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Primary Document**

**Remedial Investigation Report
for the Burial Grounds Operable Unit
at the Paducah Gaseous Diffusion Plant,
Paducah, Kentucky**



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**Remedial Investigation Report
for the Burial Grounds Operable Unit
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Paducah, Kentucky**

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Prepared for the
U.S. DEPARTMENT OF ENERGY
Office of Environmental Management

Prepared by
PADUCAH REMEDIATION SERVICES, LLC
managing the
Environmental Remediation Activities at the
Paducah Gaseous Diffusion Plant
under contract DE-AC30-06EW05001

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ACRONYMS

amsl	above mean sea level
ARAR	applicable or relevant and appropriate requirement
ASB	angled soil boring
ATSDR	Agency for Toxic Substances and Disease Registry
AT123D	Analytical Transient 1-,2-,3- Dimensional
BGOU	Burial Grounds Operable Unit
bgs	below ground surface
BHHRA	Baseline Human Health Risk Assessment
CDI	chronic daily intake
CERCLA	Comprehensive Environmental Response, Compensation, and Liability Act
<i>CFR</i>	<i>Code of Federal Regulations</i>
COC	contaminant of concern
COE	U.S. Army Corps of Engineers
COPC	chemical of potential concern
D&D	decontamination and decommissioning
DAF	dilution attenuation factor
DCE	dichloroethene
DNAPL	dense nonaqueous-phase liquid
DOE	U.S. Department of Energy
DQA	data quality assessment
DQO	data quality objective
EDD	electronic data deliverable
ELCR	excess lifetime cancer risk
EM	electromagnetic
EPA	U.S. Environmental Protection Agency
EPC	exposure point concentration
ERA	ecological risk assessment
FFA	Federal Facility Agreement
f_{oc}	organic content
FS	feasibility study
GC	gas chromatograph
GC/MS	gas chromatograph/mass spectrometer
HEAST	Health Effects Assessment Summary Tables
HI	hazard index
HQ	hazard quotient
HU	hydrostratigraphic unit
IRIS	EPA's Integrated Risk Information System
K_d	soil/water distribution coefficient
KDEP	Kentucky Department for Environmental Protection
KEEC	Kentucky Energy and Environment Cabinet
K_{oc}	organic partition coefficient
KPDES	Kentucky Pollutant Discharge Elimination System
LCS	laboratory control sample
MCL	maximum contaminant level
MDC	minimum detectable concentration
MS	matrix spike
MSA	method of standard additions

MSD	matrix spike duplicate
MW	monitoring well
NA	not applicable
NAL	no action level
NCEA	National Center for Environmental Assessment
NCP	National Contingency Plan
ND	not detected
NOAA	National Oceanic and Atmospheric Administration
NSDD	North-South Diversion Ditch
OU	operable unit
Paducah OREIS	Paducah Oak Ridge Environmental Information System
PAH	polyaromatic hydrocarbon
PCB	polychlorinated biphenyl
PEMS	Project Environmental Measurements System
PGDP	Paducah Gaseous Diffusion Plant
pH	negative logarithm of the hydrogen-ion concentration
POC	pathway of concern
POE	point of exposure
PTW	principal threat waste
QA	quality assurance
QC	quality control
RAIS	Risk Assessment Information System
RAO	remedial action objective
RCRA	Resource Conservation and Recovery Act
RfD	reference dose
RGA	Regional Gravel Aquifer
RGO	remedial goal option
RI	remedial investigation
ROD	record of decision
RPD	relative percent difference
SADA	Statistical Analysis and Decision Assistance
SESOIL	Seasonal Soil Compartment Model
SI	site investigation
SMO	Sample Management Office
SMP	site management plan
SOP	standard operating procedure
SQL	sample quantitation limit
SSL	soil screening level
SVOC	semivolatile organic compound
SWMU	solid waste management unit
TAL	target analyte list
TCE	trichloroethene
TCL	target compound list
TPU	total propagated uncertainties
TVA	Tennessee Valley Authority
UCRS	Upper Continental Recharge System
UF ₆	uranium hexafluoride
USEC	United State Enrichment Corporation
USGS	U. S. Geological Survey

VOC	volatile organic compound
WAG	Waste Area Grouping
WKWMA	West Kentucky Wildlife Management Area

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EXECUTIVE SUMMARY

The Paducah Gaseous Diffusion Plant (PGDP) is an active uranium enrichment facility that is owned by the U.S. Department of Energy (DOE). DOE is conducting environmental restoration activities at PGDP in accordance with the Paducah Federal Facility Agreement (FFA). PGDP was placed on the National Priorities List in 1994. DOE, U.S. Environmental Protection Agency (EPA), and the Commonwealth of Kentucky entered into the FFA in 1998 (EPA 1998a).

BURIAL GROUNDS OPERABLE UNIT SUMMARY

The Burial Grounds Operable Unit (BGOU) is one of five media-specific operable units (OUs) at PGDP being used to evaluate and implement remedial actions. DOE, EPA, and the Commonwealth of Kentucky have agreed upon five strategic cleanup initiatives as follows (from the Site Management Plan DOE 2009):

- BGOU Strategic Initiative,
- Decontamination and Decommissioning OU Strategic Initiative,
- Groundwater OU Strategic Initiative,
- Soils OU Strategic Initiative, and
- Surface Water OU Strategic Initiative.

The scope of the BGOU Strategic Initiative includes a remedial investigation (RI), baseline human health risk assessment, feasibility study (FS), remedy selection, and implementation of actions, as necessary, for protection of human health and the environment.

This BGOU RI assesses contamination associated with eight solid waste management units (SWMUs) that include PGDP's landfills and burial grounds; seven (SWMUs 2, 3, 4, 5, 6, 7, and 30) are located within the main PGDP secure area, and one (SWMU 145) is located within a controlled access area to the north of the main PGDP area. Two other SWMUs in the BGOU are the C-746-S (SWMU 9) and C-746-T (SWMU 10) Landfills, which are closed landfills that were not included in this RI. The following are the potential source units addressed by the BGOU RI.

- SWMU 2 C-749 Uranium Burial Ground
- SWMU 3 C-404 Low-Level Radioactive Waste Burial Ground (including the former C-404 discharge ditch)
- SWMU 4 C-747 Contaminated Burial Yard and C-748-B Burial Area
- SWMU 5 C-746-F Burial Yard
- SWMU 6 C-747-B Burial Ground
- SWMUs 7 and 30 C-747-A Burial Ground and Burn Area (which includes the area beneath SWMU 12 where it overlaps with SWMU 7)

- SWMU 145 Area P (residential/inert borrow area) and old North-South Diversion Ditch disposal trench (the area for SWMU 145 includes that beneath SWMUs 9 and 10)

Subsequent to development of the BGOU RI/FS Work Plan (DOE 2006a) and concurrent with the field investigation, an interview with a former plant operator identified potential areas of buried metal within the C-746-P and C-746-P1 Scrap Yards (SWMU 13). Assessment and remedial measures, if required, for these potential burial areas fall within the scope of the BGOU Strategic Initiative. The characterization of the potential burial areas of SWMU 13 will be addressed with a Field Sampling Plan addendum to the BGOU RI/FS Work Plan and follow-on site investigation that will be documented in a separate report. The results will be discussed with the FFA parties and, if further action is necessary, a path forward will be determined.

Remedial decisions for sediments within the BGOU SWMUs fall primarily within the scope of the Surface Water OU Strategic Initiative. Ditches of the northwest plant area that drain to the C-613 Sediment Basin will be addressed as part of the post-GDP shutdown activities for surface water.¹

The Groundwater OU Strategic Initiative will address dissolved-phase groundwater contamination in the Regional Gravel Aquifer (RGA) beneath the BGOU SWMUs; however, secondary sources of groundwater contamination that are derived from the BGOU burial grounds, such as the potential dense nonaqueous-phase liquid (DNAPL) source zones beneath SWMUs 4, 7, and 30, remain within the scope of the BGOU for assessment and remedial action, if required.

The BGOU RI/FS Work Plan (DOE 2006a) identified the following four primary goals for this RI and the follow-up FS:

- Goal 1. Characterize the nature of the source zone;
- Goal 2. Define the extent of the source zone and contamination in soil and other secondary sources at all units;
- Goal 3. Determine surface and subsurface transport mechanisms and pathways; and
- Goal 4. Support the evaluation of remedial technologies.

NATURE AND EXTENT OF CONTAMINATION (GOALS 1 AND 2)

Materials that were disposed of in each of the SWMUs of the BGOU contained hazardous substances. The conceptual model applicable to all of the BGOU SWMUs is that releases from these SWMUs have impacted soils below or adjacent to the source zones and, through vertical infiltration in the soil, have the potential to contaminate the groundwater underlying these sources. Analysis of soil and groundwater from the area of each SWMU documents the presence of metals, organic compounds, and radionuclides above screening levels. Soil and groundwater sampling results are compared with screening levels to identify the list of potential contaminants to be evaluated for the purposes of determining nature and extent of

¹ The BGOU RI risk assessment includes the available surface soils and sediments analyses for samples from within the BGOU SWMUs to complete evaluation of the exposure scenarios specified in the Work Plan (residential, industrial, and recreational).

contamination. Section 4 summarizes the characterization of the area of these SWMUs as part of the BGOU RI and as well as from previous investigation efforts.

Principal threat waste (PTW) is defined by EPA as “source materials considered to be highly toxic or highly mobile that generally cannot be reliably contained or would present a significant risk to human health or the environment should exposure occur” (EPA 1991). EPA also recognizes that “although no threshold level of risk has been established to identify principal threat waste, a general rule of thumb is to consider as a principal threat those source materials with toxicity and mobility characteristics that combine to pose a potential risk several orders of magnitude greater than the risk level that is acceptable for the current or reasonably anticipated future land use, given realistic exposure scenarios” (EPA 1997).

For the BGOU SWMUs, the trichloroethene (TCE) DNAPL at SWMUs 4, 7, and 30 and uranium at SWMUs 2 and 3 potentially are PTW. Dissolved contaminant trends in the RGA indicate that SWMU 4 and the adjoining areas of SWMUs 7 and 30 could contain TCE sources as DNAPL. The mobility and toxicity of DNAPLs make them PTW.

The uranium at SWMU 2 presents risk greater than $1E-03$ under some hypothetical exposure scenarios. Some forms of the buried uranium could be considered potential PTW where toxicity and mobility combine to pose such a risk to human health. These hypothetical exposure scenarios assume a direct contact exposure to buried waste (DOE 1997a). The uranium metal present at SWMUs 2 and 3 likely is not mobile due to its insolubility in water. The Data Summary and Interpretation Report (DOE 1997a) concluded that only some forms of uranium present may be mobile (e.g., uranyl fluoride at SWMU 2). Uncertainties concerning the risks associated with the toxicity and mobility of the uranium will be considered further during alternative evaluation in the FS.

Several analytes in subsurface soils (soils deeper than 1 ft) frequently were detected above screening criteria² used to identify contaminants for the assessment of nature and extent. Table ES.1 indicates the frequency of detection above the excavation worker no action levels (DOE 2001). All inorganic and radiological values in the table also correspond to at least one detection above background. As shown in Table ES.1, aluminum, iron, manganese, and vanadium are the most prevalent of the frequently detected contaminants in subsurface soils, often detected in more than 50% of samples representative of the nature and extent of contamination in six of the eight BGOU SWMUs. Uranium-238 was detected above screening criteria at all SWMUs except SWMU 6.

² Screening criteria for subsurface soils for the assessment of nature and extent were PGDP background levels and risk-based excavation worker no action levels. The identification of chemicals of potential concern (COPCs) for groundwater fate and transport modeling screened the subsurface soils against PGDP-specific Soil Screening Levels.

**Table ES.1. Frequency of Subsurface Soil Analytes Detected above No Action Levels
(for Analytes Detected above Background)**

Analyte	SWMU 2	SWMU 3	SWMU 4	SWMU 5	SWMU 6	SWMU 7	SWMU 30	SWMU 145
<i>Inorganics</i>								
Aluminum			94%	83%	79%	61%	88%	84%
Antimony		8%						33%
Arsenic	97%	90%	10%			72%		61%
Barium	3%		<1%	2%				2%
Beryllium	3%		5%	14%	17%	1%	8%	8%
Iron	100%		100%	97%	98%	94%	100%	
Manganese	86%		73%	75%	91%	64%	72%	
Thallium	24%							
Uranium	12%	10%				1%		13%
Vanadium	93%		99%	100%	98%		68%	
<i>Organics</i>								
Total PCB	4%		7%			3%	4%	11%
Benzo(a)pyrene							8%	
TCE	2%		3%					
1,1-DCE						3%		
Cis-1,2-DCE	3%							
Vinyl Chloride	3%		1%			1%		
<i>Radionuclides</i>								
Uranium-234	2%	3%	57%			11%	14%	19%
Uranium-235			<1%					5%
Uranium-235/236	2%					38%	20%	
Uranium-238	21%	15%	61%	21%		29%	21%	26%
Technetium-99			1%					8%
Cesium-137		3%	1%					11%
Thorium-228								85%
Thorium-230			50%			3%		9%
Thorium-232								3%
Americium-241								2%

The “no action levels” are for the Excavation Worker scenario (DOE 2001).
Blanks represent analytes which (1) did not exceed NAL or (2) exceeded NAL but not background.
Background values are from the Risk Methods Document (DOE 2001).
PCB = polychlorinated biphenyl

Metals are the most common of the frequently detected³ contaminants in both Upper Continental Recharge System (UCRS) and RGA groundwater samples. Iron and manganese are commonly present above screening levels⁴ and are the predominant contaminants in the UCRS (Table ES.2). Iron is less prevalent in the RGA (Table ES.3). In both Table ES.2 and Table ES.3, all organic analytes that were detected are listed.

³ In this section, “frequently detected” for inorganic and radiological analytes in groundwater means detected in 25% or more of the samples at levels above all screening criteria. All organics detections are listed.

⁴ While not a usable drinking water source, the screening criteria for UCRS groundwater are maximum contaminant levels (MCLs) and risk-based child resident no action levels. RGA screening criteria include PGDP background levels in addition to MCLs and risk-based child resident no action levels.

Table ES.2. Metals and Radionuclides Frequently Detected above Screening Levels and All Organic Compounds Detected in UCRS Groundwater

Source Area	Metals	Organic Compounds	Radionuclides
SWMU 2	Beryllium, Iron, Manganese, Uranium, Vanadium	1,1-DCE; TCE; <i>cis</i> -1,2-DCE; Vinyl chloride	²³⁴ U, ²³⁸ U
SWMU 3	Arsenic, Iron, Manganese, Molybdenum	TCE	⁹⁹ Tc, ²³⁴ U, ²³⁸ U
SWMU 4	Arsenic, Beryllium, Cadmium, Chromium, Copper, Iron, Lead, Manganese, Nickel, Vanadium, Zinc	<i>cis</i> -1,2-DCE; TCE; 1,1-DCE; Vinyl chloride; <i>trans</i> -1,2-DCE; Naphthalene; PCB	⁹⁹ Tc
SWMU 5	Arsenic, Beryllium, Chromium, Copper, Iron, Lead, Manganese, Molybdenum, Nickel, Vanadium, Zinc	TCE; Pyrene	--
SWMU 6	Arsenic, Chromium, Iron, Lead, Manganese, Molybdenum, Nickel, Uranium, Vanadium, Zinc	PCB-1016	⁹⁹ Tc, ²³⁴ U, ²³⁸ U
SWMU 7	Arsenic, Beryllium, Cadmium, Chromium, Iron, Lead, Manganese, Molybdenum, Nickel, Vanadium, Zinc	<i>cis</i> -1,2-DCE; TCE; Vinyl chloride ; 1,1-DCE; Benzene; Naphthalene	²²² Rn, ²³⁴ U, ²³⁸ U
SWMU 30	Arsenic, Cadmium, Iron, Lead, Manganese, Molybdenum, Nickel, Uranium, Vanadium	TCE; Vinyl chloride; Benzene; Naphthalene; PCB-1260	²³⁴ U, ²³⁸ U
SWMU 145	Arsenic, Iron, Manganese, Uranium	Chloroform	²²² Rn, ²³⁴ U, ²³⁸ U

-- = none

DCE = dichloroethene

²²²Rn = radon-222

TCE = trichloroethene

⁹⁹Tc = technetium-99

²³⁴U = uranium-234

²³⁸U = uranium-238

PCB = polychlorinated biphenyl

Table ES.3. Metals and Radionuclides Frequently Detected above Screening Levels and All Organic Compounds Detected in RGA Groundwater

Source Area	Metals	Organic Compounds	Radionuclides
SWMU 2	Arsenic, Beryllium, Cadmium, Iron, Manganese, Vanadium	1,1-DCE; TCE; <i>cis</i> -1,2-DCE; Chloroform	²³⁴ U, ²³⁸ U
SWMU 3	Arsenic, Iron, Manganese	TCE; 1,1-DCE; Chloroform	--
SWMU 4	Arsenic, Manganese, Iron, Lead	1,1-DCE; Carbon Tetrachloride; Chloroform; <i>cis</i> -1,2-DCE; TCE; Vinyl Chloride	--
SWMU 5	Iron, Lead, Manganese	TCE	--
SWMU 6	Arsenic, Iron, Lead, Manganese	TCE	--
SWMU 7	Arsenic, Beryllium, Cadmium, Iron, Lead, Manganese, Molybdenum, Nickel, Vanadium, Zinc	TCE; <i>cis</i> -1,2-DCE; Vinyl chloride; Chloroform; Carbon tetrachloride	⁹⁹ Tc, ²³⁴ U, ²³⁸ U
SWMU 30	Arsenic, Iron, Manganese	TCE; Tetrachloroethene; Chloroform	²²² Rn, ⁹⁹ Tc
SWMU 145	Arsenic, Iron, Manganese, Nickel, Vanadium	TCE; Chloroform; PCB	--

-- = none

DCE = dichloroethene

²²²Rn = radon-222

TCE = trichloroethene

⁹⁹Tc = technetium-99

²³⁴U = uranium-234

²³⁸U = uranium-238

PCB = polychlorinated biphenyl

FATE AND TRANSPORT (GOAL 3)

Modeling assessed fate and transport of contaminants for two pathways: (1) dissolved-phase transport through the aquifer and (2) vapor transport to a residential basement.⁵ Section 5 and Appendix E document the fate and transport modeling applied to the BGOU RI.

Previous work has shown that the primary pathway for groundwater flow and the site-related contaminants is predominantly vertical migration through the UCRS, followed by lateral migration in the RGA. Contaminated groundwater could migrate to the points of exposure (POEs). The POEs evaluated were at the SWMU, at the plant boundary, at the property boundary, and near the Ohio River (either at the Little Bayou Creek seeps or at the Ohio River, depending on modeled groundwater flow paths beginning at each SWMU) (Figure ES.1). Additionally, the BGOU RI includes a comprehensive evaluation of on-site risk at the SWMU that supports assessment of a SWMU boundary POE (see Table ES.4 for analytes predicted to exceed maximum contaminant levels at POEs). Not all SWMUs have transport pathways to all of the POEs. For example, SWMU 145 is located outside of the plant boundary and does not contribute to the Little Bayou seeps. SWMUs 3, 6, 7, and 30 were determined to be the only SWMUs with groundwater flow paths to the Little Bayou seeps POE, although there is some uncertainty if other SWMU flow paths are captured by the Little Bayou Creek seeps.

Table ES.4. Analytes Predicted to Exceed Maximum Contaminant Levels at the Points of Exposure

Source Area	Contaminant	SWMU	Plant Boundary	Property Boundary	Little Bayou seeps	Ohio River
SWMU 2	Arsenic	Yes ^a	No ^b	No	N/A ^c	No
	<i>cis</i> -1,2-DCE	Yes	Yes	Yes	N/A	Yes
	TCE	Yes	Yes	Yes	N/A	Yes
SWMU 3	Arsenic	Yes	No	No	No	N/A
	⁹⁹ Tc	Yes	Yes	Yes	No	N/A
	Uranium	Yes	No	No	No	N/A
SWMU 4	Arsenic	Yes	No	No	N/A	No
	<i>cis</i> -1,2-DCE	Yes	Yes	Yes	N/A	No
	⁹⁹ Tc	Yes	Yes	Yes	N/A	No
	TCE	Yes	Yes	Yes	N/A	Yes
	Vinyl Chloride	Yes	Yes	Yes	N/A	No
SWMU 5	No analytes predicted to exceed MCLs at POEs					
SWMU 6	No analytes predicted to exceed MCLs at POEs					
SWMU 7	1,1-DCE	Yes	Yes	No	No	N/A
	Arsenic	Yes	Yes	No	No	N/A
	⁹⁹ Tc	Yes	No	No	No	N/A
	TCE	Yes	Yes	No	No	N/A
	Vinyl Chloride	Yes	Yes	No	No	N/A
SWMU 30	Arsenic	Yes	Yes	No	No	N/A
SWMU 145	Antimony	Yes	N/A	No	N/A	No
	Arsenic	Yes	N/A	No	N/A	No
	⁹⁹ Tc	Yes	N/A	Yes	N/A	Yes

^a Yes = The modeled analyte concentration exceeds its maximum contaminant level
^b No = The modeled analyte concentration does not exceed its maximum contaminant
^c N/A = Not applicable: the POE does not apply to the SWMU.
DCE = dichloroethene

MCL = maximum contaminant level
POE = point of exposure
TCE = trichloroethene
⁹⁹Tc = technetium-99

⁵ Assessment of surface water runoff and sediments is principally the scope of the Surface Water OU.

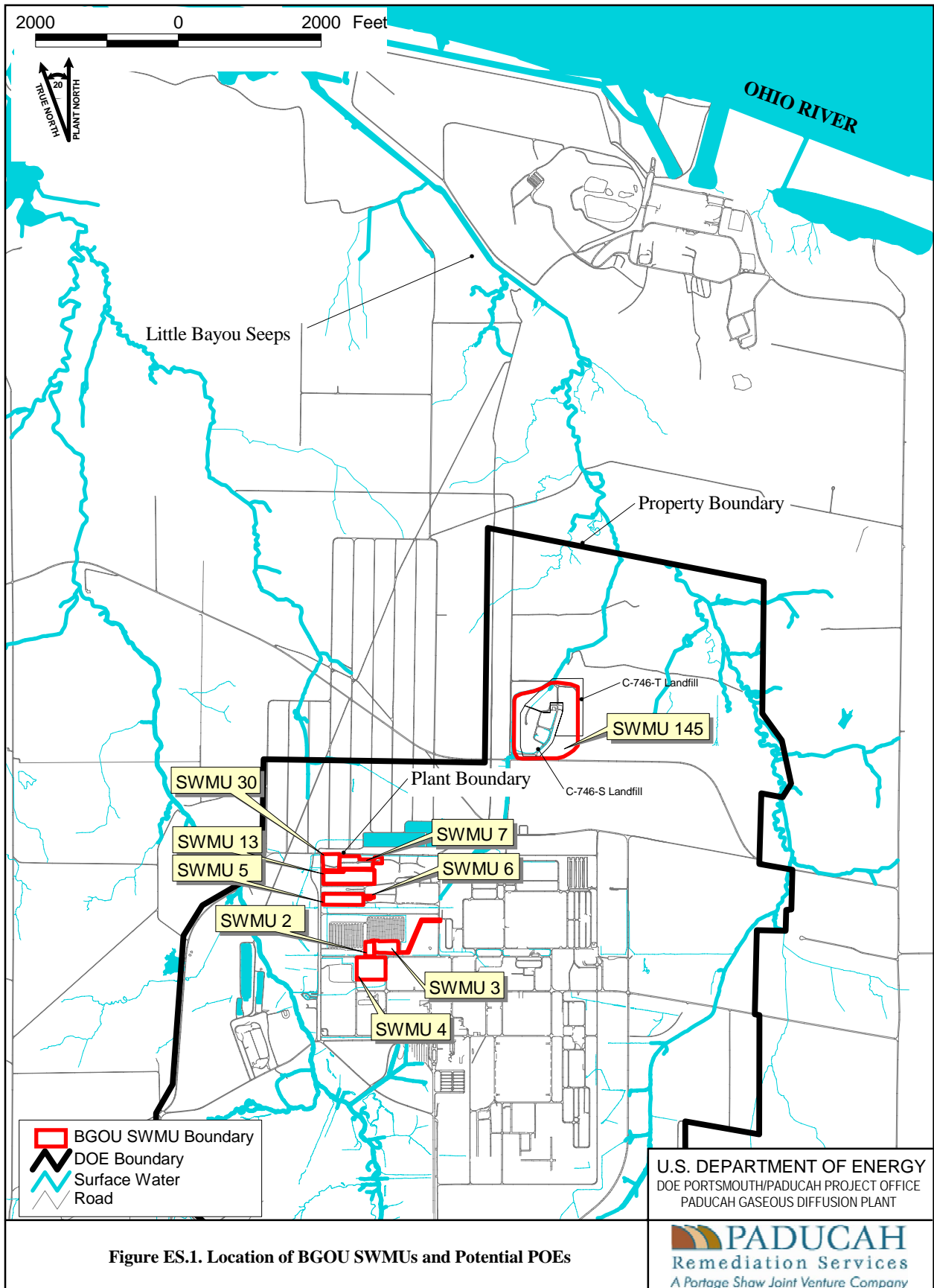


Figure ES.1. Location of BGOU SWMUs and Potential POEs

Figure No. \BGOU\d2-ri sect1.apr
DATE 09-29-09

Vapor transport modeling assessed contaminant concentrations in a hypothetical residential basement at the SWMU and in hypothetical residential basements at the plant boundary and property boundary POEs (Table ES.5). At some POEs, the excess lifetime cancer risk (ELCR) or hazard posed by hypothetical exposure to the modeled air concentration exceeded 1E-06 or 0.1, respectively. Currently there are no buildings or pipelines connected to buildings located over the contaminated material at the BGOU SWMUs. The existing buildings at PGDP are slated to be decontaminated and demolished after the plant closes; therefore, the on-site industrial worker scenario was not evaluated in the vapor modeling analysis. There are no plans to construct future facilities on the BGOU SWMUs.

Table ES.5. Analytes with Basement Air Concentrations of Concern Based on Vapor Transport Modeling Results at the Points of Exposure

Source Area	Contaminant	SWMU Boundary	Plant Boundary	Property Boundary
SWMU 2	TCE	Yes ^a	Yes	Yes
	<i>cis</i> -1,2-DCE	Yes	No ^b	No
SWMU 3	TCE	Yes	No	No
	Mercury ^c	Yes	No	No
SWMU 4	TCE	Yes	Yes	Yes
	Vinyl Chloride	Yes	Yes	No
	<i>cis</i> -1,2-DCE	Yes	No	No
SWMU 5	No analytes with basement air concentrations of concern			
SWMU 6	No analytes with basement air concentrations of concern			
SWMU 7	TCE	Yes	No	No
	Vinyl Chloride	Yes	No	No
	1,1-DCE	Yes	Yes	No
	Mercury	Yes	No	No
SWMU 30	TCE	Yes	No	No
	1,1-DCE	Yes	No	No
	Mercury	Yes	No	No
SWMU 145	Mercury	Yes	No	No

DCE = dichloroethene

SWMU = solid waste management unit

TCE = trichloroethene

^a Yes = Modeled air concentration equals or exceeds 1.0E-06 excess lifetime cancer risk (ELCR) or 0.1 hazard quotient (HQ)

^b No = Modeled air concentration is less than 1.0E-06 ELCR or 0.1 HQ

^c The vapor transport modeling for mercury was conservatively based on the metallic form, which has a Henry's Law Constant of 1.07E-02 atm-m³/mol. The rate of vaporization of mercury and certain of its inorganic compounds decrease in the sequence Hg > Hg₂Cl₂ > HgCl₂ > HgS > HgO. The Henry's Law Constant decreases dramatically down the sequence (for example, HgCl₂ has a value of 7.09E-10 atm-m³/mol).

Vapor transport modeling conducted with the Johnson and Ettinger model (1991), coded into spreadsheets by EPA (2004).

RISK ASSESSMENT (GOAL 4)

PGDP is an industrial facility. The reasonably anticipated future use of the area within the current plant boundary is expected to remain industrial. This expectation should be considered when using the risk information provided in the report to support risk management decision making.

For soil, results from previous risk assessments were used. The BGOU Work Plan did not call for additional surface (0-1 ft bgs) or subsurface (0-10 ft bgs) soil sampling at most SWMUs. The risk for the on-site resident for soil exceeds 1E-04 and the hazard index (HI) is greater than 1 at all SWMUs except for SWMUs 2, 3, and SWMU 145 (which were not evaluated for soil exposure for this scenario). The contaminants that are risk drivers for soil are aluminum, arsenic, beryllium, chromium, iron, nickel, Total

polyaromatic hydrocarbons (PAHs), uranium (as a metal), uranium-234, uranium-238, vanadium, and zinc.

Residential use of groundwater was evaluated at the SWMU boundary, plant boundary, property boundary, and Ohio River (or seeps) for all SWMUs except SWMU 6 [SWMU 6 had no groundwater chemicals of potential concern (COPCs)] and SWMU 145 (SWMU 145 was not evaluated at the plant boundary since it lies outside that boundary). At the SWMU boundary, risks and hazards from groundwater use for all evaluated SWMUs exceeded 1E-04 risk and exceeded an HI of 1. The major contaminants driving the groundwater risks and hazards at the SWMU boundary POE are arsenic; antimony; Aroclor-1260; *cis*-1,2-dichloroethene (DCE); 1-1-DCE; manganese; naphthalene; Total polychlorinated biphenyls (PCBs); TCE; technetium-99; uranium; and vinyl chloride. At the plant boundary, risks and hazards from groundwater for SWMUs 2, 3, 4, 5, 7, and 30 exceeded 1E-04 risk or exceeded an HI of 1. At the property boundary, risks and hazards from groundwater for SWMUs 2, 4, 7, 30, and 145 exceeded 1E-04 risk or exceeded an HI of 1. At the Ohio River (or seeps), risks and hazards from groundwater for SWMUs 2, 4, 7, and 30 exceeded 1E-04 risk or exceeded an HI of 1. The major contaminants driving the groundwater risks and hazards at the property boundary and Ohio River (or Little Bayou Creek seeps) POEs are arsenic, *cis*-1,2-DCE, 1,1-DCE, TCE, technetium-99, and vinyl chloride. While the migration of contamination from the potential TCE DNAPL zones at SWMU 4 and SWMUs 7 and 30 were not modeled due to uncertainties in source term development, a qualitative analysis completed considering results from the previous studies done for the PGDP (e.g., C-400 DNAPL source) indicates that TCE migration from these sources would have resulted in potential risks exceeding 1E-04 at all POEs.

For exposure to soil, at least one of the on-site receptor scenarios (industrial worker, excavation worker, or recreational user), all SWMUs (except SWMU 145 where the scenarios were not assessed) have an ELCR > 1.0E-06. For at least one of these scenarios, SWMUs 4, 5, 6, 7, and 30 have HIs > 1. Soil exposures to industrial or excavation workers are more relevant to the potential future uses of the site.

For the excavation worker who is exposed to both surface soil and subsurface soil (soil from 0 to 10 ft depth), HIs were greater than one at SWMUs 4, 5, 6, 7, and 30. Risks for the excavation worker exceeded 1E-04 at SWMUs 4, 5, 6, 7, and 30. The risk drivers for the excavation worker scenario were arsenic, beryllium, Total PAHs, uranium, uranium-235, and uranium-238.

Given the reasonably anticipated future industrial use of the areas within the plant boundary, the most likely future and current scenario is the industrial worker. The ELCR for the future industrial worker scenario exceeded 1E-04 at SWMUs 2, 3, 4, 5, 6, 7, and 30 primarily due to risk from arsenic, beryllium, Total PAHs, uranium-235, and uranium-238. The HI exceeds 1 for the industrial worker at SWMUs 4, 7, and 30; aluminum, beryllium, chromium, iron, manganese, uranium, and vanadium are the hazard drivers. Risks for the current industrial worker (at 16 days per year of exposure) were less than those for the future industrial worker; risks for the current industrial worker exceeded 1E-04 at SWMUs 4, 5, 6, 7, and 30. Table ES.6 details the exposure pathways and contaminants of concern associated with dominant risk for each SWMU for exposure to subsurface soil and groundwater.

The inclusion of beryllium as a risk driver is a result of incorporating the historical risk assessments. At the time those risk assessments were developed, beryllium still was evaluated as a carcinogen. Since then, the oral cancer slope factor for beryllium has been withdrawn and no longer is used for PGDP risk assessments. As a result, the total ELCR becomes much lower at those SWMUs where beryllium is a contaminant of concern. For SWMUs 4 and 6, removal of the contribution of beryllium to the ELCR reduces the total ELCR to within the EPA risk range for the industrial worker scenario.

Table ES.6. Exposure Routes and Exposure Pathways and Contaminants of Concern Associated with Dominant Risk for Each SWMU for Exposure to Subsurface Soil and Groundwater

Source Area	HI	ELCR
SWMU 2	– Ingestion of groundwater and household inhalation of vapors(TCE; <i>cis</i> -1,2-DCE)	– Household inhalation of vapors (TCE) – Ingestion of groundwater (TCE) – External exposure to subsurface soil (uranium-235, uranium-238)
SWMU 3	– Ingestion of groundwater (arsenic, uranium)	– Ingestion of groundwater (arsenic, ⁹⁹ Tc) – External exposure to subsurface soil (uranium-235, uranium-238)
SWMU 4	– Ingestion of groundwater (TCE) – Dermal exposure to soil (chromium, iron)	– Household inhalation of vapors and dermal exposure (TCE, vinyl chloride) – Dermal exposure to subsurface soil (beryllium)
SWMU 5	– Ingestion of RGA groundwater (arsenic, naphthalene) – Ingestion of vegetables (arsenic, aluminum)	– Ingestion of RGA groundwater (arsenic)
SWMU 6	– Ingestion of vegetables (chromium) – Dermal exposure to soil (chromium)	– Dermal exposure to subsurface soil (PAHs, beryllium) – Ingestion of vegetables (PAHs, beryllium)
SWMU 7	– Ingestion of RGA groundwater (TCE, arsenic, Aroclor-1254) – Ingestion of vegetables (iron, uranium) – Dermal exposure to soil (vanadium, iron, uranium)	– Household inhalation of vapors and ingestion of RGA groundwater (1,1-DCE) – Dermal exposure and ingestion of vegetables (beryllium, uranium-238)
SWMU 30	– Ingestion of RGA groundwater (TCE) – Ingestion of subsurface soil (uranium) – Dermal exposure to soil (vanadium, iron)	– Household inhalation of vapors (TCE) – Ingestion of vegetables (beryllium, uranium-238)
SWMU 145	– Ingestion of RGA groundwater (antimony, arsenic)	– Ingestion of RGA groundwater (Aroclor-1260)

DCE = dichloroethene
PAH = polyaromatic hydrocarbon
PCB = polychlorinated biphenyl
RGA = Regional Gravel Aquifer
⁹⁹Tc = technetium-99
TCE = trichloroethene

The BGOU RI includes a summary of previous ecological risk assessments for SWMUs 2, 4, 5, 6, 7, and 30. Neither SWMU 3 nor SWMU 145 has been assessed for ecological risk. SWMU 3 is covered by a Resource Conservation and Recovery Act (RCRA) cap, and SWMU 145 is located beneath the C-746-S and -T Landfills, which also are covered by caps. Comparison of site characterization data against No Further Action screening levels determined that all of the SWMUs have metals and organic compounds (in surface soil) that are COPCs for ecological risk to the environment, while SWMUs 7 and 30 have a radionuclide COPC (in surface soil).

CONCLUSIONS

The following are the major contaminant distribution findings for sources investigated in the BGOU RI.

- Environmental media, specifically subsurface soil and groundwater, have been impacted by releases of contaminants at all of the BGOU SWMUs.
- TCE trends in the RGA indicate that TCE DNAPL likely is present at SWMU 4 and in the vicinity of the shared border between SWMUs 7 and 30. (See Sections 3.9.4, 4.5.2, and 4.8.2.) Concentrations of TCE at SWMU 4 suggest this potential TCE DNAPL may be present both in the waste cells and underlying soils of the UCRS and in the matrix of the RGA. TCE trends at SWMUs 7 and 30 indicate that this potential TCE DNAPL source is likely constrained to the UCRS soils.
- The Baseline Human Health Risk Assessment completed as part of the BGOU RI indicates that ELCRs greater than the upper end of EPA's acceptable risk range (i.e., 1E-04) and HIs greater than 1 exist at all SWMUs; therefore, an FS is appropriate for impacted media at each SWMU. The metals arsenic, beryllium, and uranium; the organic compounds Total PAHs and Total PCBs; and the radionuclides uranium-235 and uranium-238 are common contaminants that present the dominant risks from exposure to surface and subsurface soil. The major contaminants driving the groundwater risks at the on-site POEs are arsenic, Aroclor-1260, 1,1-DCE, TCE, technetium-99, and vinyl chloride.
- Migration of contaminants through groundwater from all but SWMU 6 to locations at the SWMU boundary, the plant boundary, property boundary, and near the Ohio River also posed greater than *de minimis* risks to a hypothetical residential groundwater user. Arsenic, TCE, 1,1-DCE, technetium-99, and vinyl chloride are the primary risk drivers.
- The Screening Ecological Risk Assessment retained a number of COPCs, primarily metals, at each of the sites. Each SWMU requires further ecological evaluation.

The BGOU RI/FS Work Plan developed decision rules for the BGOU Strategic Initiative. Table ES.7 presents the decision rules (DOE 2006a).

The risk levels associated with contamination at all of the SWMUs and associated with groundwater contamination derived from all of the SWMUs meet the criteria of the decision rules to progress to evaluate actions that will mitigate risk to protect human health and the environment and to achieve applicable or relevant and appropriate requirements (ARARs); to seek an ARAR waiver in accordance with EPA guidance; or to propose alternative standards. The following are the preliminary BGOU remedial action objectives:

- Contribute to protection of current and future residential receptors from exposure to contaminated groundwater by reducing/controlling sources of groundwater contamination;
- Protect industrial workers from exposure to waste and contaminated soils; and
- Treat or remove principal threat wastes wherever practicable, consistent with 40 *CFR* § 300.430 (a)(iii)(A).

A follow-on FS will develop and evaluate remedial action alternatives for the BGOU SWMUs.

Table ES.7. Decision Rules for the BGOU Strategic Initiative

GOAL	DECISION RULE		Then statement
		If statement	
Nature of Contamination	1a	If the concentration of analytes found in the source zone could result in a cumulative ELCR greater than 1×10^{-6} or a cumulative HI greater than 1 through contact with contaminated media, or if the concentration of analytes in the source zone could result in detrimental impacts to nonhuman receptors through contact with contaminated media as indicated by exceeding ecological screening criteria, and if the concentrations of analytes in the source zone are greater than those expected to occur naturally in the environment,	then evaluate actions that will mitigate risk; otherwise pursue a “no further action” decision (see D1b and D1c).
	1b	If concentrations of analytes found in the source zone exceed ARARs,	then evaluate actions that will bring contamination within the source zone into compliance with ARARs; seek an ARAR waiver (such as technical impracticability, inconsistent application of state standards, interim measure, greater risk to human health and the environment, equivalent standard of performance) in accordance with EPA guidance; or propose/obtain alternative standards.
	1c	If contaminants found at the site are known to transform or degrade into chemicals that could lead to increased risks to human health or the environment or into chemicals for which there are ARARs, and if the concentrations of these contaminants could result in risks greater than those defined in D1a or concentrations greater than ARARs,	then evaluate actions that will mitigate potential future risk or obtain compliance with ARARs; seek an ARAR waiver (such as technical impracticability, inconsistent application of state standards, interim measure, greater risk to human health and the environment, equivalent standard of performance) in accordance with EPA guidance; or propose/obtain alternative standards.
Extent of Contamination	2a	If secondary contamination sources are found, and if the concentration of analytes within the secondary contamination source is found to potentially result in a cumulative ELCR greater than 1×10^{-6} or a cumulative HI greater than 1 through contact with contaminated media at the unit, and if the concentrations of analytes are greater than those expected to occur naturally in the environment,	then evaluate actions that will mitigate risk; otherwise, do not consider secondary contamination sources when making remedial decisions for the unit.

Table ES.7. Decision Rules for the BGOU Strategic Initiative (Continued)

GOAL	DECISION RULE		
	<i>If statement</i>	<i>Then statement</i>	
Fate and Transport	3a	<p>If contaminants are found in the source zone, or if secondary contamination sources are found, and if these contaminants are found to be migrating or may migrate from the source zone or from secondary contamination sources at concentrations that may potentially result in a cumulative ELCR greater than 1×10^{-6} or a cumulative HI greater than 1 through use of contaminated media at downgradient points of exposure, and the concentrations of analytes are greater than those expected to occur naturally in the environment,</p>	<p>then evaluate actions that will mitigate risk; otherwise, do not consider risk posed by migratory pathways when evaluating remedial alternatives for the unit (see D3b).</p>
	3b	<p>If contaminants are found in the source zone, or if secondary contamination sources are found, and if these contaminants are found to be migrating or may migrate from the source zone or from the secondary contamination source at concentrations that exceed ARARs,</p>	<p>then evaluate actions that will bring migratory concentrations into compliance with ARARs; seek an ARAR waiver (such as technical impracticability, inconsistent application of state standards, interim measure, greater risk to human health and the environment, equivalent standard of performance) in accordance with EPA guidance; or propose/obtain alternative standards; otherwise, do not consider ARARs when examining migratory pathways during the evaluation of remedial actions (see D3a).</p>
Risk Assessment	4a	<p>If Decision D1a, D1b, D1c, D2a, D3a, or D3b indicate that response actions are needed,</p>	<p>then evaluate response actions to mitigate risk in the source zone.</p>

UNCERTAINTIES/ASSUMPTIONS

The BGOU Work Plan identified data gaps on a SWMU-by-SWMU basis that were necessary to be filled in order to move forward with the FS. The Work Plan was implemented to reduce any remaining uncertainties from previous investigations regarding the nature of the source zone, extent of the source zone and secondary sources, surface and subsurface transport mechanisms, and to support evaluation of remedial technologies in the FS.

Nature of the Source Zone

A key project assumption for the upcoming FS is that the available historical documentation and soil and groundwater characterization data are sufficient relative to waste characteristics, to chemical and physical properties, and to waste volume estimates to evaluate general response actions, to screen technology types, and to conduct detailed alternative analysis for the BGOU. The potential impact of source zone uncertainties on alternatives analysis will be further documented and analyzed in the FS. While the BGOU RI field investigation sampled directly beneath the waste units using angled borings, it remains possible that the buried waste contains hazards or constituents that current sample results do not characterize (historical disposal records and waste manifests are incomplete). Additional uncertainty exists for SWMU 5 because wastes from other facilities were buried on-site through a “Work for Others” program. These waste streams may be unrelated to the typical materials used in gaseous diffusion. A related uncertainty is that the field investigation was unable to sample to the middle of a few of the larger SWMUs⁶ (SWMUs 5 and 145, particularly); therefore, there are some uncertainties in the nature and extent of the contaminant source that will need to be managed during the decision making process.

Aerial photographs were reviewed to help with locating the extent of burial pits or past activities associated with the BGOU SWMUs; however, there is some uncertainty with this approach since aerial photographs are for sporadic time periods and there were several years where photographs are not available to view to aid with delineating the extent of each burial ground.

Many of the SWMUs have been investigated previously. The BGOU RI uses a combination of historical and current sample results of soil and groundwater from the area of each SWMU. The results of previous investigations and RI sampling document and confirm the presence of metals, organic compounds, and radionuclides in the BGOU burial grounds. The associated samples were collected and analyzed over several previous and continuing investigations, as well as the BGOU RI, using several methods. Quality control/quality assurance practices at PGDP, now and previously, limit the uncertainty associated with the sampling and analysis process. Nevertheless, changes have occurred to analytical methods that limit the strict comparison of data (e.g., laboratory reporting limits have varied over time). In some cases, analytical method detection limits are above screening criteria, such as the Excavation Worker No Action Level; these data are of limited value to the RI. One potential change with large impact to the interpretation of data is the preservation of groundwater samples. Both filtered and unfiltered samples were collected in the field as “unfiltered,” with one of the fractions (the sample preserved for unfiltered analyses) being acidified in the field. The other sample was filtered and acidified in the laboratory. The unfiltered analyses represent the “whole water” chemistry (including both the dissolved and suspended fractions). These analyses may be significantly biased by turbid samples derived from temporary borings

⁶ Soil samples that best characterize releases from the SWMUs are limited to the UCRS (the shallowest 50 to 60 ft of soils). The shallowest angle of drilling/sampling that could be achieved in the deeper soil borings was 45 degrees from vertical; thus, the furthest distance under the burial grounds that could be sampled ranged from 30 to less than 60 ft, depending on the depth of the burial pit and depth to top of RGA.

and likely would lead to an overestimation of soluble transportable contaminants. Groundwater metals and radionuclide analyses would be most affected.

To minimize the potential for “age” to bias the analysis of the data, the historical sample analyses used in the BGOU RI are limited to groundwater samples collected in January 1995 and later and soil samples collected in June 1996 and later. This criterion, which was established during scoping for the BGOU RI and is documented in Section 5 of the BGOU RI Work Plan, maximizes the number of historical sample analyses available to the RI, while providing a reasonable assurance of the comparability of the data. There are limited monitoring wells in close proximity to many of the SWMUs, but temporary borings provide a snapshot of the conditions where groundwater samples could be obtained. Permanent monitoring wells would reduce the uncertainties associated with seasonal variations in water levels and contaminant trends.

Where changing conditions are present, the earlier data will be less representative of current conditions. Factors in the UCRS that might cause significant change include the degradation of organic wastes or the episodic breach of a waste container. An additional factor that might impact conditions in the RGA is an advancing contaminant plume from an upgradient source. The increasing TCE level in the RGA at SWMUs 2 and 3 is an example of conditions in the RGA being impacted by an upgradient source.

The potential for acidic leachate at each SWMU is uncertain due to the lack of disposal records. SWMUs with the greatest potential for acidic leachate are SWMU 6 (exhaust fans with perchloric acid) and SWMU 4 (records of chemicals buried are incomplete). It should be noted that angled borings beneath SWMU 6 found no evidence of acidic leachate, either from subsurface metal concentrations or groundwater pH. There are no pH measurements from shallow groundwater at SWMU 4 to allow an evaluation of the uncertainty related to acidic leachate. The potential for acid leaching at the SWMUs will be evaluated further relative to the importance of acid leaching in screening, and detailed analysis of alternatives in the FS.

For SWMUs 2 and 6, where the last disposal occurred in 1977 (32 years ago) and 1976 (33 years ago), respectively, it is reasonable to assume most, if not all, drums have failed (an Oak Ridge National Laboratory researcher estimated that failure of steel drums would be expected to occur within 18 to 36 years). At SWMU 5, where the last disposal occurred 22 years ago, it is reasonable to assume some drums still may be intact. For SWMUs 4, 7, 30, and 145, it can be assumed that drums likely are breached since they were dumped rather than having been carefully stacked. Because all drummed waste was assumed to have been released to the environment during disposal or through degradation, samples from soils surrounding the buried wastes were used to evaluate potential contaminant migration and risks associated with the SWMUs. This approach resulted in the inclusion of SWMUs with drummed waste exhibiting risk and hazard values that exceeded acceptable levels; therefore, though the integrity of buried drums is an uncertainty, the overall objectives of the RI analysis were met. The risk assessment concluded that these uncertainties related to the source zone were not estimated to have a large effect on the risk characterization and does not affect the need for taking action. Data on drummed wastes are not as critical to the decision making process, because the risk calculated from the existing data sets already indicate a need for action. The potential for unanticipated wastes is an uncertainty to be managed in the FS, because it could affect the timeline or efficacy of remedial options. This issue is discussed in more detail in Appendix F, Section F.6.4.2.

The BGOU FS may identify the need for remedial design support or additional source delineation after final selection of media-specific remedial goals and remedial alternatives. In addition, monitoring of the source zones will be evaluated in the FS.

Extent of the Source Zone and Secondary Sources

Secondary sources of groundwater contamination that are derived from the BGOU burial grounds, such as the potential DNAPL source zones beneath SWMU 4 and SWMU 7, are within the scope of the BGOU for evaluation and remedial action. The evidence for UCRS DNAPL presence is documented in previous investigations (DOE 2007a; DOE 1998a) and discussed in the RI. Collection of UCRS groundwater samples was attempted from 32 angled borings in order to detect releases or secondary sources that may be related to the SWMUs. Of the 32 attempts, 17 boring locations provided enough groundwater to collect a sample. Assessment of the secondary source in the UCRS at SWMUs 7 and 30 was based on both historical and newly generated data, while the assessment of the secondary source in both the UCRS and RGA at SWMU 4 is based primarily on historical data. Because the UCRS water samples supplement only the characterization of the BGOU SWMUs (the analysis of subsurface soil samples is the primary measure that supports the assessment of nature and extent and risk) and secondary sources, the lack of UCRS water samples from all soil borings does not limit the planned assessment of the SWMUs. There is also potential for a TCE DNAPL source at SWMU 2 based on historical disposal records; however, neither the subsurface soil nor shallow groundwater data at SWMU 2 indicate a DNAPL source. The volumetric extent of secondary source contamination has been approximated and constitutes a project assumption for evaluation of the alternatives. The impact of source volume or DNAPL uncertainties will be evaluated and further discussed in the FS.

There remains some uncertainty with regard to the boundaries of the burial pits. Geophysical surveys have not been completed across the entire area of all SWMUs (SWMU 6 had equipment in the way when it was surveyed). This uncertainty will need to be managed in the FS.

Surface and Subsurface Transport Mechanisms

Previous work has shown that the primary pathway for groundwater flow and the site-related contaminants is vertical migration through the UCRS, followed by lateral migration in the RGA. Contaminated groundwater could migrate to the POEs identified for the BGOU SWMUs at the plant boundary, property boundary, surface seeps at Little Bayou Creek, and near the Ohio River. Not all SWMUs have transport pathways to all of the POEs. For example, SWMU 145 is located outside of the plant boundary and does not contribute to the Little Bayou seeps. SWMUs 3, 6, 7, and 30 were determined to be the only SWMUs contributing to the Little Bayou seeps POE. SWMUs 2, 4, and 5 are potential contributors to the Little Bayou Creek seeps, but the numerical model used to define flowpaths indicated releases from those SWMUs would go to the Ohio River. While there is some uncertainty related to modeling in predicting whether a SWMU would contribute to the Little Bayou seeps or the Ohio River, the model results indicate this uncertainty has an insignificant effect on the modeled contaminant concentrations and should not affect remedial decisions.

The location of the water table varies in burial ground SWMUs. Most of the buried waste at SWMU 2 is saturated. The westward slope of the water table below SWMU 2 indicates that the water table must be equally shallow beneath SWMU 3. Because SWMU 3 is an aboveground facility with a RCRA multi-layered cap, all but the base of the landfill wastes likely are unsaturated. The stratigraphy of SWMU 4 is comparable to that of SWMUs 2 and 3. It appears that the hydrogeologic setting is similar, and the water table likely extends up into the waste burial pits. At SWMUs 5 and 6, even the shallowest wastes (with top near 365 ft amsl), likely are buried below the water table (at an elevation of approximately 367 ft amsl on the north side of SWMU 5). The SWMUs 7 and 30 RI (DOE 1998a) determined that a shallow water table exists approximately 5 ft bgs (Figure 3.22) and within the burial cells. UCRS piezometer and well measurements documented a strong downward gradient within the area. The elevation of the water table remains poorly documented at SWMU 145. Some buried waste at SWMU 145 likely is saturated.

There are water pipelines and electrical conduits in the proximity of some of the SWMUs. These conduits could act as pathways for contaminant transport. Because they were not specifically evaluated in the RI or in past RIs, uncertainty remains as to any contaminant transport along these pipelines and conduits. Due to the strong downward hydraulic gradient in the UCRS and the fact that the pipelines are not located below the waste cells, transport along these types of features should be limited.

Uncertainty does exist with regard to the dissolved oxygen in the UCRS at SWMUs 4 and 6 due to a lack of data. The majority of dissolved oxygen measurements from UCRS wells range from near zero to four mg/L and oxidation/reduction potential commonly ranges from -100 to 300 microVolts, with the majority of measurements greater than zero. Line plots in Figure 3.9 of the RI further demonstrate trends of dissolved oxygen (517 measurements) and oxidation/reduction potential (136 measurements) in the UCRS at the BGOU SWMUs. Due to the relative abundance of measurements for most SWMUs that demonstrate that the cumulative trend is likely to be representative of conditions at each SWMU, the oxidation/reduction potential in the UCRS at SWMUs 4 and 6 will be assumed to be similar to that in the UCRS at other BGOU SWMUs. The impact of this assumption will be evaluated further in the FS. If determined necessary to support implementation of a remedial alternative, dissolved oxygen could be measured as part of a remedial design support investigation for SWMUs 4 and 6.

Another potential pathway that exists, likely at SWMUs 7 and 30, but also possibly at other BGOU SWMUs, is lateral seepage from the burial pits into nearby ditches. The SWMU 7 and 30 RI Report (DOE 1998a) reported that water was observed emanating from the slope of a nearby ditch following a heavy rainfall. It is uncertain whether the seepage was derived from the burial pits. The RI report concluded that uranium activity concentrations in the ditch sediments suggest SWMUs 7 and 30 are contributing to contamination in the ditch, but the uranium isotope activity ratios in surface water in the ditch argued against waste burial pit waters as contributors to surface water contamination. The increased radiological activity is best explained by surface erosion carrying soil-bound radionuclides to the ditch. Likewise, some discharge of shallow groundwater in the ditch south of SWMU 2 has been observed, but the report was unclear as to the contribution of contamination to the ditch (the report concluded that contaminant migration to Outfall 015 and Bayou Creek is unlikely to exceed preliminary remediation goals) (DOE 1997a). This uncertainty will need to be managed in the FS.

Several uncertainties identified above will need to be managed in the FS. These include the following:

- Uncertainties related to risks associated with the mobility of uranium (the FS will manage this uncertainty by evaluating appropriate technologies for SWMUs where uranium is a primary contaminant);
- Uncertainties concerning the extent of source zones (burial areas) and the impact on alternative analyses (the FS will use existing knowledge and manage the uncertainties regarding the volume requiring removal or treatment);
- Uncertainties regarding the potential for acidic leachate, oxidation/reduction conditions, and degree of waste saturation (the FS will manage these uncertainties by evaluating robust technologies that are not sensitive to these types of uncertainties);
- Uncertainties regarding the extent and volume of secondary source zones such as TCE DNAPL and intact drummed wastes (the FS will manage uncertainties regarding the extent and volume of these sources for comparison);
- Uncertainties related to limited groundwater monitoring around the BGOU SWMUs (the FS will

manage this uncertainty by incorporating additional groundwater monitoring where appropriate at SWMUs where effectiveness monitoring is needed or where waste is left in place).

1. INTRODUCTION

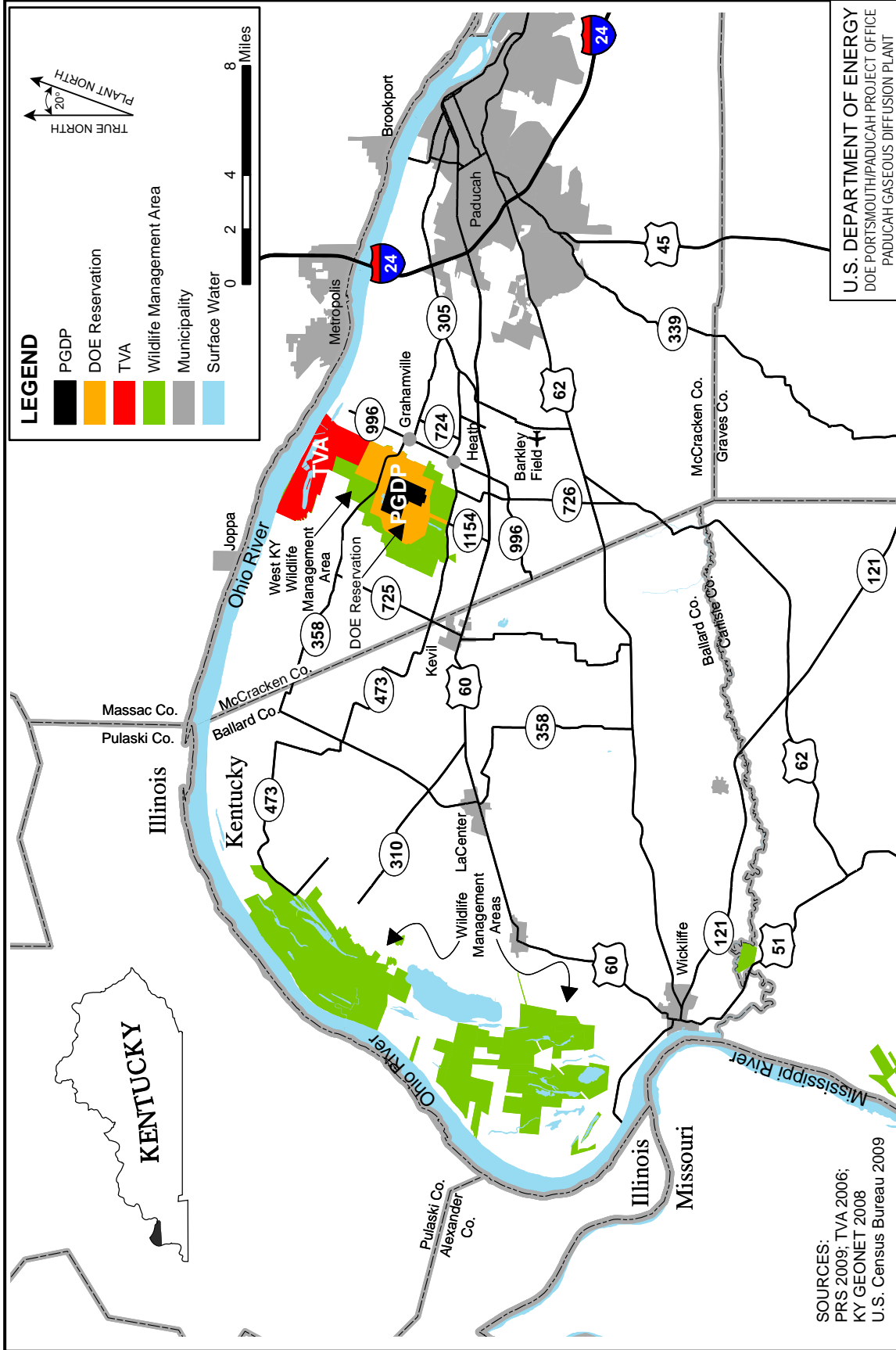
The Paducah Gaseous Diffusion Plant (PGDP), located approximately 10 miles west of Paducah, Kentucky, and 3.5 miles south of the Ohio River in the western part of McCracken County, is an active uranium enrichment facility owned by the U.S. Department of Energy (DOE). Bordering the PGDP reservation to the northeast, between the plant and the Ohio River, is a Tennessee Valley Authority (TVA) reservation on which is located the electricity generating Shawnee Steam Plant (Figure 1.1).

PGDP was owned and managed first by the Atomic Energy Commission and then the Energy Research and Development Administration, DOE's predecessors; DOE then managed PGDP until 1993. On July 1, 1993, Martin Marietta Utility Services and later the United States Enrichment Corporation (USEC) assumed management and operation of the PGDP enrichment facilities under a lease agreement with DOE. DOE still owns the enrichment complex and is responsible for environmental management activities associated with past operation of PGDP (CERCLIS # KY8-890-008-982). DOE is the lead agency for remedial actions in accordance with the Paducah Federal Facility Agreement (FFA), and the U.S. Environmental Protection Agency (EPA) and the Kentucky Energy and Environment Cabinet (KEEC) are regulatory oversight agencies (EPA 1998a).

The Burial Grounds Operable Unit (BGOU) consists of contamination associated with some of PGDP's landfills and burial grounds and known and suspected waste disposal areas in the Solid Waste Management Unit (SWMU) 13 C-746-P and C-746-P1 Scrap Yards. Burial grounds addressed by this remedial investigation (RI) are listed below and shown in Figure 1.2 (DOE 2006a).

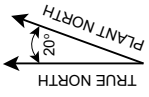
- SWMU 2—C-749 Uranium Burial Ground
- SWMU 3—C-404 Low-Level Radioactive Waste Burial Ground (plus the former C-404 discharge ditch)
- SWMU 4—C-747 Contaminated Burial Yard and C-748-B Burial Area
- SWMU 5—C-746-F Burial Yard
- SWMU 6—C-747-B Burial Ground
- SWMUs 7 and 30—C-747-A Burial Ground and Burn Area (which includes the area beneath SWMU 12 where it overlaps with SWMU 7)
- SWMU 145—Area P (the residential/inert borrow area) and old North-South Diversion Ditch (NSDD) disposal trench (the area for SWMU 145 includes that beneath SWMUs 9 and 10)

Subsequent to development of the BGOU RI/Feasibility Study (FS) Work Plan (DOE 2006a) and concurrent with the field investigation, an interview with a former plant heavy-equipment operator identified potential areas of buried metal within the C-746-P and C-746-P1 Scrap Yards (SWMU 13). Assessment and remedial measures, if required, for these potential burial areas fall within the scope of the BGOU Strategic Initiative, but are in addition to the scope defined in the BGOU RI/FS Work Plan. The characterization of the potential burial areas of SWMU 13 will be addressed with a Sampling and Analysis Plan addendum to the BGOU RI/FS Work Plan and follow-on site investigation that will be documented in the BGOU FS. The results will be discussed with the FFA parties and, if further action is necessary, a path forward will be determined.



LEGEND

- PGDP
- DOE Reservation
- TVA
- Wildlife Management Area
- Municipality
- Surface Water



SOURCES:
 PRS 2009; TVA 2006;
 KY GEONET 2008
 U.S. Census Bureau 2009

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Figure 1.1. PGDP Site Location

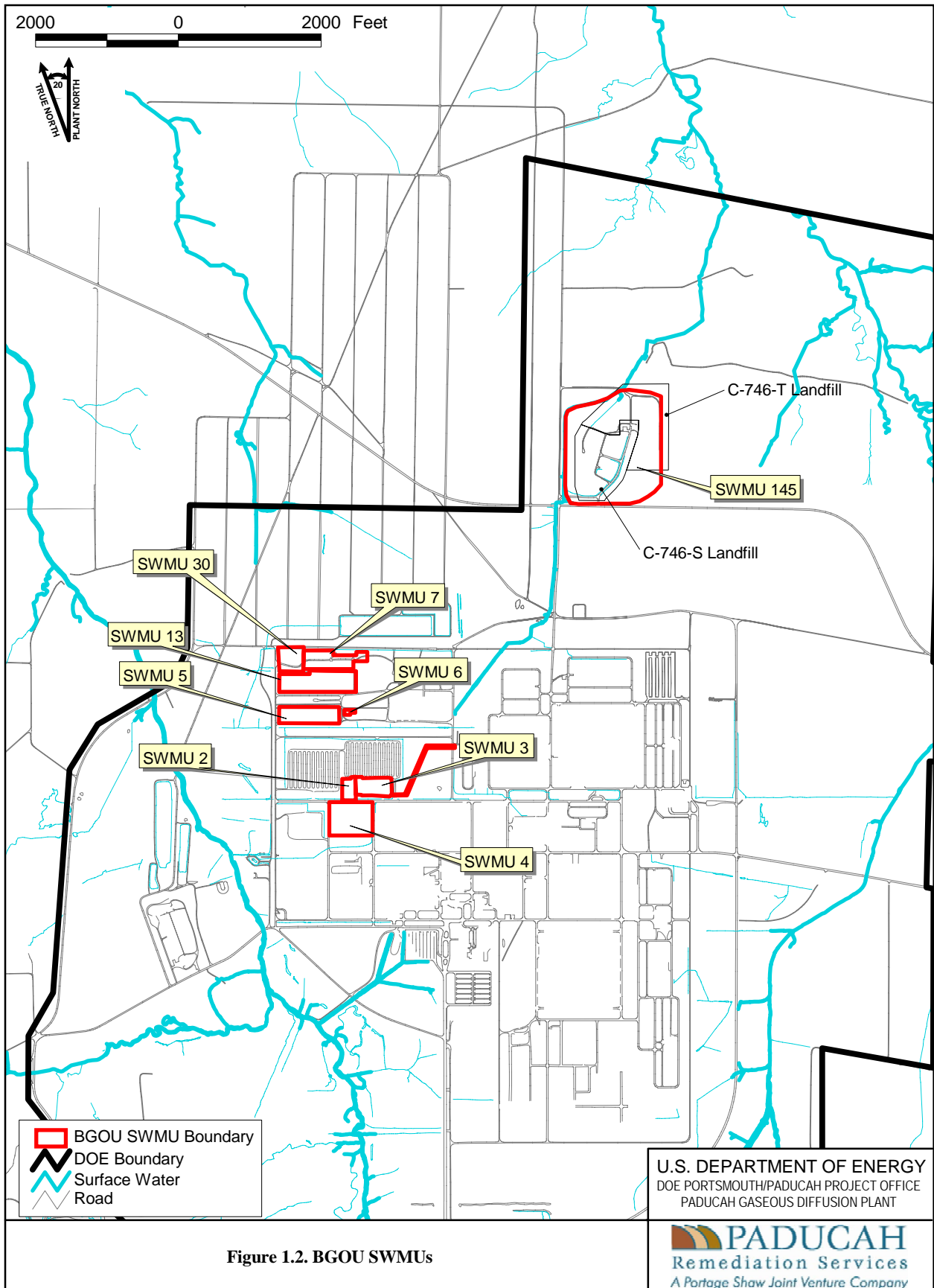


Figure 1.2. BGOU SWMUs

Figure No. \BGOU\d2-ri sect1.apr
DATE 09-29-09

1.1 PURPOSE OF REPORT

The BGOU RI followed the investigation outlined in the BGOU RI/FS Work Plan (DOE 2006a). The work plan utilized the data quality objective (DQO) process as a planning tool to assist in the identification of environmental problems and to define the data collection process needed to support decisions regarding the problem statement developed through the DQO process and documented in the BGOU RI/FS Work Plan:

Hazardous substances that have been contained in, or passed through, the BGOU SWMUs may have been released to surface water or into surrounding soil or are contained in burial cell materials. These substances may have infiltrated into groundwater below the unit and been transported through subsurface pathways. The nature and extent of contamination have been adequately defined for some SWMUs, and risk assessments have been prepared. For others, the nature and extent of contamination have not been adequately defined to assess whether potential contaminants pose unacceptable risks to human health and the environment at the SWMUs and at downgradient exposure points. Data gaps should be identified, and “closed,” so that a comprehensive RI/FS report can be prepared for the eight SWMUs within the BGOU.

The objectives of the RI included characterization of nature, extent, and magnitude of source zones and secondary sources (such as contaminated soil) at the burial ground SWMUs listed on page 1-1. Additionally, the purpose of the RI is to determine surface and subsurface transport mechanisms and to support an evaluation of remedial technologies. These goals (DOE 2006a) are listed specifically in Table 1.1.

This report documents the results of the RI and Baseline Human Health Risk Assessment (BHHRA). Recommended remedial action objectives will be presented in the forthcoming FS.

1.2 PROJECT SCOPE

The BGOU RI primarily consisted of a field investigation of the following burial grounds: C-749 (SWMU 2); C-404 (SWMU 3); C-747 and C-748-B (SWMU 4); C-746-F (SWMU 5); C-747-B (SWMU 6); C-747-A (SWMUs 7 and 30); and the residential/inert borrow area and old NSDD disposal trench (SWMU 145).

1.2.1 Scope

The BGOU RI focused on the burial grounds listed previously and the immediately affected areas adjacent to and beneath the burial cells down to the Regional Gravel Aquifer (RGA) interface to determine if the cells are contributing to groundwater contamination. As stated in the Site Management Plan (SMP), a primary objective for this project is to contribute to the protection of off-site residents by addressing sources of groundwater contamination (DOE 2009).

The Groundwater Operable Unit (OU) Strategic Initiative will address dissolved-phase groundwater contamination in the RGA beneath the BGOU SWMUs; however, secondary sources of groundwater contamination that are derived from the BGOU burial grounds, such as the potential dense nonaqueous-phase liquid (DNAPL) source zone originating from a burial ground, remain within the scope of the BGOU for assessment and remedial action, if required. The scope of the BGOU includes potential contaminant migration pathways from the burial grounds to surface water, but does not include the

ditches bounding the burial grounds. These ditches are components of the larger integrator unit of the Surface Water OU Strategic Initiative.

Table 1.1. Goals Identified for the BGOU RI

GOAL 1: CHARACTERIZE NATURE OF SOURCE ZONE
1-1: What are the suspected contaminants?
1-2: What are the plant processes that could have contributed to the contamination? When and over what duration did releases occur?
1-3: What are the concentrations and activities at the source?
1-4: What is the area and volume of the source zone?
1-5: What are the chemical and physical properties of associated material at the source areas?

GOAL 2: DEFINE EXTENT OF SOURCE ZONE AND CONTAMINATION IN SOIL AND OTHER SECONDARY SOURCES AT ALL UNITS
2-1: What are the past, current, and potential future migratory paths?
2-2: What are the past, current, and potential future release mechanisms?
2-3: What are the contaminant concentrations or activity gradients?
2-4: What is the vertical and lateral extent of contamination?
2-5: What is the relationship of the UCRS gradient to the source, to surface water bodies, and to the RGA?

GOAL 3: DETERMINE SURFACE AND SUBSURFACE TRANSPORT MECHANISMS AND PATHWAYS
3-1: What are the contaminant migration trends?
3-2: To what area is the dissolved-phase plume migrating?
3-3: What are the effects of underground utilities and plant operations on migration pathways including ditches?
3-4: What is the role of the UCRS in contaminant transport?
3-5: What are the physical and chemical properties of the formations and subsurface matrices?

GOAL 4: SUPPORT EVALUATION OF REMEDIAL ALTERNATIVES
4-1: What are the possible remedial technologies applicable for this unit?
4-2: What are the physical and chemical properties of media to be remediated?
4-3: Are cultural impediments present?
4-4: What is the extent of contamination (geologic limitations presented by the source zone or secondary contamination source)?
4-5: What would be the impact of action on and by other sources?
4-6: What would the impact of an action at the source be on the integrator units?
4-7: What are stakeholders' perceptions of contamination at or migrating from source zone or secondary contamination sources?

UCRS = Upper Continental Recharge System

The DQO process was used to focus the sampling strategy on SWMU-specific media, contamination, and migration pathways, and identifying data needs. Data collected during the BGOU RI, together with historical data presented in the BGOU RI/FS Work Plan (DOE 2006a), meet the project DQOs and are used within this RI Report.

The following list summarizes the activities that were conducted as part of the RI (not all activities were performed at each SWMU):

- Collection of geophysical data;
- Collection of surface soil, subsurface soil, and groundwater samples;

- Laboratory analysis of the samples;
- Evaluation of nature and extent of contamination related to each source unit;
- Numeric modeling of contaminant fate and transport and estimation of future exposure point concentrations at the DOE property boundary; and
- Determination of ecological and human health risks associated with each site.

Consistent with the BGOU RI/FS Work Plan, the nature and extent of surface soils (0–1 ft bgs) and sediments within the BGOU SWMUs are not included in the BGOU RI/FS.¹ Surface soils within two of the BGOU SWMUs (SWMUs 3² and 7), however, were sampled during this RI, as specified in the BGOU RI/FS Work Plan (DOE 2006a), to provide additional information. Results from these surface soil samples are presented with the BGOU analytical data.

Further, the BGOU RI sought to identify additional disposal areas that might exist beneath the scrap yards, consistent with the scope of the BGOU, as described in the SMP (DOE 2009). One such area, within the existing SWMU 13, was tentatively identified during an employee interview. A subsequent geophysical survey of SWMU 13 identified additional buried metal anomalies. The results of the geophysics survey are presented within this report in Section 2; however, this geophysical information and data available for this area will be evaluated under a separate site investigation.

To deal with uncertainties identified in the BGOU, the observational approach was used in the design of the sampling strategy for the BGOU RI/FS. The key concepts are as follows:

- The RI strategy is based on a specified “most probable site condition,” which, for the BGOU RI/FS, assumes that contamination is potentially adversely impacting human health and welfare or an impact on the environment has occurred.
- Reasonable deviations from the most probable site condition are identified. The reasonable deviation for the BGOU RI/FS is that no contamination is adversely impacting human health and welfare or the environment. Site conditions should not differ significantly from the postulated conditions shown in the conceptual models.

1.2.2 Rationale for Field Sampling

Sampling activities focused on the soils and groundwater beneath the burial pits to a depth of 60 ft bgs in order to detect any releases that may be related to the SWMUs (Section 2 includes the depths of samples). Dissolved contamination in the RGA, generally at a depth of 60 ft or greater, that is unrelated to the burial grounds is the scope of the Groundwater OU. Table 1.2 summarizes the depth of the RGA at each SWMU. Angled soil borings were utilized to collect samples for this objective. Surface² and subsurface

¹ A discussion of nature and extent of surface soils and sediment is discussed in previous investigations (DOE 1994; 1997a; 1998a; 2000a). The BGOU RI risk assessment includes results from previous RI risk assessments (for risks from direct exposure to environmental media) and groundwater modeling results from this RI (for various points of exposure). These results are used to evaluate the exposure scenarios specified in the Work Plan (residential, industrial, and recreational). The modeled points of exposure include the SWMU boundary, DOE plant boundary, DOE property boundary, Little Bayou Creek seeps, and the Ohio River.

² The SWMU 3 surface soils characterized by the BGOU RI are associated with a former pipeline located to the east of C-404.

soils adjacent to but not beneath the burial pits were not part of this investigation and will be evaluated through the Soils OU. Likewise, the RGA was not part of this investigation and will be evaluated through the Groundwater OU (with the exception of borings advanced to the RGA to evaluate upgradient and downgradient contaminant levels at SWMU 7). Assessment of the potential secondary source (TCE DNAPL) in the Upper Continental Recharge System (UCRS) at SWMUs 7 and 30 was based on both historical and newly generated data, while the assessment of the potential secondary source in both the UCRS and RGA at SWMU 4 is based primarily on historical data. Due to low groundwater yield, not all UCRS groundwater samples could be collected. Of the 32 attempts, only 18 were successful. Borings adjacent to the NSDD were advanced to a depth of 15 ft bgs to evaluate impacts from the pipeline that once discharged leachate from SWMU 3 into the NSDD. Figure 1.3 illustrates the conceptual design of the soil borings to collect these samples.

Table 1.2. Depth of the RGA at Each of the BGOU SWMUs

SWMU	Representative Lithologic Logs	Range of Depths (ft bgs) of RGA (top of HU4 or HU5)
2	SWMU 2-03, SWMU 2-05, SWMU 2-09, SWMU 2-13, SWMU 2-15, and SWMU 2-17	46.6–60
3	MW67, MW86, MW89, MW92, and MW95	55–61.5
4	004-028, 004-029, MW414, MW414A, MW415, MW416, and MW417	44–63
5	005-013, 005-026, H002, MW53, and MW54	52–61
6	006-024 and 006-025	62–63
7	GW-01	55
30	NW65 and MW66	51–52
145	145-017, 145-020, 145-021, 145-023, MW179, MW181, MW220, MW221, MW222, MW223, MW224, MW225, MW263, MW265, MW277, MW370, MW385, MW388, MW392, and MW395	37–69.5

HU = hydrogeologic unit
 MW = monitoring well
 SWMU = solid waste management unit

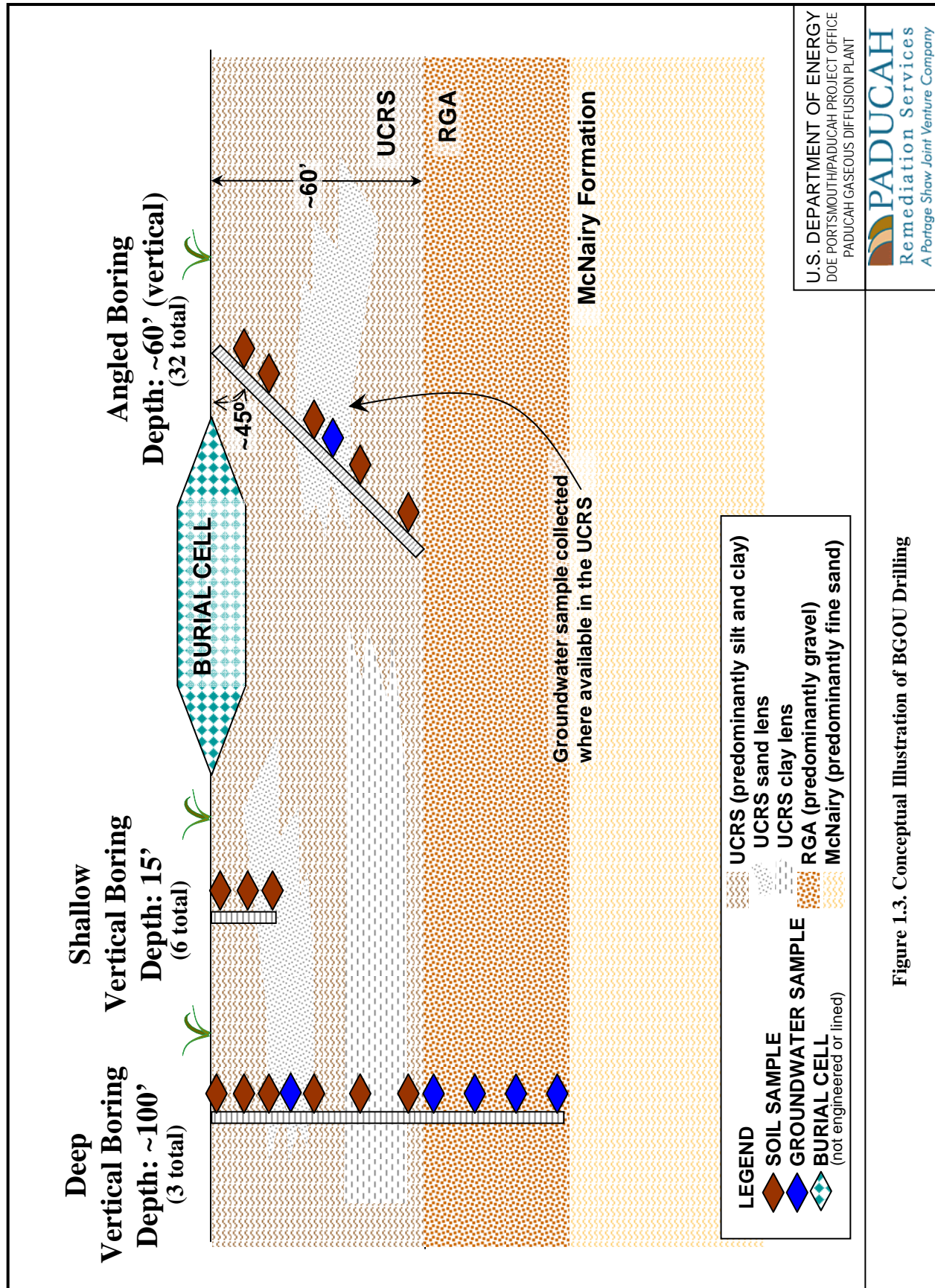


Figure 1.3. Conceptual Illustration of BGOU Drilling

1.3 SITE BACKGROUND

The burial grounds addressed by this RI are discussed in detail in the following sections. Table 1.3 summarizes this information. Much of the information regarding areas, dates of operations, and cap material is summarized from former operating contractor reports and the closure report for SWMU 3 (Union Carbide 1973; Union Carbide 1975; Union Carbide 1978; Union Carbide 1982; DOE 1987). There is some uncertainty with this information, such as burial depths, since the reports are of a general nature.

Table 1.3. Summary of BGOU SWMUs

Sub Unit	Dates of Operation	Area of Waste	Cap ^a	Known or Expected Contents (Special Hazards)
SWMU 2 C-749 Uranium Burial Ground				
	1951–1977	32,000 ft ² (7-17 ft deep)	6-inch clay 18-inch soil	Uranium (pyrophoric uranium), waste oil [polychlorinated biphenyl (PCB?)], TCE
SWMU 3 C-404 Low-Level Radioactive Waste Burial Ground				
	1952–1986	53,000 ft ² (8-12 ft deep)	RCRA multilayered cap	Uranium precipitated from aqueous solutions, uranium tetrafluoride, uranium metal, uranium oxides, degreasing sludge, and radioactively contaminated trash
SWMU 4 C-747 Burial Yard and C-748-B Burial Area				
C-747	1951 to 1958 potentially	8,300 ft ² (16 ft deep) 278,400 ft ²	2 to 3 ft soil 6-inch clay 2 to 3 ft soil	Debris (radiologically contaminated) from uranium hexafluoride feed plant
C-748-B	1973–1987	(16 ft deep)	6-inch clay	Proposed chemical landfill ^b
SWMU 5 C-746-F Burial Yard				
	1965–1987	197,400 ft ² 6-15 ft deep)	2 to 3 ft soil	Radionuclide-contaminated scrap metal, slag from nickel and aluminum smelters
SWMU 6 C-747-B Burial Ground				
Area H	1971	180 ft ² (6 ft deep)	3 ft soil	Magnesium scrap
Area I	1966	280 ft ² (8 ft deep)	5 ft soil	Exhaust fans (contaminated with perchloric acid)
Area J	Early 1960s	4,000 ft ² (6 ft deep)	3 ft soil	Contaminated aluminum
Area K	1968–1969	180 ft ² (6 ft deep)	3 ft soil	Magnesium scrap
Area L	1969	600 ft ² (6 ft deep)	3 ft soil	Modine trap
SWMU 7 C-747-A Burial Ground				
Pit B	?	10,320 ft ² (6–7 ft deep)	3 ft soil	Noncombustible trash, contaminated material and equipment
Pit C	?	10,320 ft ² (6–7 ft deep)	3 ft soil	Noncombustible trash, contaminated material and equipment, Uranium-contaminated concrete pieces of reactor tray bases
Pit D	?	1,485 ft ² (6–7 ft deep)	3 ft soil	from fluorination process of uranium tetrafluoride to uranium hexafluoride
Pit E	?	2,145 ft ² (6–7 ft deep)	3 ft soil	Uranium-contaminated concrete pieces of reactor tray bases
Pits F1–F5	?	1,600 ft ² (6–7 ft deep)	3 ft soil	Uranium-contaminated scrap metal, equipment, empty uranium/magnesium powder drums
Pit G	?	3,294 ft ² (6–7 ft deep)	3 ft soil	Noncombustible trash, contaminated material and equipment
SWMU 30 C-747-A Burn Area				
Pit A	1951–1970	128,000 ft ² (12 ft deep)	4 ft soil	Ash and debris from combustible trash, possibly uranium-contaminated

Table 1.3. Summary of BGOU SWMUs (Continued)

SWMU 145 Area P		
	44 acres (individual landfills estimated 6-10 1952–1980 ft deep)	Construction debris

^a The source material used for capping is unknown (with the exception of the SWMU 3 Resource Conservation and Recovery Act (RCRA) cap that came from the Old Hickory Clay Company).

^b The “Proposed Chemical Landfill” is the only name used to describe this burial area (Union Carbide 1973).

Several RI documents have been produced containing data pertinent to the various SWMUs within the BGOU. In most cases, the previously prepared documents grouped several SWMUs together and did not study one particular SWMU. These documents and the various monitoring wells (MWs) installed throughout PGDP provide considerable usable historical data in addition to that generated during the BGOU RI. Historical data to be used for the BGOU is documented in the BGOU RI/FS Work Plan (DOE 2006a). Additionally, the historical data set was updated to include measurements collected from monitoring wells between the periods of work plan development and RI development.

Table 1.4 identifies the previously completed reports and/or investigations primarily used.

Table 1.4. Summary of Previous Investigations of BGOU

Dates	Title	SWMU 2	SWMU 3	SWMU 4	SWMU 5	SWMU 6	SWMU 7	SWMU 30	SWMU 145
1989	Post Closure Permit Application C-404 Low-Level Radioactive Waste Burial Ground		✓						
1990– 1992	Phase II Site Investigation	✓	✓	✓	✓	✓	✓	✓	
1996	Closure Plan C-404 Low-Level Radioactive Waste Burial Ground		✓						
1996– 1997	WAG 22 SWMUs 2 and 3 Remedial Investigation and Addendum (including SWMU 2 Data Summary Report)	✓	✓						
1996– 1998	WAG 22 SWMUs 7 and 30 RI/FS						✓	✓	
1998– 2001	WAG 3 RI/FS			✓	✓	✓			
1999– 2001	Data Gaps Investigation			✓	✓		✓	✓	✓
2000– 2001	Old NSDD Sampling								✓
2002– 2003	Scrap Yards Site Characterization				✓	✓	✓	✓	
2003– 2004	C-746-S and -T Landfill Site Investigation								✓
2004	Southwest Plume Site Investigation			✓					
2006	Burial Grounds RI/FS Work Plan	✓	✓	✓	✓	✓	✓	✓	✓

In addition to the reports of previous RIs, the following documents provide significant information on the content and volume of the burial grounds:

- *The Discard of Scrap Materials by Burial at the Paducah Plant* (Union Carbide 1973) and

- *The Disposal of Solid Waste at the Paducah Gaseous Diffusion Plant* (Union Carbide 1978).

Several aerial photographs were used in researching the history of the BGOU SWMUs. These photographs are listed in Table 1.5.

Table 1.5. BGOU Aerial Photographs

Aerial Photograph (Year)	SWMUs Shown	Comments^a
1943	2, 3, 4, 5, 6, 7, 30, and 145	Prior to PGDP construction. Ground scarring shown at SWMUs 7 and 30 due to KOW operations.
1952	2, 3, 4, 5, 6, 7, 30, and 145	PGDP construction is evident.
1959	145	Plant area not shown. Ground scarring shown at SWMU 145.
1964	145	Plant area not shown. Ground scarring shown at SWMU 145.
1971	2, 3, 4, 5, 6, 7, 30, and 145	Ground scarring shown at most SWMUs.
1974/1975	2, 3, 4, 5, 6, 7, 30, and 145	Ground scarring shown at most SWMUs.
1981	2, 3, 4, 5, 6, 7, 30, and 145	Ground scarring shown at most SWMUs.
1983	2, 3, 4, 5, 6, 7, 30, and 145	Some activity possible at SWMU 5. Ground scarring shown at SWMU 145.
1988	2, 3, 4, 5, 6, 7, 30, and 145	Some activity possible at SWMUs 5 and 30. Ground scarring shown at SWMU 145.
1993	2, 3, 4, 5, 6, 7, 30, and 145	Some activity possible at SWMUs 5, 7, and 30.
1994	2, 3, 4, 5, 6, 7, 30, and 145	Some activity possible at SWMU 7.
1998	2, 3, 4, 5, 6, 7, 30, and 145	No ground scarring evident.

^a Plant records indicate that no additional disposal actions occurred at any of the burial areas beyond their identified periods of operation. The activity that caused the post-operation scarring noted at all of the SWMUs is unknown and is an uncertainty.

1.3.1 C-749 Uranium Burial Ground (SWMU 2)

1.3.1.1 Site description

The C-749 Uranium Burial Ground (SWMU 2) is located within the west-central portion of the plant. SWMU 2 encompasses an area of approximately 32,000 ft², with approximate dimensions of 160 ft by 200 ft. Records indicate that when the burial ground was in use, pits were excavated to an estimated depth of 7 to 17 ft. After the burial ground no longer was in use, the area was covered with a 6-inch thick clay cap and an 18-inch thick soil layer covered with vegetation (DOE 1995a). Figure 1.4 illustrates the burial ground, showing the historical grid layout as documented (Union Carbide 1975).

1.3.1.2 Site history

SWMU 2 was used from 1951 to 1977 for the disposal of uranium and uranium-contaminated wastes (Figure 1.5). Disposal records for SWMU 2 indicate that 270 tons of uranium, 59,000 gal of oils, and 450 gal of trichloroethene (TCE) were disposed in the unit (DOE 1999a). Disposal records also indicate that drummed wastes buried in the unit consist primarily of uranium metal from machine shop turnings, shavings, and sawdust (260 tons of the uranium is reported as being sawdust, shavings, or turnings). Table 1.6 summarizes the inventory of the wastes in SWMU 2 (DOE 1999a). Other wastes at the unit consist of drummed uranyl fluoride and TCE. Because small pieces of uranium metal may be pyrophoric (spontaneously burn in air), operating practices of that time required placing the material in drums and submerging the material in petroleum-based oil and synthetic oil to avoid contact with air.

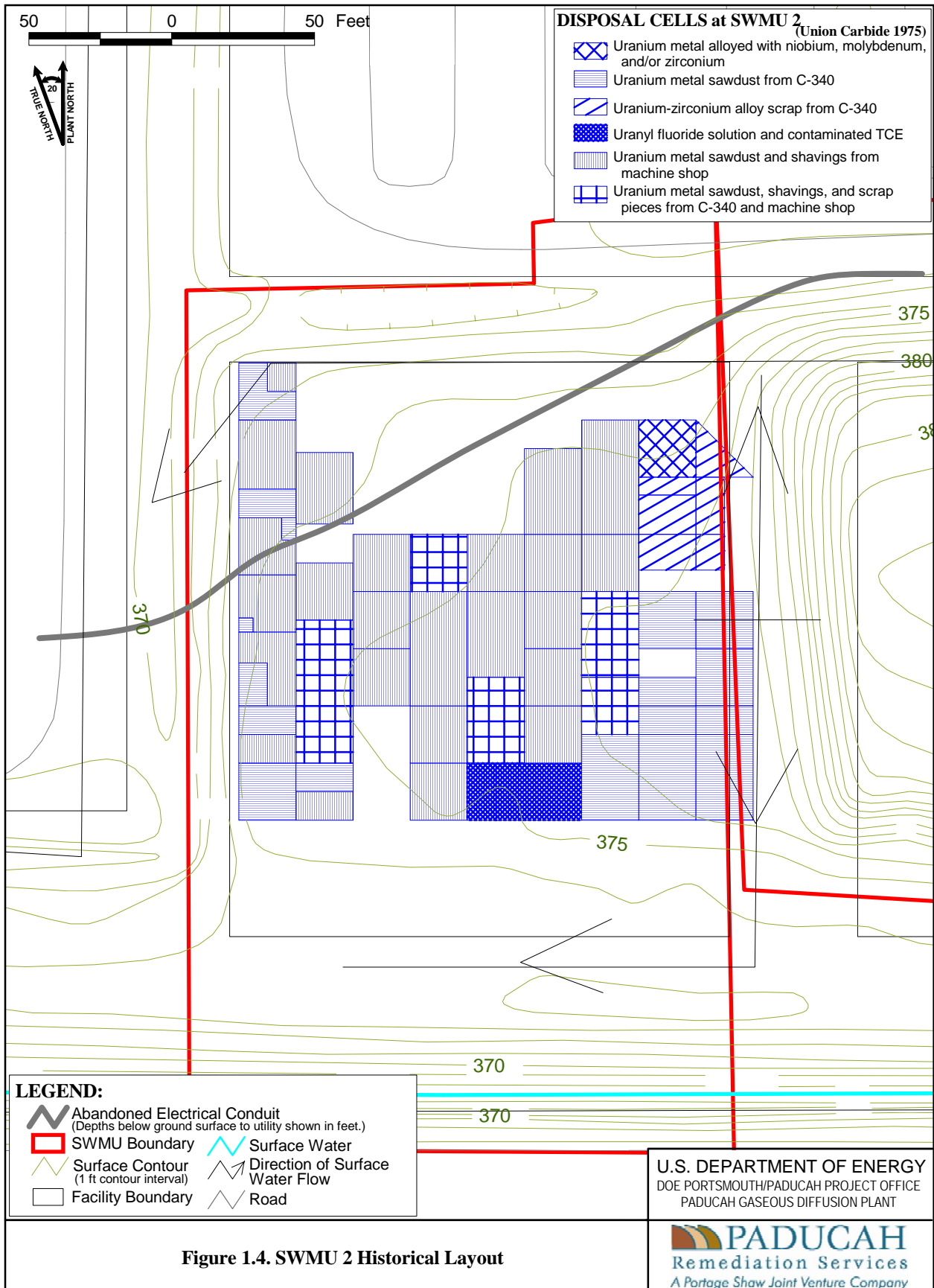


Figure 1.4. SWMU 2 Historical Layout

Figure No. 1BGOUd2-ri sect1.apr
DATE 12-16-08



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Table 1.6. Inventory of Uranium-Bearing Scrap in Solid Waste Management Unit 2

Pit No.	Date Buried	Type of Material	Number of Containers			Miscellaneous	Net Weight or Volume	Uranium (g)	Wt. % ²³⁵ U ^a	Remarks
			20-gal Drums	30-gal Drums	55-gal Drums					
1	12-16-57	U-Metal Alloy	—	4	—	Twelve 8-gal pails	4,686 lb	2,125,386 ^c	0.1400 ^d	Nb ₇ Mo-Zr alloy received from NLO
2 ^b	05-06-59	Uranium Sawdust	18	57	19		94,401b ^e	38,537,414 ^f	0.1519	From routine C-340 operations
3 ^b	06-16-59	Uranium Sawdust	—	1	—		800 lb ^e	326,585 ^f	0.1617	From routine C-340 operations
4 ^b	08-14-59	U-Metal Alloy	—	—	—	One 1-gal pail	10 lb	4,445 ^g	0.2200 ^h	Zirconium-bearing scrap from C-340
5 ^b	08-26-59	Uranium Sawdust	—	1	—		800 lb ^e	326,585 ^f	0.1675	From routine C-340 operations
5 ^b	08-31-59	Uranium Sawdust	—	1	—		800 lb ^e	326,585 ^f	0.1675	From routine C-340 operations
6 ^b	09-24-59	Slag	—	—	—	Buried loose	2,100 lb	28,576	0.2200 ^h	Zirconium-bearing scrap from C-340
7 ^b	09-30-59	Saw Chips	—	—	—	Buried loose	25 lb	10,206	0.2200 ^h	Zirconium-bearing scrap from C-340
8 ^b	10-01-59	Uranyl Fluoride Solution	—	35	—	—	6,048 liters	68,377	0.408	Contaminated waste solution
9 ^b	10-01-59	Contaminated TCE	—	15	—	—	5,843 lb	54,272	0.348	Contaminated waste solution
10 ^b	10-19-59	U-Metal Alloy	—	—	—	Buried loose	300 lb	122,469	0.2200 ^h	Zirconium-bearing scrap from C-340
10 ^b	10-19-59	Crucible Burnout	—	—	—	Buried loose	100 lb	36,287	0.2200 ^h	Zirconium-bearing scrap from C-340
10 ^b	10-19-59	Saw Chips	—	—	—	Buried loose	50 lb	20,412	0.2200 ^h	Zirconium-bearing scrap from C-340
11 ^b	11-16-59	Uranium Sawdust	—	5	—		4,000 lb ^e	1,632,924 ^f	0.1733	From routine C-340 operations
11 ^b	11-16-59	Reject "U" Scrap	—	—	—	Buried loose	3,550 lb	1,449,220	0.1733	From routine C-340 operations
12 ^b	05-27-60	Uranium Sawdust	4	—	—	—	2,400 lb ^e	979,754 ^f	0.2483	From routine C-340 operations
13	12-16-60	Uranium Sawdust	12	—	—		9,600 lb ^e	5,919,018 ^f	0.17	From routine C-340 operations
13	12-16-60	Saw Chips	—	—	—	One small box	200 lb	81,646	0.17	From routine C-340 operations
14 ^b	04-25-62	Uranium Sawdust	5	2	—		4,600 lb ^e	1,877,863 ^f	0.18	From routine C-340 operations
15 ^b	07-24-62	Sandblast Grit	—	—	—	One 1-gal pail	5 lb	Insignificant	—	Buried for safety reasons only
15 ^b	07-24-62	Zr Alloy Scrap	1	—	1		700 lb	Insignificant	—	Buried for safety reasons only
16 ^b	05-09-63	Uranium Sawdust	5	1	—		3,800 lb ^e	1,378,914 ^f	0.1800 ⁱ	From routine C-340 operations
17	11-13-63	Uranium Sawdust	13	—	—		7,800 lb ^e	3,184,202 ^f	0.2200 ⁱ	From routine C-340 operations
18 ^b	02-03-64	Uranium Sawdust and Shavings	18	39	7		4,683 lb	2,124,297 ^f	0.2335	From machining operations
19 ^b	02-03-64	Uranium Sawdust	15	16	—		25,400 lb ^e	10,369,067 ^f	0.2200 ⁱ	From routine C-340 operations
20 ^b	04-15-64	Uranium Sawdust	—	24	2		22,200 lb ^e	9,062,728 ^f	0.2200 ⁱ	From routine C-340 operations
21 ^b	04-15-64	Uranium Sawdust and Shavings	—	88	—		1,342 lb	608,717 ^f	0.2271	From machining operations
22 ^b	07-02-64	Uranium Sawdust	—	6	—		6,000 lb ^e	2,449,386 ^f	0.2200 ^j	From routine C-340 operations
23 ^b	07-02-64	Uranium Sawdust and Shavings	—	30	5		5,436 lb	2,219,416 ^f	0.3102	From machining operations

Table 1.6. Inventory of Uranium-Bearing Scrap in Solid Waste Management Unit 2 (Continued)

Pit No.	Date Buried	Type of Material	Number of Containers		Miscellaneous	Net Weight or Volume	Uranium (g)	Wt. % ²³⁵ U ⁴	Remarks
			20-gal Drums	30-gal Drums					
24 ^b	10-21-64	Uranium Sawdust	—	3	—	3,000 lb ^e	1,224,693 ^f	0.2200 ⁱ	From routine C-340 operations
25 ^b	10-21-64	Uranium Sawdust and Shavings	—	25	—	4,774 lb	2,165,439 ^f	0.2196	From machining operations
26 ^b	01-28-65	Uranium Sawdust	—	6	—	6,000 lb ^e	2,449,386 ^f	0.220	From routine C-340 operations
27 ^b	01-28-65	Uranium Sawdust and Shavings	—	25	8	1,422 lb	645,005 ^f	0.2288	From machining operations
28 ^b	04-30-65	Uranium Sawdust	—	23	—	23,000 lb ^e	9,389,313 ^f	0.2200 ⁱ	From routine C-340 operations
29 ^b	04-30-65	Uranium Sawdust and Shavings	—	5	—	1,706 lb	773,825 ^f	0.2308	From machining operations
30 ^b	10-21-65	Uranium Sawdust	—	36	—	36,000 lb	13,063,392 ^f	0.2200 ⁱ	From routine C-340 operations
31 ^b	10-21-65	Uranium Sawdust and Shavings	—	15	5	2,917 lb	1,322,940 ^f	0.22	From machining operations
32 ^b	07-20-66	Uranium Sawdust, Shavings, and Scrap Pieces	—	78	13	100,251 lb	45,472,851 ^f	0.1776	From routine C-340 and machining operations
33 ^b	11-28-66	Uranium, Sawdust, and Shavings	—	41	2	21,364 lb	9,690,703 ^f	0.2205	From machining operations
34	03-28-67	Uranium Turnings and Shavings	64	5	—	5,355 lb	2,428,931 ^f	0.1845	From machining operations
35	06-14-67	Uranium Turnings and Shavings	103	—	—	4,699 lb	2,131,542 ^f	0.2063	From machining operations
36 ^b	07-26-67	Uranium Turnings and Shavings	145	—	—	10,455 lb	4,742,294 ^f	0.2001	From machining operations
36 ^b	07-26-67	U ₃ O ₈	—	—	20	20,000 lb	7,257,440 ^f	0.1870 ⁱ	Routine C-340 operations
37 ^b	08-25-67	Uranium Turnings and Shavings	35	—	—	4,316 lb	1,957,543 ^f	0.2102	From machining operations
37 ^b	08-25-67	U ₃ O ₈	—	6	10	13,600 lb	4,935,059 ^f	0.1622 ⁱ	Routine C-340 operations
38	11-21-67	Uranium Turnings and Shavings	45	—	—	2,309 lb	1,047,394 ^f	0.2115	From machining operations
39	03-26-68	Uranium Turnings and Shavings	71	—	—	5,034 lb	2,283,564 ^f	0.1775	From machining operations
40 ^b	05-01-68	Uranium Turnings, Sawdust, and U ₃ O ₈	55	—	9	4,357 lb	1,976,130 ^f	0.154	From routine C-340 and machining operations
41	07-08-68	Uranium Turnings and Shavings	31	—	—	697 lb	307,987 ^f	0.2201	From machining operations
42	09-10-68	Uranium Turnings and Shavings	55	—	—	3,155 lb	1,412,924 ^f	0.171	From machining operations
43 ^b	11-15-68	Uranium Turnings and Shavings	83	—	—	16,297 lb	7,384,065 ^f	0.1654	From machining operations
44	03-05-69	Uranium Turnings and Shavings	82	—	—	726 lb	329,499 ^f	0.21	From machining operations
45 ^b	07-17-69	Uranium Turnings and Shavings	118	8	—	2,784 lb	1,262,602 ^f	0.207	From machining operations
45 ^b	07-17-69	U ₃ O ₈	—	—	2	2,000 lb	725,744 ^f	0.2000 ⁱ	From routine C-340 operations
46 ^b	12-22-69	Uranium Turnings and Shavings	20	—	—	288 lb	130,600 ^f	0.1983	From machining operations
46 ^b	12-22-69	U ₃ O ₈	—	—	1	1,000 lb	362,872 ^f	0.2000 ⁱ	From routine C-340 operations
47	07-21-70	Reject Uranium Metal Scrap	—	—	15	12,000 lb	5,443,080 ^f	0.2000 ⁱ	From routine C-340 operations
47	07-21-70	Uranium Turnings and Shavings	20	—	—	308 lb	139,674 ^f	0.2200	From routine C-340 operations
48	03-05-71	Uranium Turnings and Shavings	93	—	—	9,283 lb	4,210,526 ^f	0.1626	From machining operations
49	08-23-71	Uranium Turnings and Shavings	107	—	—	10,680 lb	4,844,368 ^f	0.1626	From machining operations
50	03-16-72	Uranium Turnings and Shavings	200	—	—	19,575 lb	8,879,024 ^f	0.199	From machining operations
51 ^b	11-08-72	Uranium Turnings and Shavings	143	—	—	10,803 lb	4,900,000 ^f	0.201	From machining operations

Table 1.6. Inventory of Uranium-Bearing Scrap in Solid Waste Management Unit 2 (Continued)

Pit No.	Date Buried	Type of Material	Number of Containers			Miscellaneous	Net Weight or Volume	Uranium (g)	Wt. % ²³⁵ U ^a	Remarks
			20-gal Drums	30-gal Drums	55-gal Drums					
52 ^b	10-19-73	Uranium Turnings and Shavings	111	—	16		7,057 lb	0.198	From machining operations	
53	12-12-74	Uranium Turnings and Shavings	86	—	2		7,151 lb	0.198	From machining operations	

^a Weight percent ²³⁵U of all material shown as originating from routine C-340 operations was derived from the average assay of uranium metal produced during the approximate period the scrap was accumulated.

^b Denotes only partial use of the location for the specified burial.

^c Derived from application of shipper's analyses to Paducah net weight.

^d Received into the plant on the basis of the current metals plant assay at the time of receipt (in accordance with letter from K.C. Brooks to R.G. Jordan on September 9, 1957).

^e Based on an average net weight of sawdust plus oil obtained from a limited number of weighted drums.

^f Calculated on basis of the net weight (sawdust plus oil) and an estimated average of 80% uranium.

^g Calculated on a basis of 98% uranium.

^h Estimated.

ⁱ Derived from the average assay of uranium metal processed during the period the sawdust was accumulated.

^j Derived from a net balance of receipts, shipments, and inventories.

^k Uranium sawdust from C-340 buried on top of material from shop in Location B-1.

^l Calculated on basis of the net weight and an estimated average analysis of 80% uranium.

Most of the waste in the unit is believed to consist of pyrophoric uranium metal in the form of machine shop turnings, shavings, and sawdust. Pyrophoric uranium metal usually was placed in 20-, 30-, or 55-gal drums. Occasionally, underground fires were reported as a result of oxidation of pyrophoric uranium metal, but no documentation of these fires is available. No subsidence has been observed as a result of volume reductions due to the fires. It is possible that the oils used in the pyrophoric uranium drums may have included some polychlorinated biphenyl (PCB)-contaminated oils. Other forms of uranium, including oxides of uranium (solid and dissolved in aqueous solutions), uranyl-fluoride solutions, uranium-zirconium alloy, slag, and uranium tetrafluoride, were buried in small quantities (DOE 1996).

The most likely scenario is that the uranium buried at PGDP is in the metallic state or is coated with uranium (IV) oxide. Neither of these forms of uranium is very susceptible to leaching. The kinetics of dissolution of the buried metal and uranium (IV) oxide is controlled by the amount of oxygen and carbon dioxide that leaches through the waste. Site records show that much of the metal was coated with oil, possibly PCB oil. Such oils are resistant to chemical and biological degradation and from leaching by percolating waters. In addition, oils, as they slowly degrade, consume oxygen, which lowers the oxidation-reduction potential. Under such conditions, uranium dissolution is negligible (ORNL 1998).

No documentation of technetium-99 disposal at SWMU 2 exists; however, during the years of feed plant operation from 1953 to 1964 and from 1968 intermittently through 1977, recycled uranium feed material from nuclear reactors was reprocessed through the feed plant, resulting in the introduction of reactor-produced radioactive impurities, such as technetium-99, into the enrichment process. It is possible that a portion of the uranium-contaminated wastes disposed of in burial grounds at PGDP contains technetium-99 from reprocessing activities (DOE 1994).

Materials contaminated with TCE also are known to have been disposed of at SWMU 2. In August 1984, the western portion of the area designated as containing uranyl fluoride solution and contaminated TCE on Figure 1.4 was excavated with the intent of removing TCE in the soil or drums due to concern about the integrity of TCE-containing drums (15 30-gal drums) reportedly disposed of in this area. It is reported that during excavation, four 30-gal drums (one of these drums contained a uranium and TCE sludge and the others were of such poor integrity that the contents could not be ascertained) and 35 55-gal drums (30 of these drums contained uranium sludges, not TCE, one drum contained TCE, and the rest were of such poor integrity their contents could not be ascertained) were recovered. The 55-gal drum containing TCE was placed in an overpack for proper disposal. None of the 15 30-gal drums containing TCE was found intact. Additionally, the liquid portion of the uranium solutions found in the other drums was transferred to new drums for proper disposal (Ashburn 1984). The remaining materials (everything except the 55-gal drum containing TCE that was overpacked and the liquid portion of the uranium solutions that was transferred to new drums) were returned to the pit and covered with soil. It is important to note the grid and inventory from records did not match what was found during this excavation, which means there is additional uncertainty with the quality of the disposal records.

An FS (DOE 1995a), proposed plan (DOE 1995b), and record of decision (ROD) (DOE 1995c) for SWMUs 2 and 3 were put in place to implement an interim action at SWMU 2 to mitigate risks posed to groundwater and the potential for direct contact. The ROD also included a remedial design investigation to address some data gaps that could affect the effectiveness of the proposed action (i.e., installation of a cap to limit infiltration of water into the waste). The additional information was used to determine that the waste at SWMU 2 was mostly saturated, indicating that a cap would not effectively reduce infiltration. Subsequently, construction of the cap for SWMU 2 was cancelled. A second FS and proposed plan (DOE 1999a) were issued, but a second ROD never was signed or implemented.

1.3.2 C-404 Low-Level Radioactive Waste Burial Ground (SWMU 3)

1.3.2.1 Site description

The C-404 Low-Level Radioactive Waste Burial Ground (SWMU 3) is 1.2 acres located in the west-central portion of the PGDP secured area. The unit originally was constructed as a rectangular, aboveground surface impoundment measuring 387 ft by 137 ft, with a floor area of approximately 53,000 ft². The floor of the surface impoundment was constructed of well-tamped earth, surrounded by clay dikes to a height of 6 ft. The C-404 impoundment was designed with an overflow weir at its southwest corner. When the impoundment overflowed, the effluent flowed west in a ditch (not the NSDD) and eventually discharged through what is now Kentucky Pollutant Discharge Elimination System (KPDES) Outfall 015. Figure 1.6 shows C-404 along with a schematic of this design. Historic effluent/leachate discharges later were rerouted to the NSDD via what is now an abandoned pipeline and ditch leading from the northeast corner of the landfill. This ditch is now filled mostly with a rail loading facility and building being constructed above the course of the former discharge ditch.

1.3.2.2 Site history

SWMU 3 operated as a surface impoundment from approximately 1952 until early 1957. During this time, all influents to the impoundment originated from C-400. From 1957 through 1976, the C-404 surface impoundment was converted to a solid waste disposal facility for solid uranium-contaminated wastes. The waste consists of uranium precipitated from aqueous solutions, uranium tetrafluoride, uranium metal, uranium oxides, degreasing sludge, and radioactively contaminated trash. Uranium-contaminated magnesium-fluoride slag from the metal reduction plant and rejected UF₄ constituted much of the disposal volume. The net weight of uranium committed to the area from 1957 through 1976 is reported to be approximately 3,200 tons. There are no records documenting the cleanout of sludges and sediments from the pond when it was converted to a landfill. When the C-404 impoundment was converted into a disposal facility, a sump was installed at the weir. Leachate was pumped from the sump through an underground transfer line. The transfer line discharged into a northeast-southwest ditch just east of C-404. From this ditch, the leachate flowed into the NSDD. NSDD historically carried PGDP effluents north to Little Bayou Creek. The date of termination of the leachate discharge through the underground transfer line into the NSDD has not been determined. It is known that, prior to landfill closure in 1986, this underground transfer line to the NSDD was not in operation, and leachate from the C-404 Landfill was being collected in the sump for treatment at the C-400-D Lime Precipitation Unit in the C-400 Facility. At some time following closure of the C-404 Landfill, treatment of leachate from C-404 at C-400 was discontinued, and treatment of the leachate was transferred to the C-752 Remedial Action Waste Holding Facility. Some of the constituents found in the leachate and their ranges have included fluoride (4.8-10.0 mg/L); TCE (1-22 µg/L); PCBs (0.41-1.18µg/L); neptunium-237 (0.42-11.7 pCi/L); technetium-99 (90.6-365 pCi/L); and uranium-238 (2,160-37,900 pCi/L).

The upper tier of waste within C-404 contains approximately 450 drums of waste similar to that collected in the impoundment plus smelter furnace liners and drums of Extraction-Procedure-Toxicity, characteristically hazardous, [Resource Conservation and Recovery Act (RCRA) waste codes D006 (for cadmium), D008 (for lead), and D010 (for selenium)]. The drums of extraction-procedure were produced in C-400 during treatment of wastes including sodium bisulfate solution, hydrochloric acid, chromic acid, nickel stripper solution, miscellaneous acids and alkalies, and aqueous solutions containing metals. Table 1.7 presents a list of the waste constituents in the C-404 facility (DOE 1995a). A partial clay cap was installed on the eastern end of the landfill in 1982 (DOE 1987).

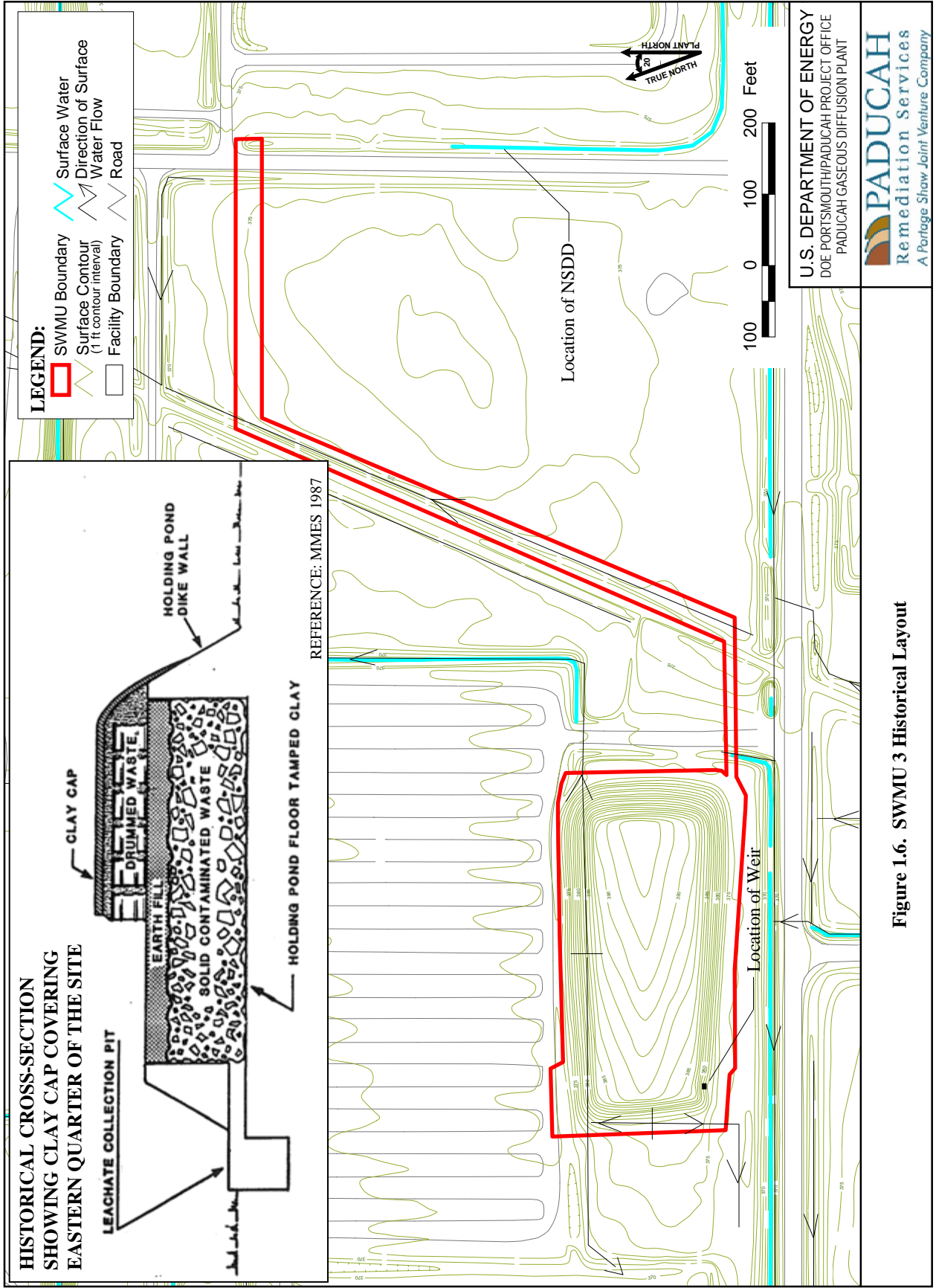


Figure 1.6. SWMU 3 Historical Layout

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Table 1.7. C-404 Waste Constituents

Waste	Constituents	
Gold Dissolver Precipitate ^{1,2}	Cadmium Calcium Copper Lead Magnesium Nickel	Potassium Selenium Sodium Uranium Zinc
Calcium Carbonate Trap Material ¹	Calcium Magnesium	Uranium
Furnace Liners ¹	Aluminum Calcium Iron	Magnesium Uranium
Decontamination Precipitate ¹	Calcium Copper Magnesium Nickel	Sodium Uranium Zinc
Alumina Trap Mix ¹	Aluminum	Uranium
Smelter Dust ¹	Aluminum Nickel	Sodium Uranium
Sodium Fluoride Trap Material ¹	Sodium Fluoride	Uranium
Magnesium Fluoride Slag ³	Magnesium Fluoride	Uranium
Calcium Fluoride Slag ³	Calcium Fluoride	Uranium
Uranium Metal ³	Uranium	
Uranium Oxides ³	Uranium	Fluoride
Concrete ^{3,4}	Uranium	
Uranium Tetrafluoride ³	Uranium	Fluoride
Roofing Materials ^{3,4}	Uranium	
Cleanup Debris ^{3,4}	Uranium	
Radioactive Sources ^{3,4}	Cobalt	Cesium

¹Data based on spectrochemical and EP toxicity data.

²Hazardous waste.

³No analytical data available. Major constituents listed based on knowledge of type of waste.

⁴Possible source of organic constituents, most likely petroleum-based products.

Some of the chemicals of potential concern (COPCs) known to be associated with C-400 include TCE; nitric acid; sulfuric acid; radionuclides (Americium-241, cesium, thorium-230, neptunium-237, plutonium-239, technetium-99, uranium-234, -235, and -238; hexavalent chromium discharge; fluoride/fluorine; lime/sodium hydroxide; heavy metals from cleaning; and PCBs.

Approximately 6,615,000 lb of uranium-contaminated wastes was disposed of at SWMU 3. The total volume is approximately 260,000 ft³. Some uranium contaminated waste also is contaminated with TCE, radionuclides, and metals. In 1986, the disposal of waste at C-404 Landfill was halted, and a portion of the disposed waste was found to be RCRA-hazardous. The landfill was covered with a RCRA multilayered cap and certified closed in 1987. It currently is regulated under RCRA as a land disposal unit and compliance is required by a RCRA postclosure permit issued in 1992. The cap continues to be maintained according to the closure plan, and leachate is pumped from the leachate pit when needed. Based on the amount of leachate requiring removal, the cap appears to be functioning as intended to reduce infiltration. The closure plan required continued groundwater monitoring (DOE 1989). A permit

modification was submitted in May 2008, revising the MW network for the unit (DOE 2008a) to add a new upgradient well, MW420. MW420 is screened in the upper RGA.

An FS (DOE 1995a), proposed plan (DOE 1995b), and ROD (DOE 1995c) for SWMUs 2 and 3 were put in place to implement an interim action at SWMU 2. Since SWMU 3 had undergone RCRA closure in 1987, the ROD did not require additional remedial or corrective measures.

1.3.3 C-747 Contaminated Burial Yard and C-748-B Burial Area (SWMU 4)

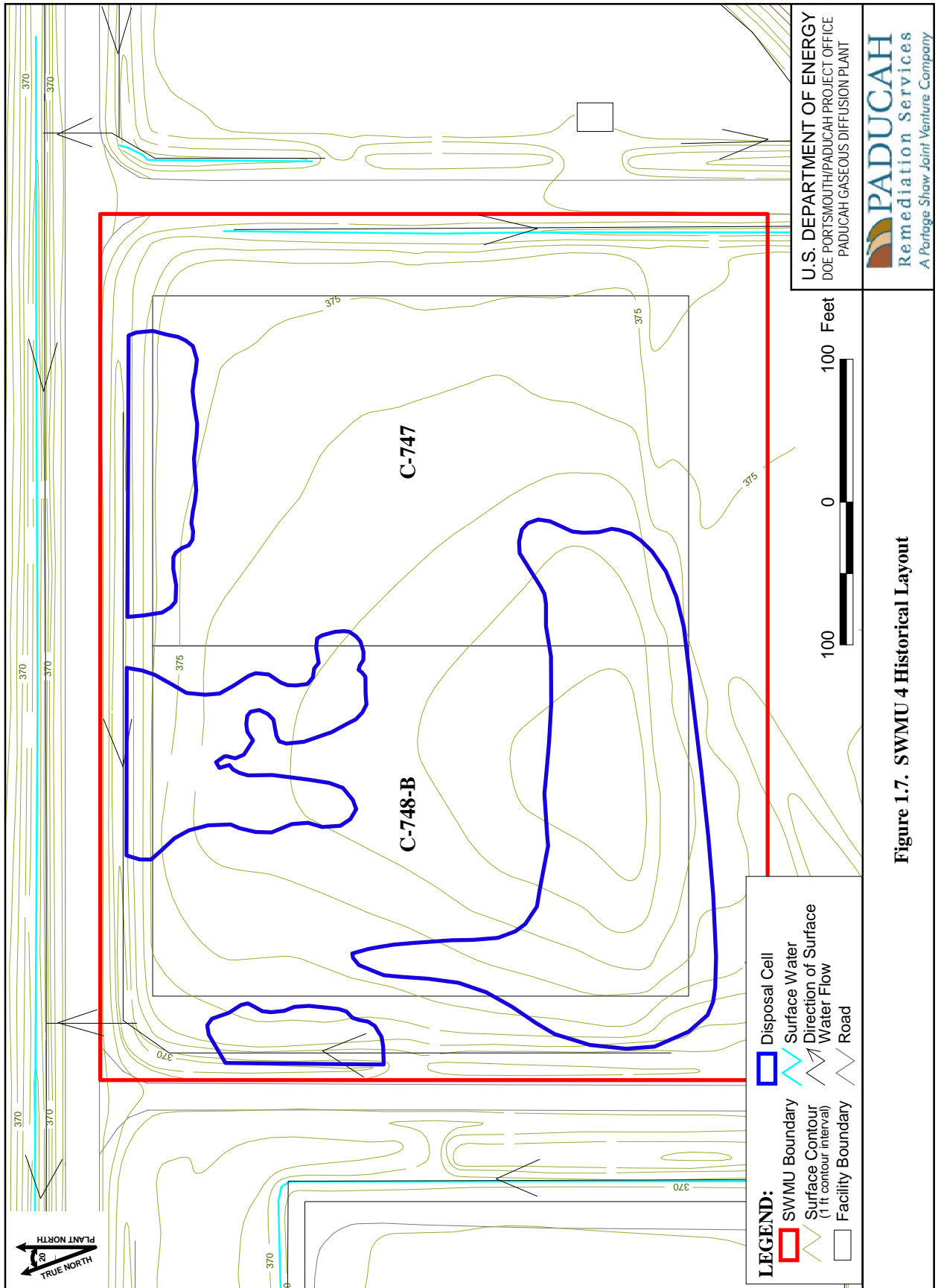
1.3.3.1 Site description

The C-747 Contaminated Burial Yard and the C-748-B Burial Area (SWMU 4) is located in the western section of the PGDP secured area. SWMU 4 (which covers an area of approximately 286,700 ft²) is bounded on the north, east, and west by plant roads and on the south by an active railroad spur (Figure 1.7). This SWMU is an open field that, at one time, was used for the burial and disposal of various waste materials in designated burial cells. A short, narrow, gravel road that enters from the west is nearly completely grass-covered. Except for this rarely used road, the entire site is covered with a variety of field grasses and clovers. The site typically is mowed once a month from April through September. SWMU 4 is bounded on three sides (north, east, and west) by shallow drainage swales that direct surface runoff to the northwest corner of the site. There is an elevation difference of approximately 10 ft between the highest point in the SWMU to the adjacent drainage swales. Both burial yards, C-747 and C-748-B, were covered with 2 to 3 ft of soil material and a 6-inch clay cap was placed over the area in 1982 (DOE 1998b).

1.3.3.2 Site history

The C-747 Burial Yard was in operation from 1951 to 1958 for the disposal of radiologically contaminated and uncontaminated debris originating from the C-410 uranium hexafluoride (UF₆) feed plant. The area originally consisted of two pits covering an area of approximately 8,300 ft² (50 ft by 15 ft and 50 ft by 150 ft) (Union Carbide 1978).

Some of the trash was burned before burial. According to PGDP personnel, a majority of the contaminated metal was buried in the northern part of the yard. When the yard was closed, a smaller pit was reported to have been excavated for the disposal of radiologically contaminated scrap metal.



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Figure 1.7. SWMU 4 Historical Layout

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The C-748-B Burial Area, located on the west side of C-747, is identified as a Proposed Chemical Landfill Site in the 1973 Union Carbide document on waste disposal (however it is unknown if chemicals were ever buried here). Geophysical surveys indicate the area of C-748-B does contain disposal cells, which are shown in Figure 1.7. The waste is assumed to be similar to that buried in C-747. The original SWMU Assessment Report dated August 24, 1987, for SWMU 4 included only the C-747 Contaminated Burial Yard. The C-748-B Burial Area was incorporated into various descriptions of SWMU 4 starting in the mid-1990s as a result of a geophysical survey. As a result of this addition, the area of the SWMU was changed from 8,300 ft² to 286,700 ft² (DOE 2007b).

SWMU 4 also may have received sludges designated for disposal at the C-404 Burial Ground. The source of these sludges is unknown, but the Waste Area Grouping (WAG) 3 RI Work Plan (DOE 1998b) indicated that the sludges potentially included uranium-contaminated solid waste and ⁹⁹Tc-contaminated magnesium fluoride. The total volume of material disposed at this site is unknown. Potential contaminants associated with this SWMU include uranium, technetium-99, metals, and TCE.

During the summer of 1996, a small sinkhole (approximately 3 ft across and 3 ft deep) developed in the southern burial cell, apparently from settling of material within the SWMU. The sinkhole was backfilled with soil. This hole previously had been reported in the WAG 3 RI Report and the BGOU Work Plan as having developed in the fall of 1999.

In the fall of 1999, employee interviews led to designating the C-747 Burial Yard as a classified area. Access subsequently was restricted based on security considerations.

1.3.4 C-746-F Burial Yard (SWMU 5)

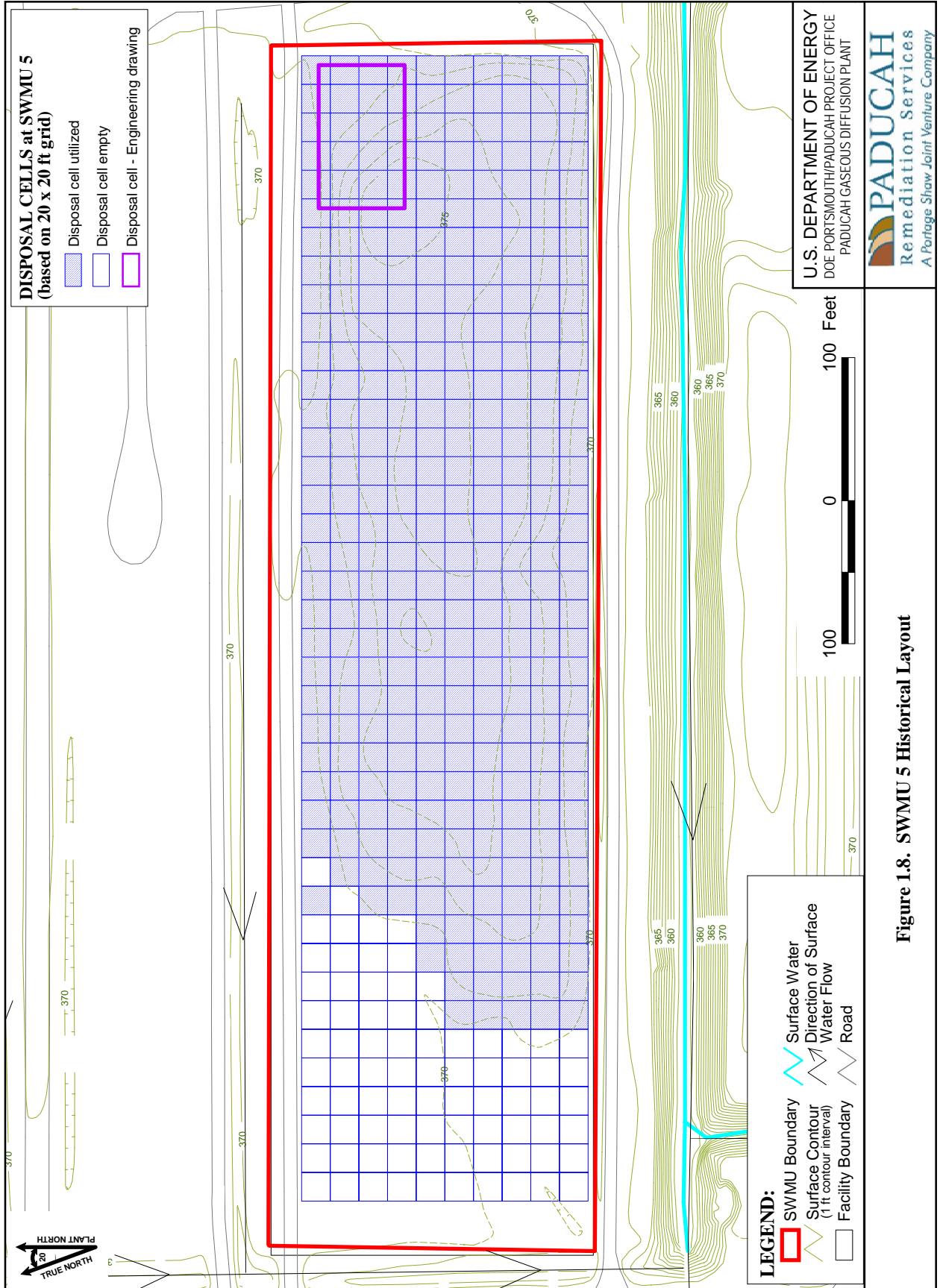
1.3.4.1 Site description

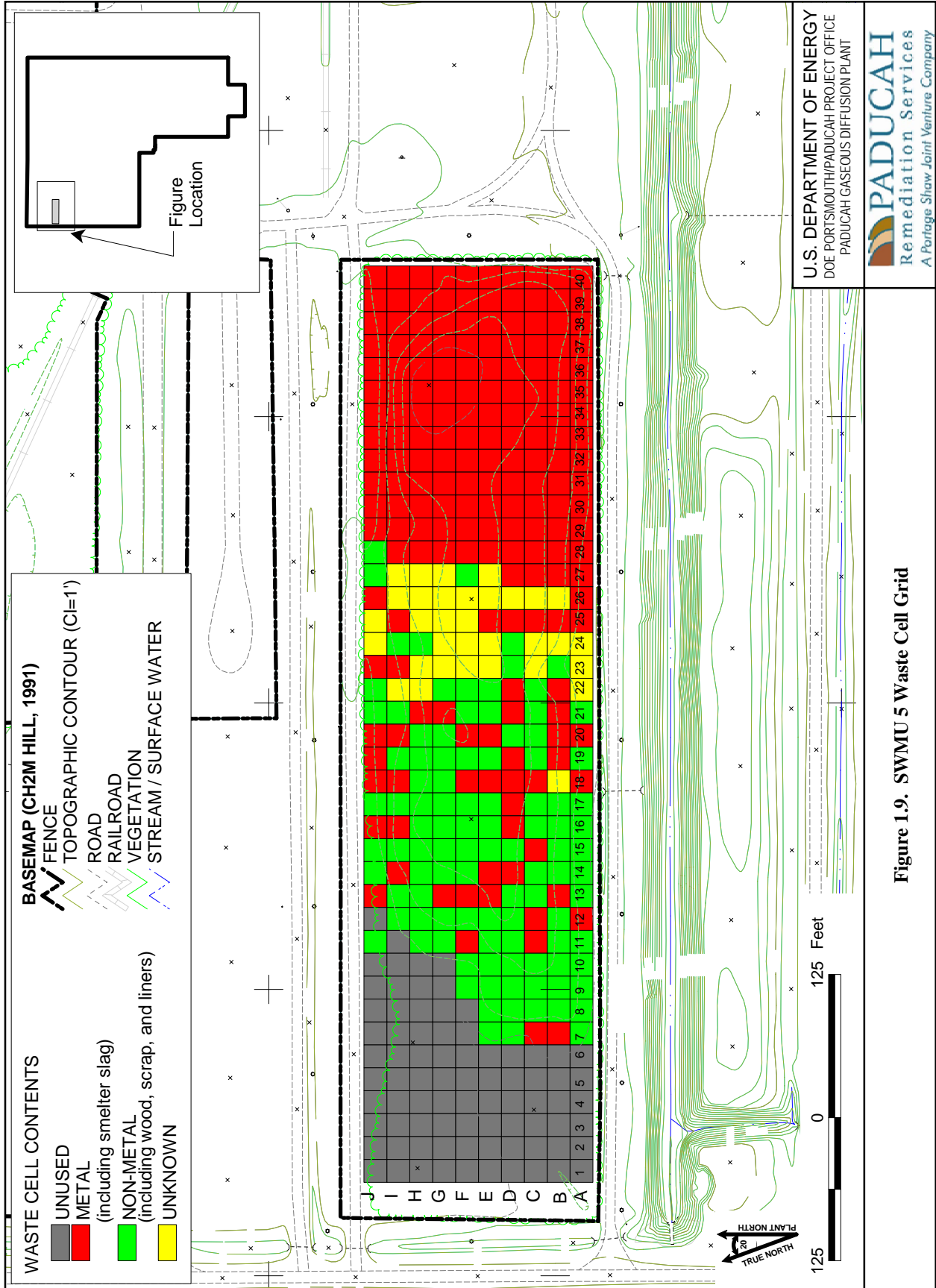
The C-746-F Burial Yard is located in the northwestern section of the PGDP secured area. SWMU 5 (which covers an area of approximately 197,400 ft²) is located adjacent to a scrap yard to the north (C-746-P/P1) and SWMU 6 to the east. Disposal pits were located on a grid system. Documentation of the size of these grids ranges from 10 by 10 ft cells to 20 by 20 ft cells excavated to a depth of 6 to 15 ft bgs. Figure 1.8 shows these cells as 20 by 20 ft. Worker interviews indicate this spacing is roughly accurate; however, historical aerial photographs indicate the earliest grid spacing may have been smaller. The fence around SWMU 5 has regularly spaced reflectors, which may have been used by workers as a reference in defining the waste cell grid in the field.

Waste placed in the yard disposal pits was covered with 2 to 3 ft of soil. SWMU 5 is fenced to limit access to authorized personnel only. The ground surface is covered with short grasses and various flowering herbaceous plants (DOE 1998b).

1.3.4.2 Site history

SWMU 5 was in operation from 1965 to 1987. The burial pits were used for the burial of components from the “Work for Others” activities, some radionuclide-contaminated scrap metal, and slag from the nickel and aluminum smelters (Figure 1.9). Table 1.8 shows examples of composition of material for what may have been placed in SWMU 5. Metals and radioisotopes are the primary potential contaminants of interest at this SWMU. The total quantity of wastes buried at the yard could be up to 896,000 ft³, assuming an average quantity of 2,800 ft³ waste placed in each cell and 320 cells receiving waste. Chemically unstable or incompatible compound/metal wastes are thought to have been placed here





also. This conclusion is supported by the occurrence of an underground fire (thought to have occurred circa 1975–1976) in the southeast corner of the yard. This fire burned for several weeks, and individuals observing the fire reported that the ground surface appeared to become unstable. The source and/or cause were never determined; however, subsequent worker interviews indicate the fire was thought to be a reaction from hot slag in contact with water, producing acetylene gas. The fire extinguished itself without intervention, and no testing was performed to prove or disprove this theory. Common practice following this incident was to allow slag to cool before placing it in the burial yard. No data are available related to contaminant releases from the fire. It was common practice in the 1960s and 1970s to collect materials such as scrap metal and slag from smelters and bury the waste without containerization.

Table 1.8. SWMU 5 Examples of Materials of Composition

Waste	Possible Constituents		
Metal	Aluminum	Gold	Steel
	Beryllium	Iron	Tantalum
	Boron	Lead	Tin
	Brass	Magnesium	Titanium
	Bronze	Nickel	Uranium
	Cadmium	Platinum	Zinc
	Chrome	Silver	Zirconium
	Copper		
Non-metal	Asbestos	Fiberglass	Plastics
	Carbon	Glass	Rubber
	Epoxies	Paint	Tritium

1.3.5 C-747-B Burial Ground (SWMU 6)

1.3.5.1 Site description

The C-747-B Burial Ground is located in the northwestern section of the plant area east of SWMU 5. The entire burial area covers an area of approximately 13,500 ft², which is divided into five separate burial cells (Figure 1.10). The following are the dimensions of each of the cells.

- Area H—This disposal site covers an area of about 12 by 15 ft and is about 6 ft deep. A 3 ft cover of soil was placed on top of the buried drums.
- Area I—This discard pit is approximately 8 by 35 ft and is about 8 ft deep. The waste was covered with about 5 ft of soil. A smaller pit located near the northwest corner of Area I, designated I-2 on Figure 1.8, is approximately 6 ft by 6 ft.
- Area J—This burial site is about 4,000 ft² (37 by 110 ft) and was excavated to a depth of about 6 ft. The area was covered with about 3 ft of soil.
- Area K—This disposal site consists of an area of about 12 by 15 ft and is about 6 ft deep. A 3 ft cover of soil was placed on top of the buried drums.
- Area L—This burial area is about 20 by 30 ft and about 6 ft deep. The disposed waste was covered with about 3 ft of soil.

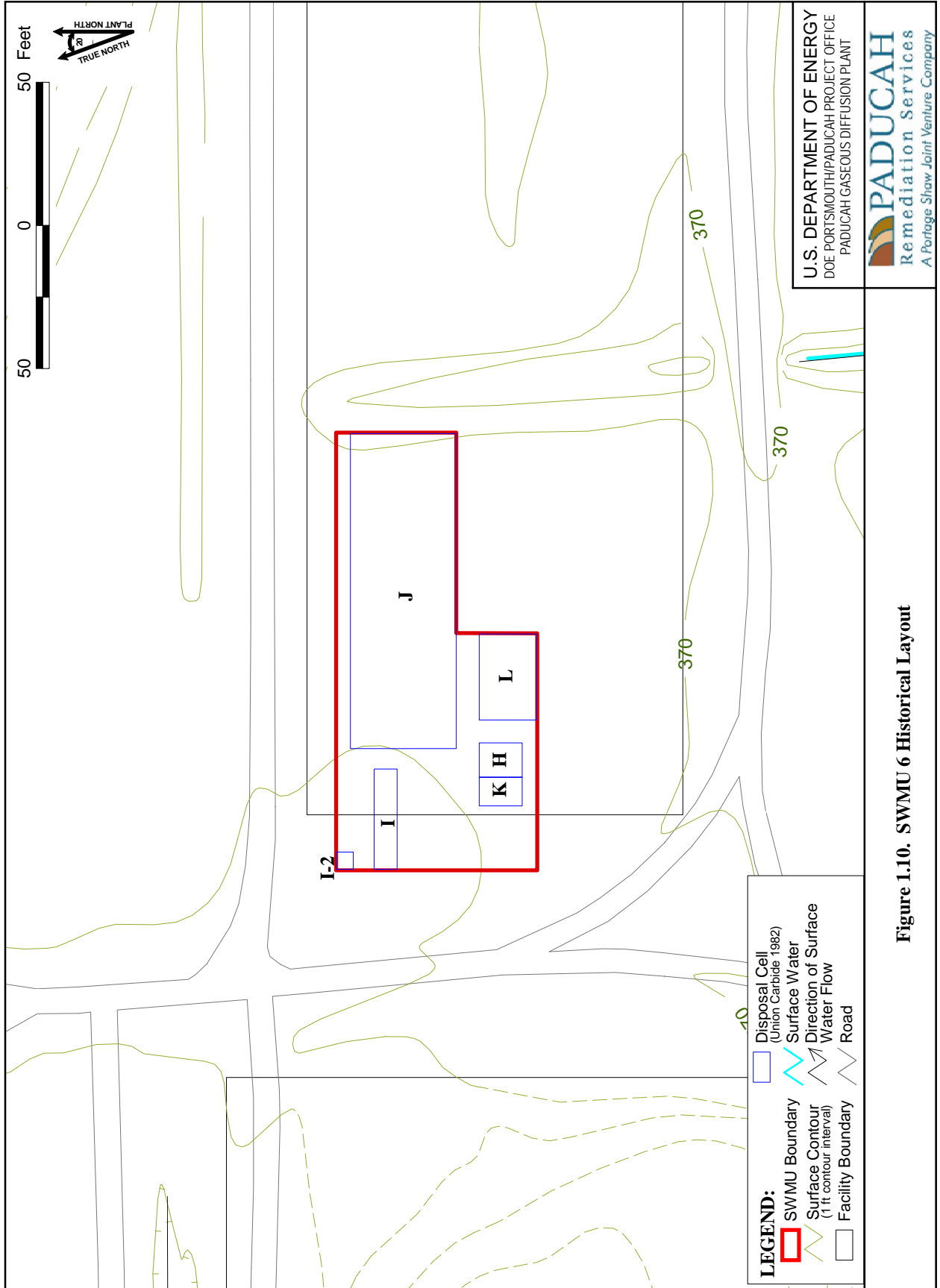


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SWMU 6 is relatively flat and is bounded to the north by a set of abandoned railroad tracks, to the east by a 5-ft wide by 4-ft deep drainage ditch that drains into Ditch 001, and unnamed gravel roads to the west and south. The ground surface is medium to tall grasses (up to 3 ft high) with occasional pockets of young trees and shrubs (DOE 1998b).

1.3.5.2 Site history

SWMU 6 was in operation from 1960 to 1976. Each of the burial cells was used for the disposal of a different waste. Each cell and its contents were identified in the WAG 3 RI Report (DOE 2000a) as follows:

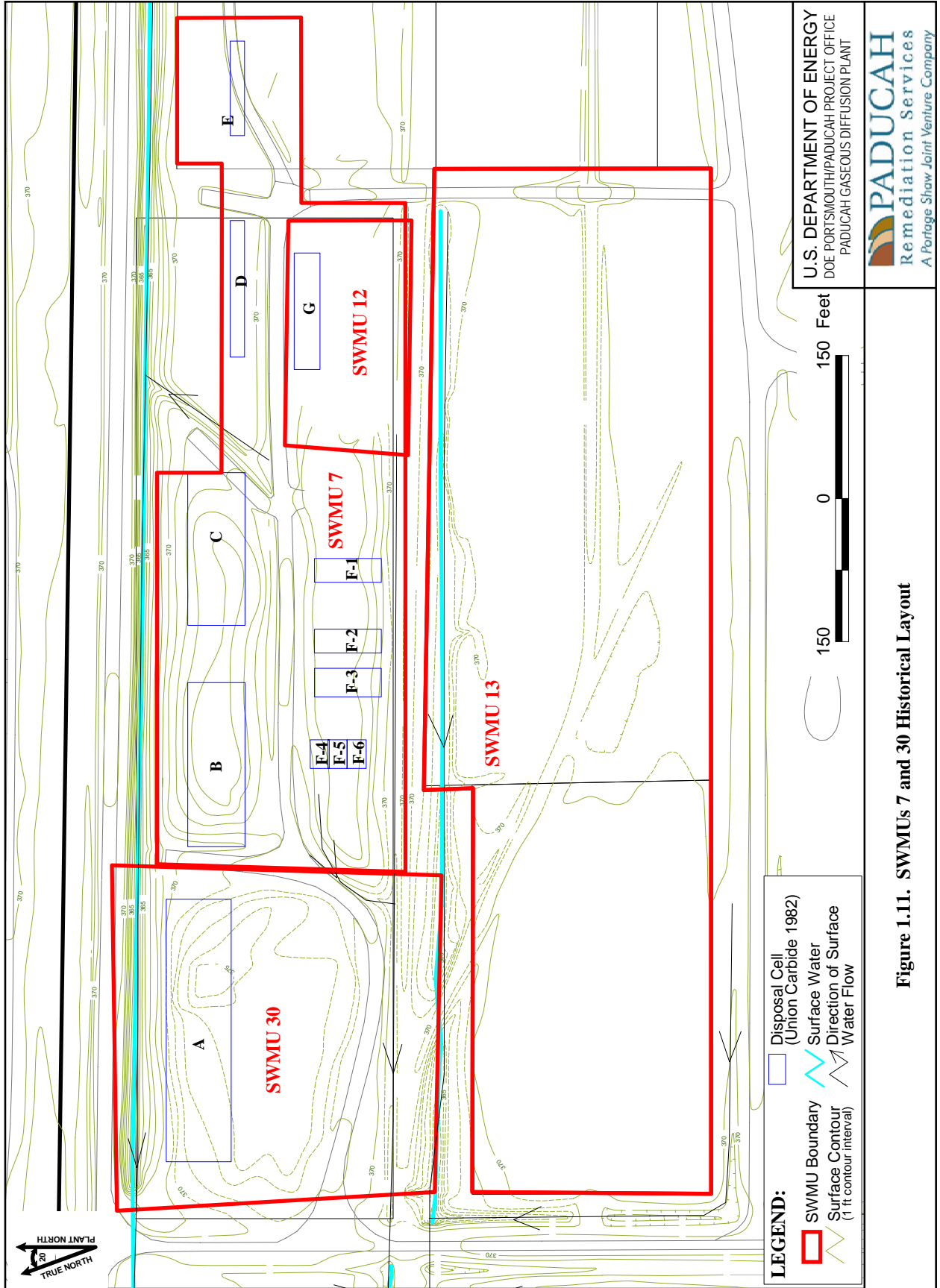
- Area H—Magnesium Scrap Burial Area. The scrap buried at this location is magnesium, in various shapes, generated in the machine shop. A total of about ten drums of scrap was buried during midsummer 1971.
- Area I—Exhaust Fan Burial Area. Eight exhaust hood blowers removed from C-710 were discarded to this pit. These blowers, which were about 15 inch in diameter and weighed about 100 lb each, were discarded in 1966 because of contamination with perchloric acid. Each blower was spaced about 4 ft apart in the hole. In 1976, additional exhaust fans from C-710 were buried in pit I-2.
- Area J—Contaminated Aluminum Burial Area. The contaminated scrap buried in this hole involved about 100 to 150 drums of aluminum scrap in the form of nuts, bolts, plates, trimmings, etc., that were generated in the converter and compressor shop. This scrap was buried in the early 1960s.
- Area K—Magnesium Scrap Burial Area. The scrap buried at this location is magnesium in various shapes generated in the machine shop. A total of about 20 drums of scrap was buried on September 3, 1968, and December 23, 1969.
- Area L—Modine Trap Burial Area. A single contaminated modine trap was buried in this area. The cold trap was about 4 ft in diameter, approximately 15 ft long, and weighed about 5,000 lb. This equipment was buried on March 5, 1969.

The WAG 3 RI Report (DOE 2000a), stated that approximately 50% of the surface area of SWMU 6 was used to store radioactively contaminated equipment and materials. These items include industrial forklifts and transport carts, flatbed trailers, generators, concrete pipes, and other miscellaneous items. This stored equipment has been removed. The area no longer is used for storage (DOE 2007c).

1.3.6 C-747-A Burial Ground (SWMU 7)

1.3.6.1 Site description

The C-747-A area is located in the northwest corner of the PGDP secured area. SWMU 7 comprises the eastern two-thirds of C-747-A. The SWMU is bounded on the north and south sides by perimeter ditches, on the west side by the C-747-A Burn Area (SWMU 30), and on the east side by the C-746-E Contaminated Scrap Yard. SWMU 7 covers approximately 240,900 ft² and includes six discrete burial pit areas described below and illustrated in Figure 1.11 (DOE 1998c). A soils operable unit SWMU, SWMU 12, overlaps a portion of SWMU 7. Any buried material beneath SWMU 12 is considered part of SWMU 7.



- Pit B—This pit is approximately 60 by 172 ft. According to the Phase II Site Investigation (SI) geophysical survey, the actual excavation extends beyond the designated boundaries and may connect with the adjacent burial pit (Pit C). A geophysical survey conducted for this RI interprets B and C as separate pits.
- Pit C—This pit is approximately the same size as Pit B. Based on the Phase II geophysical survey, Pit C and Pit B may be one continuous pit; however, a geophysical survey conducted for this RI interprets B and C as separate pits.
- Pit D—This pit is approximately 15 by 143 ft.
- Pit E (outside the eastern boundary of SWMU 7 and within the C-746-E Contaminated Scrap Yard)—This pit is approximately 15 by 99 ft.
- Pits F1–F5—These pits are all small (average size of each pit is approximately 20 by 80 ft). Engineering drawings indicate a sixth “F” pit that was not labeled.
- Pit G—This pit was documented as approximately 27 by 122 ft.

Records indicate the burial pits, in general, were excavated to a depth of 6 to 7 ft bgs, filled with wastes, and covered with approximately 3 ft of earth (Union Carbide 1978); however, geophysical surveys during the Phase II SI indicated waste in pits to a depth of 8-15 ft (CH2M HILL 1992).

A stockpile of radiologically contaminated scrap drums, locally known as Drum Mountain, formerly was located on the southeast corner covering Pit G. Interviews with a former heavy-equipment operator who worked in the SWMU 7 area indicate Drum Mountain was created only after the area between the F Pits and Pit G had been filled with similar material. This interview was corroborated by geophysical evidence (see Section 2.1).

The land surface slopes within SWMU 7. Burial Pits B and C form a slight hill on the north side of SWMU 7, and Burial Pit F forms a lesser mound on the south side of the SWMU. Pit D underlies a level area north of where Drum Mountain once was located. Shallow drainage swales occur on the west side of Burial Pit B, between Burial Pits C and D. The ground surface of the west half of the SWMU is covered by grassy vegetation, except where gravel roads extend through the site. A PGDP scrap metal project covered the west half of the SWMU with 1 to 2 ft of gravel as a working base for truck and tractor traffic. This gravel also prevents exposure to contaminated soils resulting from the earlier removal of scrap material in Drum Mountain.

Infrastructure has been placed in the area in support of the Scrap Metal Removal Action project. This infrastructure includes an extensive gravel pad constructed to support a truck scale in the area of Burial Pit G.

The upper 20 ft of soils at SWMU 7 consist of surface soil, fill, and loess, alternatively described as silt or clay, in the area boreholes. Surface soils, to a depth of 6 inches, were sampled and described during the Phase II SI. Soil textures range from sand with gravel to lean clay with gravel. During the Phase II SI, double-ring infiltrometer tests were conducted on surface soils at SWMU 7. Average long-term infiltration rates ranged less than 5.7 ft/day (CH2M HILL 1992). Logs of deeper soil borings demonstrate that coarse textures generally are limited to the upper 2 ft, with the exception of the burial pits that are now known to be as much as 10 ft deep.

The surface water that drains from SWMU 7 into the surrounding ditches is carried west through Outfall 001 into Bayou Creek. In 2002, a sedimentation basin was constructed to contain runoff from PGDP scrap yards. Runoff now flows into the sedimentation basin and is released periodically into Outfall 001.

1.3.6.2 Site history

PGDP used the burial pits for disposal of wastes from 1957 to 1979. Burial Pits B, C, and G were used for disposal of noncombustible, contaminated and uncontaminated trash, material, and equipment. Contaminated concrete removed from the C-410 Feed Plant during May and June 1960 was placed in Burial Pits D and E. Burial Pit F was used for disposal of uranium-contaminated scrap metal and equipment. Empty uranium and magnesium powder drums also were reported to have been buried in Burial Pit F (Union Carbide 1978).

The following summarizes what is known about the disposed waste in the burial pits.

- Pit B—Buried material includes noncombustible trash and contaminated and noncombustible material and equipment (however, no specific disposal records exist).
- Pit C—Historic records indicate that both Pit B and C received the same material.
- Pit D—Documented buried material consists of uranium-contaminated concrete pieces of reactor tray bases from C-410 used during the fluorination process of uranium tetrafluoride to uranium hexafluoride.
- Pit E—Documented buried material consists of uranium-contaminated concrete pieces of reactor tray bases.
- Pits F1–F5—Documented buried material consists of uranium-contaminated scrap metal and equipment and empty uranium and magnesium powder drums (engineering drawings indicate there was a sixth “F” pit that was not numbered).
- Pit G—Documented buried material consists of noncombustible trash and contaminated and noncombustible material and equipment.

In addition to these burial pits, the Phase II SI geophysical investigation also identified another anomaly in the shape of a rough circular area (15 ft diameter) between SWMU 30 and SWMU 7, west of the F-4 and F-5 Pits (see Section 2). There is no information confirming the presence or the nature of any buried wastes associated with this anomaly.

An FS was prepared for SWMUs 7 and 30 in 1998 (DOE 1998c). The purpose of the FS was to develop and evaluate alternatives including (1) no action, (2) a limited action, (3) installation of a soil cover and additional groundwater monitoring, and (4) surface soil removal and additional groundwater monitoring. The proposed plan and ROD never were issued.

1.3.7 C-747-A Burn Area (SWMU 30)

1.3.7.1 Site description

SWMU 30 includes the western one-third of C-747-A. It consists of an historical burn-and-burial pit (Burial Pit A) and the location of a former incinerator. The SWMU is bounded on the north and south sides by ditches, on the west side by a plant road, and on the east side by SWMU 7 (Figure 1.11). The unit encompasses approximately 117,600 ft². The pit is reported to have been excavated to a depth of 12 ft and covered with 4 ft of earth. The land surface slopes gently, and a slight mound rises over Burial Pit A. SWMU 30 is bordered by drainage ditches on the north and south side. Grassy vegetation covers the ground, except where gravel roads extend through the site.

Phase II SI surface soil sample sites H-361 through H-366, H-370, and H-373 provide characterization of surface soil texture from eight locations across SWMU 30. The upper 6 inches of soil ranges from lean clay to sand. Surface soil samples from the Burial Pit A area tend to be lean clay with gravel, whereas surface soil textures from the south side of SWMU 7 range from lean clay to silty sand with gravel (DOE 1998c). The Phase II SI included double-ring infiltrometer tests on surface soils at three locations. Average long-term infiltration rates were less than 6×10^{-3} ft/day for two of the tests. All deeper soil borings, including Phase II SI borings H-211 and H-212, MW 66, and boring S-2, encountered surficial fill materials to depths of 2 to 12 ft.

1.3.7.2 Site history

SWMU 30 was used from 1951 to 1970 to burn combustible trash, which may have contained uranium contamination. An incinerator was constructed for use at SWMU 30, but the exact time frame is uncertain. The incinerator was a steel mesh, “tee pee” shaped structure primarily used to burn paper, wood, cardboard, and other combustibles. Ash and debris were buried below ground in Burial Pit A beginning in 1962, when use of an on-site incinerator was discontinued. It is assumed ash from incineration was buried at SWMU 30 rather than taken elsewhere at the site. Site maps and a surface electromagnetic geophysical survey of the Phase II SI identify the location of Burial Pit A. Prior to identification by Phase II SI surface geophysics testing; it was believed that remnants of the former incinerator were not present. Further research identified images of the incinerator at the location. This disposal site covers an area of about 250 ft by 50 ft. Geophysical data from the Phase II SI indicate that the actual area of excavation does not exactly match the rectangular outline and extends beyond the rectangular outline to the north and east. Material disposed in Pit A included contaminated and uncontaminated trash, ash, and debris.

In addition to Pit A, the Phase II SI geophysical investigation also identified another anomaly in the shape of a rough circle approximately 43 ft in diameter (see Section 2). The SWMUs 7 and 30 RI confirmed this anomaly likely was the metal reinforcement within the footer and retaining walls of the former incinerator and/or parts of the unit buried there upon decommissioning (DOE 1998c).

An FS was prepared for SWMUs 7 and 30 in 1998 (DOE 1998c). The purpose of the FS was to develop and evaluate alternatives including (1) no action, (2) a limited action, (3) installation of a soil cover and additional groundwater monitoring, and (4) surface soil removal and additional groundwater monitoring. The proposed plan and ROD never were issued.

1.3.8 Area P (SWMU 145)

1.3.8.1 Site description

Area P (SWMU 145) is located north of the PGDP security area and is defined by encompassing the area underneath SWMUs 9 and 10 (the C-746-S and -T Landfills, respectively). The SWMU is approximately 44 acres and began operation in the early 1950s. Currently, the C-746-S and -T Landfills are located on top of SWMU 145, but are not included in SWMU 145 (DOE 1999b), as illustrated in the conceptual drawing, Figure 1.12. The boundaries of the area previously had not been well defined outside of the area utilized by the C-746-S and -T Landfills.

1.3.8.2 Site history

SWMU 145 began operation in the early 1950s. A 1973 document *The Discard of Scrap Materials by Burial at the Paducah Plant* (Union Carbide 1973), states this area was used by the contractor during the construction of PGDP to discard all types of scrap and waste materials. Use of the area for discarding of scrap and waste by subcontractors was continued until the early 1980s. Construction debris, such as concrete, roofing materials, wire, wood, shingles with asbestos, and welding rods, are expected to have been disposed in the area (during drilling for the RI, glass was recovered that appears to have come from a laboratory). Approximately once a year, the accumulated scrap piles were moved by plant personnel into piles or earth depressions and, whenever practicable, covered with dirt. The area was later permitted for the construction and operation of the C-746-S and -T Landfills (BJC 2001a). The C-746-S Landfill began operation in 1981. Figure 1.13 shows historical aerial photographs of the area and depicts evident ground scarring, likely indicating disposal areas.

Several monitoring wells are present in the area for permit-related monitoring. Since 2003, these wells have indicated the presence of PCBs in the RGA (see Section 4).

1.4 REPORT ORGANIZATION

This RI report was prepared following the guidance found in Appendix D of the FFA for PGDP (EPA 1998a). The outline of this report followed the guidance presented in Appendix D of the *Work Plan for the Burial Grounds Operable Unit Remedial Investigation/Feasibility Study at the Paducah Gaseous Diffusion Plant, Paducah, Kentucky* (DOE 2006a).

These sections are consistent with the FFA. The following are their locations within this report.

- Chapter 1—Introduction
- Chapter 2—Study Area Investigation
- Chapter 3—Physical Characteristics of the Study Area
- Chapter 4—Nature and Extent of Contamination
- Chapter 5—Fate and Transport
- Chapter 6—Baseline Human Health Risk Assessment
- Chapter 7—Summary and Conclusions
- Chapter 8—References

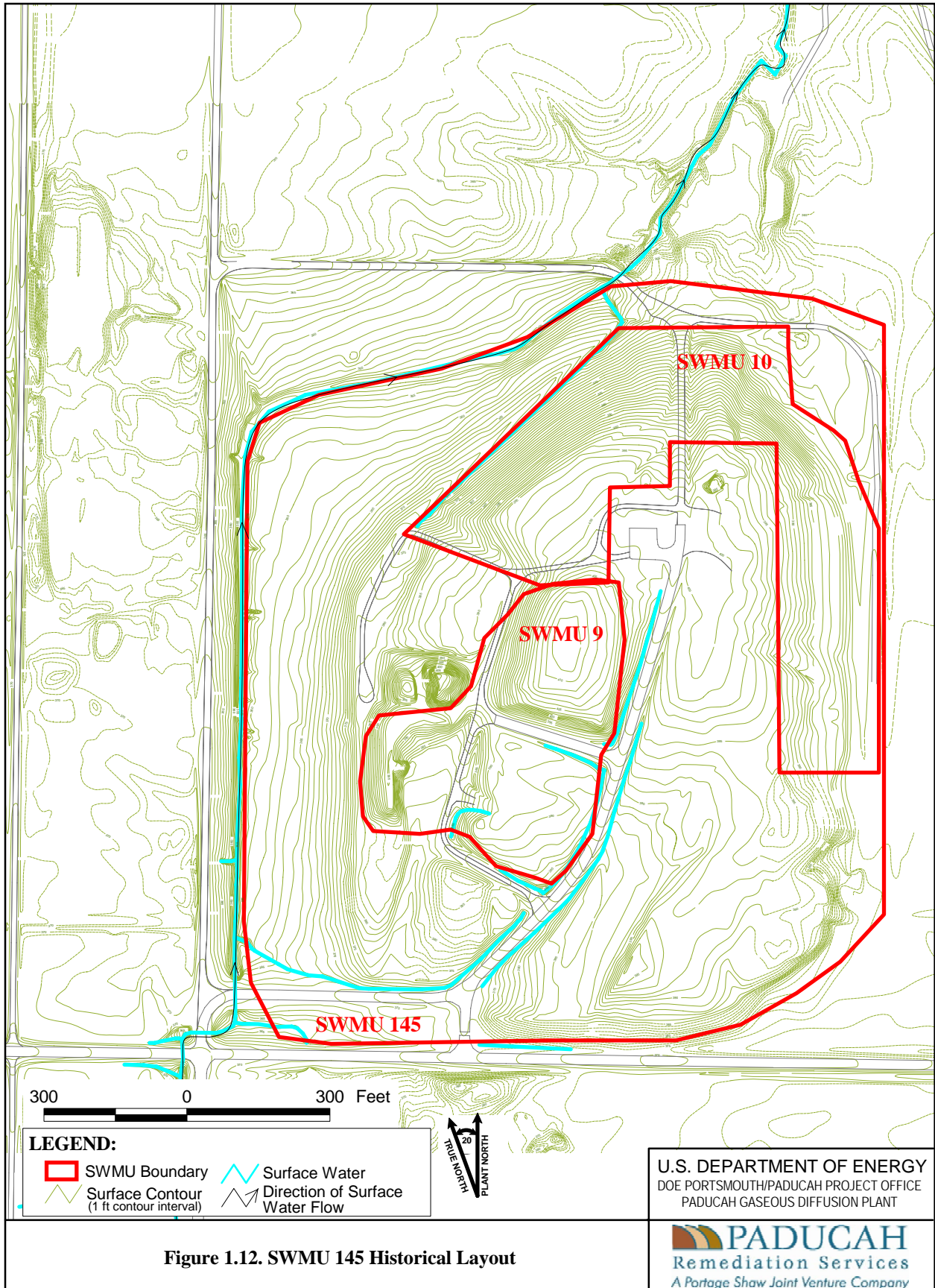


Figure 1.12. SWMU 145 Historical Layout

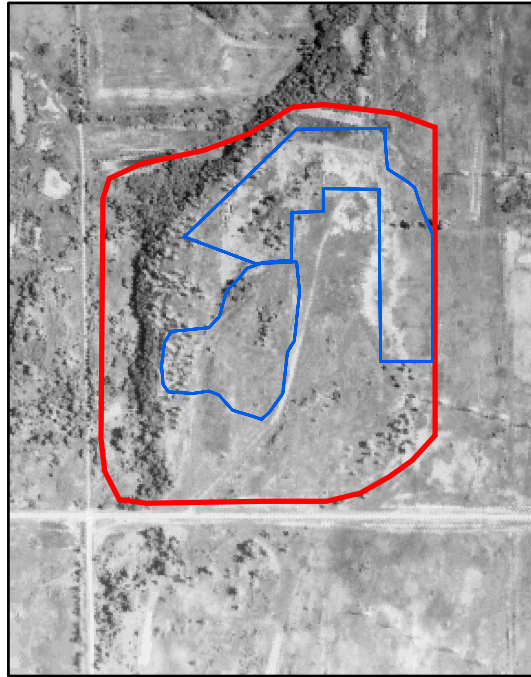
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Figure No. 1BGOU1.R1.apr
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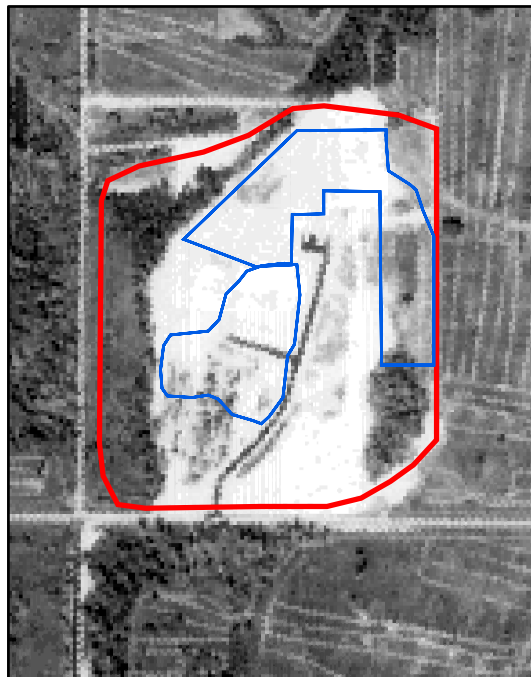
1959 Aerial Photograph



1964 Aerial Photograph



1975 Aerial Photograph



1981 Aerial Photograph

LEGEND

- CURRENT SWMU 145 BOUNDARY
- CURRENT SWMUs 9 and 10 BOUNDARIES

0 300 600 1,200 Feet



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Figure 1.13. Historical Aerial Photographs of SWMU 145

Additionally, the following appendices are included to support the information presented in the text.

- Appendix A—Technical Memorandum for Field Activities
- Appendix B—Lithologic Logs and Well Construction Diagrams, Groundwater Stabilization Logs, and Well Development Logs
- Appendix C—Analytical Data and Quality Assurance (QA)/Quality Control (QC) Evaluation Results
- Appendix D—Three-Dimensional Visualization Figures
- Appendix E—Fate and Transport Modeling
- Appendix F—Baseline Human Health Risk Assessment
- Appendix G—Review of Ecological Risk Assessments

2. STUDY AREA INVESTIGATION

Section 2 includes all field activities associated with site characterization of the BGOU. Technical memoranda documenting details of field activities are included in Appendix A.

2.1 GEOPHYSICAL INVESTIGATIONS

Current geophysical investigations were combined with historical geophysical information to create a more complete picture of the burial area.

As part of the RI field activities, geophysical surveys of SWMUs 2, 5, 7, 30, and 145 were conducted prior to sampling activities. The BGOU represented a difficult target for geophysical characterization because the SWMUs contain a heterogeneous collection of wastes and backfill soils, and some of these SWMUs consist of multiple burial pits of various depths. Geophysical surveys were not planned to be conducted at SWMUs 2 and 5 during scoping of the work plan; however, in support of the excavation/penetration permits procedure, geophysical surveys were conducted in proposed drilling locations where there was uncertainty in waste boundaries to ensure waste cells were not encountered.

An electromagnetic (EM)-61 magnetometer survey was conducted at the surface of these SWMUs to delineate the location and extent of the burial pits. The EM-61 survey was implemented for the most part along continuous lines primarily spaced 5 ft apart in a grid layout. A data logger was employed for data acquisition, and resultant geophysical anomalies were marked in the field and plotted using plant coordinates to an electronic overlay.

In addition to the current geophysical surveys, historical geophysical information is available for SWMUs 2, 4, 5, 6, 7, and 30. Results of the geophysical surveys conducted for the BGOU RI and historical information gathered during this RI is presented in Figures 2.1 through 2.6. The geophysical anomalies to the north and east of the SWMU 6 boundary shown in Figure 2.4 represent equipment (forklifts, mowers, metal debris) that was stored or parked on the surface. While the historical geophysical survey did not cover all of the waste cells (H, I, K, and L) at SWMU 6, engineering drawings were adequate to indicate where buried material was located. The geophysical anomaly identified on Figure 2.6 was mapped during a previous investigation. Limits of the geophysical anomaly, mapping the Old North-South Diversion Ditch (NSDD), were determined by the previous investigation objectives. The filled ditch channel extends beyond the limits of the geophysical survey.

The area within SWMU 13 identified by an employee interview was confirmed by a geophysics survey. The results of this survey are presented in Figure 2.7. After the site investigation, the geophysical information and data for this area will be evaluated and discussed with the FFA parties and, if further action is necessary, a path forward will be determined.

2.2 CONTAMINANT SOURCE INVESTIGATIONS

In order to evaluate migration of contaminants from sources (burial cells), angled soil borings were utilized to collect samples from the soils and groundwater beneath the burial pits. Available information from aerial photographs, historical and current geophysics, engineering drawings, and previous RIs was used to determine the most probable location of the burial pit. Angled soil borings then were placed to collect samples from beneath the burial pits and/or cells. These temporary borings provide a “snapshot” of

the current conditions at the time of sampling. Figures 2.8 through 2.14 show the locations of these angled borings.

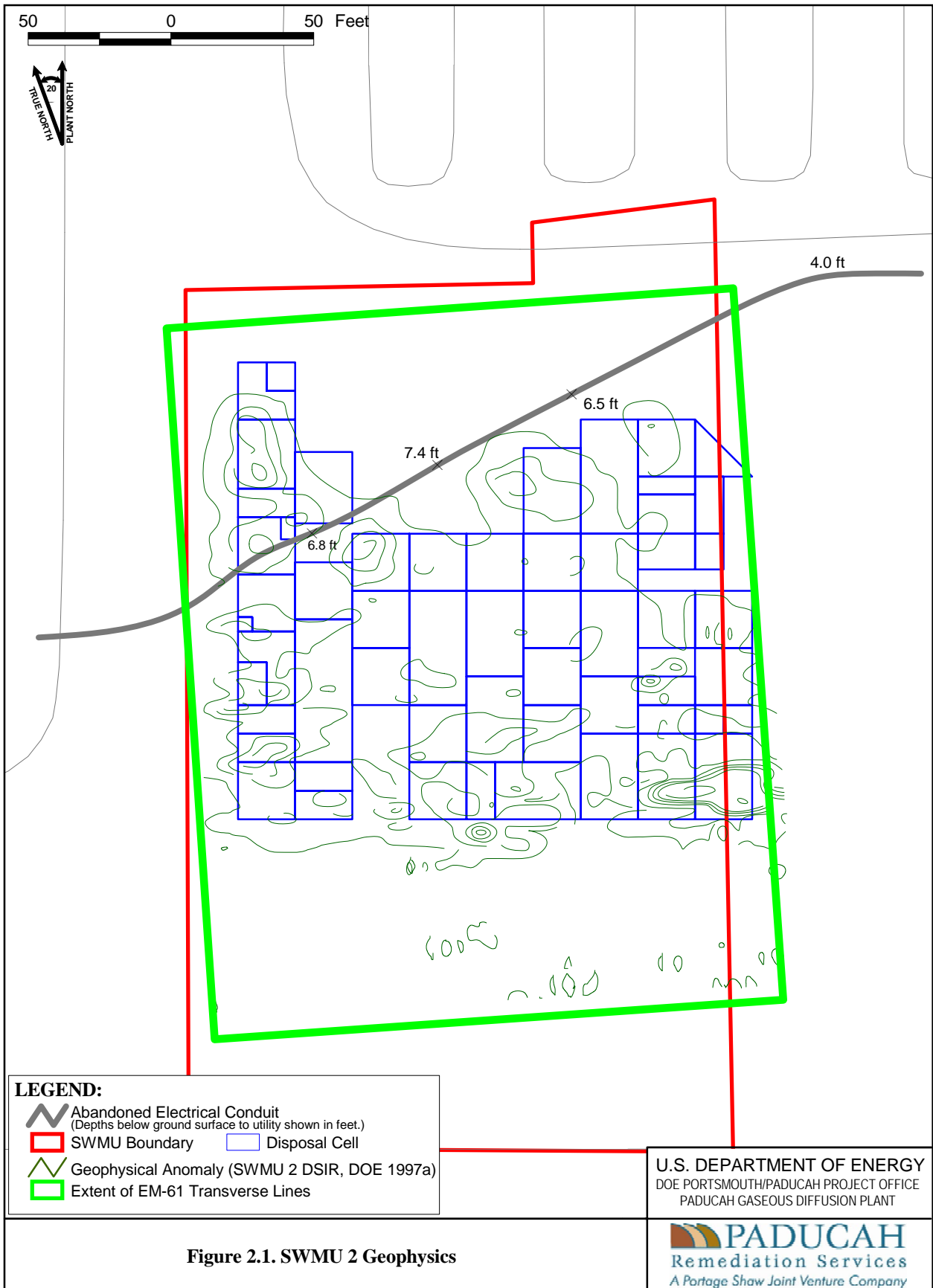
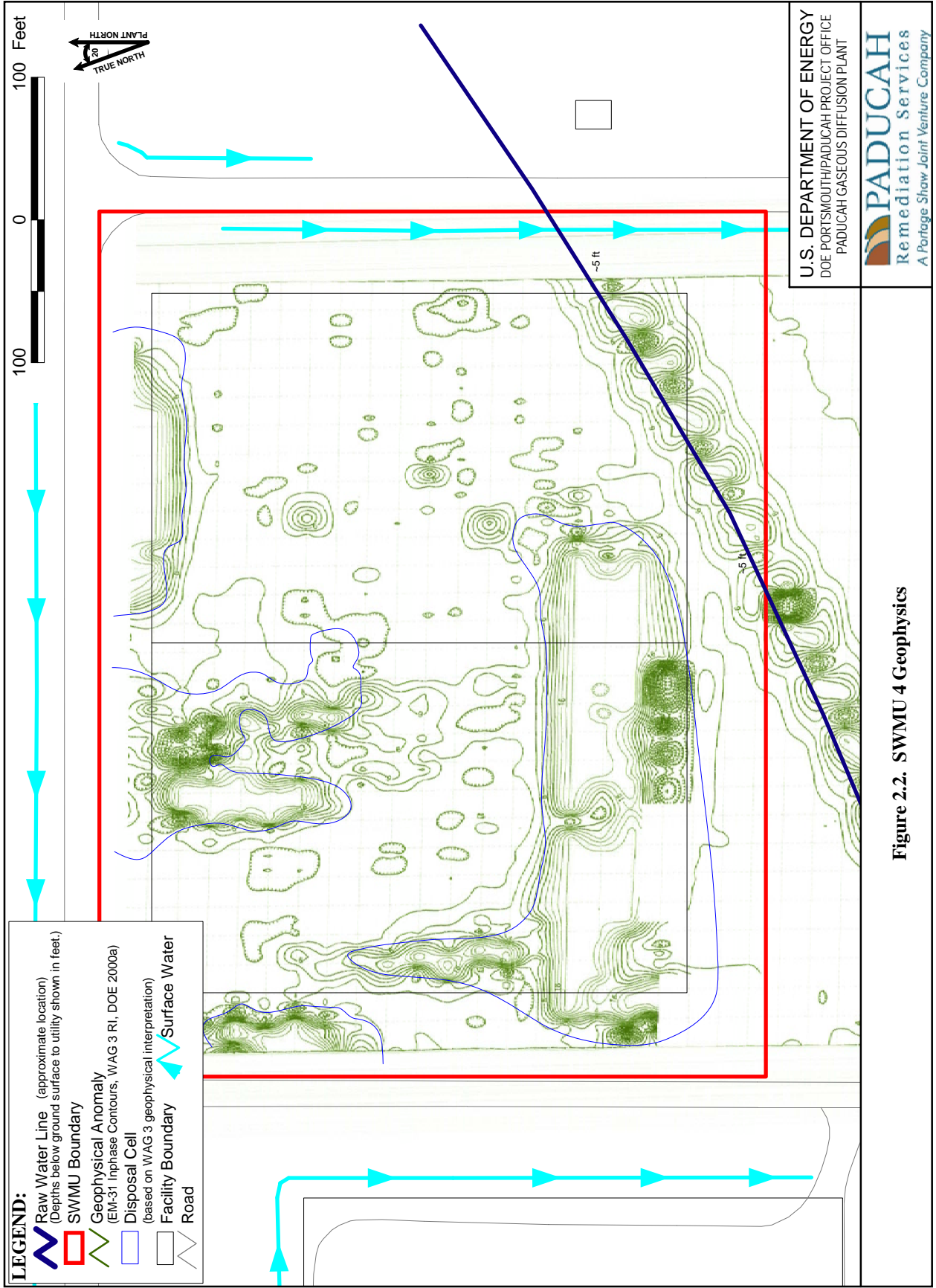
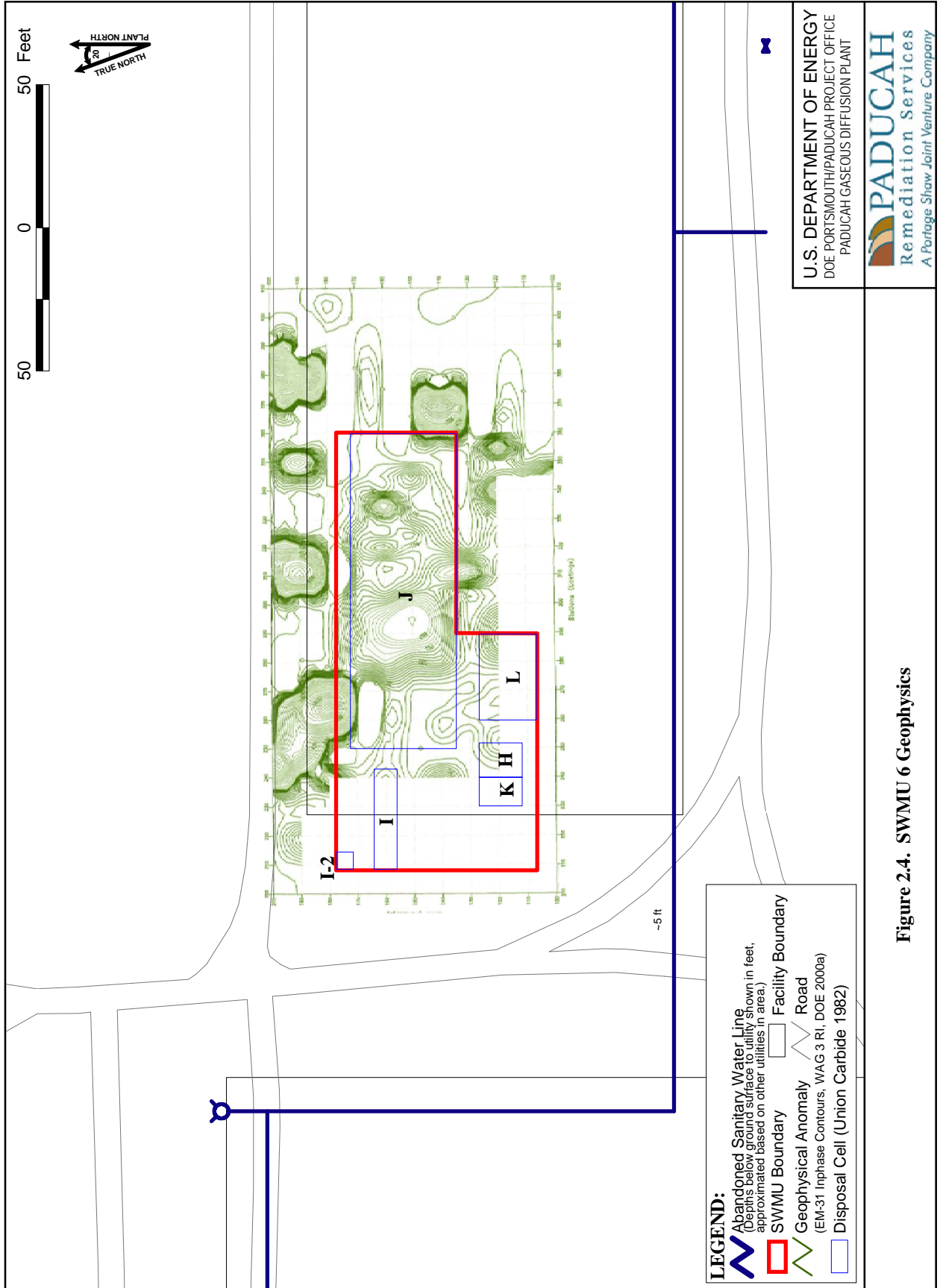


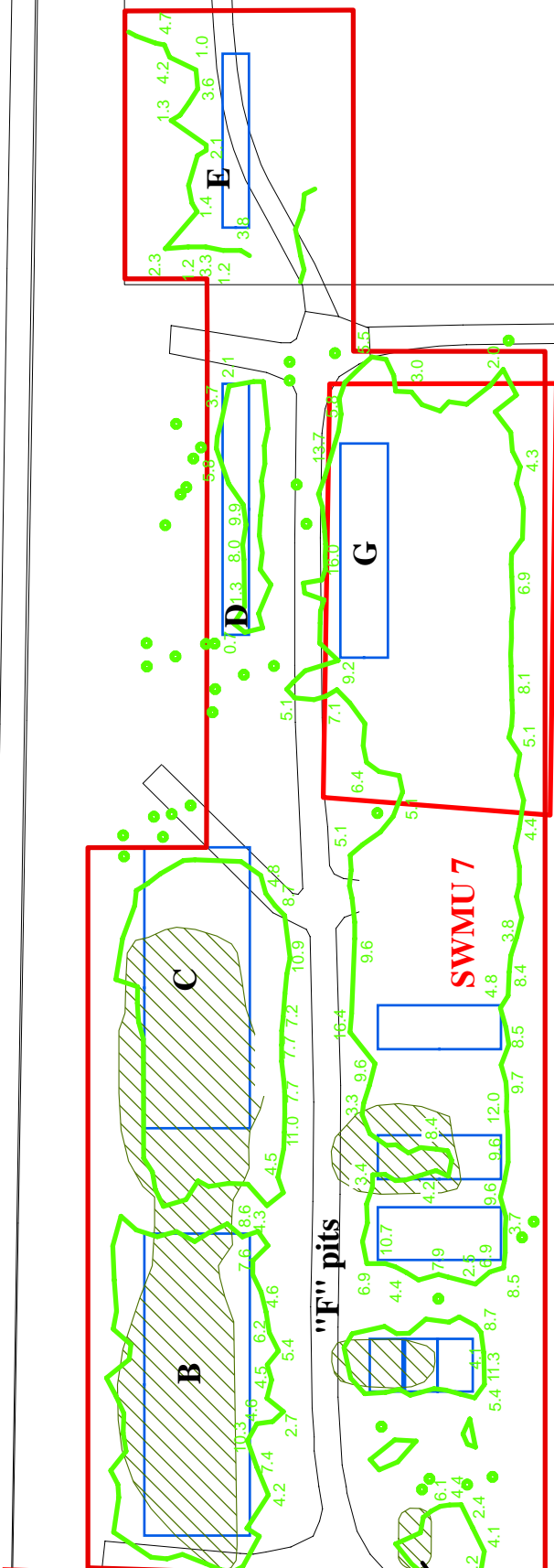
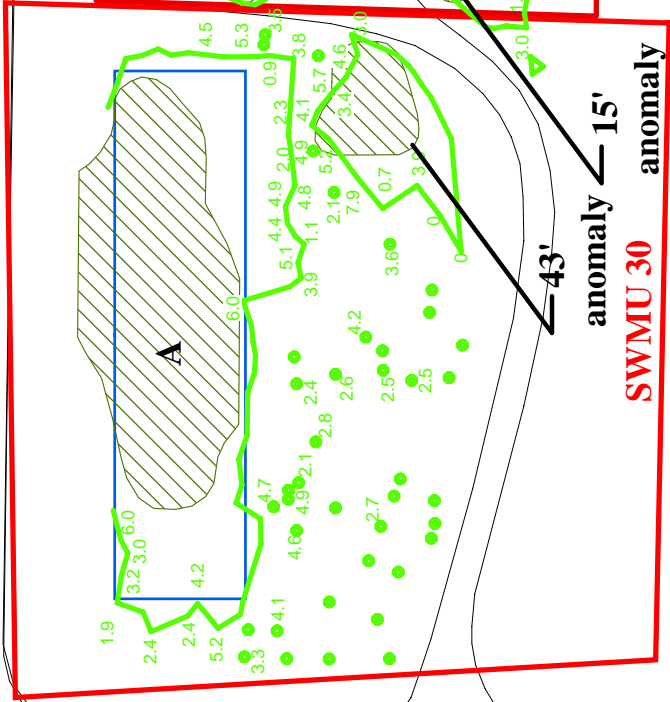
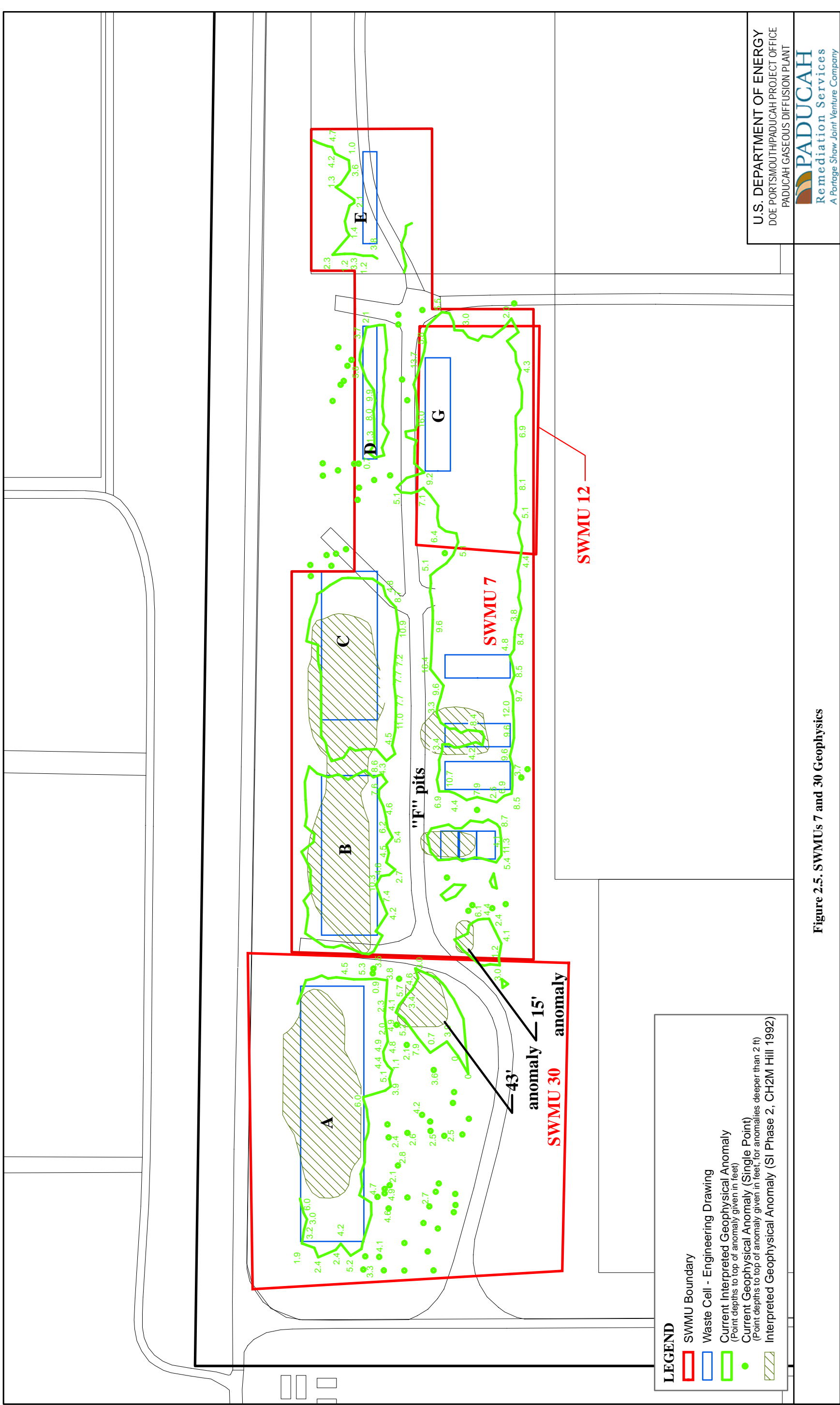
Figure 2.1. SWMU 2 Geophysics

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LEGEND

- SWMU Boundary
- Waste Cell - Engineering Drawing
- Current Interpreted Geophysical Anomaly (Point depths to top of anomaly given in feet)
- Current Geophysical Anomaly (Single Point) (Point depths to top of anomaly given in feet, for anomalies deeper than 2 ft)
- Interpreted Geophysical Anomaly (SI Phase 2, CH2M Hill 1992)

Figure 2.5. SWMUs 7 and 30 Geophysics

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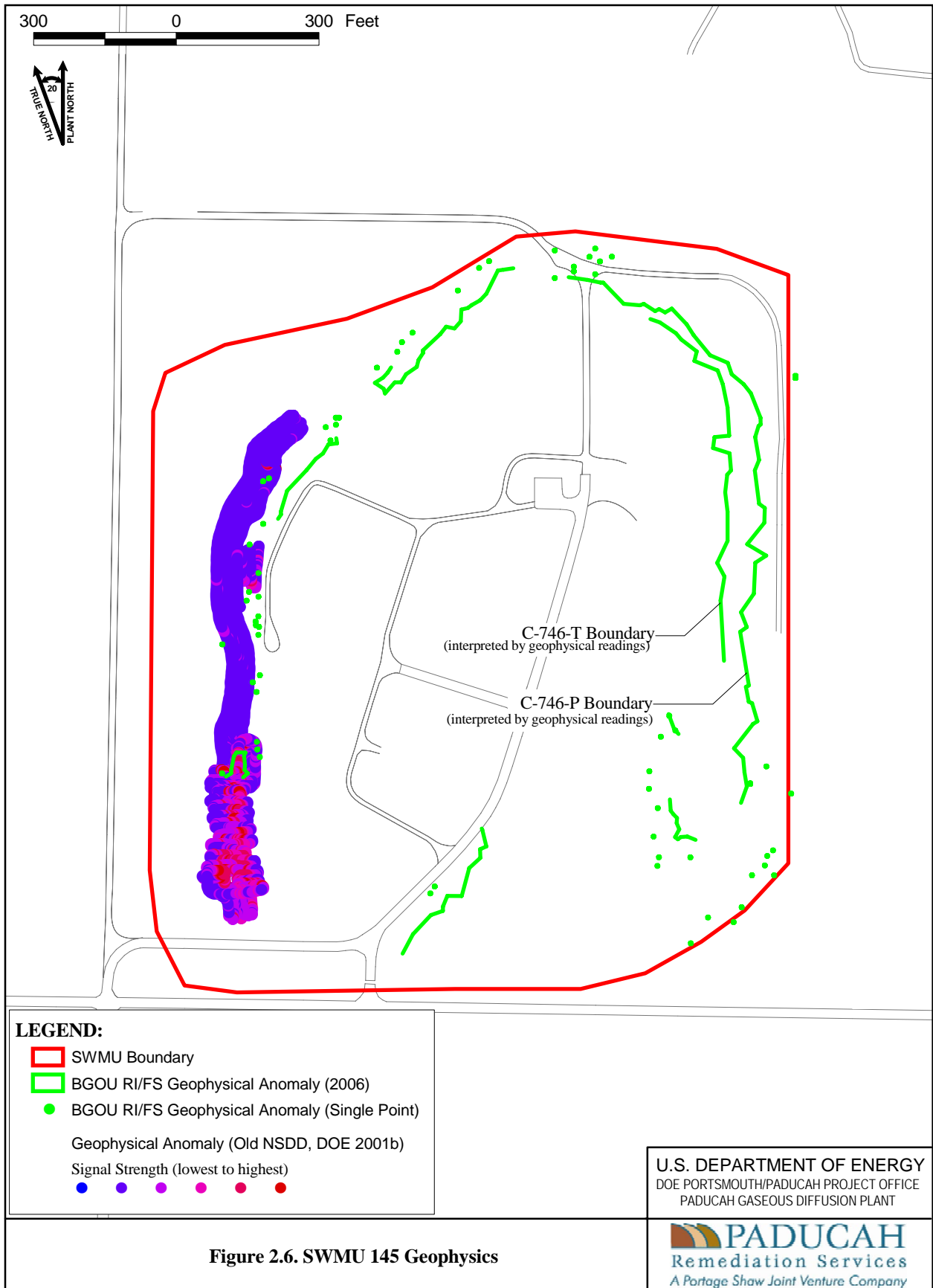
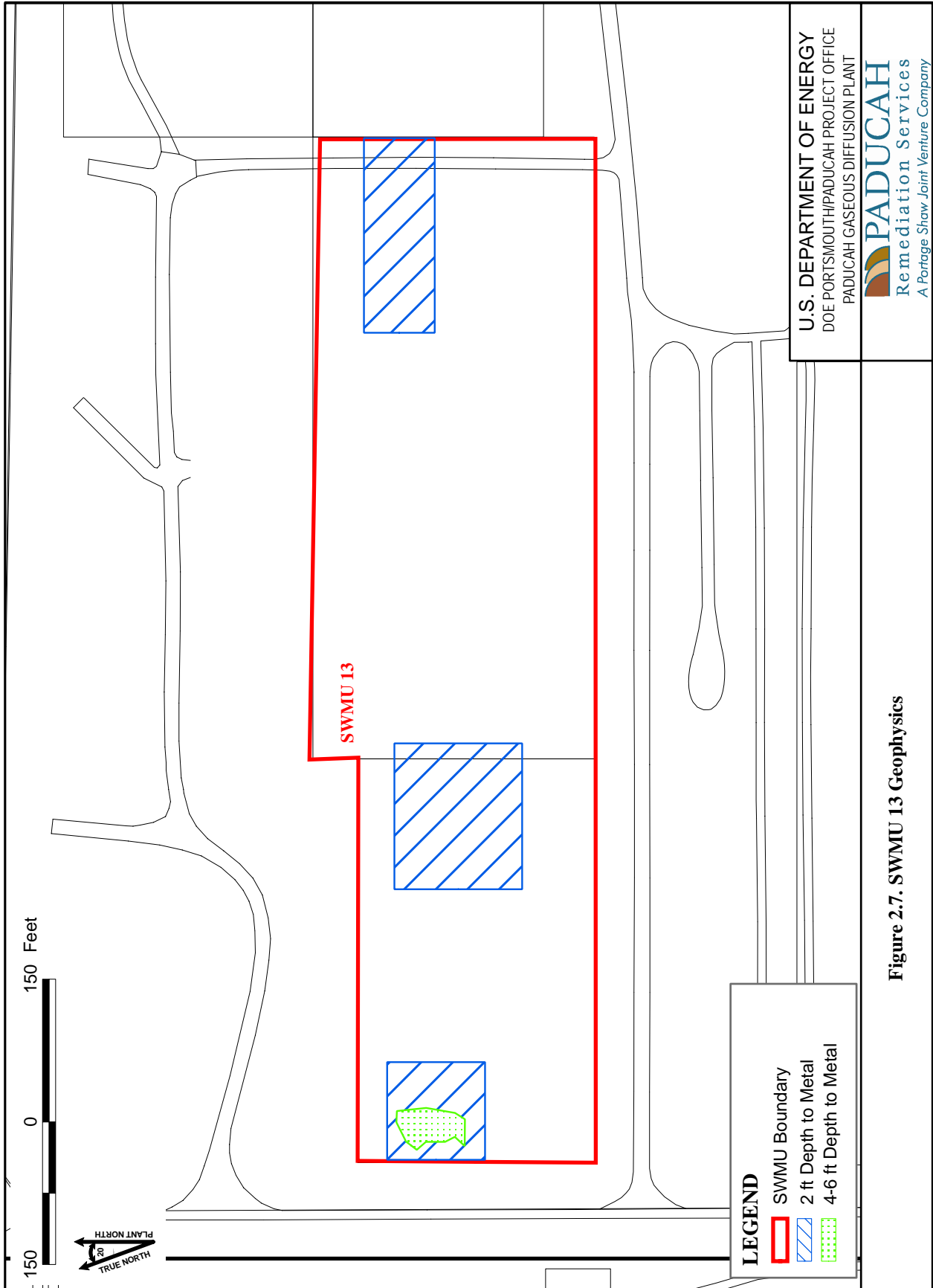


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Figure No. bgoufri swmu 13.mxd
 DATE 12-18-07

Figure 2.7. SWMU 13 Geophysics

LEGEND

- SWMU Boundary
- 2 ft Depth to Metal
- 4-6 ft Depth to Metal

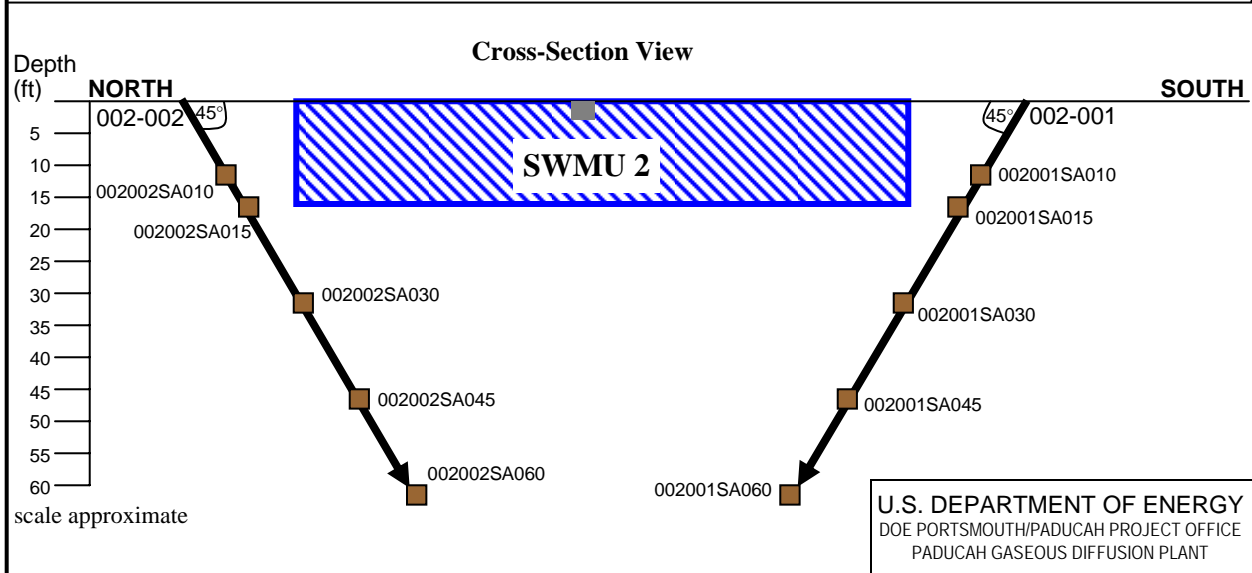
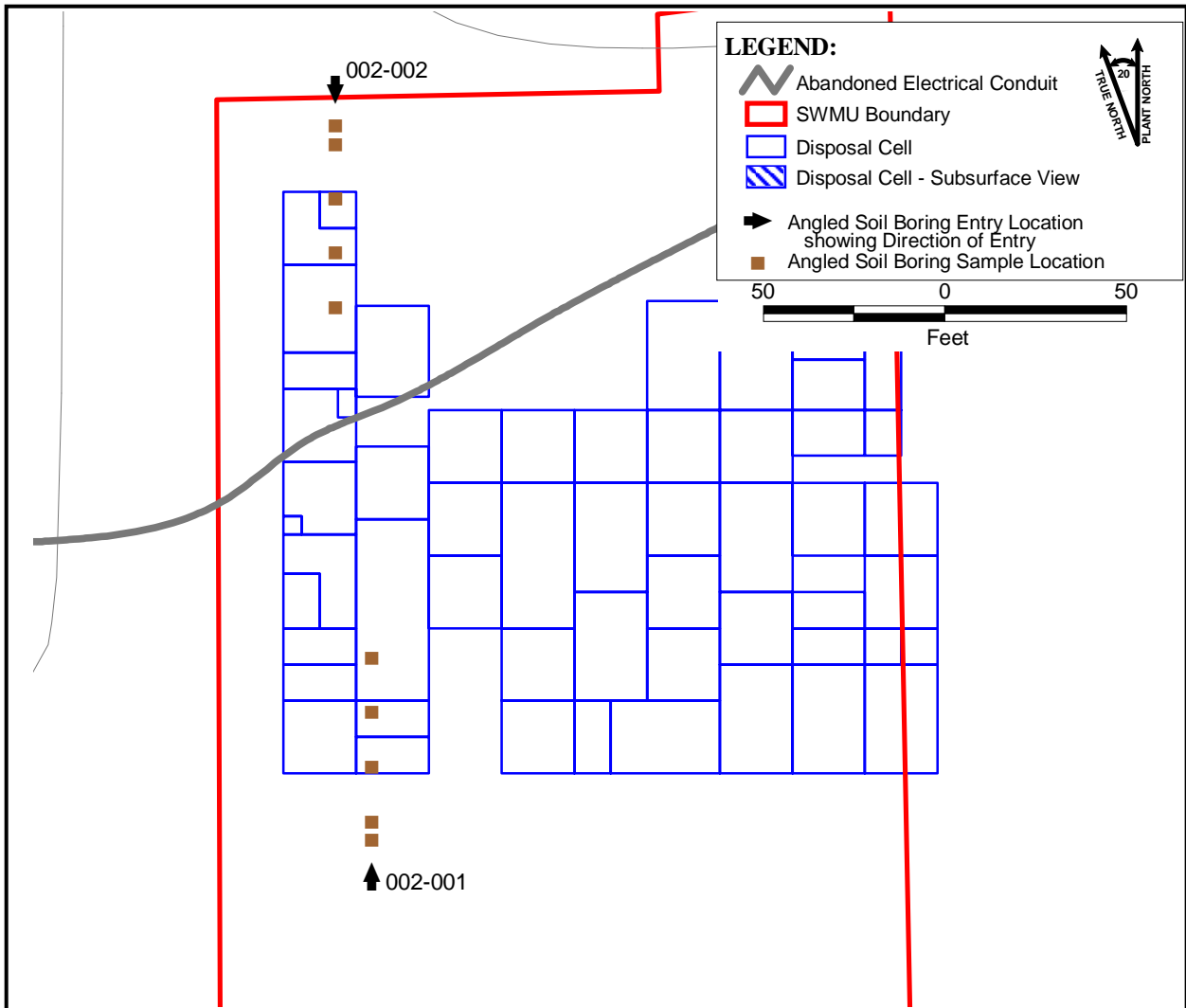
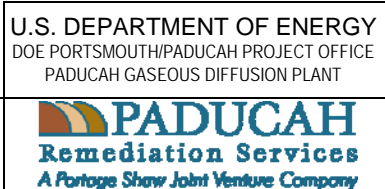


Figure 2.8. SWMU 2 Angled Boring Locations



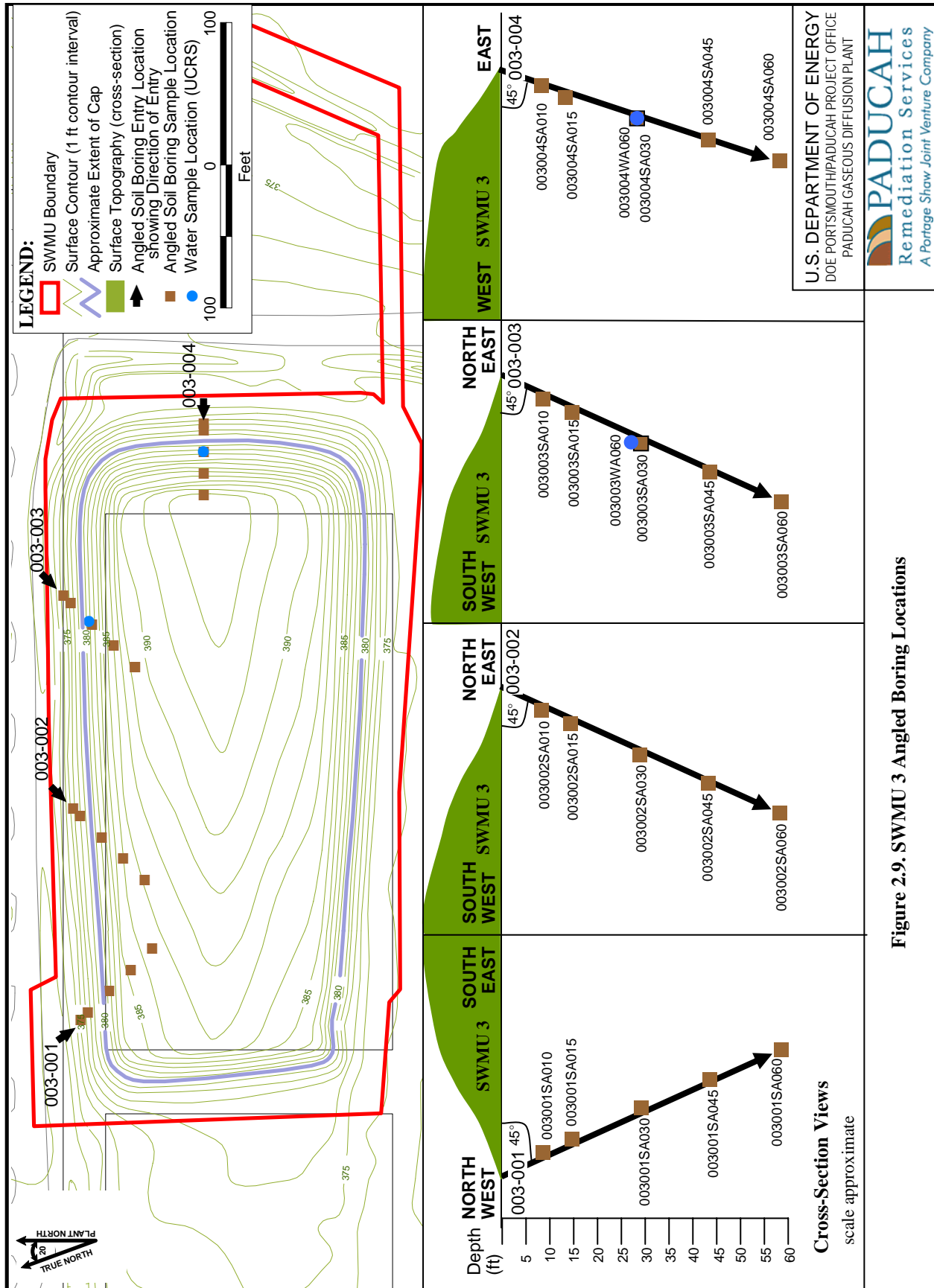
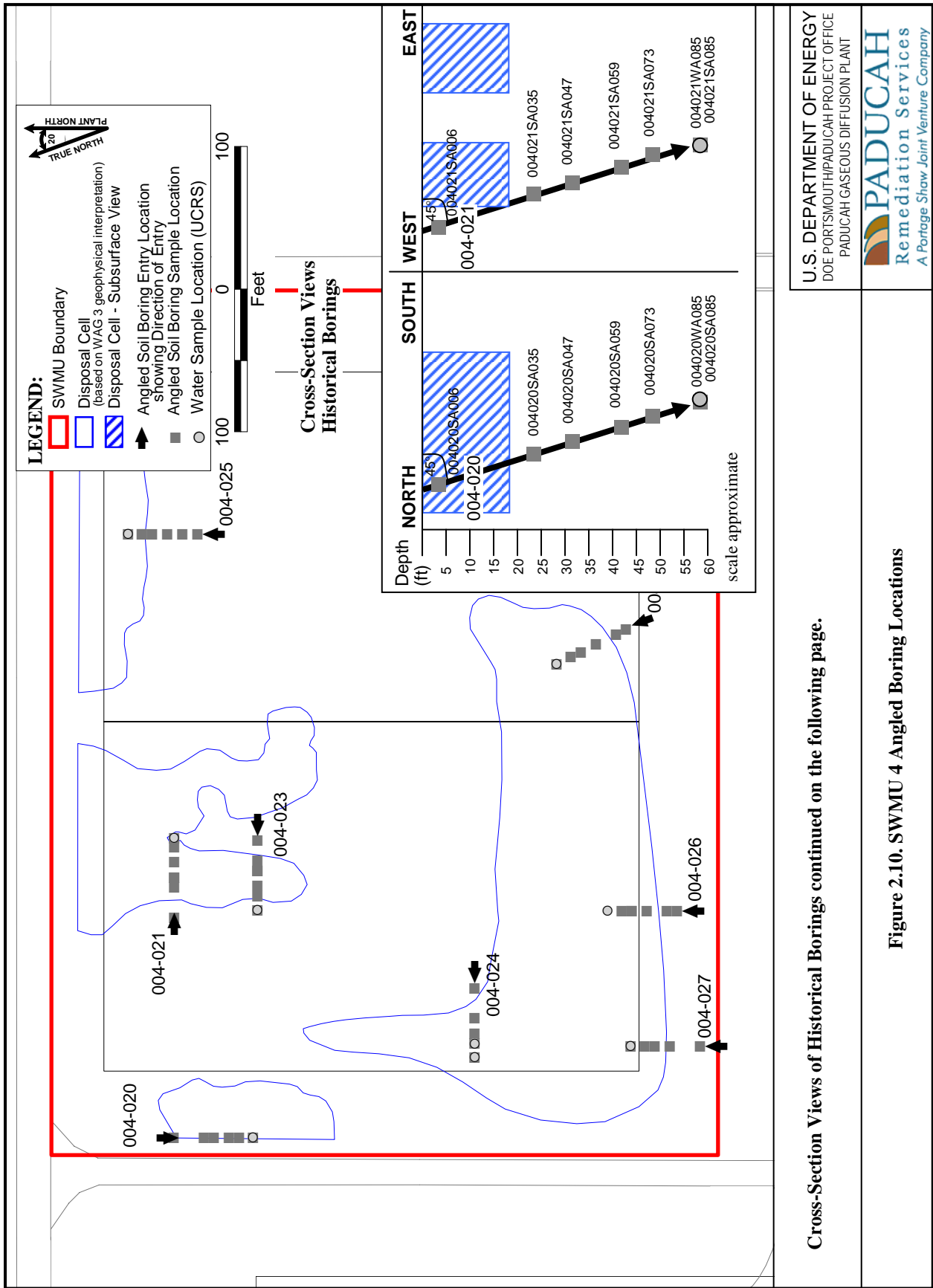


Figure 2.9. SWMU 3 Angled Boring Locations

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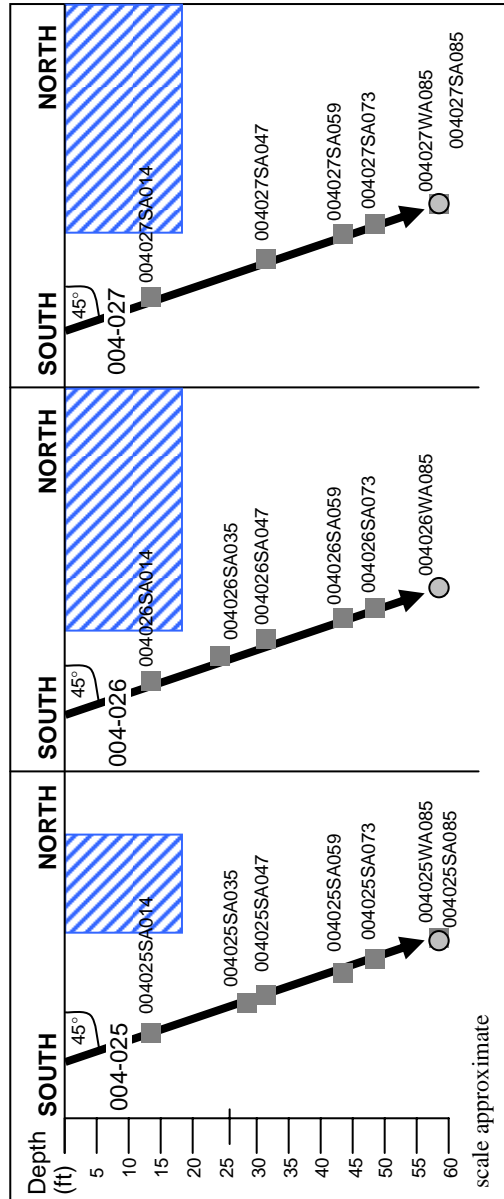
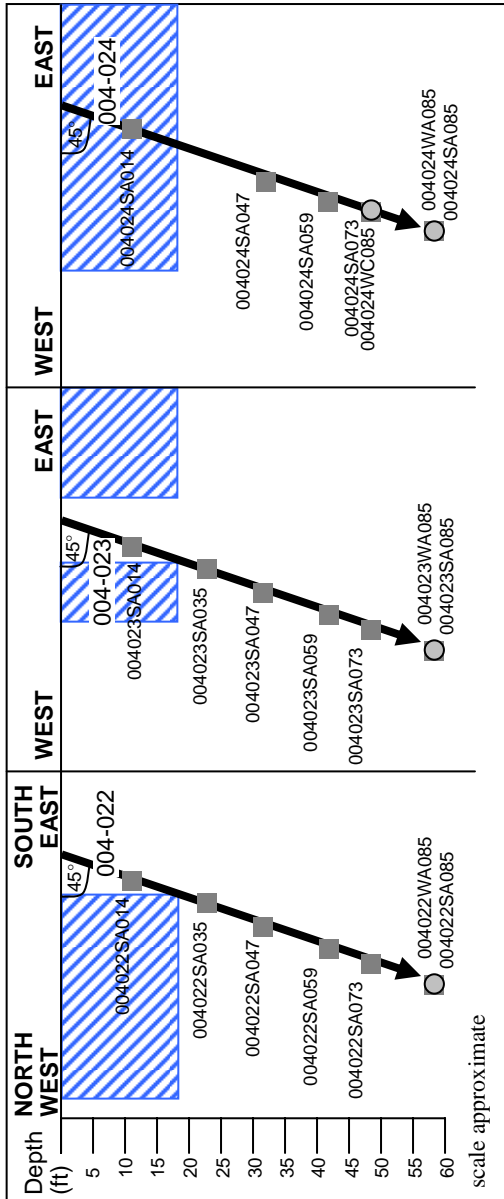
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Cross-Section Views of Historical Borings continued on the following page.

Figure 2.10. SWMU 4 Angled Boring Locations

**Cross-Section Views
Historical Borings**



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Figure 2.10. SWMU 4 Angled Boring Locations (Continued)

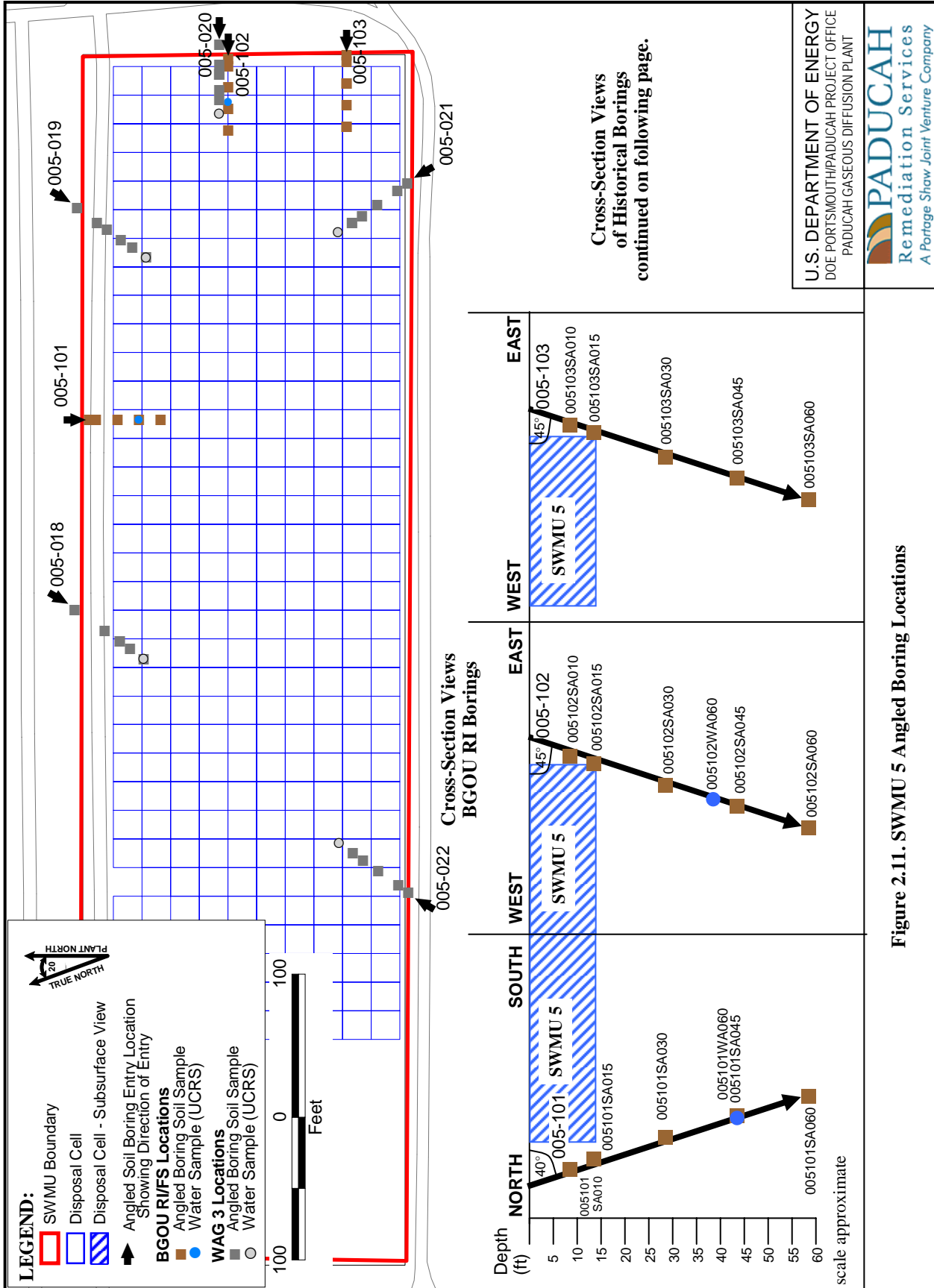
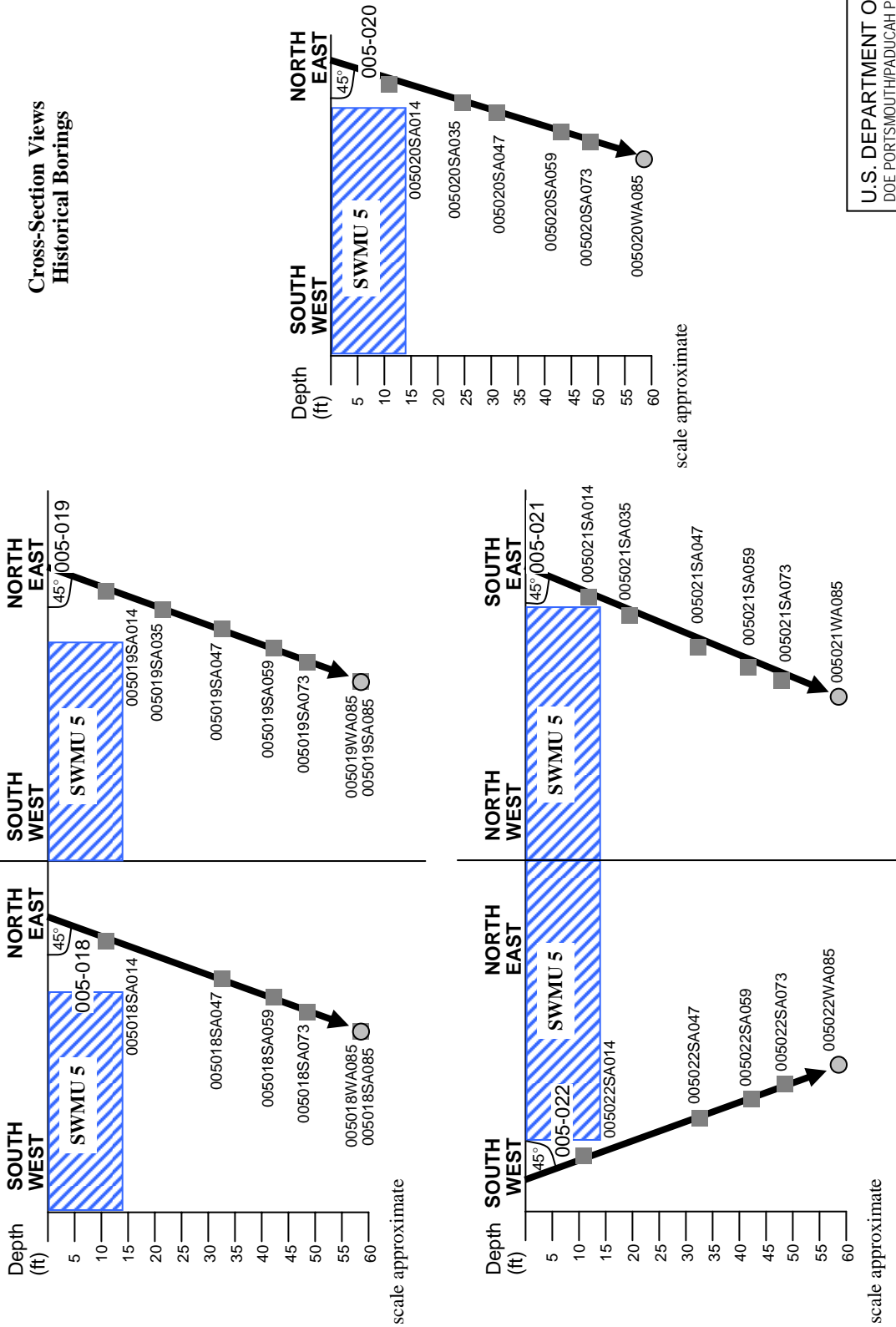


Figure 2.11. SWMU 5 Angled Boring Locations

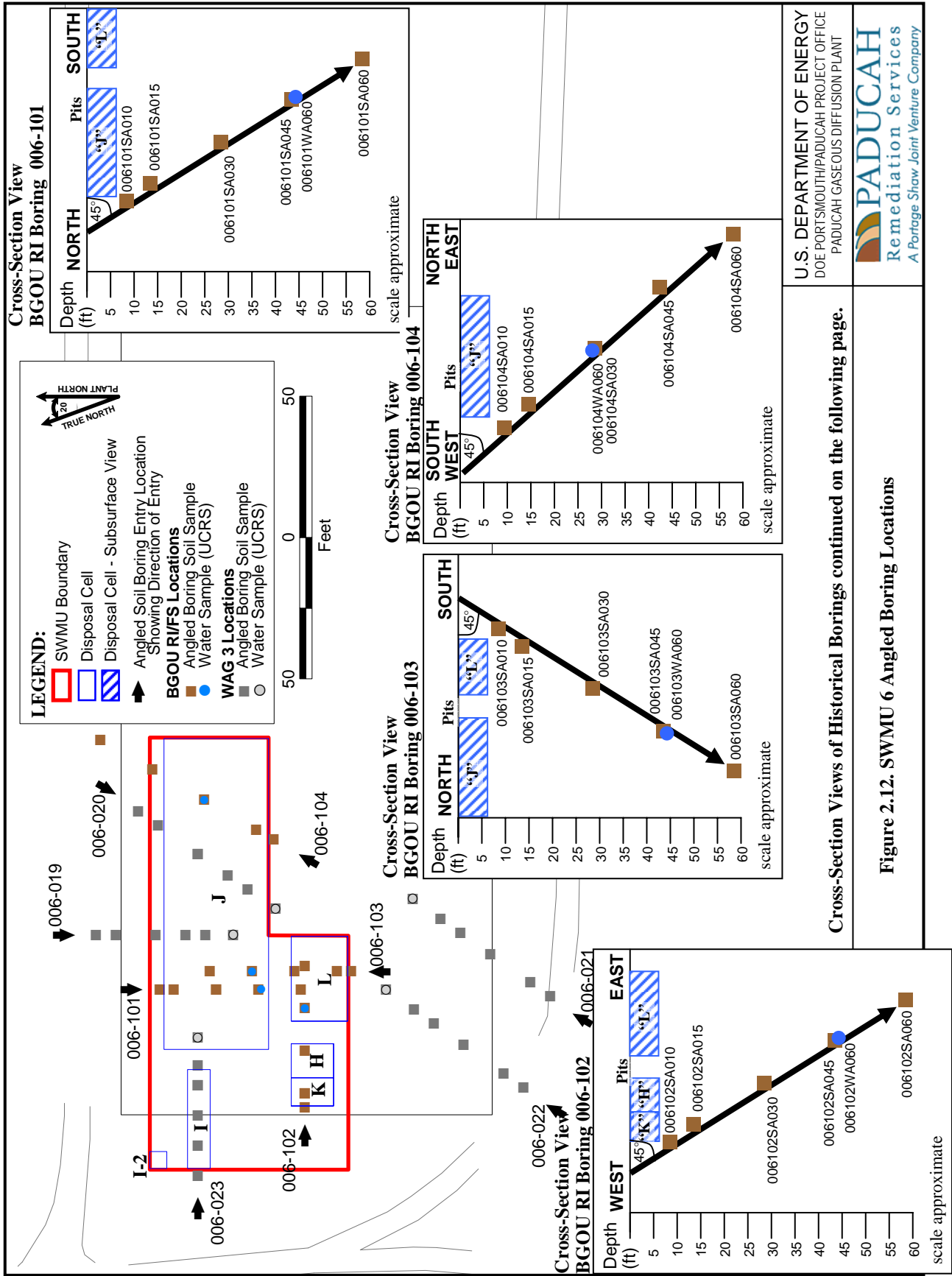
**Cross-Section Views
Historical Borings**



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Figure 2.11. SWMU 5 Angled Boring Locations (Continued)



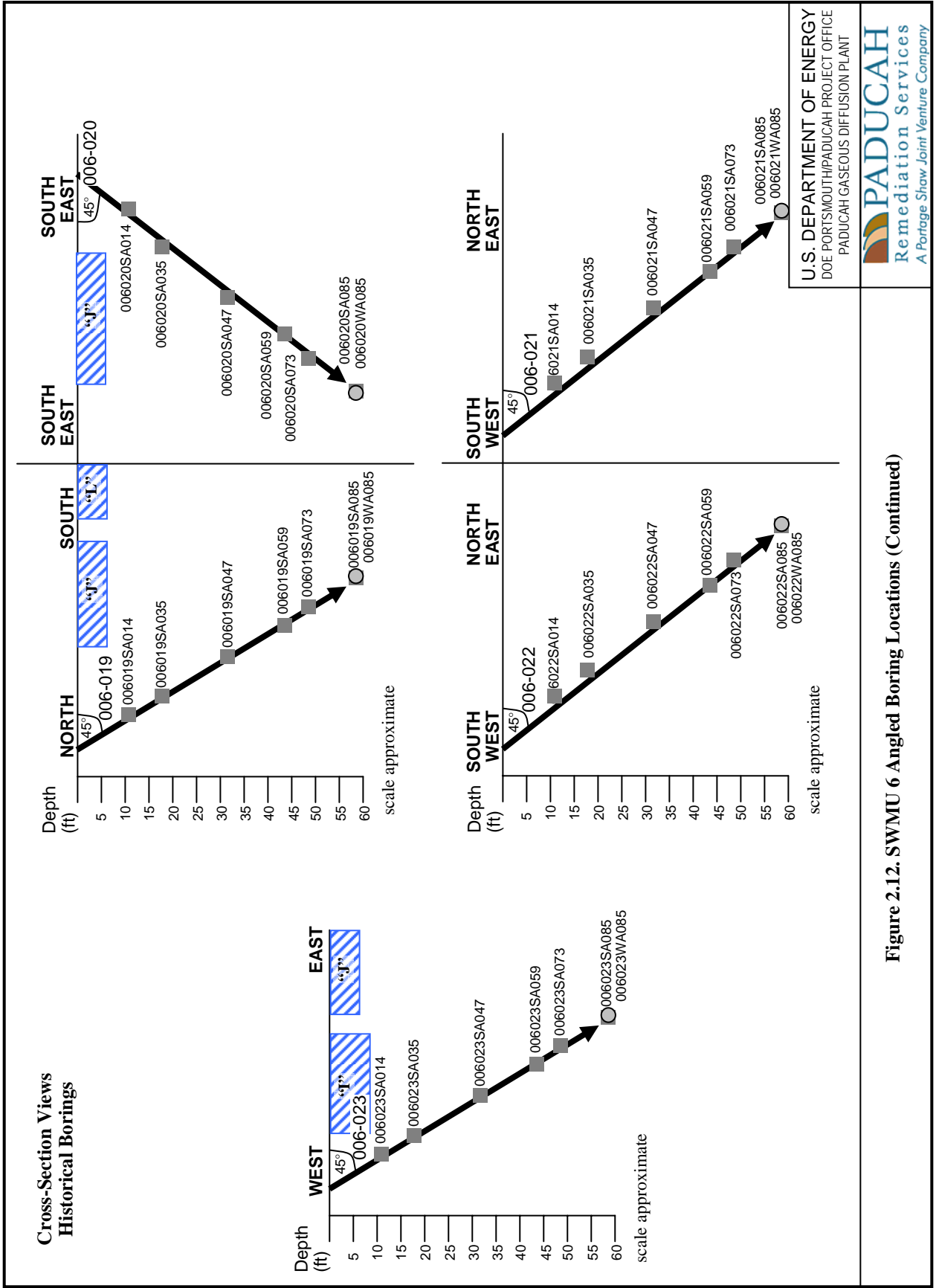


Figure 2.12. SWMU 6 Angled Boring Locations (Continued)

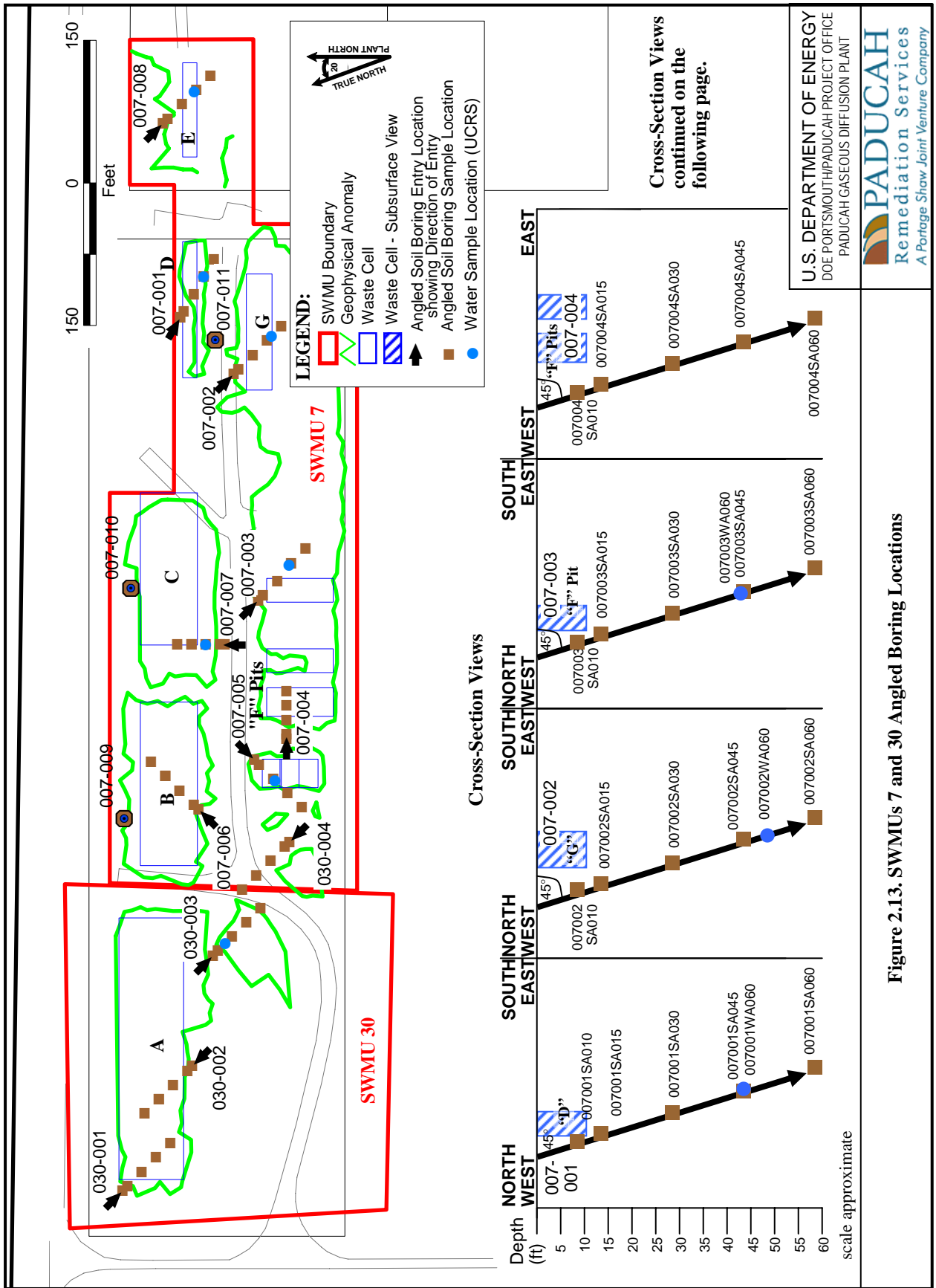
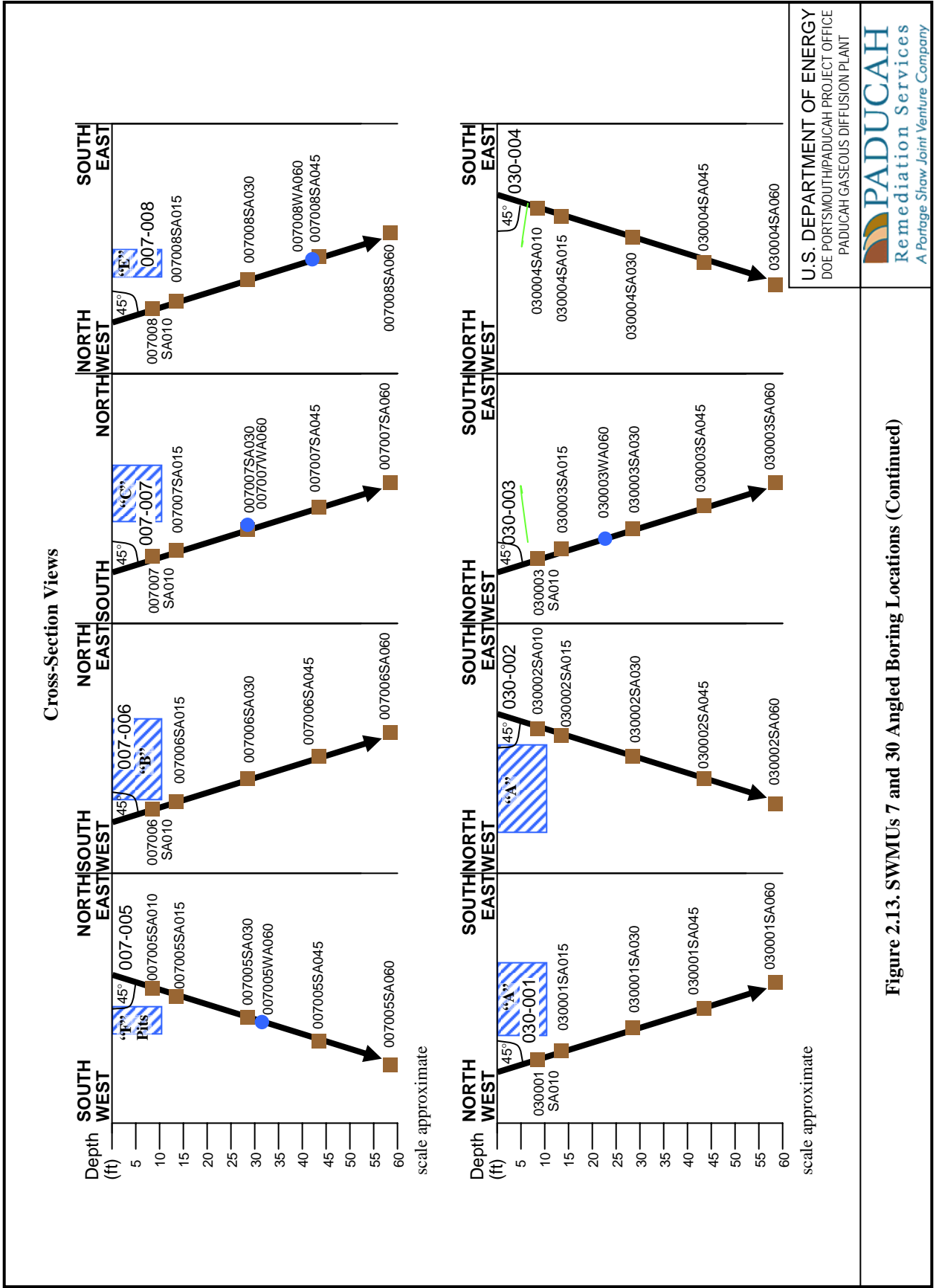


Figure 2.13. SWMUs 7 and 30 Angled Boring Locations



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Figure 2.13. SWMUs 7 and 30 Angled Boring Locations (Continued)

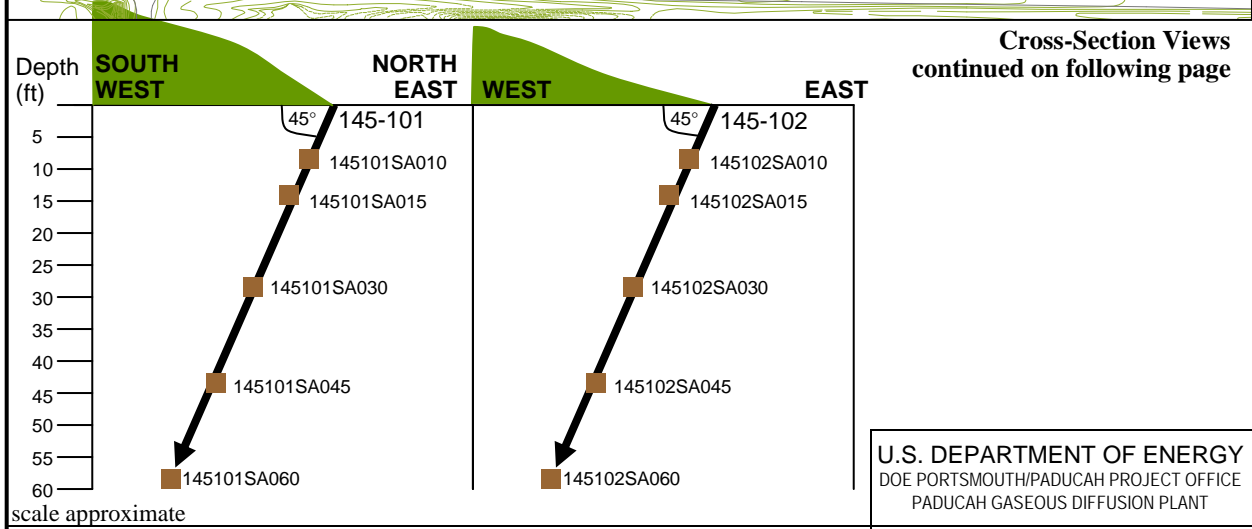
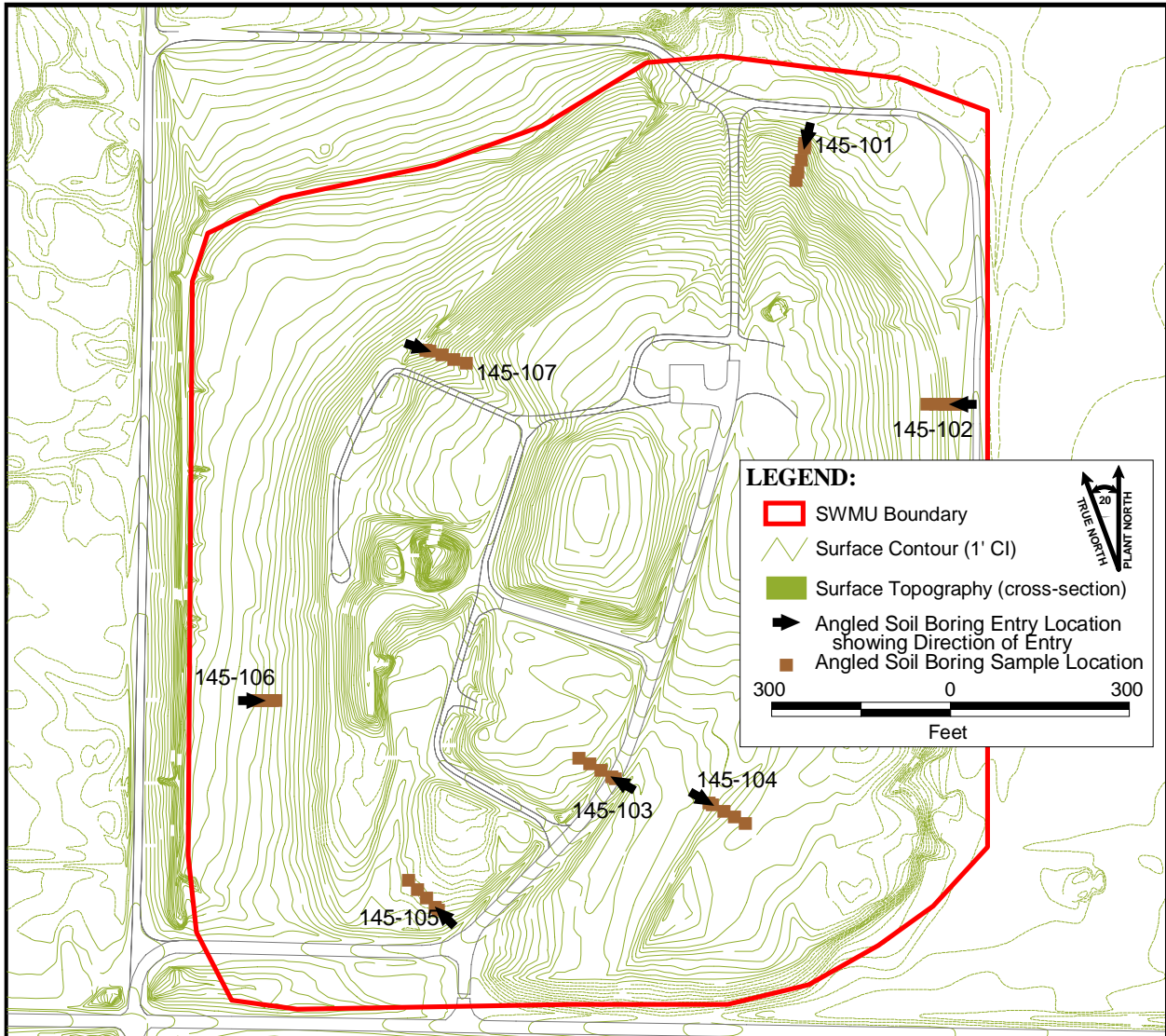
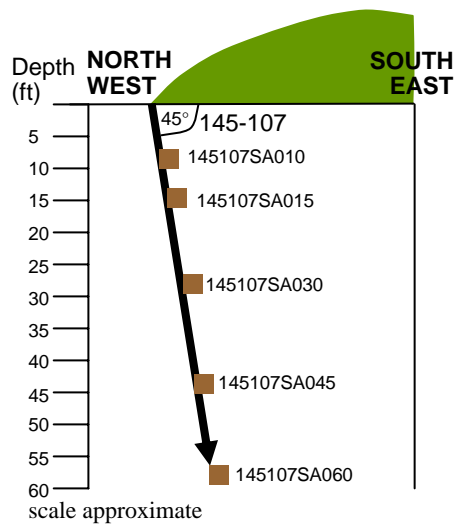
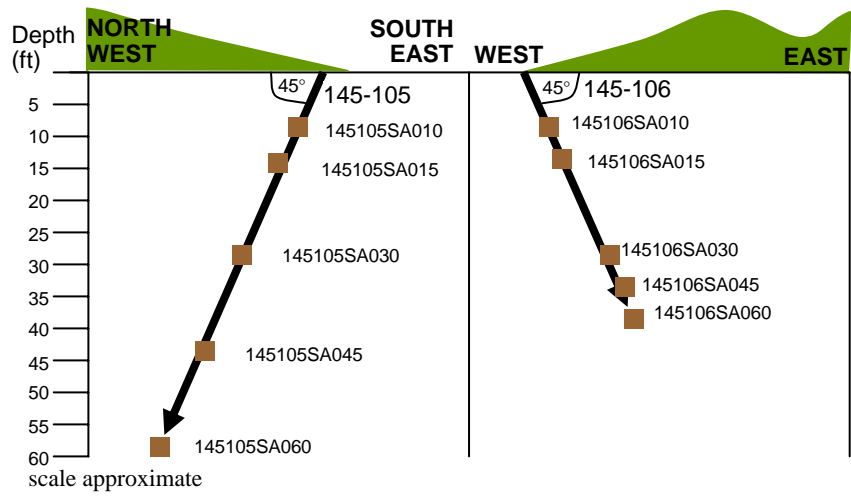
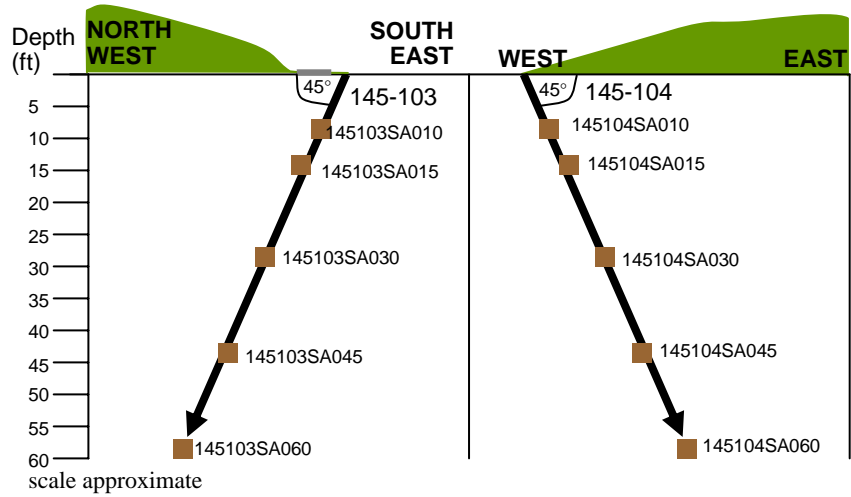


Figure 2.14. SWMU 145 Angled Boring Locations



Cross-Section Views



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Figure 2.14. SWMU 145 Angled Boring Locations (Continued)



The locations of the angled soil borings and deep vertical soil borings were determined in order to avoid drilling into any burial cells. Set-back calculations, the use of geophysics, and historical process knowledge were utilized to determine pit boundaries and depths. During the drilling of Boring 104 at SWMU 145, the field effort was stopped when a safety meter detected a gas coming from the drill stem near the lower explosive limit. According to the recollection of a landfill operator who had worked at the site for several years, roofing material had been disposed of and covered with soil in the area, though not expected to be the source of the gas. Gas samples were collected and analyzed immediately by the USEC laboratory. The gas was determined to be methane. It is believed the methane was migrating from buried material in the area. The borehole was allowed to vent and sampling was completed during the next two weeks.

2.3 SOIL INVESTIGATIONS

Subsurface soil samples from the angled borings were collected generally from 7 to 11 ft, 11 to 14 ft, 28 to 32 ft, 42 to 46 ft, and 57 to 60 ft bgs (a total of five sets of soil samples per boring) in order to effectively identify probable and potential contaminant migration and exposure pathways, as directed by the BGOU Work Plan (DOE 2006a). Soil samples were not collected at or near the surface in angled borings because these borings were installed at a given distance from the burial cell, outside the influence of a burial pit. Locations of these soil samples, relative to their surface penetration, are shown in Figures 2.8 through 2.14. Table 2.1 summarizes soil sampling and analysis from the BGOU RI. Appendix C provides the soil and groundwater analytical results in a searchable database on compact disk.

In addition to the angled borings, subsurface soil samples were collected from both shallow and deep vertical borings (see Figures 2.8 through 2.14). Ten shallow borings were installed along a former drainage ditch that connected the C-404 Landfill and the NSDD. Samples from these borings were collected at the surface and from 1 to 5 ft, 5 to 10 ft, and 10 to 15 ft. Three deep vertical borings were installed within SWMU 7. Samples from these borings were collected at the surface and from 3 to 5 ft, 8 to 10 ft, 13 to 15 ft, 28 to 30 ft, 43 to 45 ft, and 58 to 60 ft.

Table 2.1. Summary of BGOU RI Soil Sampling and Analysis

Location	Activity	Number of Borings	Sampling Interval (ft bgs) ^a	Analyses per Sampling Interval	Sampling Rationale
SWMU 2	Angled Borings	2	7-11 11-14 28-32 42-46 57-60	Metals PCBs Radionuclides VOCs	Sampling intended to characterize soils beneath typical waste cell.
SWMU 3	Angled Borings	4	7-11 11-14 28-32 42-46 57-60	Metals PCBs Radionuclides VOCs	Sampling intended to characterize soils beneath waste cell.
	Shallow Vertical Borings	10	0-1 1-5 5-10 10-15	Metals PCBs Radionuclides	Sampling intended to characterize soils along former discharge ditch.

Table 2.1. Summary of BGOU RI Soil Sampling and Analysis (Continued)

Location	Activity	Number of Borings	Sampling Interval (ft bgs) ^a	Analyses per Sampling Interval	Sampling Rationale
SWMU 5	Angled Borings	3	7-11 11-14 28-32 42-46 57-60	Metals PCBs Radionuclides	Sampling intended to characterize soils beneath typical waste cell.
SWMU 6	Angled Borings	4	7-11 11-14 28-32 42-46 57-60	Metals PCBs Radionuclides	Sampling intended to characterize soils beneath typical waste cell.
SWMU 7	Angled Borings	8	7-11 11-14 28-32 42-46 57-60	Metals PCBs Radionuclides SVOCs VOCs	Sampling intended to characterize soils beneath geophysics-defined waste cells.
	Deep Vertical Borings	3	0-1 3-5 8-10 13-15 28-30 43-45 58-60	Metals PCBs Radionuclides SVOCs VOCs	Sampling intended to characterize soils downgradient of typical waste cell.
SWMU 30	Angled Borings	4	7-11 11-14 28-32 42-46 57-60	Metals PCBs Radionuclides SVOCs VOCs	Sampling intended to characterize soils beneath geophysics-defined waste cells.
SWMU 145	Angled Borings	7	7-11 11-14 28-32 42-46 57-60	Metals PCBs Radionuclides VOCs	Sampling intended to characterize soils beneath geophysics-defined waste cell boundary and areas of disturbance defined in historical photos.

^a Sampling Interval reported in vertical depth.

PCB = polychlorinated biphenyl

SVOC = semivolatile organic compound

VOC = volatile organic compound

2.4 GROUNDWATER INVESTIGATIONS

Collection of an UCRS groundwater sample was attempted for each angled boring. Of the 32 attempts, 18 boring locations provided enough groundwater to collect a sample. Locations of these samples are shown in Figures 2.8 through 2.14.

RGA groundwater samples in addition to UCRS groundwater samples were collected from the deep vertical borings. Generally, UCRS samples were collected from 30 to 45 ft bgs; while RGA samples were collected at 10 ft intervals beginning at 60 ft bgs to the base of the RGA. RGA groundwater samples were collected only at SWMU 7 during this RI. Table 2.2 summarizes groundwater sampling and analysis from the BGOU RI. In the field, the BGOU RI collected only unfiltered water samples. Water samples

collected for metals analysis, however, were filtered in the laboratory prior to preserving the samples with acid. Samples collected in the field for “total” or unfiltered analyses were preserved in acidified bottles while in the field.

Table 2.2. Summary of BGOU RI Groundwater Sampling and Analysis

Location	Activity	Boring	Sampling Interval (ft bgs) ^a	Analyses per Sampling Interval	Comments
SWMU 2	Angled Borings	002-001 002-002	None collected None collected	Metals PCBs, VOCs Radionuclides	Groundwater not present in sufficient quantity for samples.
SWMU 3	Angled Borings	003-001 003-002 003-003 003-004	None collected None collected 28 30	Metals PCBs Radionuclides VOCs	Groundwater present in two of four borings in sufficient quantity for samples.
SWMU 5	Angled Borings	005-101 005-102 005-103	45 40 None collected	Metals PCBs Radionuclides	Groundwater present in two of three borings in sufficient quantity for samples.
SWMU 6	Angled Borings	006-101 006-102 006-103 006-104	46 45 45 45	Metals PCBs Radionuclides	Groundwater present in all four borings in sufficient quantity for samples.
SWMU 7	Angled Borings	007-001 007-002 007-003 007-004 007-005 007-006 007-007 007-008	45 50 43 None collected 45 None collected 45 43	Metals PCBs Radionuclides SVOCs VOCs	Groundwater present in six of eight borings in sufficient quantity for samples.
	Deep Vertical Borings	007-009 007-010 007-011	50 69 ^b 80 ^b 90 ^b 45 60 ^b 66 ^b 80 ^b 90 ^b 100 ^c 45 60 ^b 70 ^b 80 ^b 90 ^b	Metals PCBs Radionuclides SVOCs VOCs	Groundwater samples collected in UCRS and at 10 ft intervals within the RGA.
SWMU 30	Angled Borings	030-001 030-002 030-003 030-004	None collected None collected 23 None collected	Metals PCBs Radionuclides SVOCs, VOCs	Groundwater present in one of four borings in sufficient quantity for samples.
SWMU 145	Angled Borings	145-101 145-102 145-103 145-104 145-105 145-106 145-107	None collected None collected None collected None collected None collected None collected None collected	Metals PCBs Radionuclides VOCs	Groundwater not present in sufficient quantity for samples.

^a Sampling interval reported in vertical depth, not drilled length.

^b RGA

^c Suspected McNairy

PCB = polychlorinated biphenyl; SVOC = semivolatle organic compound; VOC = volatile organic compound

Groundwater-productive intervals are uncommon in the UCRS. The following steps were followed to identify target sample depths and ensure the quality of the groundwater samples.

- (1) Prior to drilling, identify likely depths of saturated, permeable horizons in the UCRS (typically a sand unit or soil unit containing appreciable sand content) from soil boring logs of area boreholes.
- (2) As drilling proceeds, examine soil samples to determine the presence of saturated, permeable soils and monitor indications of water [i.e., drilling progress, drill cuttings (when using augers), and indications of water within the borehole (e.g., wet center rods)] to identify target horizons for UCRS groundwater samples.
- (3) Upon drilling into a target groundwater-producing horizon in the UCRS, pull back the drill string several ft to expose the walls of the borehole and allow groundwater to flow into the open drill string.
- (4) Measure depth to groundwater to determine the presence of water and the rate of rise of water within the drill string.
- (5) If the target soils are water-productive, lower a clean sampling pump within the drill string. Position the sampling pump at least five ft above the base of the drill string, if possible, to minimize the potential of “sand locking” the pump inside the drill string.¹
- (6) Purge up to two to three gal² of water, as necessary, to reduce the turbidity of the discharge water.
- (7) Upon completion of the purge of initial, turbid water, route the discharge water through a flow-through cell equipped with a water quality monitor and document the stability of water quality criteria over time.
- (8) Upon documenting stable water quality in the discharge stream, route the discharge stream through a sampling port, isolating the water in the sample stream from the flow-through cell.
- (9) Collect the water sample. Because the discharge water often remained turbid and the analytical laboratory was not able to analyze volatile organic and semivolatile organic samples containing excessive turbidity, the field crew often filled bottles for these analyses last in an attempt to collect water samples with less turbidity.
- (10) After collecting the UCRS groundwater sample and recovery of the sample pump, measure the depth to water in the borehole and then resume drilling and collection of soil samples. For the three vertical boreholes at SWMU 7, continue the borehole into the RGA, collecting groundwater samples at 10 ft depth intervals.

Comparison of Whole Water and Filtered Water Samples

The BGOU RI collected only whole water (unfiltered) groundwater samples in the field. To support an assessment of the inorganic contaminants that may be migrating through groundwater, as required in the project work plan, the laboratory split all samples for analyses of metals and the dominant uranium

¹ Field experience revealed that sand settling out of the water column, as the soil borings were pumped, often accumulated in the base of the drill string. Later attempts to recover the pump could wedge the pump inside the drill string.

² For most soil borings that did not produce clear water, two to three gal of initial purge water was adequate to assess the potential of producing a better quality water sample.

isotopes into whole water and filtered water (0.45 µ filter) samples. The filtered water sample analyses are more representative of the inorganic dissolved species and colloids that are migrating through groundwater. Unless otherwise designated, all results in Section 4 tables are unfiltered analyses (e.g., a filtered analysis result is designated “Aluminum, dissolved”).

With the exception of all but the shallowest groundwater sample from locations 007-010 and 007-011, the BGOU RI collected all of the groundwater samples from the UCRS. Per the work plan, the BGOU RI collected RGA samples from 007-010 (5 deepest samples) and 007-011 (4 deepest samples).

The laboratory analyzed the groundwater samples for the following 24 metals:

aluminum	calcium	magnesium	silver
antimony	chromium	manganese	sodium
arsenic	cobalt	mercury	thallium
barium	copper	molybdenum	uranium
beryllium	iron	nickel	vanadium
cadmium	lead	selenium	zinc

Three of the metals (calcium, magnesium, and sodium) are essential nutrients and are known to be toxic only at extremely high concentrations. These three metals were not assessed further. Of the remaining 21 metals, 12 of them either were rarely detected or were not present in detectable levels in the filtered water samples, as follows:

aluminum	cadmium	lead	thallium
antimony	chromium	mercury	uranium
beryllium	copper	silver	vanadium

These metals are not being transported in groundwater at levels of human health concern.

The following remaining nine metals were frequently detected in the filtered water samples:

arsenic	iron	nickel
barium	manganese	selenium
cobalt	molybdenum	zinc

These metals are potentially mobile as dissolved species or colloids in groundwater at levels of human health concern.

UCRS: Among the UCRS groundwater samples, sample locations 006-101, 006-103, and 006-104 yielded whole water groundwater samples with the highest levels of arsenic, barium, cobalt, manganese, nickel, and zinc. (Note that 006-101 and 006-103 were excessively turbid, such that the laboratory could not produce a filtered water sample.) These analyses (whole water samples for locations 006-101 and 006-103 and whole and filtered water samples for location 006-104) are biased by the turbidity of the water samples and likely are not representative of area groundwater.

The levels of metals are almost always higher in the whole water samples, reflecting the presence of metals in the suspended fraction of the groundwater samples. Most analyses of these metals in the whole water fraction are less than 10X the metals analyses of the corresponding filtered water samples. These levels of metals in the filtered water samples likely represent naturally occurring conditions. Iron was the most common metal in the whole water analyses to exceed 10X the dissolved water analysis (in 15 of 19 samples). Iron commonly occurs as a coating on the gravels in the UCRS. The higher iron levels in the filtered water samples likely are an artifact of the disturbance of the gravels during drilling.

Of the nine potentially mobile metals, only arsenic, barium, and selenium have maximum contaminant levels (MCLs) for comparison. Only cobalt and zinc do not have an associated Child Resident No Action Level (NAL). The frequency of occurrence above the MCL and NAL for each metal is as follows:

<u>Metal</u>	<u>MCL (mg/L)</u>	<u>% Exceeding MCL</u>	<u>Child Resident NAL</u>	<u>% Exceeding Child Resident NAL</u>
arsenic	1.00×10^{-2}	6 of 20	3.50×10^{-5}	19 of 19 ^a
barium	2.00×10^0	1 of 20	1.04×10^{-1}	16 of 20
cobalt	NA ^b	NA	9.06E-02	1 of 16
iron	NA	NA	4.49×10^{-1}	20 of 20
manganese	NA	NA	3.50×10^{-2}	20 of 20
molybdenum	NA	NA	7.53×10^{-3}	17 of 20
nickel	NA	NA	3.01×10^{-2}	14 of 20
selenium	5.00×10^{-2}	1 of 20	7.54×10^{-3}	8 of 20
zinc	NA	NA	4.50E-01	3 of 15

^a The method detection limit for 1 analysis exceeded the MCL.

^b NA = primary MCL not available or not applicable.

Analyses of uranium-234, uranium-235, and uranium-238 in whole water commonly were approximately 10X the analyses of filtered water except in samples of location 030-003 (filtered water analyses \approx 2X whole water analyses) and uranium-238 analyses of 007-003B and 007-007 (filtered water analyses \approx whole water analyses). The uranium isotope abundance at location 030-003 is markedly higher than the other UCRS sample locations, suggesting the presence of contamination. Uranium-238 levels at locations 007-003B and 007-007 are within the range of levels seen in other UCRS samples.

RGA: The BGOU RI sampled RGA groundwater at locations 007-010 and 007-011. Levels of the metals barium, cobalt, iron, nickel, and zinc in the whole water samples of 007-011 are greater than 10X the levels of the metals in the corresponding filtered water samples and are elevated compared to the metals levels in the whole water samples of 007-010. This relationship suggests that metals are present in elevated levels in the RGA matrix (disturbed during drilling) at location 007-011. One potential explanation is that a release of metals contamination previously migrated through the 007-011 location, but no longer is present. The metals either became sorbed to the RGA matrix or precipitated within the aquifer.

The analyses compare to MCLs and NALs as follows:

<u>Metal</u>	<u>MCL (mg/L)</u>	<u>% Exceeding MCL</u>	<u>Child Resident NAL</u>	<u>% Exceeding Child Resident NAL</u>
arsenic	1.00×10^{-2}	3 of 9	3.50×10^{-5}	9 of 9
barium	2.00×10^0	None	1.04×10^{-1}	7 of 9
cobalt	NA ^a	NA	9.06E-02	2 of 24
iron	NA	NA	4.49×10^{-1}	7 of 9
manganese	NA	NA	3.50×10^{-2}	9 of 9
molybdenum	NA	NA	7.53×10^{-3}	3 of 9
nickel	NA	NA	3.01×10^{-2}	3 of 7 ^b
selenium	5.00×10^{-2}	None	7.54×10^{-3}	4 of 9
zinc	NA	NA	4.50E-01	2 of 23

^a NA = primary MCL not available or not applicable.

^b The method detection limit for 2 analyses exceeded the MCL.

The lone exceedances of an MCL (3 of 9 analyses for barium exceeded the MCL) and the NAL exceedances for molybdenum and nickel derive from samples of location 007-010. Analyses of location 007-011 account for 3 of the 4 NAL exceedances for selenium.

2.5 DEVIATIONS FROM ORIGINALLY PLANNED SAMPLE LOCATIONS

To deal with uncertainties identified in the BGOU, the observational approach was used in the design of the sampling strategy for the BGOU RI/FS. Site conditions and results of the geophysical investigation necessitated movement of some of the RI borings from their originally planned location. Movement of these locations was communicated among the parties and agreed to beforehand. All boring relocations were successful from the standpoint that no unexpected materials were encountered during drilling.

2.5.1 SWMU 2

Site conditions, results of a geophysical investigation, and a historical records search necessitated moving final placement of both SWMU 2 borings.

The angled boring 002-002 was formerly proposed at the center, northern edge of the unit. The boring was moved in order to place it north of burial areas (see Figure 2.8), which reportedly contain uranium sawdust and shavings from routine C-340 operations and machining operations. Buried material in this area is expected to be representative of other waste buried at C-749 (DOE 1995b).

The angled boring 002-001 initially was proposed to be moved slightly south to avoid penetrating the abandoned electrical conduit; however, the setup location for the drilling was too close to the cylinder yard to the west of SWMU 2. Historical sampling of the waste itself is available in this area [boring location SWMU2-12 of the SWMU 2 Interim Remedial Design Investigation (DOE 1997a)]; therefore, 002-001 was moved to the southern edge of the unit, angling to the north.

2.5.2 SWMU 3

Movement of planned boring locations to their final placement varied only slightly to allow for set-back to avoid penetrating the C-404 cap or the bottom of the burial cell.

2.5.3 SWMU 5

Final locations of these borings were moved only slightly in consultation with KEEC personnel. Angled boring 005-101 was relocated to the east to split the difference between historical WAG 3 borings 005-018 and 005-019. Angled boring 005-102 was relocated to the eastern boundary from the northern boundary and angled to the west to target a specific burial pit found on engineering drawings. The angle of angled boring 005-103 was changed from southwest to due west in order to drill in a perpendicular orientation to the adjacent waste cell.

2.5.4 SWMU 6

Site conditions and results of the geophysical investigation necessitated movement of two borings as described below.

The originally planned location for Boring 006-101 was adjacent to Burial Area I, angling to the east, beneath Burial Area J. Area I is reported to contain exhaust fans contaminated with perchloric acid buried

in 1966. Since perchloric acid presents a serious explosion hazard, “Danger” signs had been placed at the edges of the burial pit. In order to avoid disturbance of the acid, 006-101 was relocated to the north, angling to the south under the western third of Area J.

Boring 006-102 was adjusted southward slightly to better intersect the center of Burial Areas K, H, and L. In the area of the available geophysical survey, the location of the burial areas as indicated on the engineering drawing compared favorably to the areas delineated by geophysics in the field.

2.5.5 SWMUs 7 and 30

Site conditions and results of the geophysical investigation necessitated moving final placement of several of these borings.

Boring 030-001 was impossible to complete as originally planned under the current site conditions. The ditch to the north of the planned boring location is wide and filled with water. Between the ditch and the “Pit A” is a silt fence placed as part of an interim corrective measure, preventing the drill rig from setting up in the planned location. Boring 030-001 was moved approximately 62 ft west and angled under the pit to the southeast.

Although the previous geophysical survey did not indicate buried items in the originally planned location, the current survey showed Boring 030-002 within the boundary of buried material. Boring 030-002 was moved approximately 50 ft south and 70 ft west. The revised location was adequate to provide equivalent information regarding potential contamination leaching from Pit A.

Borings 007-011 (45 ft north and 15 ft west) and 007-002 (10 ft east and 60 ft north) were moved from the originally planned location because the previous geophysical survey did not indicate the apparent large burial area, connecting the F Pit area and Pit G. When the previous geophysical survey took place, SWMU 12 (Drum Mountain) had not been removed and electromagnetic survey near the area was not possible. The new location provided equivalent information regarding potential contamination migrating from Pit G.

Boring 007-003-ASB (angled soil boring) was moved 50 ft north. The previous geophysical survey did not indicate the apparent large burial area, connecting the F Pit area and Pit G. When the previous geophysical survey took place, SWMU 12 (Drum Mountain) had not been removed and electromagnetic survey near the area was not possible. The new location provided equivalent information regarding potential contamination leaching from the F Pits.

Boring 007-005-ASB was moved 25 ft east and 80 ft north to the opposite side of the pits, allowing the boring to be closer to the geophysical anomalies.

2.5.6 SWMU 145

Sampling locations for SWMU 145 were better defined in a revision to the Work Plan (DOE 2006b) issued in November 2006. Historical aerial photographs from 1959, 1964, 1975, and 1981 were used to locate soil borings in areas that appeared to be waste disposal locations. Site conditions and results of the geophysical investigation necessitated movement of these borings’ final placement only slightly.

Boring 145-101 encountered a shallower than expected RGA. (The planned samples for SWMU 145 were limited to UCRS soil and groundwater.) Adjustments were made to soil sample intervals at 145-106 and 145-107 to account for the shallow RGA at this location.

2.6 QUALITY ASSURANCE/QUALITY CONTROL

QC was monitored throughout the RI process. QC included field sampling, laboratory analysis, and data management.

2.6.1 Field QC

Field QC samples were collected to assess data quality. Appendix C provides the data from the field QC samples in a searchable database on compact disk. Table 2.3 lists the QC samples collected for each SWMU. The target frequency of collection for QC samples for the entire project was 1 in 20 for equipment rinseates, field blanks, and field duplicates. Overall, this target was met for the project. Trip blanks were collected at a frequency of one per sample cooler containing volatile organic compound (VOC) samples.

Table 2.3. Summary of BGOU RI QC Sampling

Location	QC Sample Type	Frequency of Collection ^a
SWMU 2	Equipment Rinseates	1/10
	Trip Blanks	4/10
	Field Blanks	1/10
	Field Duplicates	1/10
SWMU 3	Equipment Rinseates	3/46
	Trip Blanks	10/46
	Field Blanks	3/46
	Field Duplicates	2/46
SWMU 5	Equipment Rinseates	1/18
	Trip Blanks	N/A
	Field Blanks	1/18
	Field Duplicates	1/18
SWMU 6	Equipment Rinseates	2/24
	Trip Blanks	N/A
	Field Blanks	2/24
	Field Duplicates	2/24
SWMU 7	Equipment Rinseates	4/82
	Trip Blanks	24/82
	Field Blanks	4/82
	Field Duplicates	5/82
SWMU 30	Equipment Rinseates	1/21
	Trip Blanks	5/21
	Field Blanks	1/21
	Field Duplicates	1/21
SWMU 145	Equipment Rinseates	2/35
	Trip Blanks	10/35
	Field Blanks	2/35
	Field Duplicates	2/35
BGOU Summary	Equipment Rinseates	14/236
	Trip Blanks	53/194
	Field Blanks	14/236
	Field Duplicates	14/236

N/A = not applicable (no VOCs collected)

^a Frequency of collection is the number of QA samples collected per number of regular samples collected.

2.6.2 Laboratory QC

The USEC Paducah laboratory performed all of the laboratory analyses of soil and groundwater samples for the BGOU RI. The laboratory was contracted through the DOE Sample Management Office (SMO) and is DOE-approved and Nuclear Regulatory Commission licensed. The USEC laboratory is a DOE approved laboratory audited annually for compliance with requirements. Approved SW-846 methods were used for all samples, except those parameters for which other methods are necessary. The analysis followed SW-846 protocols, and Level C and Level D data packages were provided along with electronic data deliverables (EDDs). Filtered and unfiltered analyses were performed on metals and uranium isotopes. All other analyses were performed using unfiltered samples.

The following data qualifiers were used for reporting fixed-base laboratory results:

Inorganic Analysis

- B This flag is used when the analyte is found in the associated blank as well as in the sample.
- U The analyte was analyzed for, but not detected.
- J Indicates an estimated value.
- E The reported value is estimated because of the presence of interference. An explanatory note must be included under comments on the cover page (if the problem applies to all samples) or on the specific Form I (if it is an isolated problem).
- M Duplicate injection precision was not met.
- N Spiked sample recovery was not within control limits.
- S The reported value was determined by the method of standard additions (MSA).
- W Postdigestion spike for furnace atomic absorption analysis is out of control limits (85-115%), while sample absorbance is less than 50% of spike absorbance.
- X Other specific flags may be required to properly define the results.
- * Duplicate analysis was not within control limits.
- + Correlation coefficient for the MSA is less than 0.995.

Organic Analysis

- U Indicates compound was analyzed for, but not detected.
- J Indicates an estimated value. This flag is used under the following circumstances: (1) when estimating a concentration for tentatively identified compounds where a 1:1 response is assumed and (2) when the mass spectral and retention time data indicate the presence of a compound that meets the pesticide/PCB identification criteria, and the result is less than the contract-required quantitation limit, but greater than zero.
- P This flag is used for a pesticide/PCB target analyte when there is greater than 25% difference for detected concentrations between the two gas chromatograph (GC) columns.
- C This flag applies to pesticide results where the identification has been confirmed by gas chromatograph/mass spectrometer (GC/MS).
- B This flag is used when the analyte is found in the associated blank as well as in the sample.
- E This flag identifies compounds whose concentrations exceed the calibration range of the GC/MS instrument for that specific analysis.
- D This flag identifies all compounds identified in an analysis at a secondary dilution factor.
- X Other specific flags may be required to properly define the results.
- Y Indicates matrix spike (MS)/matrix spike duplicate (MSD) recovery and/or relative percent difference (RPD) failed to meet acceptance criteria.

Radionuclide Analysis

- B Method blank not statistically different from sample at 95% level of confidence.
- D Sample is statistically different from duplicate at 95% level of confidence.

- L Expected and measured value for laboratory control sample (LCS) is statistically different at 95% level of confidence.
- M Expected and measured value for MS is statistically different at 95% level of confidence.
- T Tracer recovery is < 20% or > 105%.
- U Indicates compound was analyzed for, but not detected.
- X Other specific flags may be required to properly define the results.

Precision, accuracy, and completeness objectives were presented in Table 11.3 of the BGOU RI Work Plan (DOE 2006a). An assessment of these objectives for laboratory analytical data was performed. The results of this assessment are provided in Table 2.4.

Precision refers to the level of agreement among repeated measurements of the same characteristic, usually under a given set of conditions. To determine the precision of the laboratory analysis, a routine program of replicate analyses is performed. The absolute difference between the two values calculated is referred to as the RPD. Precision was determined for this RI by reviewing laboratory-applied qualifiers that pertain to laboratory duplicates (i.e., “M” and “*” for inorganic analyses, “Y” for organic analyses, and “D” for radionuclide analyses) over all analyses. QA objectives for precision given in the RI Work Plan are performance based, with RPDs that ranged from 13 to 50%. These objectives were met by the data collected during this RI.

Table 2.4. QA Assessment for Laboratory Measurements of RI Data

Parameter	Method	Matrix	Precision	Accuracy	Completeness ^c
TCL volatiles	SW-846 ^a 8260	Soil	RI Data: 99%	99%	99% ^d
		Water	RI Data: 99%	99%	100% ^d
TCL semivolatiles	SW-846 8270	Soil	RI Data: 99%	99%	99%
		Water	RI Data: 100%	100%	100%
TAL metals	SW-846 6010, 6020, and 7000 series	Soil	RI Data: 95%	92%	94% ^e
		Water	RI Data: 94%	81%	97% ^f
TCL PCBs	SW-846 8082	Soil	RI Data: 97%	97%	99%
		Water	RI Data: 100%	100%	67% ^g
Gross alpha	SW-846 9310	Soil	RI Data: 100%	100%	99%
Gross beta	SW-846 9310	Soil	RI Data: 100%	100%	99%
²³⁴ U, ²³⁵ U, and ²³⁸ U	RL-7128 ^b	Soil	RI Data: 100%	100%	99%
		Water	RI Data: 100%	100%	100%
⁹⁹ Tc, ²³⁰ Th, ²³⁹ Pu, ¹³⁷ Cs, and ²³⁷ Np	RL-7100, RL-7124, and RL-7128 ^b	Soil	RI Data: 100%	100%	99%
		Water	RI Data: 96%	88%	93%

^aEPA 1996. *Test Methods for Evaluating Solid Waste, Physical/Chemical Methods*, Third Edition, SW-846, December.

^bLaboratory-specific method, derived from DOE guidance.

^cCompleteness for groundwater samples calculated based on locations where groundwater was available for sampling.

^dCompleteness for acrolein analysis in soil samples was 45% and 77% in groundwater samples.

^eCompleteness for silver analysis in soil samples was 80%.

^fCompleteness for antimony, chromium, and nickel analyses in groundwater samples was 77%, 84%, and 87%, respectively.

^gCompleteness for PCBs in groundwater was less than the 90% objective due to eleven results being rejected by validation. Of the ten results, nine were from SWMU 7 and two were from SWMU 6. Section 2.6.4 contains additional discussion.

TAL = Target Analyte List

TCL = Target Compound List

¹³⁷Cs = cesium-137

²³⁷Np = neptunium-237

²³⁹Pu = plutonium-239

PCB = polychlorinated biphenyl

⁹⁹Tc = technetium-99

²³⁰Th = thorium-230

²³⁴U = uranium-234

²³⁵U = uranium-235

²³⁸U = uranium-238

Accuracy refers to the nearness of a measurement to an accepted reference or true value. To determine the accuracy of an analytical method and/or the laboratory analysis, a periodic program of sample spiking is conducted. Accuracy for this RI was determined by reviewing laboratory-applied qualifiers that pertain to laboratory spikes over all analyses (i.e., “N” and “W” for inorganic analyses; “Y” for organic analyses; and “B,” “M,” and “L” for radionuclide analyses). The accuracy range objective specified in the RI Work Plan was 80–100%. These objectives were met by the data collected during this RI.

Representativeness is the degree to which discrete samples accurately and precisely reflect a characteristic of a population, variations at a sampling location, or an environmental condition. Representativeness is a qualitative parameter and will be achieved through careful, informed selection of sampling sites, drilling sites, drilling depths, and analytical parameters and through the proper collection and handling of samples to avoid interference and minimize contamination and sample loss.

Completeness is a measure of the percentage of valid, viable data obtained from a measurement system compared with the amount expected under normal conditions. The goal of completeness is to generate a sufficient amount of valid data to satisfy project needs. A summary of RI data rejected by validation for each SWMU is shown in Table 2.5.

Table 2.5. Remedial Investigation Data Rejected during Validation

SWMU	Rejected Analyte (number of rejects)	
	Soil	Groundwater
SWMU 2	Acrolein (11)	
SWMU 3	Acrolein (34); Silver (10)	
SWMU 5	Silver (12)	
SWMU 6		Total PCB (2); PCB-1016 (2); PCB-1221 (2); PCB-1232 (2); PCB-1242 (2); PCB-1248 (2); PCB-1254 (2); PCB-1260 (2); PCB-1268 (2); Technetium-99 (2)
SWMU 7	Acrolein (19); Silver (14)	Acrolein (6); Antimony (7); Chromium (5); Mercury (1); Nickel (4); Total PCB (9); PCB-1016 (9); PCB-1221 (9); PCB-1232 (9); PCB-1242 (9); PCB-1248 (9); PCB-1254 (9); PCB-1260 (9); PCB-1268 (9)
SWMU 30		Acrolein (1)
SWMU 145	Acrolein (37); Silver (10); Vanadium (10)	

No data was collected at SWMU 4 during this RI

For this project, the completeness objective for laboratory measurements was 90%. This objective was met as intended by the RI data with the exception of PCBs in groundwater. Completeness for PCBs in groundwater was less than the 90% objective due to 11 PCB results being rejected by validation. Of the 11 results, 9 were from SWMU 7 and two were from SWMU 6. Section 2.6.4 contains additional discussion. These measurements were not used for decision calculations in this RI Report.³

³ UCRS groundwater analyses of the BGOU RI soil borings are used primarily to supplement the assessment of soil analyses to identify significant sources of groundwater contamination. The BGOU RI risk assessment uses only analyses of UCRS groundwater collected from MWs within and adjacent to the SWMUs to calculate a conservative reasonable maximum risk estimate (see Attachment F2 of Appendix F). The low completion rate of UCRS groundwater PCB analyses does not limit significantly the nature and extent and risk assessments for the BGOU RI.

Completeness also is a measure of samples collected during the field effort with respect to those targeted for collection in the work plan. All soil samples targeted for collection during this RI were collected with the exception of one surface soil sample at SWMU 7. Additionally, a sufficient volume of soils was not available from two intended locations for all analyses planned (metals were not analyzed from location 145-104 at 15 ft bgs and radionuclides were not analyzed from location 007-001 at 60 ft bgs). For boring 007-001, the entire sampling interval was not recovered (the sampling interval from 57 to 60 ft stated minimal sample recovered). For boring 145-104, the sample included wood fragments, which limited the amount of soil in the sample.

Groundwater sample objectives were fulfilled. The Work Plan (DOE 2006a) strategy for sampling groundwater in the UCRS was to sample water-bearing zones as they were available. Where sand and gravel zones were encountered in the UCRS that would yield sufficient water for the collection of a quality water sample, the field crew collected water samples for analysis. Of the 35 soil borings of the BGOU RI deeper than 15 ft (those targeted for groundwater sampling in the Work Plan), the RI collected UCRS groundwater samples from 18. Because the UCRS water samples only supplement the characterization of the BGOU SWMUs (the analysis of subsurface soil samples is the primary measure that supports the assessment of nature and extent and risk), the lack of UCRS water samples from all soil borings does not limit the planned assessment of the SWMUs.

For those borings that extended through the RGA, water samples were collected as scheduled in 10-ft depth increments throughout the thickness of the RGA. Table 2.6 summarizes the water samples that were collected and analyzed for the BGOU RI/FS. Of all of the water samples, the only sample to be partially collected was the UCRS sample from soil boring 006-103-ASB. This boring provided insufficient water for the collection of samples for analysis of dissolved metals and dissolved radionuclides levels.

Table 2.6. Summary of BGOU RI Water Samples

Location	Soil Boring	Water Sample Depth Interval (ft)
UCRS Groundwater Samples		
SWMU 2	-- ¹	NA ²
SWMU 3	003-003-ASB	28-32
	003-004-ASB	28-32
SWMU 4	none planned	NA
SWMU 5	005-101-ASB	42-46
	005-102-ASB	40-41
SWMU 6	006-101-ASB	42-46
	006-102-ASB	18-19
	006-103-ASB	50-51
	006-104-ASB	42-46
	007-001-ASB	42-46
	007-002-ASB	50-51
	007-003-ASB	42-46
SWMU 7	007-005-ASB	42-46
	007-007-ASB	42-46 ³
	007-008-ASB	42-46
	007-009-VSB	45-55
	007-010-VSB	40-45 and 55-60
	007-011-VSB	40-45 and 55-60
SWMU 30	030-003-ASB	23
SWMU 145	-- ¹	NA ²

**Table 2.6. Summary of BGOU RI Water Samples
(Continued)**

Location	Soil Boring	Water Sample Depth Interval (ft)
RGA Groundwater Samples		
	007-009-VSB	69, 80, 90 ³
SWMU 7	007-010-VSB	66, 80, 90, 100
	007-011-VSB	70, 80, 90

¹ None of the soil borings at this SWMU yielded UCRS water samples.

² No water sample was collected.

³ A duplicate water sample was collected from this depth interval.

Comparability is the extent to which comparisons among different measurements of the same quantity or quality will yield valid conclusions. Comparability will be assessed in terms of field standard operating procedures (SOPs), analytical methods, QC, and data reporting. In addition, data validation assesses the processes employed by the laboratory that affect data comparability.

Historical data also was evaluated for precision and accuracy as described previously. This assessment was performed over all measurements for the projects associated with the BGOU SWMUs. Multiple laboratories analyzed samples for these projects. The comparison for the precision and accuracy of historical results encompassed the entire historical data set and did not differentiate between projects or laboratories. A summary of this assessment is provided in Table 2.7.

Table 2.7. Assessment for Laboratory Measurements of Historical Data Used in RI

Parameter	Method	Matrix	Precision ^a	Accuracy ^a
TCL volatiles	SW-846 8260	Soil	94%	94%
		Water	99%	99%
TCL semivolatiles	SW-846 8270	Soil	99%	99%
		Water	99%	99%
TAL metals	SW-846 6010, 6020, and 7000 series	Soil	81%	48%
		Water	99%	83%
TCL PCBs	SW-846 8082	Soil	100%	100%
		Water	100%	100%
²³⁴ U, ²³⁵ U, and ²³⁸ U	Various methods	Soil	100%	100%
		Water	100%	100%
⁹⁹ Tc, ²³⁰ Th, ²³⁹ Pu, ¹³⁷ Cs, and ²³⁷ Np	Various methods	Soil	99%	100%
		Water	99%	99%

^a Values indicated are for the percent of valid values.

¹³⁷Cs = cesium-137

²³⁷Np = neptunium-237

²³⁹Pu = plutonium-239

⁹⁹Tc = technetium-99

²³⁰Th = thorium-230

²³⁴U = uranium-234

²³⁵U = uranium-235

²³⁸U = uranium-238

All historical analyses were within the criteria established by the RI Work Plan for RI data, with the exception of accuracy of metals analyses in soil.

2.6.3 Surveillances

A surveillance was completed during fieldwork to verify adherence to project specific plans and procedures. Surveillance results are documented and filed in the Document Management Center. The DOE SMO conducts routine laboratory surveillances of the laboratory through the Consolidated Audit Program. These surveillances of the BGOU laboratory were conducted in February 2006 and February 2007.

2.6.4 Data Management

The BGOU Project Environmental Measurements System (PEMS) was used to manage field-generated data; import laboratory-generated data; add data qualifiers based on data verification, validation, and assessment; and transfer data to the Paducah Oak Ridge Environmental Information System (Paducah OREIS). PEMS included a tracking system to identify, track, and monitor each sample and associated data from point of collection through final data reporting. The system included field measurements, chain-of-custody information, and a tracking system for tracking hard-copy data packages and EDDs. PEMS also included information for field planning and data evaluation.

All data packages and EDDs received from the laboratory were tracked, reviewed, and maintained in a secure environment. When first received, data packages were assigned a document control number and then logged into a tracking system. The following information was tracked: sample delivery group numbers, date received, document control number, number of samples, sample analyses, receipt of EDDs, and comments.

The data verification processes for laboratory data were implemented for both hard-copy data and EDDs. The data packages were reviewed to confirm that all samples had been analyzed for the requested parameters. Discrepancies were reported to the laboratory and the data validators. As part of a series of internal integrity checks within PEMS, a check was run to identify which of the requested samples and analyses were not received in an EDD. Hard copy data packages were checked to confirm agreement with the associated EDD. Integrity checks in PEMS also were used to check the list of compounds generated by the laboratory to confirm that data were provided for all requested analytes. Discrepancies were reported to the laboratories for responses and/or correction and to the data validators.

Data verification within PEMS included standardization of analytical methods, chemical names and units, as well as checks for holding time violations and detections above background values.

PEMS system requirements included backups, security, change control, and interfacing with other data management systems. PEMS was housed on the Paducah network. System backups were performed nightly following standard Paducah network protocol. Updates made to the files were copied to a computer backup tape each night, and an entire backup was performed each week.

Security of PEMS and data used for the data management effort was considered essential to the success of the project. The security protocol followed by the data management team was consistent with that of the Paducah network. Access to the network is password-protected. Access to PEMS was limited, on an as-needed basis, to the data management personnel. Read-write, graded access to PEMS was limited to the data management team, which consisted of the PEMS Coordinator and the supporting data entry staff. The data management staff assisted other project members with data needs from PEMS by running requested queries.

A large volume of data was generated during the BGOU RI. To confirm that the data set could be used in the decision making process, the RI team performed various checks and reviews during and after the fieldwork to maintain data consistency and identify problem areas. These checks and reviews included electronic verification and manual assessments by the RI team, as well as independent validation of fixed-base laboratory data. Approximately 36,000 records were reviewed during the BGOU RI data assessment.

Data validation is a process performed for a data set by a qualified individual independent from sampling, laboratory, project management, and other decision making personnel for the project. Data validation is performed in accordance with EPA guidance. In the data validation process, the laboratory adherence to analytical method requirements is evaluated. Portage Environmental, Inc., validated data collected for this RI at a frequency of 100%.

As part of the data review process, findings were qualified as necessary to reflect data validation results. The following qualifiers were assigned by the data validators:

- U Analyte or compound considered not detected above the reported detection limit.
- J Analyte or compound identified; the associated numerical value is approximated.
- UJ Analyte or compound not detected above the reported detection limit, and the reported detection limit is approximated due to quality deficiency.
- R Result is not usable for its intended purpose, so data are of “information only” quality and should be supplemented with additional data for decision-making.
- = Data were validated; however, no qualifier was added.

The majority of the data rejected by validation was acrolein analyses in soil (101 of the 177 sample results collected) due to initial and continuing calibration relative response factors less than 0.05. Also rejected by validation in soil were silver and vanadium analyses (46 and 10 of 216 sample results, respectively). These analyses were rejected due to the interference check sample, the MS and/or MSD, lab control sample, and post digestion spike recoveries being below the lower control limit and the MS/MSD pair exceeding the RPD limit.

The risk assessment does not identify either acrolein or silver as a COPC for the BGOU RI (nor have they been recognized commonly as site-related contaminants); thus, the rejection of these sample results likely has little importance. In contrast, vanadium is a COPC at several BGOU SWMUs. All soil samples with rejected vanadium analyses were collected from SWMU 145 (Figure 2.15). A significant percentage of the vanadium analyses of subsurface soils at SWMU 145 was rejected (10 of 34). Although the risk assessment does not identify vanadium as a COPC for SWMU 145, some uncertainty remains. There are 576 analyses of vanadium (non-rejected) among the historical data and RI data for the BGOU SWMUs. Comparison of these data with PGDP background values (37 mg/kg for subsurface soil and 38 mg/kg for surface soil) demonstrates that the vanadium is naturally occurring; 94% of the vanadium analyses are equal or less to the PGDP background values. Since most vanadium analyses are less than background, and vanadium exceedances were not concentrated in one area, but detected at five separate SWMUs, the rejected data should have little impact on remedial decisions to be made for SWMU 145.

Analyses of groundwater samples resulted in the following number of rejections: 7 of 26 acrolein sample results; 7 antimony results, 5 chromium results, 1 mercury result, and 4 nickel results of 31 samples; 2 of 31 technetium-99 sample results; and 11 of 31 PCB (total) and congeners sample results. Acrolein was rejected because initial and continuing calibration relative response factors were <0.05. Metals were rejected for recoveries being below the lower control limits on one or more of the following controls: continuing calibration verification, interference check sample, lab control sample, the MS and/or MSD, and/or post digestion spike. Additionally, in some cases, the MS/MSD pair exceeded the RPD limit.

Technetium-99 was rejected for unacceptable laboratory control sample bias, significant difference between the sample and the duplicate, and the MS and/or MSD recovery being below the lower control limit. PCBs were rejected because the decachlorobiphenyl surrogate was recovered at lower than acceptable QC limits and the results were nondetect; however, the tetrachloro-m-xylene surrogate also was used and recovered within acceptable QC limits.

The presence of the metals antimony, chromium, mercury, and nickel largely are naturally occurring in the PGDP groundwater. PCBs typically are not mobile in groundwater through soils with small pores and clay or organic matter. Thus, remedial decisions for the BGOU SWMUs should not be impacted by the rejected metals analyses. Because fate and transport modeling results using soil data, and not UCRS groundwater, were used to assess risk, the rejected metals analyses have no impact in the assessment of leachability for metals in soils related to the SWMUs.

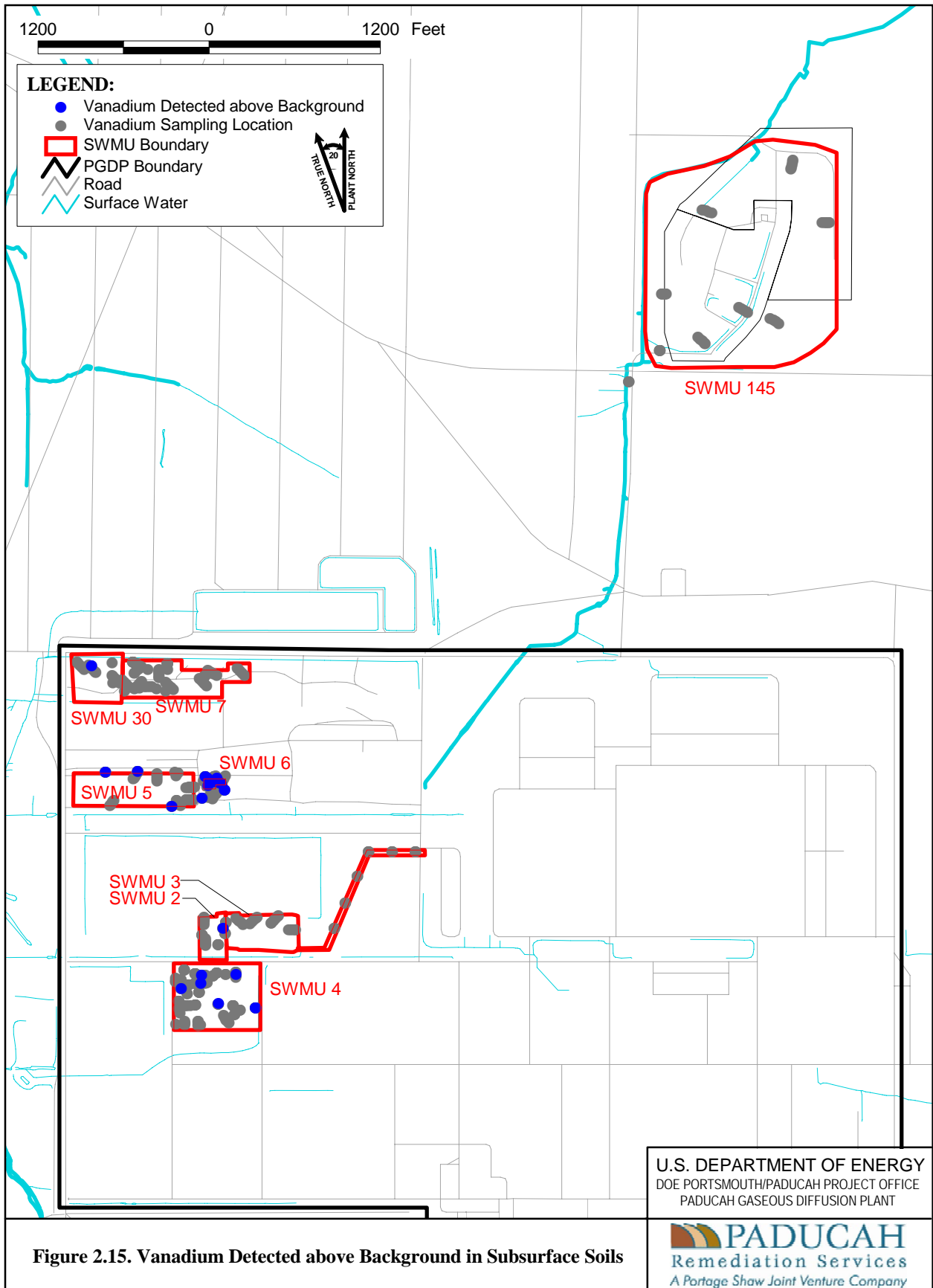


Figure 2.15. Vanadium Detected above Background in Subsurface Soils

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3. PHYSICAL CHARACTERISTICS OF THE STUDY AREA

This chapter presents the physical and ecological characteristics of PGDP, in general, and of the BGOU SWMUs, in particular, that bear on contaminant release and migration. The discussion focuses from region- and PGDP-wide characteristics to SWMU-specific characteristics in sufficient detail to support subsequent evaluations of the nature and extent and the fate and transport of contaminants exiting the SWMUs and entering the external environment.

Numerous investigations detail physical characteristics of PGDP that are pertinent to the BGOU. In addition to the BGOU SWMU investigations identified in Table 1.2, the primary references include the following:

- *Results of the Site Investigation, Phase I, at the Paducah Gaseous Diffusion Plant, Paducah, Kentucky* (CH2M HILL 1991)
- *Results of the Site Investigation, Phase II, at the Paducah Gaseous Diffusion Plant, Paducah, Kentucky* (CH2M HILL 1992)
- *Report of the Paducah Gaseous Diffusion Plant Groundwater Investigation Phase III* (Clausen *et al.* 1992)
- *Environmental Investigations at the Paducah Gaseous Diffusion Plant, and Surrounding Area, McCracken County, Kentucky* (COE 1994)
- *Groundwater Monitoring Plan for the C-746-S Residential Landfill, Paducah Gaseous Diffusion Plant, Paducah, Kentucky* (BJC 2001b)

This RI field effort focused on collection and analysis of soil and groundwater samples to address deficiencies in the existing characterization of the nature and extent of contamination. These field activities yielded additional analyses of the subsurface soils and groundwater that are incorporated into the SWMU-specific discussions. Other than the area of the historical ditch that routed effluent from SWMU 3 to the NSDD and the vertical borehole locations of SWMU 7, the BGOU RI did not include additional characterization of surface soils (see Section 1.2.1).

3.1 SURFACE FEATURES

PGDP is located on a 3,556-acre DOE site approximately 10 miles west of Paducah, Kentucky, and 3.5 miles south of the Ohio River in the western part of McCracken County (Figure 1.1). The PGDP industrial area occupies approximately 650 acres of the DOE site, surrounded by an additional 689-acre buffer zone. DOE licenses most of the remaining acreage to the Commonwealth of Kentucky as part of the West Kentucky Wildlife Management Area (WKWMA). TVA's Shawnee Steam Plant borders the DOE site to the northeast, between the plant and the Ohio River.

Three small communities are situated within three miles of the DOE property boundary: Heath and Grahamville to the east and Kevil to the southwest. The next closest municipality is Metropolis, Illinois, five miles to the northeast of PGDP on the north side of the Ohio River.

The dominant topographic features in the area of PGDP are nearly level to gently sloping dissected plains and the flood plain of the Ohio River. Local elevations range from 290 ft above mean sea level (amsl) along the Ohio River to 450 ft amsl southwest of PGDP. Ground surface elevations vary from 360 to 390 ft amsl within the PGDP plant boundary, where most of the BGOU SWMUs are located, and

from 360 to 410 ft at SWMU 145. Generally, the topography in the PGDP area slopes toward the Ohio River at an approximate gradient of 27 ft per mile (CH2M HILL 1992).

3.2 METEOROLOGY

The National Weather Service office at Barkley Regional Airport (located four miles to the southeast of PGDP) documents hourly meteorological measurements. Current and historical meteorological information regarding temperature, precipitation, and wind speed/direction are available from the National Oceanic and Atmospheric Administration’s National Climatic Data Center.

The climate of the PGDP region is humid-continental. Summers are warm (July averages 79 °F) and winters are moderately cold (January averages 35 °F). PGDP experiences a yearly surplus of precipitation versus evapotranspiration. The 30-year average monthly precipitation for the period 1961 through 1990 is 4.11 inches,¹ varying from an average of 3.00 inches in October (the monthly average low) to an average of 5.01 inches in April (the monthly average high). Monthly estimates of evapotranspiration using the Thornthwaite method (Thornthwaite and Mather 1957) equal or exceed average rainfall for the period May through September (season of no net infiltration).

Heavy rainfall associated with thunderstorms or low-pressure systems occurs occasionally at PGDP. Table 3.1 presents the predicted storm recurrence intervals for PGDP (Hershfield 1963; Johnson *et al.* 1993; DOE 1997a).

Table 3.1. Precipitation as a Function of Recurrence Interval and Storm Duration for the Site

Recurrence Interval (years)	Storm Duration (hours)						
	0.5	1	2	3	6	12	24
	Precipitation (inches)						
1	1.08	1.30	1.66	1.85	2.23	2.65	3.06
2	1.26	1.56	1.91	2.14	2.61	3.08	3.53
5	1.55	1.98	2.38	2.67	3.20	3.69	4.38
10	1.80	2.23	2.75	3.02	3.66	4.33	4.97
25	1.99	2.57	3.13	3.44	4.18	4.83	5.71
50	2.23	2.83	3.46	3.83	4.62	5.53	6.42
100	2.45	3.13	3.83	4.24	5.02	5.97	6.88
10,000*	3.80	4.94	5.99	6.59	7.85	9.32	10.85

* Extrapolated values calculated using least-squares methodology

The prevailing wind is from the south-southwest at approximately 10 miles per hour. Historically, stronger winds are recorded when the winds are from the southwest.

3.3 SURFACE WATER HYDROLOGY

PGDP is situated in the western portion of the Ohio River basin, 15 miles downstream of the confluence of the Ohio River with the Tennessee River and 35 miles upstream of the confluence of the Ohio River with the Mississippi River. The Ohio River is located approximately 3.5 miles north of PGDP. It is the most significant surface-water feature in the region, carrying over 25 billion gal/day of water through its

¹ For the recent five-year period June 2002 through May 2007, average monthly precipitation was slightly less (3.90 inches), ranging from 3.25 inches in October (monthly average low) to 4.94 inches in September (monthly average high).

channel. A U. S. Geological Survey (USGS) gaging station at Metropolis, Illinois (USGS 03611500), monitors the Ohio River stage near PGDP. River stage typically varies between 290 and 328 ft amsl over the course of a year. Water levels on the lower Ohio River generally are highest in winter and early spring and lowest in late summer and early fall. The entire PGDP is above the historical high water floodplain of the Ohio River (CH2M HILL 1991) and above the local 100-year flood elevation of the Ohio River (333 ft). [The highest Ohio River stage recorded at Metropolis, Illinois (February 2, 1937) was 343 ft.]

The plant overlies the divide between Little Bayou and Bayou Creeks (Figure 3.1). Bayou Creek is a perennial stream on the western boundary of the plant that flows generally northward, from approximately 2.5 miles south of the plant site to the Ohio River along a 9 mile course. Little Bayou Creek is an intermittent stream located on the eastern boundary of the plant; its drainage originates within WKWMA and extends northward along a 6.5 mile course, which joins Bayou Creek near the Ohio River. Most of the flow within Bayou and Little Bayou Creeks is from process effluents or surface water runoff from PGDP. Networks of ditches discharge effluent and surface water runoff from PGDP to the creeks. Contributions from PGDP comprise approximately 85% of the base flow within Bayou Creek and 100% of the base flow within Little Bayou Creek.

Multiple groundwater aquifers underlie PGDP. (See Section 3.6 for a discussion of PGDP hydrogeology.) The shallowest aquifers occur in the Continental Deposits and the McNairy Formation, both of which discharge into the Ohio River north of PGDP. A large, downward, vertical hydraulic gradient within the Upper Continental Deposits, which represents an aquitard, typically limits the amount of groundwater discharge to the ditches of PGDP and adjacent creeks. Gaining reaches in the creeks are found on Bayou Creek south of PGDP and on Little Bayou Creek to the north of PGDP where it meets the Ohio River flood plain. Both creeks have gaining reaches adjacent to the Ohio River.

Other surface water bodies in the vicinity of PGDP include several small ponds, inactive clay and gravel pits, and settling basins scattered throughout the PGDP plant area; a marshy area just south of the confluence of Bayou Creek and Little Bayou Creek; ash settling ponds of the Shawnee Steam Plant; and Metropolis Lake, located east of the Shawnee Steam Plant.

3.4 GEOLOGY

PGDP lies within the Jackson Purchase region of western Kentucky, which represents the northern tip of the Mississippi Embayment portion of the Coastal Plain Province. The stratigraphic sequence in the region consists of Cretaceous, Tertiary, and Quaternary sediments unconformably overlying Paleozoic bedrock (Figure 3.2). The following sections describe the primary geologic units of the PGDP region. Section 3.9 presents the shallow geology specific to each of the BGOU SWMUs.

3.4.1 Bedrock

Mississippian carbonates, composed of dark gray limestone, with some interbedded chert and shale underlie the entire PGDP area at an approximate depth of 300 to 340 ft.

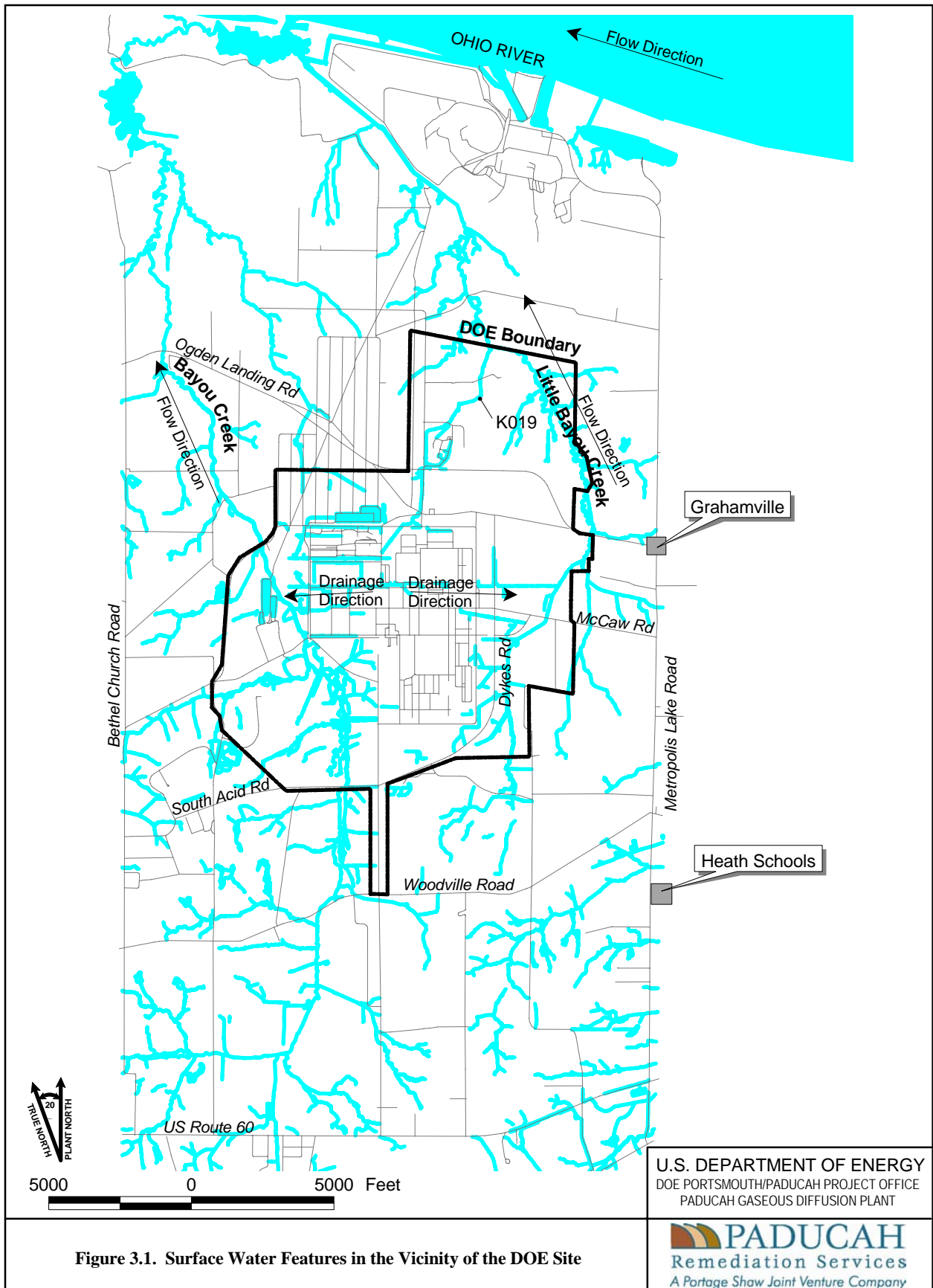
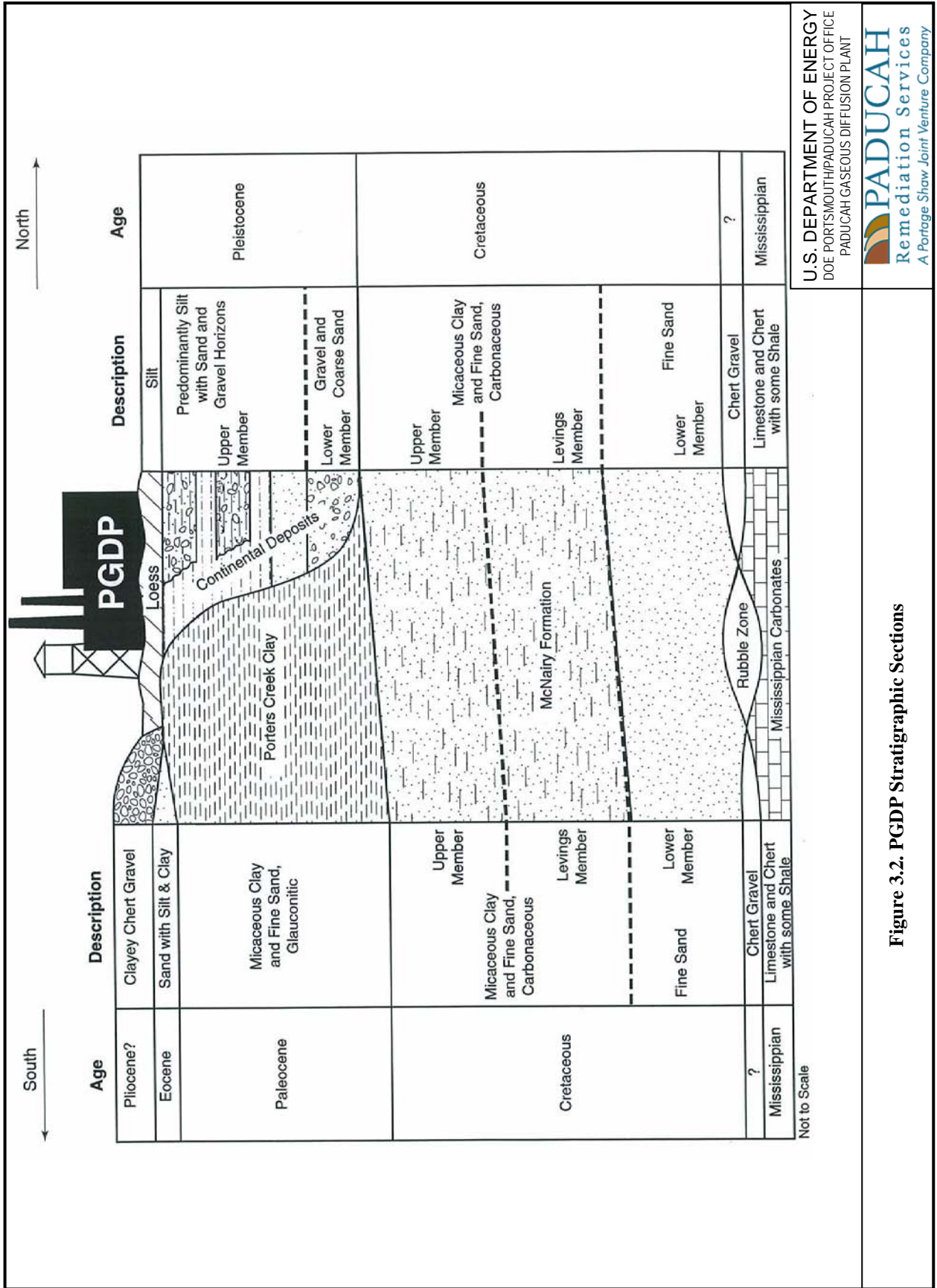


Figure 3.1. Surface Water Features in the Vicinity of the DOE Site

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PADUCAH GASEOUS DIFFUSION PLANT



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PADUCAH GASEOUS DIFFUSION PLANT



Figure 3.2. PGDP Stratigraphic Sections

3.4.2 Rubble Zone

Deep soil borings at PGDP commonly encounter a rubble zone of chert gravel at the top of the bedrock. The age and continuity of the rubble zone remain undetermined.

3.4.3 McNairy Formation

The McNairy Formation consists of Upper Cretaceous, fine clastic sediments. At PGDP the upper and middle members of the McNairy Formation are typically grayish-white to dark-gray, micaceous silt and clay interbedded with gray to yellow, very fine- to fine-grained sand. The middle (Levings) member tends to contain fewer sand interbeds. The basal McNairy member at PGDP is primarily a light gray, very fine to fine sand.

3.4.4 Porters Creek Clay/Porters Creek Terrace Slope

Paleocene age Porters Creek Clay underlies the southern portions of the DOE site and consists of dark gray to black silt with varying amounts of clay and fine-grained, micaceous, commonly glauconitic, sand. The Porters Creek Clay subcrops along a buried terrace slope that extends east–west under the south end of the PGDP industrial area. This subcrop is the northern limit of Porters Creek Clay and the southern limit of the Pleistocene Lower Continental Deposits under PGDP.

3.4.5 Eocene Sands

Eocene sands occur south of PGDP above the Porters Creek Clay. This unit includes undifferentiated quartz sands and interbedded and interlensing silts and clays of the Claiborne Group and Wilcox Formation (Olive 1980). The Eocene sands thicken to the south of PGDP.

3.4.6 Continental Deposits

Continental sediments [Pliocene(?)² to Pleistocene age] unconformably overlie the Cretaceous through Eocene strata throughout the area. These continental sediments were deposited on an irregular erosional surface consisting of several terraces. The thicker Continental Deposits sections represent Pleistocene valley fill sediments that comprise a fining-upward cycle. The continental sediments have been divided into the two distinct facies described below.

- (1) Lower Continental Deposits. The Lower Continental Deposits is a gravel facies consisting of chert, ranging from pebbles to cobbles, in a matrix of poorly sorted sand and silt. Gravels of the Lower Continental Deposits overlie three distinct terraces in the PGDP area.
 - The upper terrace Lower Continental Deposits consists of Pliocene(?) gravel units, ranging in thickness from near 0 to 30 ft, occurring in the southern portion of the DOE site at elevations greater than 350 ft amsl. This gravel unit overlies the Eocene sands and Porters Creek Clay (where the Eocene sands are missing).
 - Pliocene(?) gravels of the Lower Continental Deposits also occur on an intermediate terrace eroded into the Porters Creek Clay at an elevation of approximately 320 to 345 ft amsl in the southeastern and eastern portions of the DOE site. The thickness of this unit typically ranges from 15 to 20 ft.

² A question mark indicates uncertain age.

- The Lower Continental Deposits of the upper and intermediate terraces are collectively referred to as the Terrace Gravel.
- The third and most prominent of the three Lower Continental Deposits members consists of a Pleistocene gravel deposit resting on an erosional surface at an elevation of approximately 280 ft amsl. This gravel underlies most of the plant area and the region to the north, but pinches out under the south side of PGDP along the subcrop of the Porters Creek Clay. The Pleistocene member of the Lower Continental Deposits averages approximately 30 ft in thickness. Trends of greater thickness, as much as 50 ft, fill deeper scour channels that trend east–west beneath the site.

(2) Upper Continental Deposits. The Upper Continental Deposits is a Pleistocene age, fine-grained clastics facies that commonly overlies the Lower Continental Deposits. This unit ranges in thickness from 15 to 55 ft. The Upper Continental Deposits includes three general horizons beneath PGDP: (1) an upper silt and clay interval, (2) an intermediate interval of common sand and gravel lenses (sand and gravel content generally diminishes northward), and (3) a lower silt and clay interval. The upper silt and clay interval consists of the Peoria Loess and Roxana Silt (DOE 2003; WLA 2006). The Peoria Loess and Roxana Silt blanket the entire PGDP area.

3.5 SOILS

The surficial deposits found in the vicinity of PGDP are Pleistocene loess and Holocene alluvium. Both units commonly consist of clayey silt or silty clay and range in color from yellowish-brown to brownish-gray or tan, making field differentiation difficult. The general soil map for Ballard and McCracken Counties delineates three soil associations within the vicinity of PGDP: the Rosebloom-Wheeling-Dubbs association, the Grenada-Calloway association, and the Calloway-Henry association (USDA 1976).

In the immediate PGDP area, the predominant soil is the Henry soil series of the Calloway-Henry association, which consists of nearly level, somewhat poorly- to poorly-drained, medium-textured soils on upland positions. The Henry soil series contains poorly drained, acidic soils that have a fragipan. Henry soils typically have moderate permeability above the fragipan and low permeability within the fragipan. Permeability in the fragipan is less than 0.4 ft/day (DOE 1998c). It should be noted that soils within the industrial area of PGDP could be classified as “urban” since they have been impacted by human influence and many of the original characteristics have been lost.

Several other soil groups also occur in limited areas of the region, including the Grenada, Falaya-Collins, Waverly, Vicksburg, and Loring.

The soils in the vicinity of PGDP tend to have a low buffering capacity, with a pH ranging from 4.5 to 5.5. Measurements of the cation exchange capacity of site soils range from 8.92 to 69.8 milliequivalents per liter (DOE 1999c). Under background conditions, the cation exchange capacity is sufficient to bind metals in the soils; however, acidic leachate will significantly increase metal solubility and mobility. The potential for acidic leachate at each SWMU is uncertain due to the lack of disposal records. SWMUs with the greatest potential for acidic leachate are SWMU 6 (exhaust fans with perchloric acid) and SWMU 4 (poor records of chemicals buried). There were no indications of acidic leachate or mobilized contaminants found at SWMU 6 during this RI (UCRS groundwater pH values ranged from 5.04 to 6.75). There likely was minimal perchloric acid residue on the exhaust fans that has degraded over time. On the other hand, wastes that went into SWMU 3 were neutralized at C-400 prior to disposal and should present a low potential for generation of acidic leachate (leachate samples from SWMU 3, from 2003 to 2008, have a pH between 8.0 to 8.5).

3.6 HYDROGEOLOGY

The significant geologic units relative to shallow groundwater flow at PGDP include the Terrace Gravel and Porters Creek Clay (south sector of the DOE site) and the Pleistocene Continental Deposits and McNairy Formation (underlying PGDP and adjacent areas to the north). Groundwater flow in the Pleistocene Continental Deposits is a primary pathway for transport of dissolved contamination from PGDP. The following paragraphs provide the framework of the shallow groundwater flow system at PGDP.

- (1) Terrace Gravel Flow System. The Porters Creek Clay is a confining unit to downward groundwater flow south of PGDP. A shallow water table flow system is developed in the Terrace Gravel, where it overlies the Porters Creek Clay south of PGDP. Discharge from this water table flow system provides baseflow to Bayou Creek and underflow to the Pleistocene Continental Deposits to the east of PGDP.

The elevation of the top of the Porters Creek Clay is an important control to the area's groundwater flow trends. A distinct groundwater divide is centered in hills located approximately 9,000 ft southwest of PGDP, where the Terrace Gravel and Eocene sands overlie a "high" on the top of the Porters Creek Clay. In adjacent areas where the top of the Porters Creek Clay approaches land surface, as it does south of PGDP and near the subcrop of the Porters Creek Clay to the west of the industrial complex, the majority of groundwater flow is forced to discharge into surface streams (gaining reaches) and little underflow occurs into the Pleistocene Continental Deposits. To the east of PGDP, the Terrace Gravel overlies a lower terrace eroded into the top of the Porters Creek Clay. In this area, a thick sequence of Terrace Gravel occurs adjacent to the Pleistocene Continental Deposits, allowing significant underflow from the Terrace Gravel. Surface drainages in this area are typically losing reaches. Figure 3.3 presents hydraulic potential trends for the Terrace Gravel flow system.

- (2) UCRS. The upper strata, where infiltration of water from the surface occurs and where the uppermost zone of saturation exists, in the Upper Continental Deposits (beneath PGDP and the contiguous land to the north) is called the UCRS. Groundwater flow is primarily downward in the Upper Continental Deposits. A plot of elevation of water level versus midpoint of well screen for UCRS wells at PGDP (Figure 3.4) demonstrates that steep vertical hydraulic gradients are characteristic of the UCRS. Vertical hydraulic gradients generally range from 0.5 to 1 ft/ft where measured by wells completed at different depths in the UCRS. Vertical gradients are 1 to 2 orders of magnitude greater than lateral hydraulic gradients. While groundwater flow is predominantly downward, there will be some lateral flow due to heterogeneities in the shallow soils.

Direct measurements of the UCRS water table elevation are available only for the south-central PGDP industrial area, where water levels commonly occur in the screen interval of the wells, and the location of two source unit investigations (the SWMU 2 Interim Remedial Design Investigation and the SWMUs 7 and 30 RI) in the west PGDP industrial area. All other well measurements, where water levels occur above the well screen interval, provide lower bounds to the elevation of the water table. Figure 3.5 illustrates the general features of the UCRS water table for the PGDP industrial area (DOE 1997b). While Figure 3.5 shows data from November and December 1995, hydrographs of UCRS monitoring wells on-site indicate fluctuations of only a few ft over the past 10 years; therefore, this figure still provides an adequate representation of the UCRS water table. The main features of the water table are a broad trough in the northeast and central areas, a linear discharge area associated with a ditch in the northwest, and a lateral hydraulic gradient toward Bayou Creek on

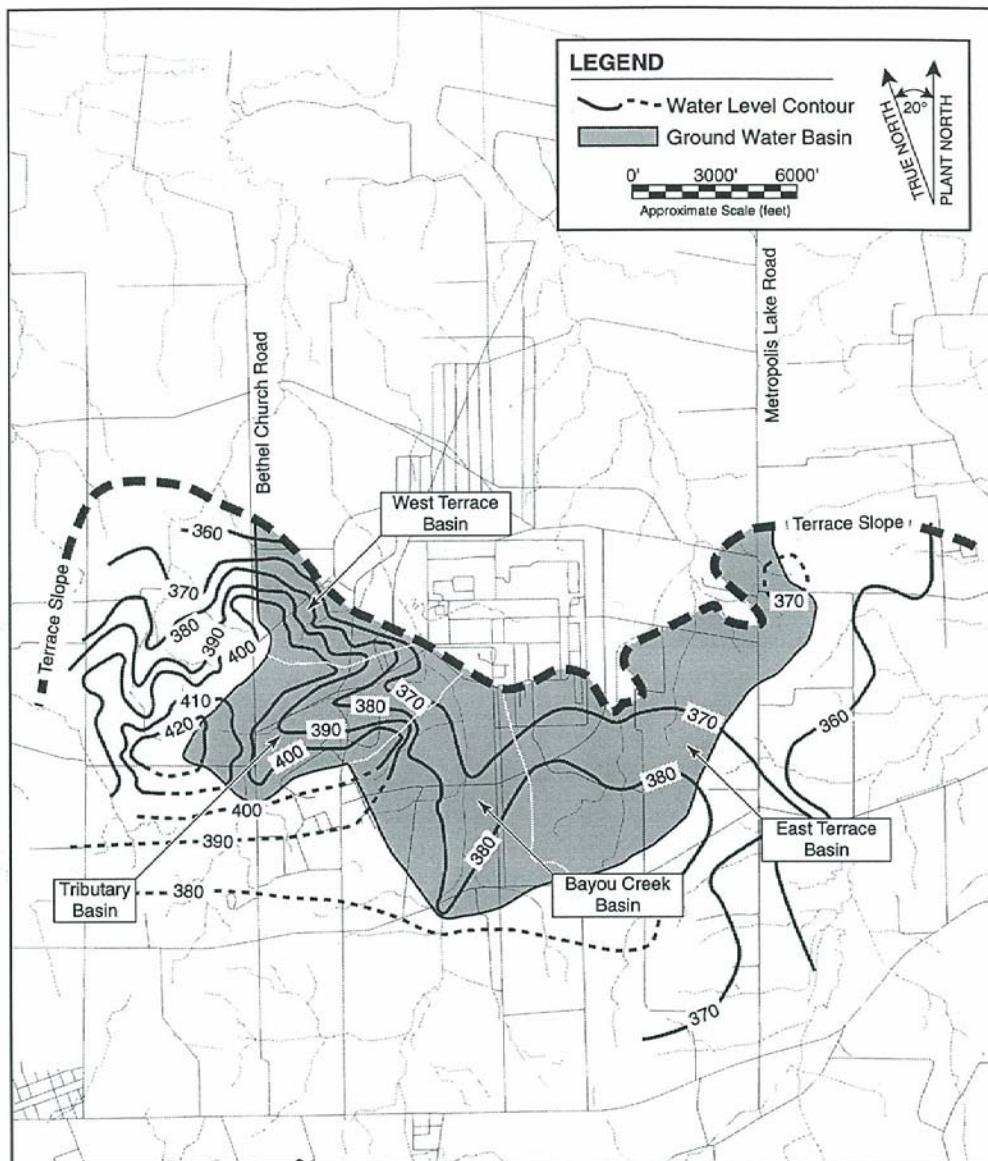
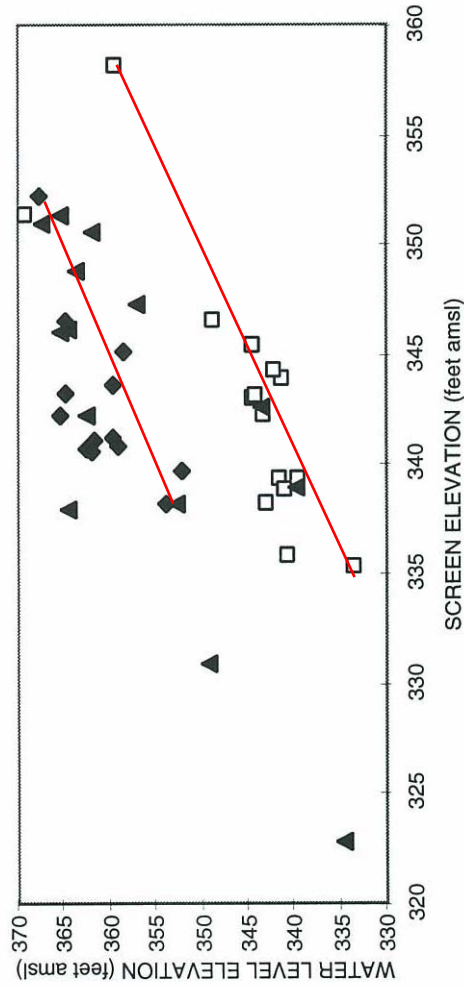


Figure 3.3. Water Table Trends in the Terrace Deposits South of the PGDP

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◆ WEST CENTRAL WELLS □ SOUTH CENTRAL WELLS ▲ OTHER PGDP AREA WELLS

Figure 3.4. Plot of Water Level Versus Well Screen for Upper Continental Recharge System Wells

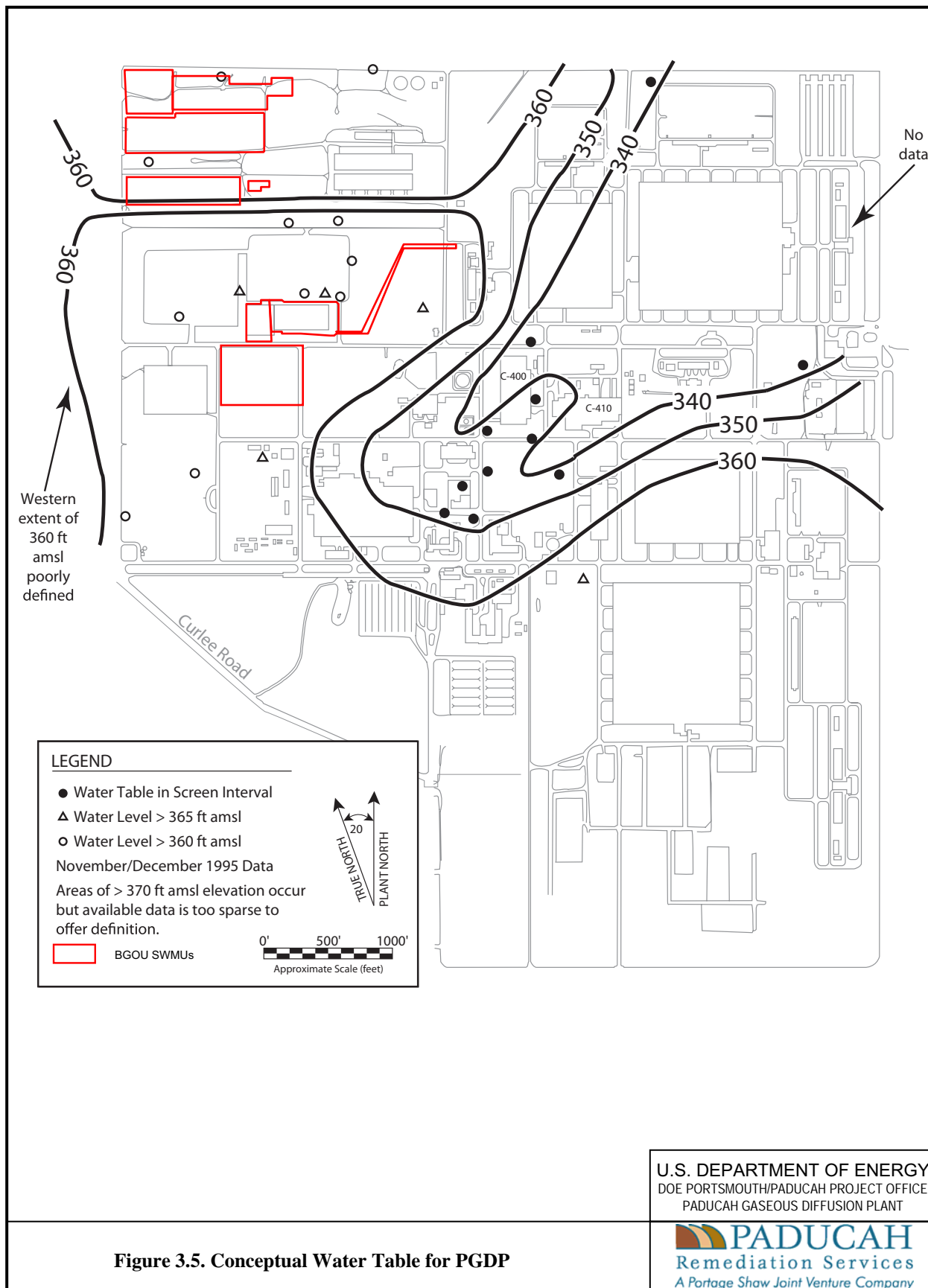


Figure 3.5. Conceptual Water Table for PGDP

the west side. In general, the water table is less than 20 ft deep in the western half of PGDP and as much as 40 ft deep in the northeastern corner.

The infiltration rate for the PGDP area is approximately 6.6 inches/yr based on site-specific groundwater modeling. This 6.6 inches/yr applied over the area of the industrial area of the plant yields approximately 0.4 mgd of recharge to the shallow groundwater system. Leakage from plant water utilities, ditches, lagoons, and cooling tower basins is suspected to be another important source of infiltration at PGDP. Water use for PGDP for calendar year 2006 averaged 13 mgd. Municipal water systems lose as much as 24% of their daily conveyance (Jowitt and Xu 1990). A similar loss of the PGDP system would equal 3.1 mgd. Since the UCRS groundwater flow is predominantly downward, areas with higher anthropogenic recharge create mounding of hydraulic head in the RGA that can affect contaminant transport. Because the hydraulic conductivity in the RGA on-site is relatively large, the mounding is only slight (often less than 1ft) and difficult to measure.

- (3) RGA. Vertically infiltrating water from the UCRS moves downward into a basal sand member of the Upper Continental Deposits and the Pleistocene gravel member of the Lower Continental Deposits and then laterally north toward the Ohio River. This lateral flow system is called the RGA. The RGA is the shallow aquifer beneath PGDP and contiguous lands to the north. Groundwater of the RGA meets requirements of a Class II groundwater as delineated in *Guidelines for Ground-Water Classification under the EPA Ground-Water Protection Strategy* (EPA 1988).

Hydraulic potential in the RGA declines toward the Ohio River, which is the control of base level of the region's surface water and groundwater systems. The RGA potentiometric surface gradient beneath PGDP is commonly 10^{-4} ft/ft, but increases by an order of magnitude near the Ohio River. (Vertical gradients are not well documented, but small.)

The hydraulic conductivity of the RGA varies spatially. Pumping tests have documented the hydraulic conductivity of the RGA ranges from 53 ft/day to 5,700 ft/day. East-to-west flow of the ancestral Tennessee River, which laid down the Pleistocene Continental Deposits gravel member, tended to orient permeable gravel and sand lenses east-west. Thus, with the hydraulic head in the RGA generally decreasing northward toward the Ohio River, groundwater flow trends to the northeast and northwest from PGDP in response to the anisotropy of the hydraulic conductivity as well as the anthropogenic recharge, which is greatest in the industrial portion of the plant. Anthropogenic recharge from waterline leaks, lagoons, cooling tower basins, and other sources provides the primary driving force in moving groundwater in northeastern and northwestern flow directions from the industrial plant area. Ambient groundwater flow rates in the more permeable pathways of the RGA commonly range from 1 to 3 ft/day.

- (4) McNairy Flow System. Groundwater flow in the fine sands and silts of the McNairy Formation is called the McNairy Flow System. The overall McNairy groundwater flow direction in the area of PGDP is northward to the Ohio River, similar to that of the RGA. Hydraulic potential is greater in the RGA than in the McNairy Flow System beneath PGDP. Area monitoring well clusters document an average downward vertical gradient of 0.03 ft/ft. Because the RGA has a steeper hydraulic potential slope toward the Ohio River than does the McNairy Flow System, the vertical gradient reverses nearer the Ohio River. [The "hinge line," which is where the vertical hydraulic gradient between the RGA and McNairy Flow System changes from a downward vertical gradient to an upward vertical gradient, parallels the Ohio River near the northern DOE property boundary (LMES 1996).]

The contact between the Lower Continental Deposits and the McNairy Formation is a marked hydraulic properties boundary. Representative lateral and vertical hydraulic conductivities of the upper McNairy Formation in the area of PGDP are approximately 0.02 ft/day and 0.0005 ft/day,

respectively. Vertical infiltration of groundwater into the McNairy Formation beneath PGDP is on the order of 0.1 inch per year. (Lateral flow in the McNairy Formation beneath PGDP is on the order of 0.03 inch per year.) As a result, little interchange occurs between the RGA and McNairy Flow System.

3.6.1 Hydrogeologic Units

Five hydrogeologic units (HUs) are commonly used to discuss the shallow groundwater flow system beneath the DOE site and the contiguous lands to the north (Figure 3.6). In descending order, the HUs are described below:

- Upper Continental Deposits
 - HU 1 (UCRS): Loess that covers the entire site.
 - HU 2 (UCRS): Discontinuous, sand and gravel lenses in a clayey silt matrix. In some areas of the plant, the HU2 interval consists of an upper sand and gravel member (HU2A) and a lower sand and gravel member (HU2B) separated by a thin silt unit.
 - HU 3 (UCRS): Relatively impermeable unit that acts as the upper semiconfining-to-confining layer for the RGA. The lithologic composition of HU3 varies from clay to fine sand, but is predominantly silt and clay.
 - HU 4 (RGA): Near-continuous sand unit with a clayey silt matrix that forms the top of the RGA.
- Lower Continental Deposits
 - HU 5 (RGA): Gravel, sand, and silt.

3.7 DEMOGRAPHY AND LAND USE

The WKWMA and some sparsely populated agricultural lands surround PGDP. Historically, the economy of western Kentucky has been based on agriculture, although there has been increased industrial development in recent years. PGDP's operations contractor, USEC, employs approximately 1,400 people, while the TVA Shawnee Steam Plant employs an additional 260 people. According to the 2000 U.S. Census, the total population within the 32 counties that lie within a 50-mile radius of PGDP is approximately 731,500; and approximately 88,500 people live within the three counties that contain the 10 mile radius of the plant (Massac County, Illinois and Ballard and McCracken Counties, Kentucky). The estimated population of Paducah, Kentucky, (2006) is approximately 25,600. Metropolis, Illinois, has an estimated population (2006) of approximately 6,400 (U.S. Census Bureau 2007).

In addition to the residential population surrounding the plant, WKWMA draws thousands of visitors each year for recreational purposes. Visitors primarily use the area for hunting and fishing, but other activities include horseback riding, hiking, and bird watching. An estimated 5,000 fishermen visit the area each year.

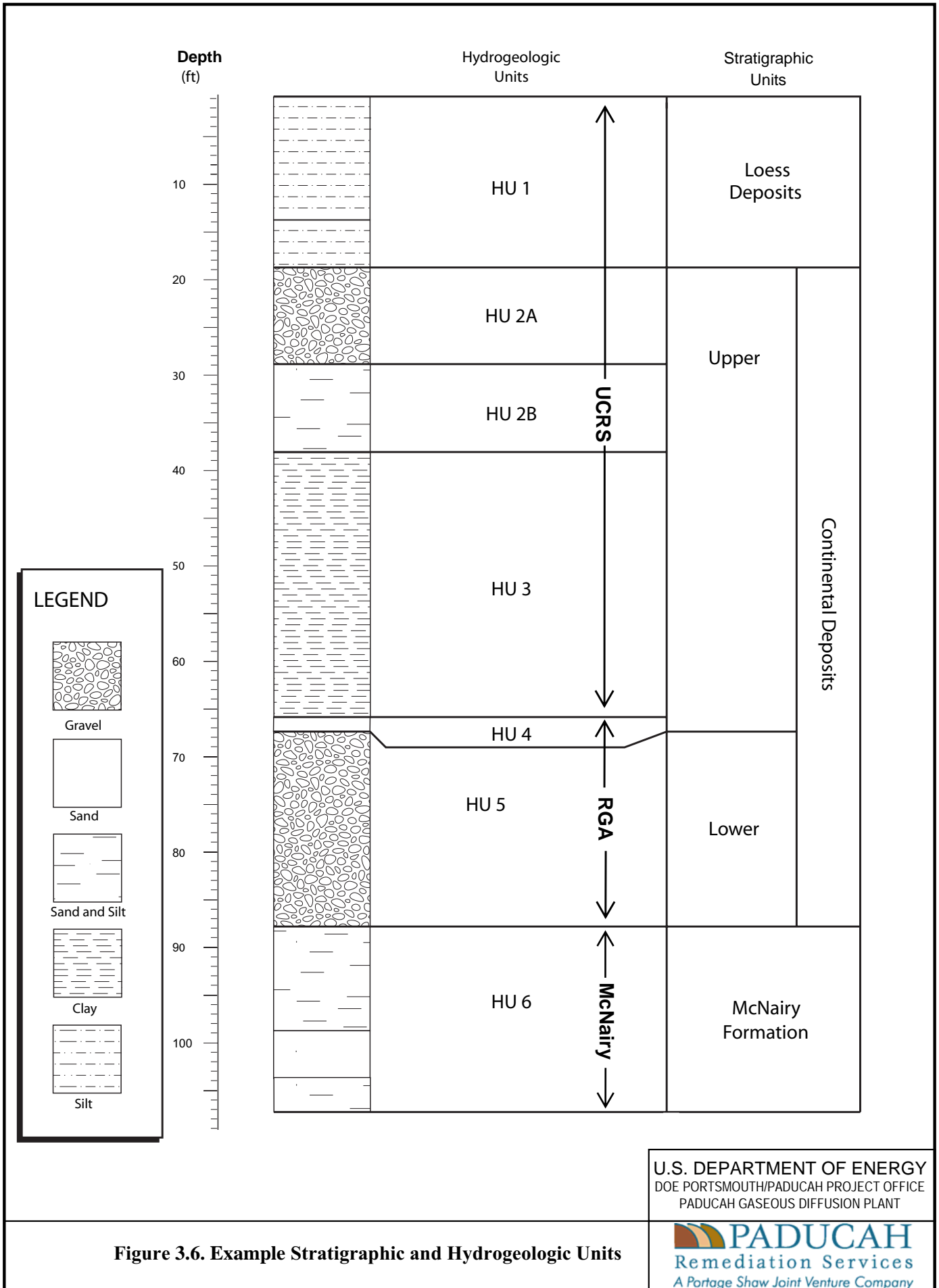


Figure 3.6. Example Stratigraphic and Hydrogeologic Units

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3.8 ECOLOGY

The following sections give a brief overview of the terrestrial and aquatic systems at PGDP. A more detailed description, including identification and discussion of sensitive habitats and threatened/endangered species, is contained in the *Investigation of Sensitive Ecological Resources Inside the Paducah Gaseous Diffusion Plant, Paducah, Kentucky* (CDM Federal 1994) and *Environmental Investigations at the Paducah Gaseous Diffusion Plant and Surrounding Area, McCracken County, Kentucky, Volume V: Floodplain Investigation, Part A: Results of Field Survey* (COE 1994).

3.8.1 Terrestrial Systems

The terrestrial component of the PGDP ecosystem includes the plants and animals that use the upland habitats for food, reproduction, and protection. Upland vegetative communities in the vicinity of PGDP consist primarily of grassland, forest, and thicket habitats with agricultural areas. The main crops grown in the PGDP area include soybeans, corn, tobacco, and sorghum.

Most of the area in the vicinity of PGDP has been cleared of vegetation at some time. PGDP mows much of the grassland habitat adjacent to the plant. The Kentucky Department of Fish and Wildlife Resources manages a large percentage of the adjacent WKWMA to promote native prairie vegetation by burning, mowing, and various other techniques.

Dominant overstory species of the forested areas include oaks, hickories, maples, elms, and sweetgum. Understory species include snowberry, poison ivy, trumpet creeper, Virginia creeper, and Solomon's seal. Thicket areas consist predominantly of maples, black locust, sumac, persimmon, and forest species in the sapling stage with herbaceous ground cover similar to that of the forest understory.

Wildlife commonly found in the PGDP area consists of species indigenous to open grassland, thicket, and forest habitats. Small mammal surveys conducted on WKWMA documented the presence of southern short-tailed shrew, prairie vole, house mouse, rice rat, and deer mouse (KSNPC 1991). Large mammals commonly present in the area include coyote, eastern cottontail, opossum, groundhog, whitetail deer, raccoon, and gray squirrel. Mist netting activities in the area have captured red bat, little brown bat, Indiana bat, northern long-eared bat, evening bat, and eastern pipistrelle (KSNPC 1991).

The typical birds of the area are European starling, cardinal, red-winged blackbird, mourning dove, bobwhite quail, turkey, killdeer, American robin, eastern meadowlark, eastern bluebird, bluejay, red-tail hawk, and great horned owl.

Amphibians and reptiles present in the PGDP area include cricket frog, Fowler's toad, common snapping turtle, green tree frog, chorus frog, southern leopard frog, eastern fence lizard, and red-eared slider (KSNPC 1991).

3.8.2 Aquatic Systems

The aquatic communities in and around the PGDP area that could be impacted by plant discharges include two perennial streams (Bayou Creek and Little Bayou Creek), the NSDD (a former ditch for the discharge of plant effluents to Little Bayou Creek), a marsh located at the confluence of Bayou Creek and Little Bayou Creek, and other smaller drainage areas. The dominant taxa in all surface waters include several species of sunfish, especially bluegill and green sunfish, as well as bass and catfish. Shallow streams, characteristic of the two main area creeks, are commonly dominated by bluegill, green and longear sunfish, and stonerollers.

3.8.3 Wetlands and Floodplains

The wetlands of the PGDP vicinity include a swamp covering 165 acres immediately south of the confluence of Bayou and Little Bayou Creeks. A 1994 study of the PGDP area by the U.S. Corps of Engineers (COE) (1994) groups the area wetlands into 16 vegetative cover types encompassing forested, scrub/shrub, and emergent wetlands. Wetland vegetation consists of species such as sedges, rushes, spikerushes, and various other grasses and forbs in the emergent portions; red maple, sweet gum, oaks, and hickories in the forested portions; and black willow and various other saplings of forested species in the thicket portions. Wetlands inside the plant security fence are confined to portions of drainage ditches traversing the site (CDM Federal 1994).

At PGDP, three bodies of water cause most area flooding: the Ohio River, Bayou Creek, and Little Bayou Creek. The floodplain analysis performed by the COE (COE 1994) found that much of the built-up portions of the plant lie outside the 100- and 500-year floodplains of these streams. In addition, this analysis determined that ditches within the plant area can contain the expected 100- and 500-year discharges. It should be noted that precipitation frequency estimates for the 100- and 500-year events were updated in 2004 in the National Oceanic and Atmospheric Administration's (NOAA) Atlas 14 (NOAA 2004). In the updated report, the mean precipitation estimate for the 100-year, 24-hour event in Atlas 14 for the Paducah area is 10.1% to 15% greater than the mean estimate in previous publications. As stated in Atlas 14, in many cases, the mean precipitation estimate used previously still is within the confidence limits provided in Atlas 14; therefore, it is likely the plant ditches still will contain the 100- and 500-year discharges.

3.9 BGOU PHYSICAL CHARACTERISTICS

The following sections present the settings and physical characteristics of the BGOU SWMUs that govern contaminant migration.

3.9.1 BGOU Surface Features

The PGDP facility generally consists of three land uses, (1) areas of permanent structures and paved roads that are engineered to promote drainage, (2) UF₆ cylinder storage yards and scrap yards, and (3) former burial grounds and aboveground landfills. All of the SWMUs for the BGOU are former burial grounds or aboveground landfills. Drainage ditches that discharge into KPDES outfalls and then to Bayou Creek west of the plant skirt all of the BGOU SWMUs, except SWMU 145 (Figure 3.7). Runoff from SWMU 145 flows through the NSDD to Little Bayou Creek.

SWMU 2 is a uranium burial ground located immediately west of SWMU 3, in the west-central portion of the plant (Figure 1.4). Graveled storage yards bound SWMU 2, to the north and west, respectively. The main drainage ditch to KPDES Outfall 015 passes between SWMU 2 and Virginia Avenue, to the south. SWMU 2 is grass covered. The land surface at SWMU 2 is relatively flat (with a slight mound on the east side); surface elevations range from 370 to 375 ft amsl. PGDP maintains SWMU 2 as a Radioactive Materials Area, with applicable boundary access controls.

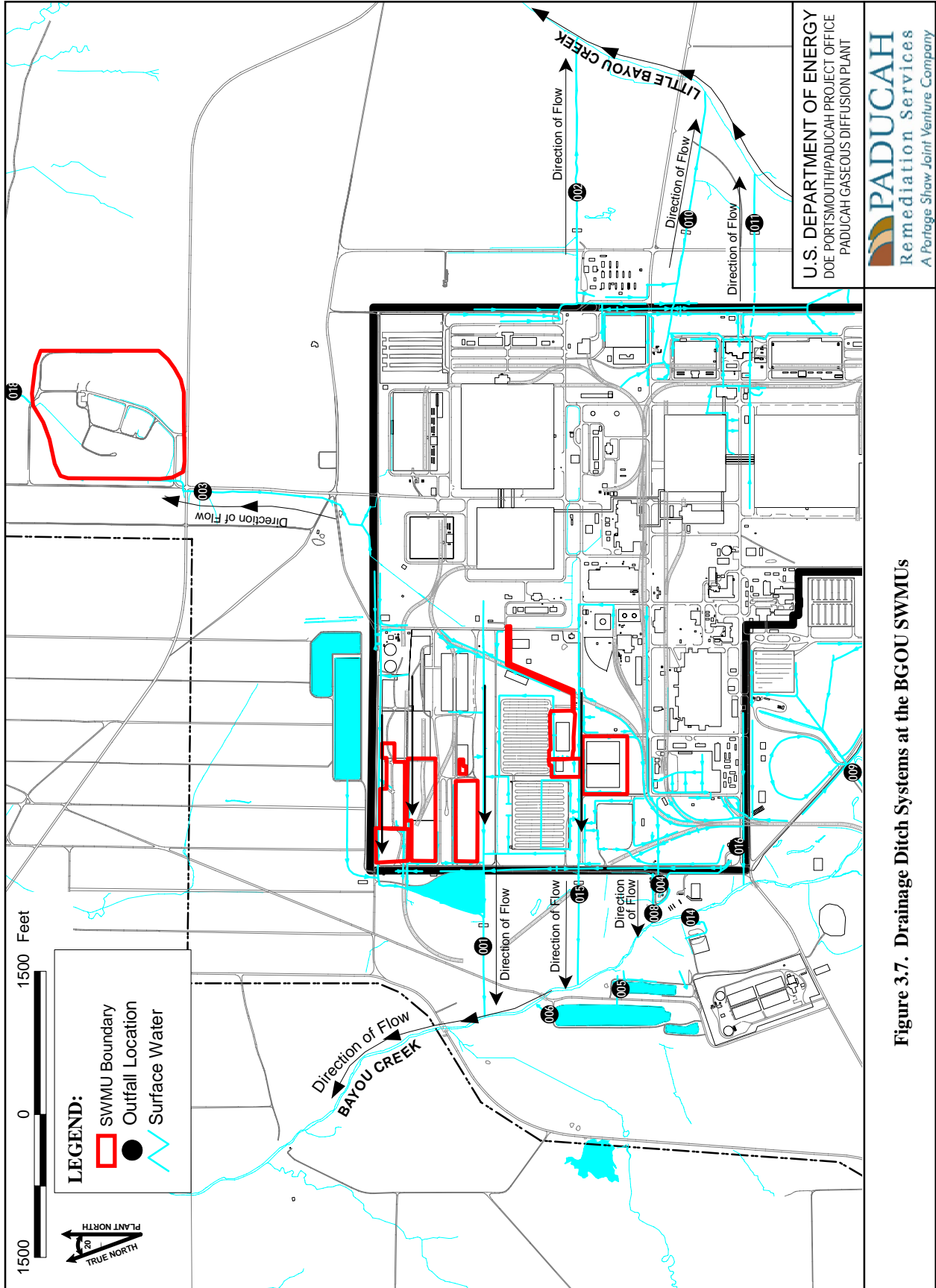


Figure 3.7. Drainage Ditch Systems at the BGOU SWMUs

SWMU 3 (Figure 1.6) consists of an aboveground surface impoundment that was converted to a solid waste disposal facility (C-404) and a field to the east where a northeast-southwest ditch drained the C-404 surface impoundment to the NSDD. C-404 is a grass covered mound with steep, 10-ft high sides and a gently sloping cap (highest on the east side). Elevations at C-404 range from 375 to 392 ft amsl. An empty, graveled, cylinder storage yard borders C-404 to the north. The same main drainage ditch to KPDES Outfall 015 passes between C-404 and Virginia Avenue to the south.

Gravel roads provide limited access to the east and south sides of C-404. PGDP maintains C-404 as a Radioactive Materials Area.

SWMU 4 is an open grass field that was used for the burial and disposal of waste materials. This SWMU is bounded on the north by Virginia Avenue, on the east by 6th Street, on the west by 4th Street, and on the south by an active railroad spur (Figure 1.7). Shallow drainage swales that direct surface runoff to the northwest corner of the site bound SWMU 4 on three sides (north, east, and west). Surface runoff passes beneath Virginia Avenue through a drainage culvert where it discharges into the main drainage ditch to KPDES Outfall 015. The ground surface of the burial area is graded so that surface runoff is directed toward the surrounding drainage swales. There is an elevation difference of approximately 10 ft between the highest point in the SWMU to the adjacent drainage swales. PGDP maintains a fence around SWMU 4 to control access.

SWMU 5 is a burial area in the northwest quadrant of the plant (Figure 1.8). Unnamed gravel roads parallel the north, south, and east sides, while a paved road lies to the west. Shallow drainage swales bordering the SWMU direct surface runoff to a settling pond (C-613) and then to KPDES Outfall 001. The ground surface is grass-covered with no significant surface structures. Approximately five ft of topographic relief exists between the mound of the burial area, which is offset to the east, and the sides of the SWMU. The SWMU is fenced to limit access to authorized personnel only.

The SWMU 6 burial plots (Figure 1.10) are located due east of SWMU 5. This area is relatively flat and is bounded by unnamed gravel roads to the west and south and to the north by a ditch that drains through the C-613 settling pond to KPDES Outfall 001. PGDP maintains the area as a grassed field with occasional shrubs. SWMU 6 is a Radioactive Materials Area with boundary chains to mark limited access.

SWMU 7 is a burial pit area in the northwest corner of the plant (Figure 1.11). Ditches of the KPDES Outfall 001 drainage system border SWMU 7 to the north and south. A scrap yard lies to the east. SWMU 30 adjoins SWMU 7 to the west. The earthen cover over the burial pits form slight hills (two ft high) on the north and south sides of SWMU 7. A gravel pad covers the east end of SWMU 7. PGDP maintains grass cover over the west burial pits. Boundary chains limit access to the west burial pits, which are delimited Radioactive Materials Areas and High Radioactive Materials Areas.

The same KPDES Outfall 001 drainage ditches bound SWMU 30 on the north and south sides. A paved road borders SWMU 30 on the west side. The surface of the SWMU 30 earthen cover ranges from an elevation of 375 ft at its highest point near the northeast corner of the SWMU to 371 ft near the edges of the burial pit. As at SWMU 7, PGDP maintains a grass cover over the burial pit and boundary chains limit access (Radioactive Materials Area).

SWMU 145 (Figure 1.12) is located to the north of the plant, beneath the C-746-S and -T Landfills. Boundaries of the waste fill are not well defined. The BGOU RI used review of historical aerial photography and geophysical surveys to delineate areas for characterization of the historic waste fill (see Section 2.1). Ogden Landing Road (Kentucky Highway 358) borders the south side of SWMU 145. PGDP's currently operating landfill (C-746-U) lies to the north of SWMU 145. The present trace of the

NSDD passes on the west and north sides. Grasslands of the WKWMA adjoin SWMU 145 to the east. Area runoff drains through the NSDD. Fencing for the C-746-S and -T Landfills limits access to SWMU 145.

3.9.2 Underground Utilities and Plant Operations

Underground utilities are sparse in the area of the BGOU SWMUs. Because they are sparse in this area, it is likely that they have had no impact on contaminant migration from or into the SWMU areas. A raw water line that supplies the PGDP water treatment plant underlies the southeast corner of SWMU 4. This raw water line is the most extensive of the underground utilities located near the BGOU RI burial areas. At its closest approach, the raw water line crosses approximately 30 ft southeast of the SWMU 4 south burial pit. (See Figure 2.2) The base of the raw water line occurs at 367 ft amsl; the base of the south burial pit at SWMU 4 (closest pit) is approximately 362 ft; thus, some buried waste likely is present above the elevation of the raw water line. Contaminants potentially could migrate from the shallower buried waste to the raw water line, which, in turn, might be a pathway for transport away from SWMU 4.

An abandoned water line (six-inch diameter) installed for the Kentucky Ordnance Works (pre PGDP) underlies the north and east ends of SWMU 5 and the south side of SWMU 6. As at SWMU 4, the water line is located outside the nearest burial pits. This water line is unlikely to act as an avenue for contaminant migration from the burial pits because the water line occurs at depths of four to seven ft, above most of the buried waste.

The only other utility in the vicinity of the burial grounds is an abandoned electrical conduit that underlies the north end of SWMU 2 at depths of six to seven ft. This conduit directly overlies areas of buried waste; however, it likely does not intercept the waste because of the shallow depth of the conduit. Like the water line near SWMUs 5 and 6, the conduit is unlikely to offer a pathway of contaminant migration because it lies above the buried waste.

Plant operations subsequent to waste operations at each of the SWMUs have contaminated surface soils. The common presence of polycyclic aromatic hydrocarbons (PAHs) and uranium in surface soil are related directly to past and on-going plant operation. Ditches bound all of the BGOU SWMUs and provide a potential pathway for contaminant migration. The Surface Water Operable Unit SI assesses the nature and extent of this contamination in most areas addressed by the BGOU RI. Ditches of the northwest plant area that drain to the C-613 Sedimentation Basin will be addressed as part of the post-GDP shutdown for surface water.

3.9.3 BGOU Hydrogeology

The scope of the BGOU RI focused on contaminant migration in the soils of the Pleistocene Continental Deposits and in the groundwater of the UCRS and RGA flow systems. Appendix B provides the lithologic logs of the boreholes drilled for the BGOU RI. The following sections summarize the general characteristics of the UCRS and RGA and present hydrogeologic data for each SWMU based on field information obtained during the BGOU RI and previous studies. This presentation of the site hydrogeology uses the framework of the five HUs as summarized in Section 3.6.1.

Sorption. Cation exchange capacity and total organic carbon content are common measures of the sorption capacity of soils. The SWMU 2 Interim Remedial Design Investigation (DOE 1997a) characterized cation exchange capacity and total organic carbon content for each HU. Table 3.2 presents the data. Cation exchange capacity values for UCRS soils range from 15 to 26 milliequivalents per 100 g (meq/100 g). These values are typical of silty soils with some clay. Only three values are available for the

RGA HU5 interval. The two lowest values of 9 and 10 meq/100 g are most representative of the overall RGA.

Table 3.2. Sorption Measurements from SWMUs 7 and 30 Remedial Investigation Report and the SWMU 2 Interim Remedial Design Investigation

	SWMUs 7 and 30	SWMU 2			
HU	Uranium Distribution Ratio (mL/g)	Sample ID	Uranium Distribution Ratio (mL/g)	Cation Exchange Capacity (meq/100 g)	Total Organic Carbon Content (%)
HU1	253 ± 10.0	S03211	3,200	NA	NA
		S05211	1,530	17.54	0.1020
		S17211	NA	20.80	0.0819
HU2 Sand	1,170 ± 264	S03212	9,080	NA	0.0701
		S05212	8,070	15.11	0.0869
		S13211	NA	17.87	0.2400
HU2 Silt		S03213	13,100	17.78	0.0465
		S05213	72,200	21.04	0.0968
		S17212	NA	18.18	0.0566
HU3	3,640 ± 2,060	S03214	93,900	NA	NA
		S05214	7,020	21.94	0.0862
		S09213	NA	23.02	0.0807
		S17213	NA	25.63	0.0720
HU4	761 ± 172	S13212	NA	23.80	0.1060
		S17214	NA	NA	0.0464
HU5	66.8 ± 3.82	S03215	4,950	NA	0.2530
		S05215	49,900	9.98	0.0453
		S09215	NA	NA	0.0394
		S13214	NA	23.72	0.0796
		S13215	NA	9.40	0.0321
		S17215	NA	NA	0.0199

NA = not available (not measured)

Total organic carbon content for the SWMU 2 data is similar for the UCRS and RGA. Values range from 0.05 to 0.24% (with a median value of 0.08%) for the UCRS measurements and 0.02 to 0.25% (with a median value of 0.05%) for the RGA measurements. The WAG 6 RI (DOE 1999c) also measured total organic carbon content of UCRS and RGA soils. Total organic carbon content measurements ranged from 0.002 to 0.2% (median of 0.04% for 20 measurements) in UCRS samples and from 0.003 to 0.3% (median of 0.02% for 38 measurements) in RGA samples.

The SWMUs 7 and 30 RI (DOE 1998a) and SWMU 2 Interim Remedial Design Investigation also characterized the uranium distribution ratio for all HUs because of the significance of potential uranium transport from the burial cells (Table 3.2). All measurements of the uranium distribution ratio are greater than 1, which means that uranium will preferentially partition from groundwater to the soils. The magnitude of the UCRS values (253 to 93,900 mL/g) indicates that common forms of uranium leachate are not likely to migrate from the burial grounds to the RGA. Even the sands and gravel units of the RGA would provide significant retention of uranium [uranium distribution coefficient (K_d) of 66.8 mL/g]. The fate and transport modeling for this RI, as documented in Appendix E, uses a K_d of 66.8 mL/g to minimize the potential of eliminating uranium as a contaminant of concern (COC) so that it can be properly addressed in the BGOU FS.

Groundwater Geochemistry. In areas that are not heavily influenced by dissolved contaminants, both UCRS and RGA groundwater tends to be mildly acidic and well buffered. As the groundwater migrates through the UCRS, bicarbonate replaces sulfate as the dominant anion (Clausen *et al.* 1992; DOE 1997a) with depth. The dominant cations in both UCRS and RGA water are commonly sodium followed by calcium.

Data are available to document the nature of dissolved oxygen levels and oxidation/reduction potential applicable to the BGOU SWMUs for both the UCRS and RGA. The dissolved oxygen data includes 517 measurements of UCRS well water and 1,799 measurements of RGA well water. Oxidation/reduction potential of the SWMUs is represented by 136 measurements of UCRS well water and 574 measurements of RGA well water. Table 3.3 summarizes the available analyses of dissolved oxygen and oxidation/reduction potential for all UCRS groundwater samples (collected from wells and temporary borings) for the BGOU SWMUs. These UCRS groundwater samples characterize areas adjacent to the disposal cells as well as areas directly below the disposal cells. The samples from directly below the disposal cells are representative of constituents migrating from the disposal cells.

The distribution and cumulative trends of dissolved oxygen measurements (collected *ex situ*) are similar for the UCRS and RGA monitoring well samples (Figure 3.8) (groundwater flows from the UCRS into the RGA). In both the UCRS and RGA, more measurements of dissolved oxygen occur in the 1.00-1.99 mg/L range than any other 1 mg/L range (25% of the measurements in both populations.) The majority of dissolved oxygen measurements from UCRS wells (75%) range from near zero³ to 4 mg/L; nearly half of the measurements (45%) are less than 2 mg/L. In the RGA, the majority of measurements (68%) range between 1 and 5 mg/L. These general trends are consistent with the site conceptual model of groundwater flow because groundwater travels relatively slowly in the UCRS and microenvironments occur where biotic and abiotic activity consume the dissolved oxygen as compared to the RGA where groundwater flow is relatively rapid. There, oxygenated water mixes with water depleted in dissolved oxygen and dissolved oxygen diffuses into depleted waters, tending to equalize the dissolved oxygen content. Consequently, the RGA does not appear to have the microenvironments where biotic or abiotic activity is consuming the dissolved oxygen.

Figures 3.9 and 3.10 show the dissolved oxygen measurements in the UCRS and RGA, respectively, for each of the BGOU SWMUs where data are available. Previous investigations of SWMU 2 (DOE 1997a) and SWMUs 7 and 30 (DOE 1998a) identified high levels of reductive dechlorination byproducts of TCE within and below some waste disposal areas. These byproducts [principally *cis*-1,2-dichloroethene (DCE) and vinyl chloride] are evidence that reducing conditions (little to no dissolved oxygen) have been present, and may continue to be present, locally within some of the burial cells where other organic wastes, such as oils, have been co-located. Low dissolved oxygen concentrations in UCRS MW186 at SWMUs 7 and 30 confirm the presence of reducing conditions. There is uncertainty with regard to the dissolved oxygen in the UCRS at SWMUs 4 and 6 due to a lack of data, but the range can be anticipated from the general UCRS trends. The presence of TCE degradation products in the UCRS at SWMU 4 provides some evidence of low dissolved oxygen at that unit.

Figure 3.11 compares oxidation/reduction potential measurements in the UCRS and RGA for the BGOU SWMUs as distribution and cumulative trend curves. The oxidation/reduction potential in the UCRS commonly ranges from -100 to 300 microVolts, with the majority of measurements greater than zero. Oxidation/reduction potential of the RGA is mostly confined in the 100 to 300 microVolts range. Plots of the data for each SWMU (as available), overlaid on the cumulative trend plots, (Figures 3.12 and 3.13)

³ It should be noted that measurements of dissolved oxygen determined with flow-through cells, such as the method used at PGDP, are not accurate below 1 mg/L, which leads to uncertainty in the distribution; however, most measurements of dissolved oxygen obtained at the site (>75%) are greater than 1 mg/L.

illustrate the range and relative abundance of measurements of oxidation/reduction potential in the UCRS and RGA for most of the SWMUs.

The data summaries in Table 3.3 and Figures 3.8 through 3.13 demonstrate that a wide range of dissolved oxygen and oxidation/reduction potential conditions are applicable at each disposal area.

RGA Hydraulic Potential. The potentiometric surface of the RGA trends north-northeast toward the regional hydraulic base level represented by the Ohio River (see Figure 3.14). Representative values for hydraulic gradient at PGDP and to the north commonly range between 10^{-4} ft/ft and 10^{-3} ft/ft. In the area of the plant, the potentiometric surface remains relatively flat throughout the year. The area north of the DOE property boundary tends to be an area of higher hydraulic gradient, except following an extended rise in the Ohio River stage.

The hydraulic potential of the RGA near the center of the plant averages 328 ft amsl and commonly fluctuates five ft over a yearly high-and-low cycle. RGA water levels near the Ohio River are often 10 ft lower. Low pool elevation of the Ohio River north of PGDP is 290 ft amsl.

**Table 3.3. Summary of Dissolved Oxygen and Oxidation/Reduction Potential Data of the UCRS
(Samples from 64 ft depth or less) for the BGOU RI**

SWMU	Sample Location	Sample Depth (ft)	Dissolved Oxygen		Oxidation/Reduction Potential	
			(mg/L)	Data Type	(mV)	Data Type
2	SWMU 2-3	63	6.3	C	112	C
	SWMU 2-9	24	11.5	C	174	C
		43	9.4	C	--	--
	SWMU 2-10	22	3.0	C	--	--
	SWMU 2-17	22	7.8	C	46	C
	PZ74	32-42	5.0	B	240	B
	MW154	16-18	4.5	A	303	C
	PZ334	8-10	5.7	A	224	B
PZ335	8-10	2.8	B	254	B	
PZ336	8-10	5.5	B	244	B	
3	MW85	30-40	8.2	A	225	A
	MW88	29-40	2.0	A	228	A
	MW91	29-39	6.8	A	223	A
	MW94	29-39	2.5	A	192	A
4	004-020	60	--	--	156	C
	004-021	60	--	--	183	C
	004-022	60	--	--	241	C
	004-105	64	--	--	144	C
	004-107	64	--	--	168	C
	004-108	64	--	--	84	C
	004-110	64	--	--	134	C
5	005-015	55-60	--	--	179	C
	005-018	60	--	--	220	C
	MW190	18-22	2.9	A	--	--
6	006-016	32-37	--	--	171	C
	006-018	22-27	--	--	175	C
	006-019	60	--	--	207	C
7	WBP-9A	11	1.6	B	164	B
	WBP-12A	9	2.3	B	64*	B
	MW186	18-23	0.8	A	157	A
30	WBP-4A	8	2.6	B	7*	B
	MW64	28-33	6.3	A	--	--
	MW187	22-26	2.0	A	198	A
145	145-021	59	--	A	24	A
	MW16	20-40	5.8	A	--	A
	MW18	35-55	0.7	A	--	A
	MW180	22-27	2.3	A	179	A
	MW182	15-20	1.5	A	-7*	A
	MW386	20-30	1.3	A	21	A
	MW390	28-38	2.7	A	197	A
	MW393	28-38	1.2	A	-13	A
MW396	34-44	1.0	A	36	A	

Data Type:

A = median of measurements of four or more sample events

B = average of measurements of two or three sample events

C = single value available

* Range includes negative and positive oxidation/reduction potential values

-- indicates no data available.

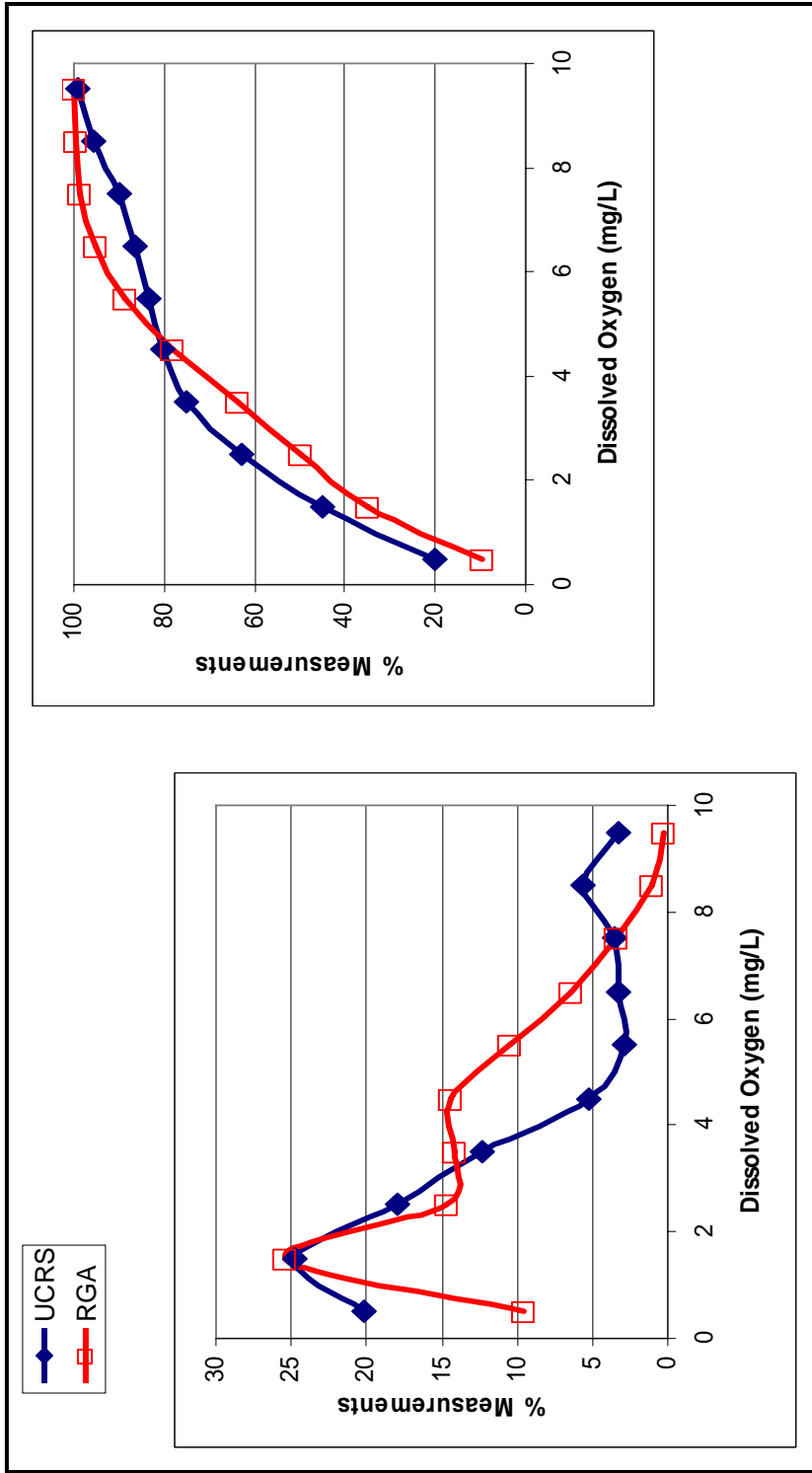
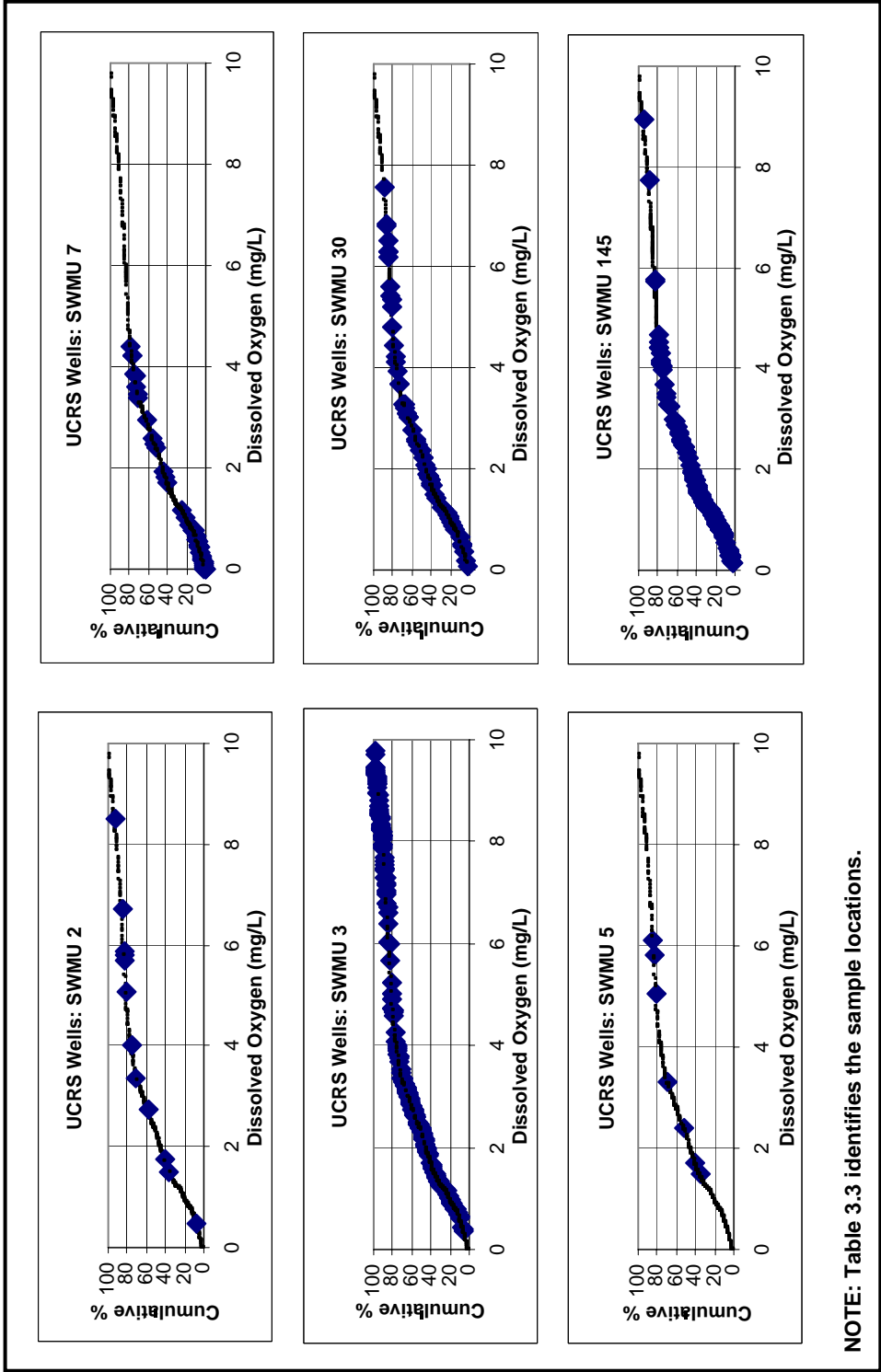
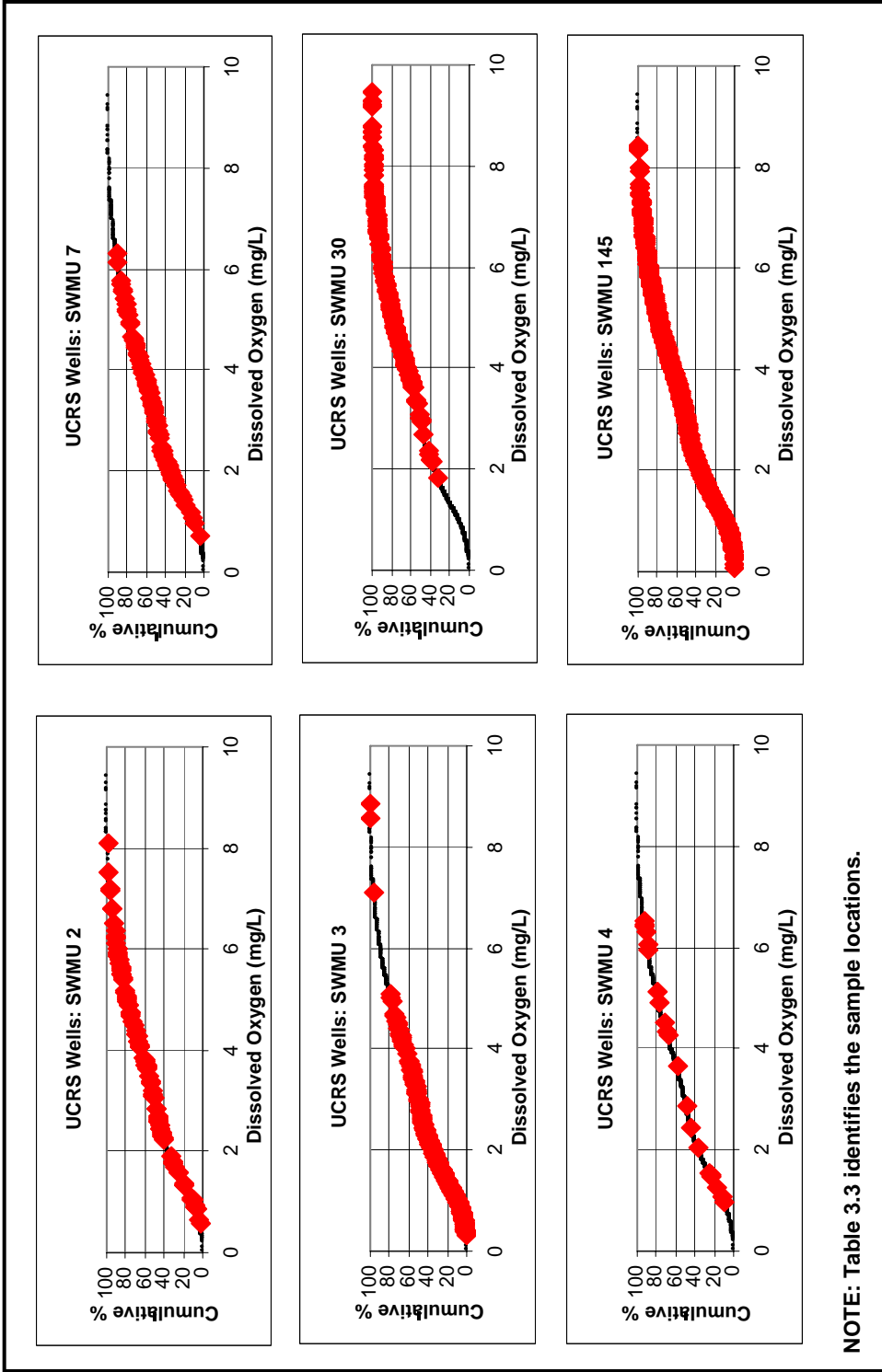


Figure 3.8. Distribution of Dissolved Oxygen in the UCRS and RGA for the BGOU SWMUs



NOTE: Table 3.3 identifies the sample locations.

Figure 3.9. Dissolved Oxygen Measurements in the UCRS for Each BGOU SWMU



NOTE: Table 3.3 identifies the sample locations.

Figure 3.10. Dissolved Oxygen Measurements in the UCRS for Each BGOU SWMU

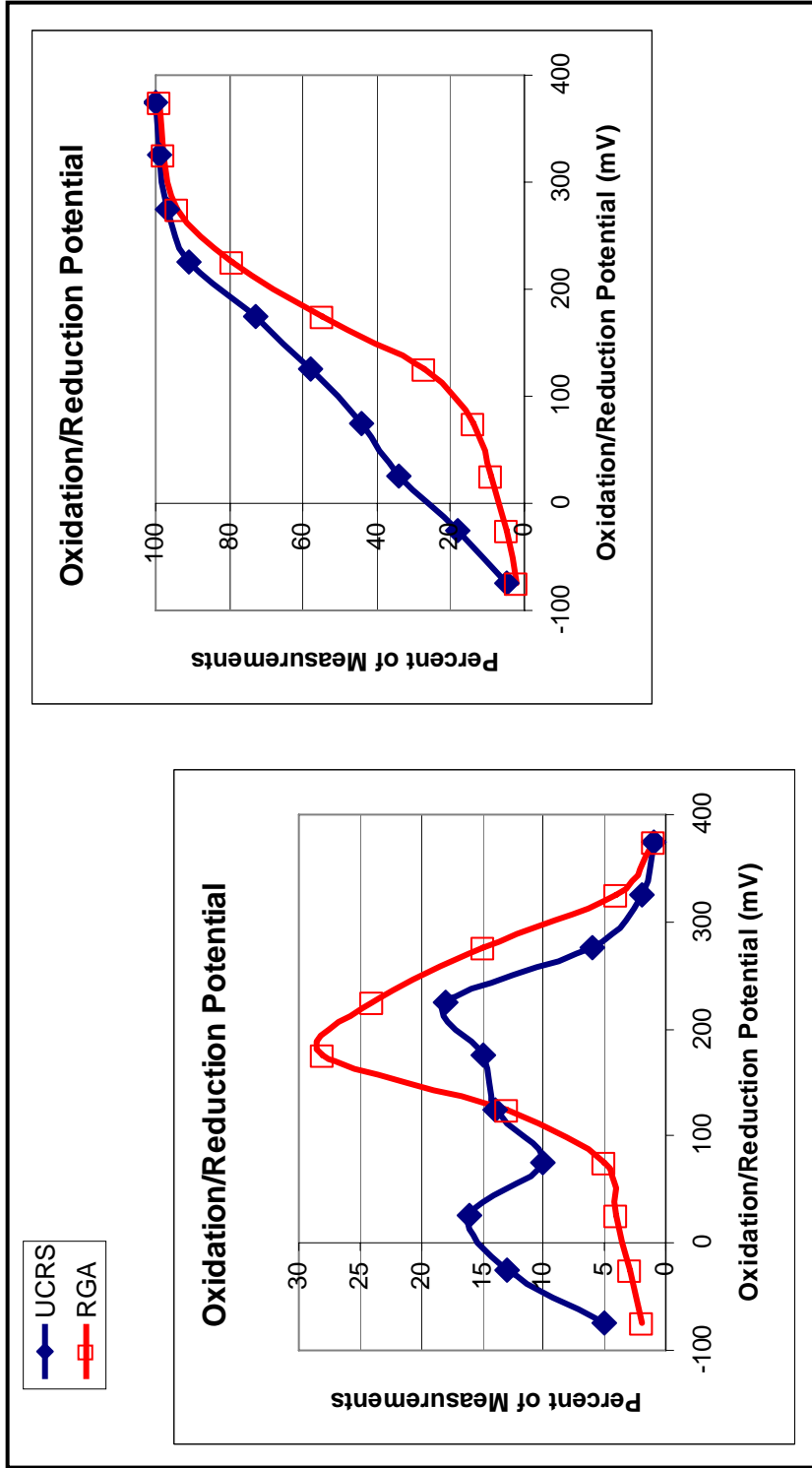
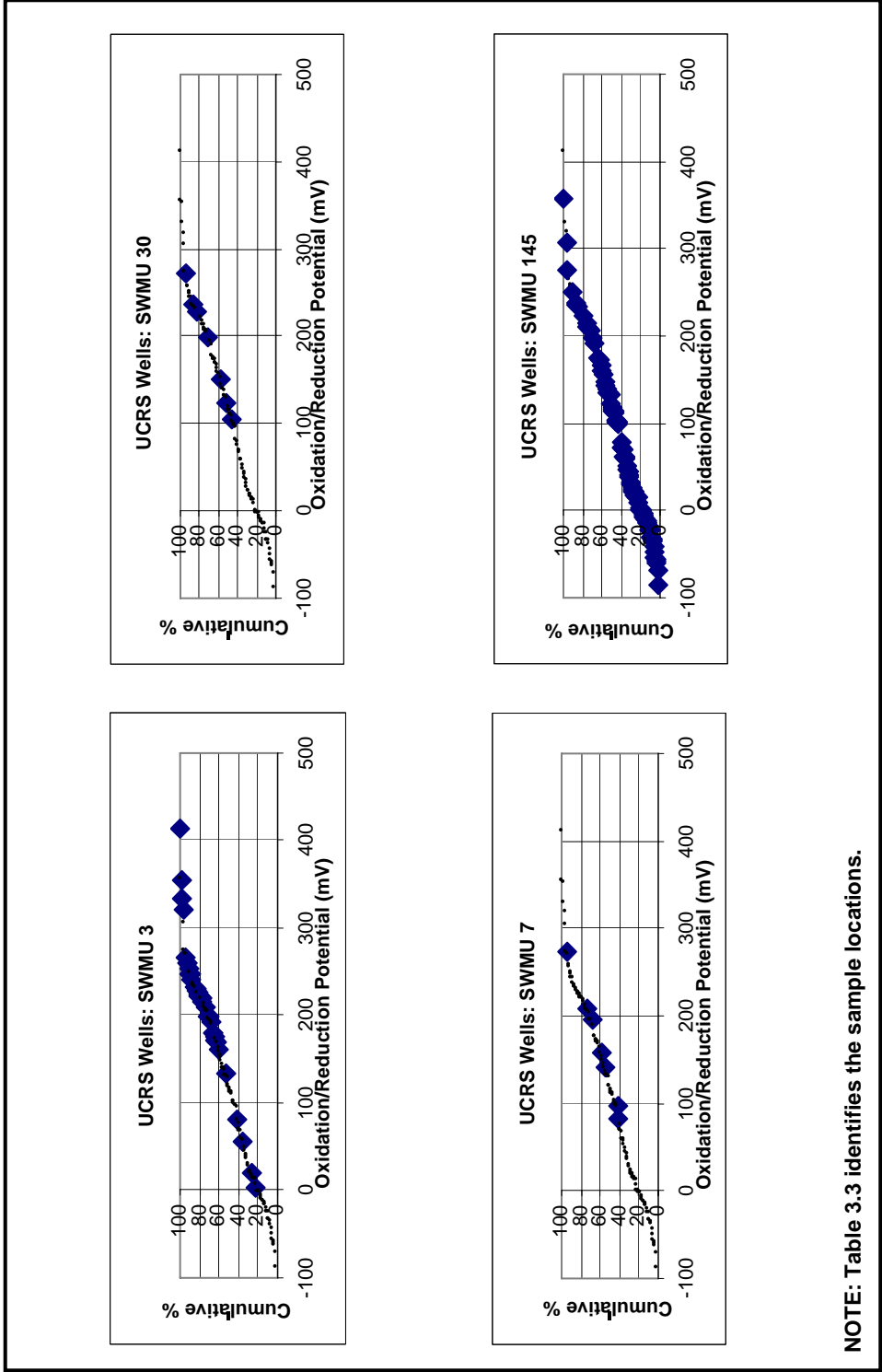
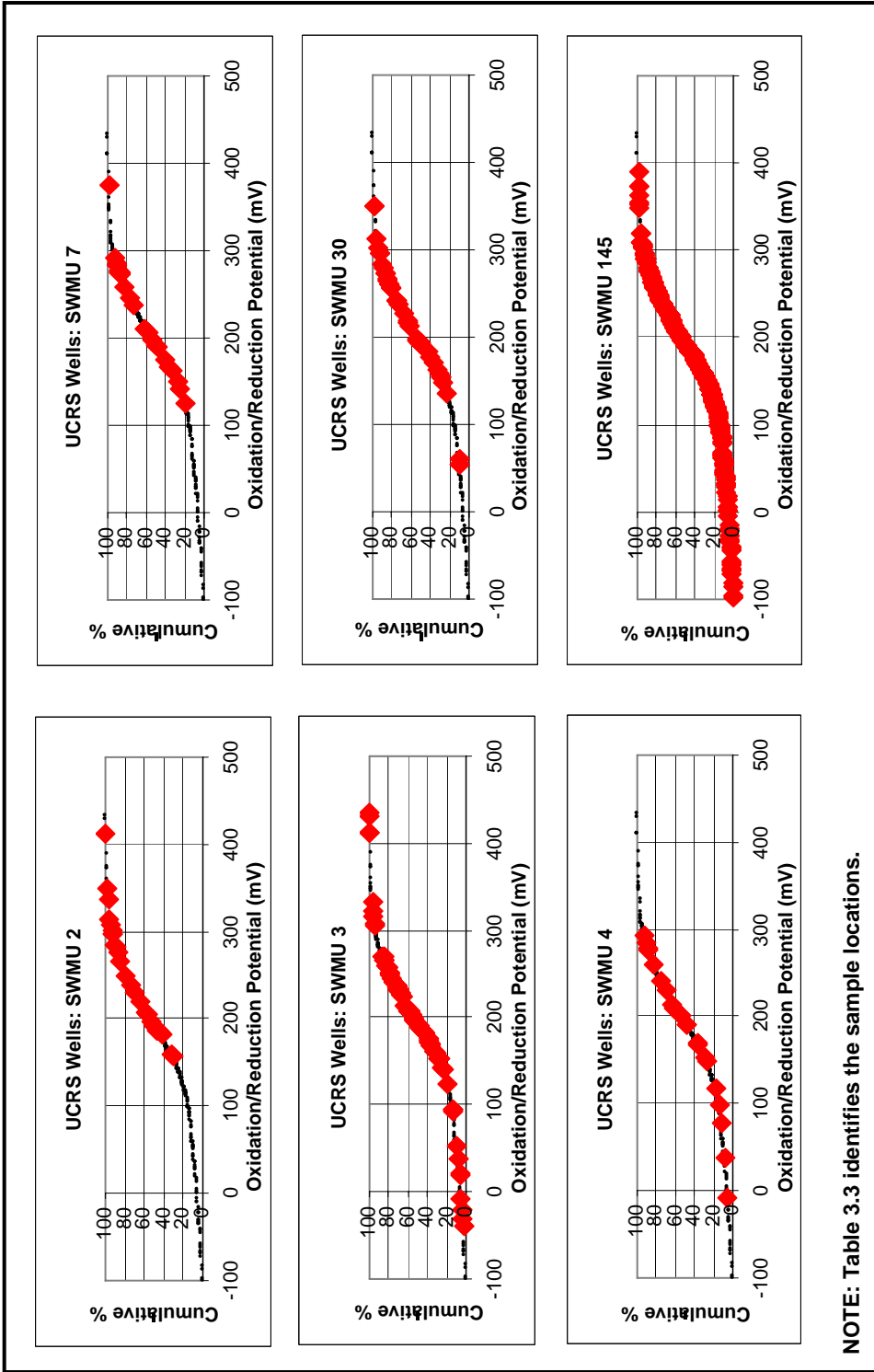


Figure 3.11. Distribution of Oxidation/Reduction Potential in the UCRS and RGA for the BGOU SWMIUS



NOTE: Table 3.3 identifies the sample locations.

Figure 3.12. Oxidation/Reduction Potential Measurements in the UCRS for Each BGOU SWMU



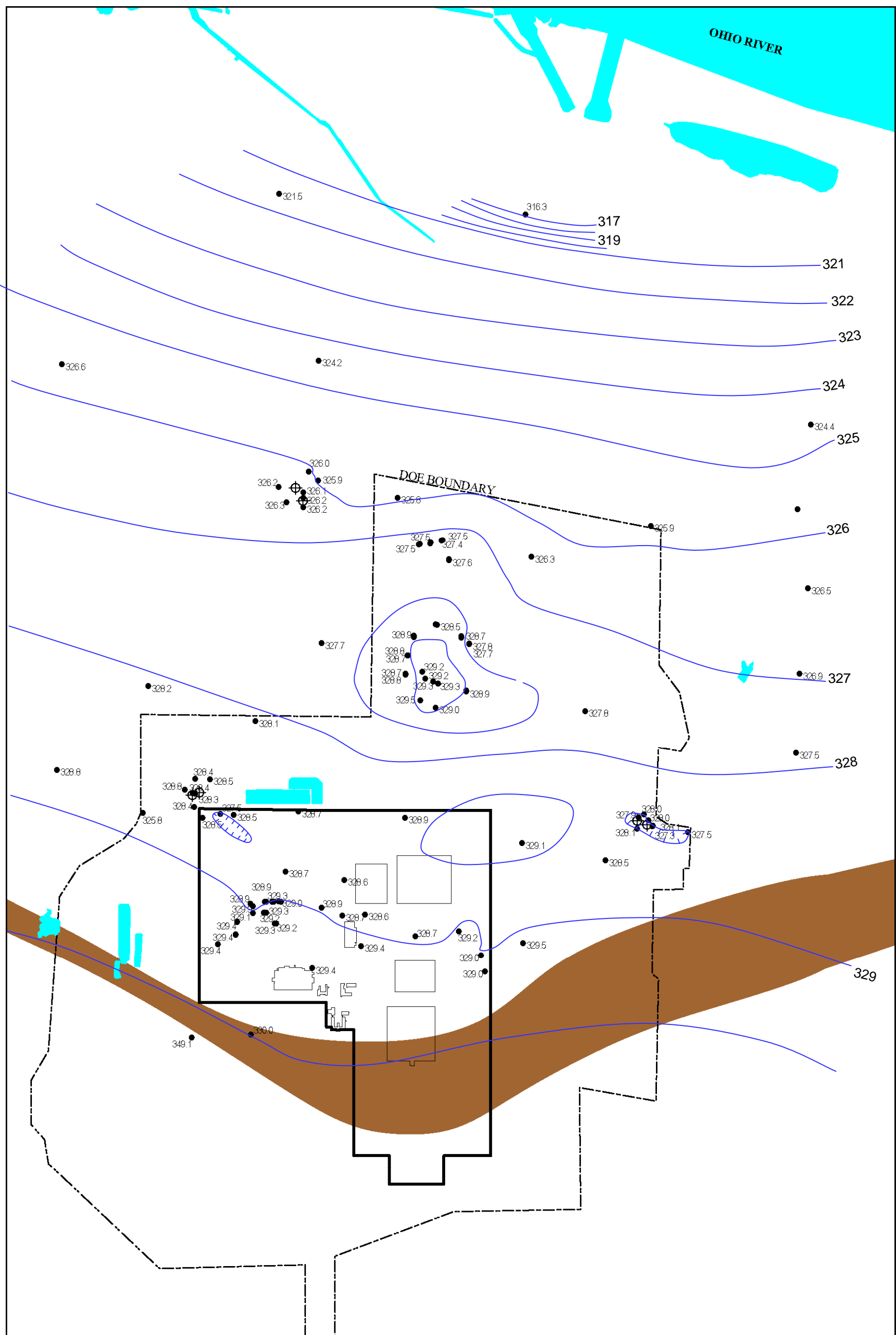
NOTE: Table 3.3 identifies the sample locations.

Figure 3.13. Oxidation/Reduction Potential Measurements in the UCRS for Each BGOU SWMU





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OHIO RIVER

Figure 3.14. RGA Potentiometric Surface, April - June 2007



RGA Potentiometric Surface: April - June 2007

-  POTENTIOMETRIC CONTOUR
-  MONITORING WELL
-  EXTRACTION WELL
-  TERRACE SLOPE

2000 0 2000 Feet



U.S. DEPARTMENT OF ENERGY
 DOE PORTSMOUTH/PADUCAH PROJECT OFFICE
 PADUCAH GASEOUS DIFFUSION PLANT
 COORDINATE SYSTEM: PGDP
 FILE NAME: WaterLev.ppt
 DATE: 10/17/2008
 AUTHOR: Paducah Remediation Services, Inc.
 SOURCES: Paducah OREIS

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3.9.3.1 SWMUs 2 and 3 hydrogeologic interpretation

Waste Disposal Background. SWMUs 2 and 3 are adjacent waste disposal facilities located in the west-central portion of the plant. PGDP buried uranium and uranium-contaminated waste in cells excavated to depths of 7 to 17 ft at SWMU 2. SWMU 3 (C-404) operated as a rectangular, aboveground, surface impoundment from approximately 1952 until 1957, when PGDP converted the surface impoundment to a solid waste disposal facility for uranium-contaminated wastes. (See Sections 1.3.1 and 1.3.2.)

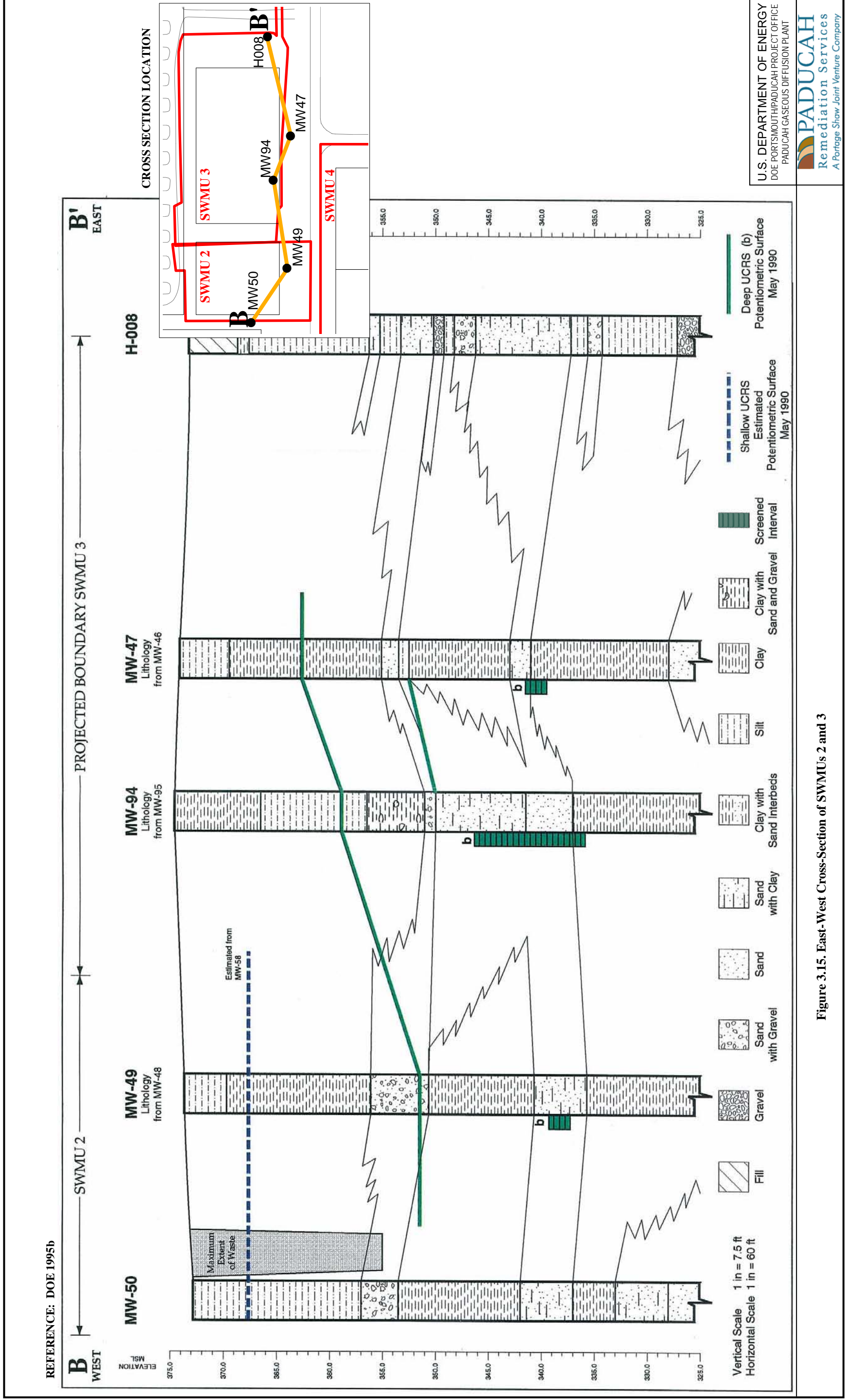
Stratigraphy. The burial cells of SWMU 2 are excavated into the HU1 loess member (silt with some clay) of the Upper Continental Deposits. Some waste cells likely extend to near the base of the HU1 unit, at a depth of 18.5 ft. The underlying HU2 interval consists of upper and lower sand and gravel horizons, separated by an intervening clayey silt unit, to a depth of 40 ft. A nine-ft-thick silty clay interval (HU3) separates the HU2 sand and gravel horizons from the basal HU4 sand and the sands and gravels of the Lower Continental Deposits (HU5). SWMU 3 rests upon the top of the Upper Continental Deposits. East-west cross sections of the stratigraphy below SWMUs 2 and 3 (Figure 3.15, DOE 1995b) demonstrate the relative continuity of the HU2 sand and gravel intervals.

UCRS Groundwater Flow and Hydraulic Potential. Figure 3.16 (DOE 1997a) summarizes the key hydrogeologic parameters that govern groundwater flow through the UCRS at SWMU 2 (the parameters also are applicable to SWMU 3, but SWMU 3 doesn't have burial pits as shown in this figure). The SWMU 2 Interim Remedial Design Investigation Report (DOE 1997a) documents the depth and gradient of the water table using measurements from shallow monitoring wells and piezometers. Four rounds of measurements of water level during a one-week period in August, 1996 consistently demonstrate that the water table occurred within 10 ft of land surface, sloping toward a ditch on the west side. Most of the buried waste at SWMU 2 is saturated. The SWMU 2 report (DOE 1997a) also reported some shallow groundwater discharged into the ditch just southwest of the burial cells. The westward slope of the water table below SWMU 2 indicates that the water table must be equally shallow beneath SWMU 3. Because SWMU 3 is an aboveground facility with a RCRA multi-layered cap, the actual saturation level within the waste is unknown; however, with the shallow water table and generation of leachate, it is assumed that all but the base of the landfill wastes are likely unsaturated.⁴ The annual average volume of leachate removed from the SWMU 3 leachate sump from CY 2005 through 2008 was approximately 2,200 gal (this amount of leachate is equivalent to less than 0.1 inches of infiltration per year, which indicates the cap is functioning as intended to reduce infiltration).

Dr. Cary McConnell of the University of Missouri at Rolla modeled groundwater flow at C-404 (McConnell 1992), using the results of a C-404 aquifer test (Terran 1990) and other measures of hydraulic conductivity in the C-404 vicinity (slug tests of wells and permeameter tests of soil core). The groundwater flow model determined the vertical hydraulic conductivity of the upper confining unit to the RGA (HU3) to be 1.2×10^{-3} ft/day and confirmed observations of a steep vertical gradient (approximately 1 ft/ft) in the area's UCRS wells (Figure 3.17).

⁴ The continuing recovery of leachate from the facility indicates that some infiltration occurs and the base of the disposal cell must be saturated.

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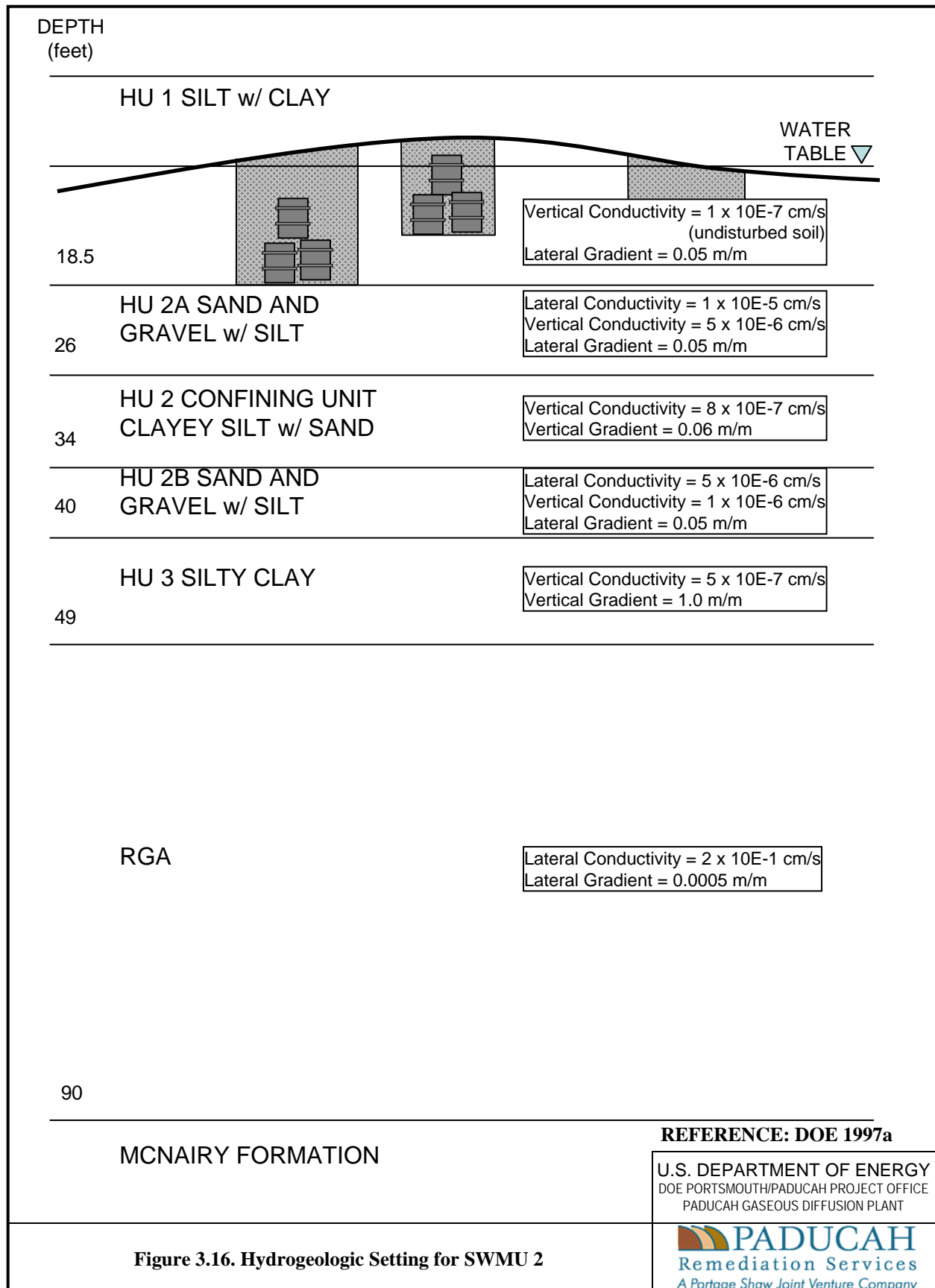
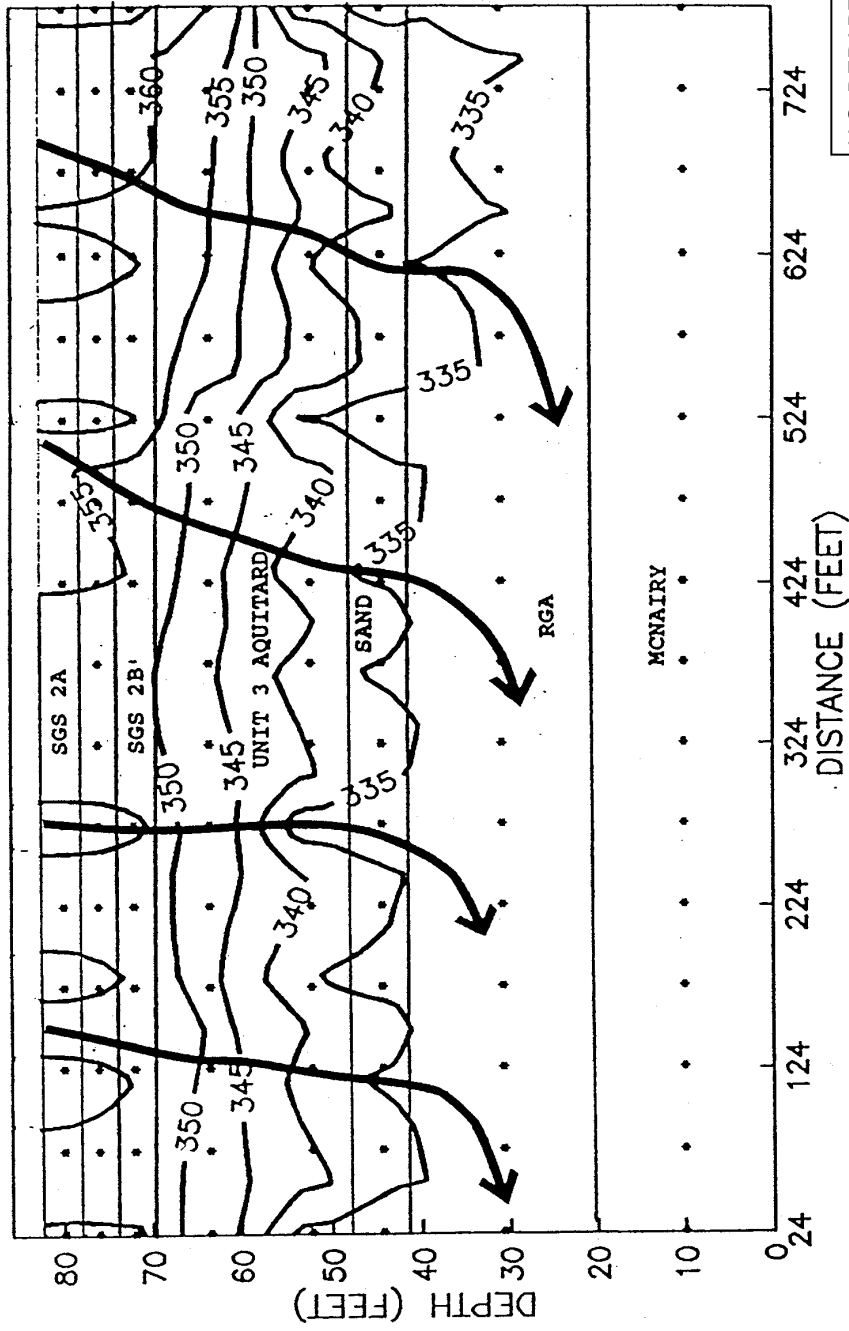


Figure 3.16. Hydrogeologic Setting for SWMU 2

SCALE 1 inch = 75.25 FEET

PLANT NORTH

CROSS SECTION OF HEADS ALONG SIMULATION ROW 9



U.S. DEPARTMENT OF ENERGY
DOE PORTSMOUTH/PADUCAH PROJECT OFFICE
PADUCAH GASEOUS DIFFUSION PLANT



Figure 3.17. Model of the UCRS Vertical Hydraulic Gradient at SWMU 3 (from McConnell 1992)

RGA Groundwater Flow and Hydraulic Potential. The Terran Corporation performed an aquifer test in the C-404 area in September 1989 (Terran 1990) as part of characterization for the Post-Closure Groundwater Monitoring System. Pumping at MW79, located approximately 200 ft east of the northeast corner of C-404, proceeded 48 hours before it was interrupted by a heavy rainfall. The aquifer test documented drawdown as far as 300 ft from the pumping well (extending under C-404); the rate of drawdown indicated that conditions were near steady-state when the aquifer test was terminated.

Test results were consistent with a leaky aquifer setting in which leakance was primarily derived from the underlying McNairy Formation. The 20 ft of drawdown in MW79 induced upward recharge during the aquifer test. Normally, the higher hydraulic potential of the RGA in the C-404 area prevents upward groundwater flow from the McNairy Formation into the RGA.

Analysis of the test results determined that the hydraulic conductivity of the RGA in the C-404 vicinity ranges between 53 and 107 ft/day. Attachment E of the Hazardous Waste Facility Permit, KY8-890-008-982, Modification 4, Effective November 20, 2008, attributes a hydraulic conductivity to the RGA of 21 to 140 ft/day, based on the MW79 aquifer test, another RGA aquifer test from the east side of PGDP (CH2M HILL 1992), and slug tests of RGA wells distributed across the PGDP.

Summaries of measurements of RGA hydraulic potential for C-404 are available from the permit required annual groundwater flow rate and direction within the November version of the C-404 Hazardous Waste Landfill Semiannual Groundwater Reports (C-404 GW Reports). During the 2006—2009 period, the RGA hydraulic potential beneath C-404 has ranged from 323 to 329 ft amsl. Potentiometric maps associated with these reports document a hydraulic gradient that consistently slopes northward, but varies from northeast to northwest (Figure 3.18). A northward-migrating TCE plume that has crossed under the west end of C-404, beginning in 2005, further supports a northward groundwater flow direction.

Each annual report presents the average hydraulic gradient of the RGA at C-404 for the year.

Year	RGA Hydraulic Gradient	
	Upper RGA	Lower RGA
2006	5.89×10^{-3}	8.35×10^{-3}
2007	2.82×10^{-3}	1.25×10^{-3}
2008	8.38×10^{-4}	8.36×10^{-4}
2009	5.60×10^{-4}	NA*

*Beginning with the 2009 C-404 reports, modification of the permit limits characterization of the RGA to upper RGA wells.

The RGA hydraulic gradient at C-400 ranges from 5.60×10^{-4} to 8.35×10^{-3} ft/ft, with an average of 2.93×10^{-3} ft/ft and a median of the gradient measurements of 1.25×10^{-3} ft/ft.

The C-404 Groundwater reports calculate an average yearly groundwater flow velocity as the product of hydraulic conductivity and gradient, divided by the porosity. The RGA is considered to have a porosity of 25%, a typical value for a sandy gravel.

Year	Groundwater Flow Velocity			
	Upper RGA		Lower RGA	
	k* = 21 ft/day	k = 140 ft/day	k = 21 ft/day	k = 140 ft/day
2006	0.49	3.29	0.70	4.68
2007	0.24	1.58	0.10	0.70
2008	0.07	0.46	0.07	0.47
2009	0.05	0.31		

*k = hydraulic conductivity

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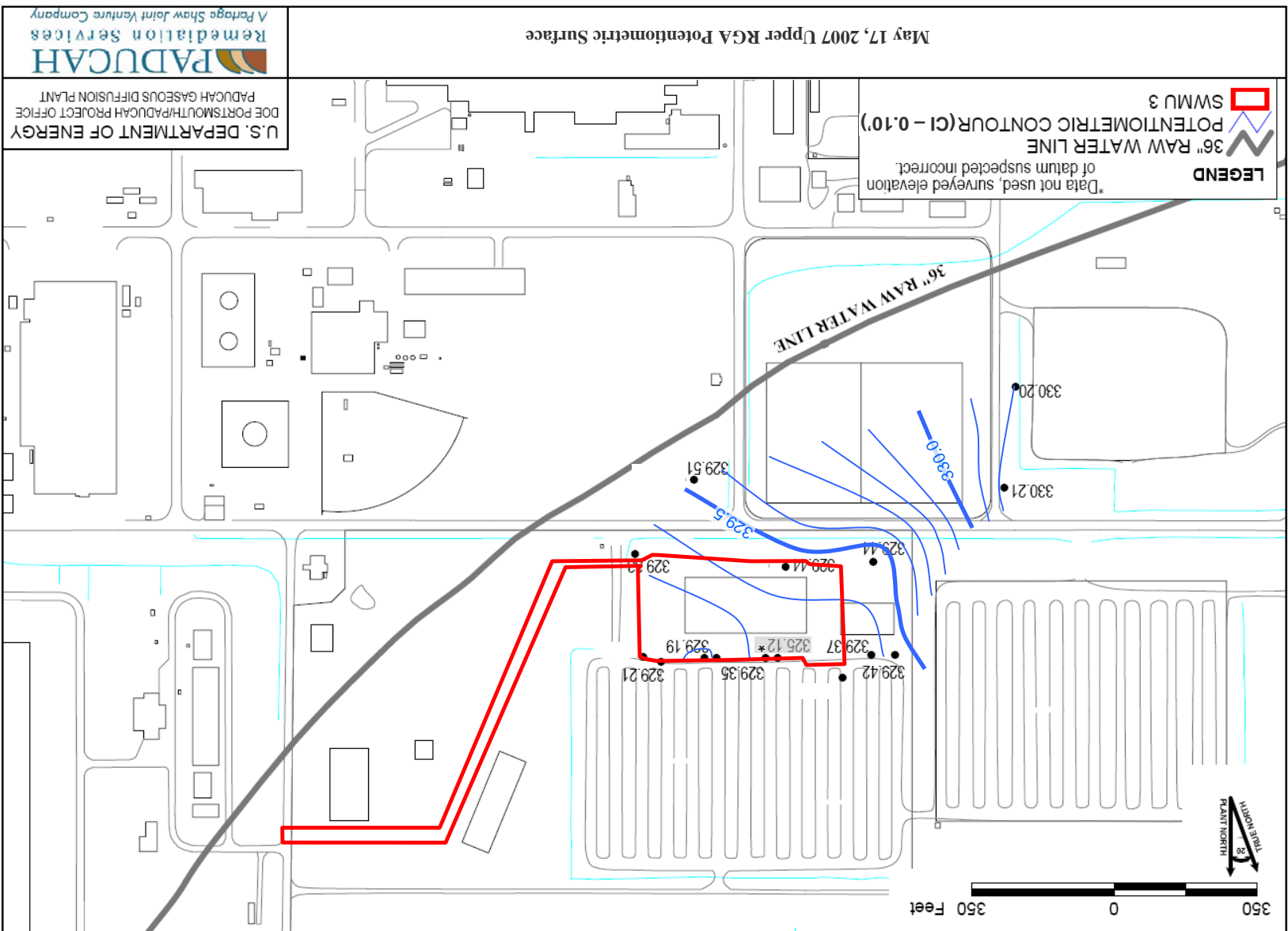
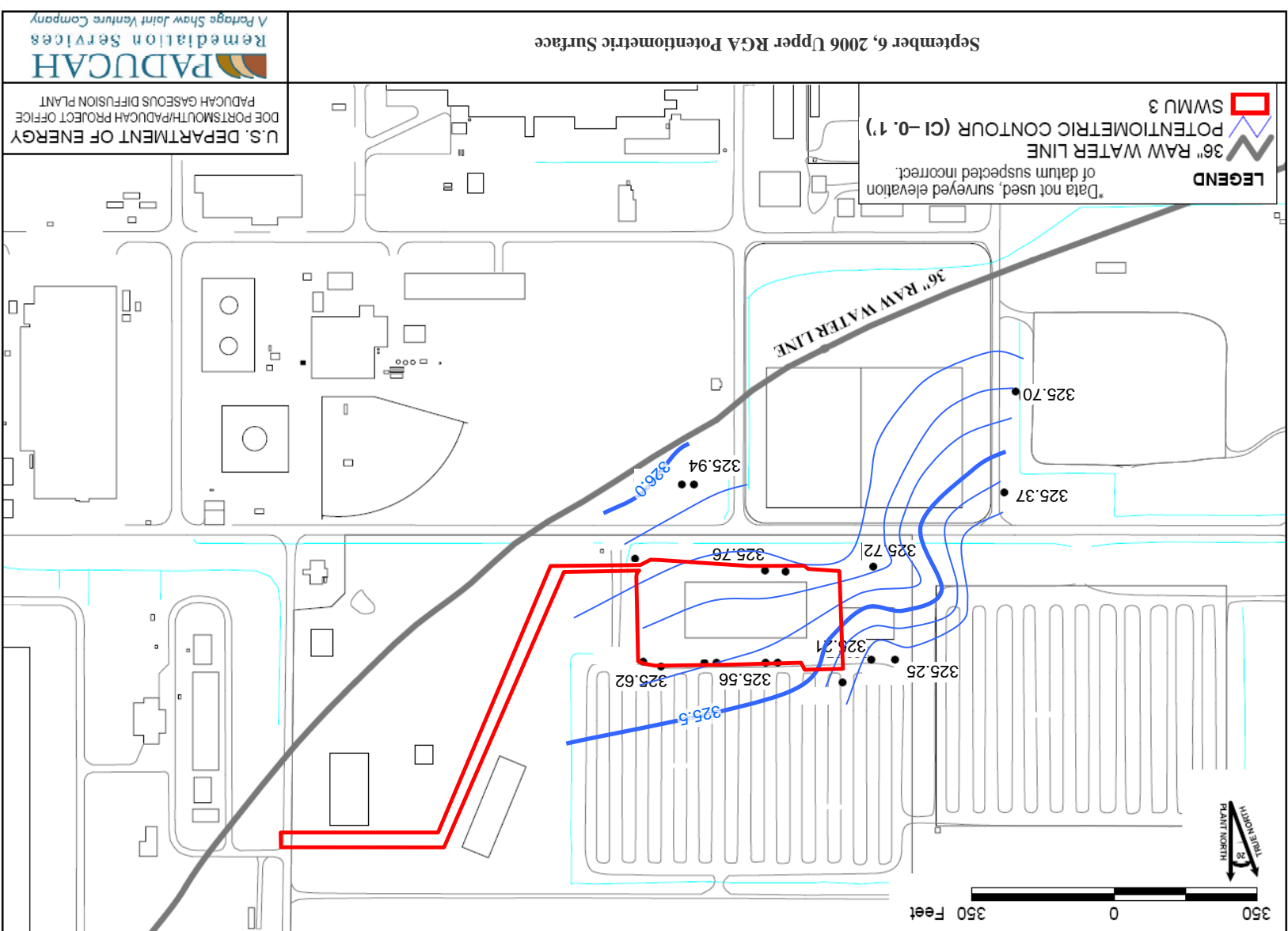


Figure 3.18, Upper RGA Potentiometric Surface at SWMU 3



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The calculated RGA groundwater flow velocities for the 2006—2009 period ranged from 0.05 to 4.68 ft/day, with an average of 0.95 ft/day and a median value of 0.47 ft/day. For comparison, the average RGA groundwater flow velocity in the areas of the contaminant plumes is typically 1 to 3 ft/day.

3.9.3.2 SWMU 4 hydrogeologic interpretation

Waste Disposal Background. SWMU 4 includes four burial pit areas to the south of SWMUs 2 and 3, excavated to a depth of approximately 15 ft for the disposal of various wastes (Section 1.3.3).

Stratigraphy. Like SWMU 2, the burial cells of SWMU 4 penetrate into the HU1 loess member (predominately silt) of the Upper Continental Deposits. These burial cells likely extend to near the base of HU1, at a depth of 15 to 20 ft. Lithologic logs of wells MW415 and MW417 document the presence of an upper and lower HU2 sand horizon, separated by an intervening silt member beneath SWMU 4. The HU2 occurs over the approximate depths of 20 to 40 ft. This, in turn, is underlain by the HU3 silt interval down to a depth of 50 ft. The HU4 sand is approximately 15 ft thick at SWMU 4. Sand and gravelly sand members of the Lower Continental Deposits (HU5) extend down to a depth of approximately 100 ft. The underlying McNairy Formation consists of fine sands and clays. Cross sections based on the numerous soil borings of the WAG 3 RI demonstrate the lateral continuity of these units beneath SWMU 4 (Figure 3.19, taken from DOE 2000a).

UCRS Groundwater Flow and Hydraulic Potential. The depth to the water table at SWMU 4 is uncertain since there are no direct measurements of the depth of the water table beneath SWMU 4. Since the stratigraphy and hydrogeology is comparable to that of SWMUs 2 and 3, and SWMU 4 is in close proximity to those SWMUs, it is reasonable to assume a similar depth to water in the UCRS (10 to 15 ft bgs). The water table likely extends up into the waste burial pits.

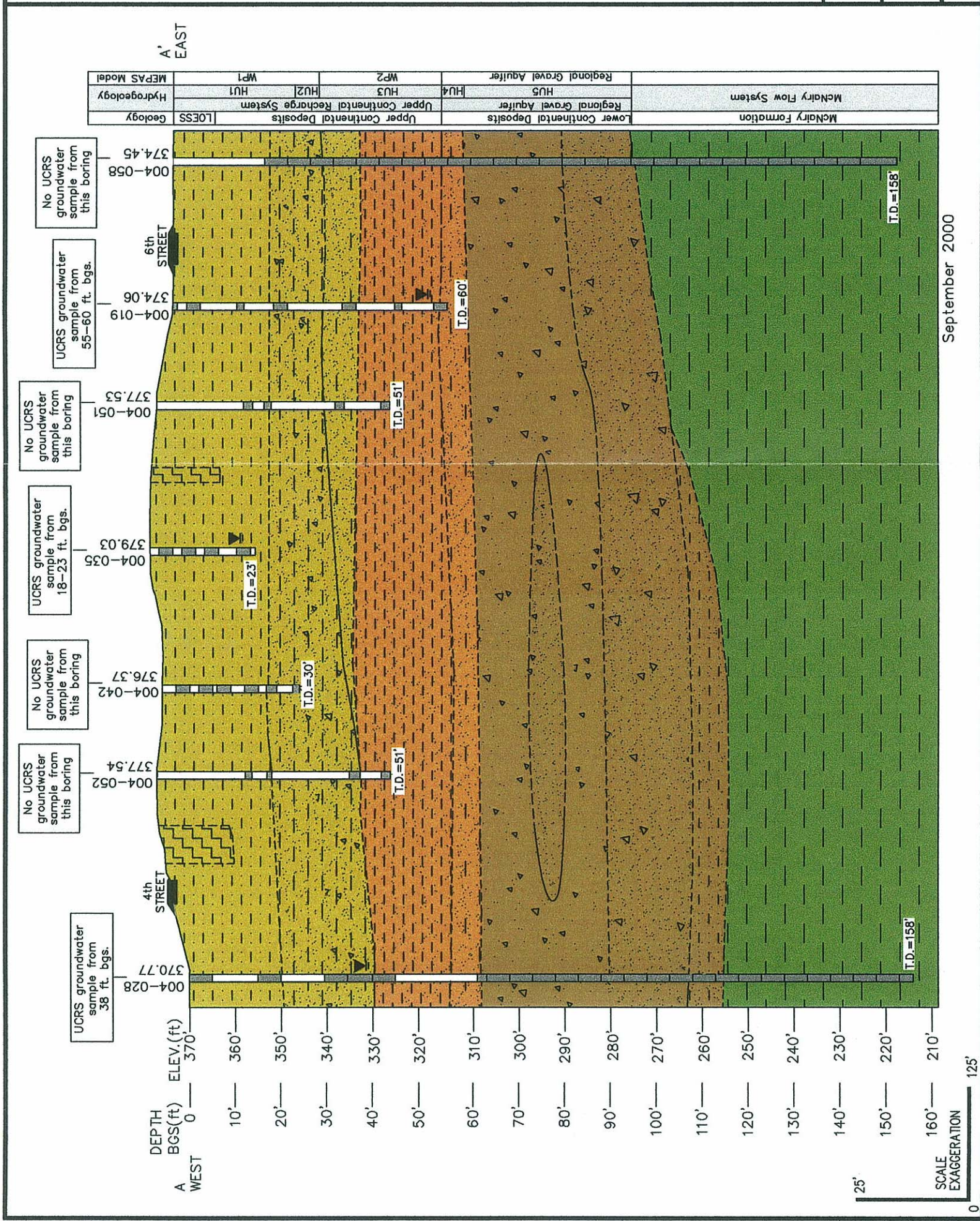
RGA Groundwater Flow and Hydraulic Potential. The northwest flow direction demonstrated for the immediate area to the south of SWMU 3 and the general west-northwest trend of the Southwest Plume define the dominant flow paths in the RGA beneath SWMU 4. It is anticipated that the hydraulic conductivity of the RGA is similar to that of other on-site areas containing the main contaminant plumes, 1,200 to 1,300 ft/day. Average RGA groundwater flow velocity in the areas of the contaminant plumes is commonly 1 to 3 ft/day.

3.9.3.3 SWMUs 5 and 6 hydrogeologic interpretation

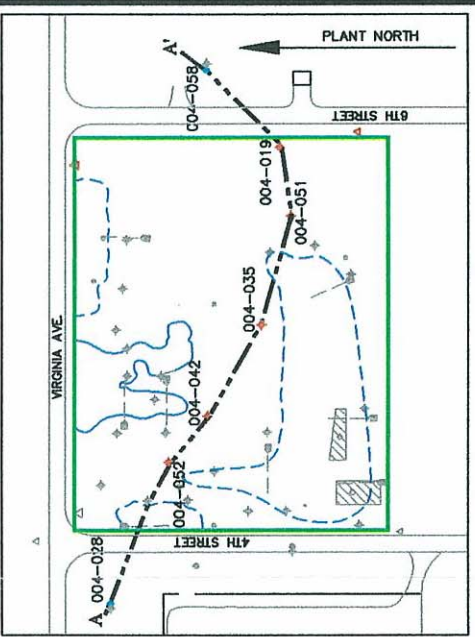
Waste Disposal Background. SWMUs 5 and 6 are adjacent waste disposal facilities near the northwest corner of the PGDP industrial area. Both are burial grounds. The disposal pits of SWMU 5 extend 6 to 15 ft deep (Section 1.3.4). Those of SWMU 6 range from 6 to 8 ft deep (Section 1.3.5).

Stratigraphy. The burial cells of SWMUs 5 and 6 are excavated into the HU1 loess member (silt with some clay) of the Upper Continental Deposits. Only the deeper SWMU 5 pits likely extend to near the base of the HU1 unit, at a depth of 18 to 20 ft. Soil borings of the WAG 3 RI (Figure 3.20, taken from DOE 2000a) document that the HU2 interval in this area is a silty clay with sand and gravel lenses, to a depth of 30 ft below SWMU 6 and 40 ft below SWMU 5. The bottom of the HU3 interval, clay with variable amounts of silt and sand, occurs uniformly at depths of 58 to 60 ft. Soil borings infrequently identified a thin (5 to 7 ft thick) sand interval at the top of the RGA (HU4). In most soil borings, the RGA is a mix of sand and gravel deposits. In the area of SWMUs 5 and 6, the upper McNairy consists primarily of clay, beginning at depths of 100 to 105 ft.

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- LITHOLOGY**
- SILT/CLAY
 - GRAVEL
 - CLAY
 - WASTE CELL
 - SAND AND SILTY SAND AND/OR GRAVEL
- HYDROGEOLOGY**
- HU1 - THE UPPER FLUVIAL/ALLUVIAL SILTY CLAYS RUST BROWN TO GREY-BROWN; PLEISTOCENE AGE.
 - HU2 - LOESS, CLAYEY SILTY SAND TO SANDY SILT. SAND LENSES ARE DISCONTINUOUS BUT CORRELATABLE, RUST BROWN TO YELLOW-BROWN; PLEISTOCENE AGE.
 - HU3 - CLAY TO SILTY CLAY, VERY IMPERMEABLE CLAY THAT ACTS AS A SEMICONFINING TO CONFINING LAYER, PRIMARY CONSTITUENT OF THE UPPER CONTINENTAL DEPOSITS; LOWER PLEISTOCENE AGE.
 - HU4 - SAND WITH GRAVEL AND SILT, CONSIDERED THE UPPER PART OF THE RGA AS PART OF THE LOWER CONTINENTAL DEPOSITS; PLEISTOCENE AGE.
 - HU5 - GRAVEL WITH SAND AND SILT, CONSIDERED THE LOWER PART OF THE RGA AS PART OF THE LOWER CONTINENTAL DEPOSITS; PLEISTOCENE AGE.
- MCNAIRY FORMATION - GREYISH-WHITE TO DARK MICA-FLOW SYSTEM - CEOLUS CLAY, INTERBEDDED WITH LIGHT GREY TO YELLOW-BROWN VERY FINE SAND, CRETACEOUS AGE.**
- NOTE:**
 GRAY INSERTS IN WELLS ARE LOCATIONS WHERE SAMPLES FOR LITHOLOGIC CHARACTERIZATION WERE COLLECTED.
 ▽ GROUNDWATER SAMPLE

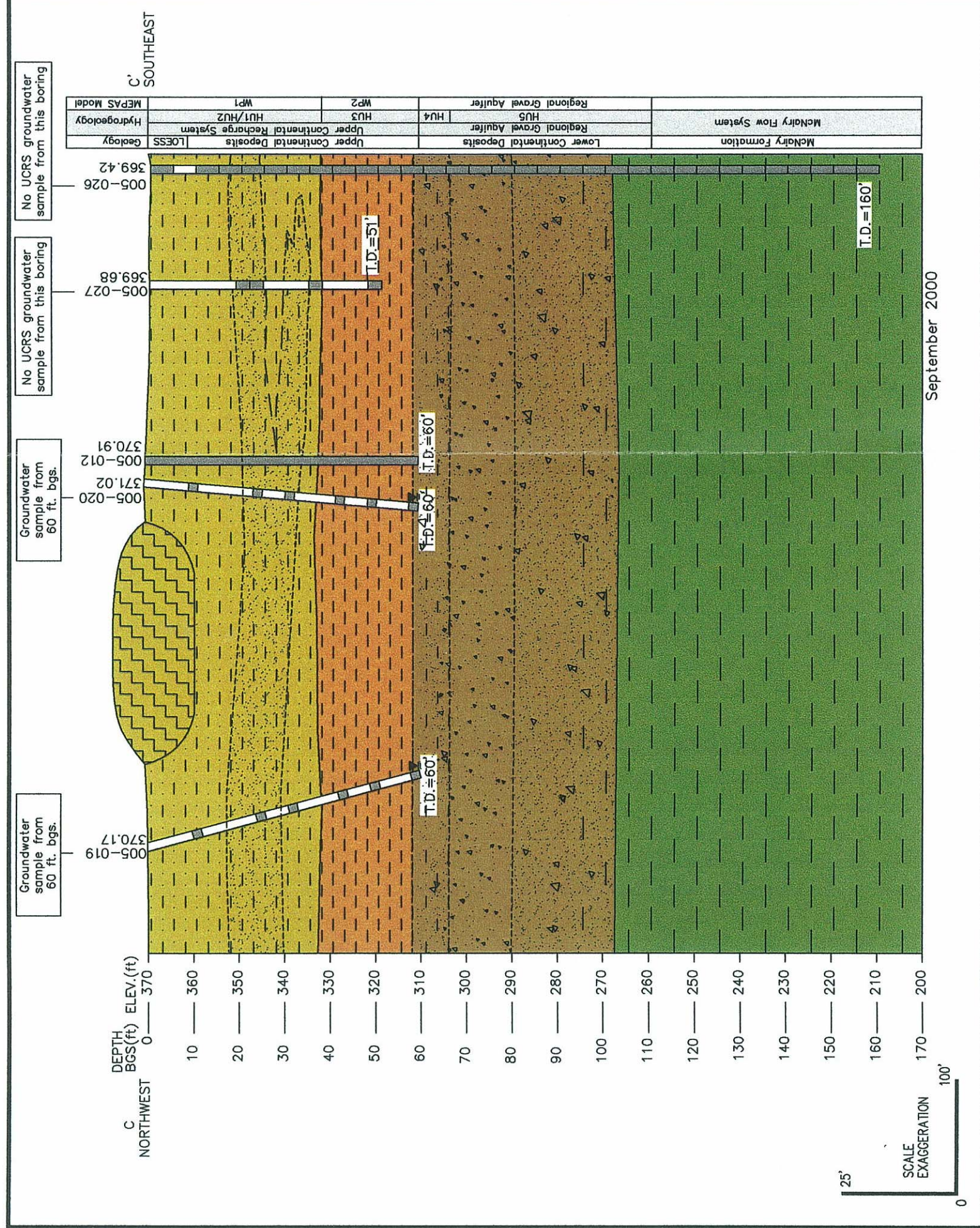


September 2000

SCALE EXAGGERATION 125'

Figure 3.19. WAG 3, SWMU 4 Lithologic Cross-Section A-A'

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Groundwater sample from 60 ft. bgs.

Groundwater sample from 60 ft. bgs.

Groundwater sample from 60 ft. bgs.

No UCRS groundwater sample from this boring

No UCRS groundwater sample from this boring

No UCRS groundwater sample from this boring

McNairy Formation	HU5	Regional Gravel Aquifer	MPAS Model
Lower Continental Deposits	HU4	Regional Gravel Aquifer	MP1
Upper Continental Deposits	HU3	Upper Continental Recharge System	HU1/HU2
Upper Continental Deposits	HU2	Upper Continental Recharge System	Hydrogeology
Upper Continental Deposits	HU1	Upper Continental Recharge System	Geology

LITHOLOGY

	SILT/CLAY		WASTE CELL
	GRAVEL		SAND AND SILTY SAND AND/OR GRAVEL
	CLAY		

HYDROGEOLOGY

HU1 THE UPPER FLUVIAL/ALLOUVIAL SILTY CLAYS RUST BROWN TO GREY-BROWN; PLEISTOCENE AGE.

HU2 LOESS, CLAYEY SILTY SAND TO SANDY SILT SAND LENSES ARE DISCONTINUOUS BUT CORRELATABLE, RUST BROWN TO YELLOW-BROWN; PLEISTOCENE AGE.

HU3 CLAY TO SILTY CLAY, VERY IMPERMEABLE CLAY THAT ACTS AS A SEMICONFINING TO CONFINING LAYER, PRIMARY CONSTITUENT OF THE UPPER CONTINENTAL DEPOSITS; LOWER PLEISTOCENE AGE.

HU4 SAND WITH GRAVEL AND SILT, CONSIDERED THE UPPER PART OF THE RGA AS PART OF THE LOWER CONTINENTAL DEPOSITS; PLEISTOCENE AGE.

HU5 GRAVEL WITH SAND AND SILT, CONSIDERED THE LOWER PART OF THE RGA AS PART OF THE LOWER CONTINENTAL DEPOSITS; PLEISTOCENE AGE.

MCNAIRY MCNAIRY FORMATION - GREYISH WHITE TO DARK MICA-CEOUS CLAY, INTERBEDDED WITH LIGHT GREY FLOW SYSTEM TO YELLOW-BROWN VERY FINE SAND. CRETACEOUS AGE.

NOTE:
GRAY INSERTS IN WELLS ARE LOCATIONS WHERE SAMPLES FOR LITHOLOGIC CHARACTERIZATION WERE COLLECTED.

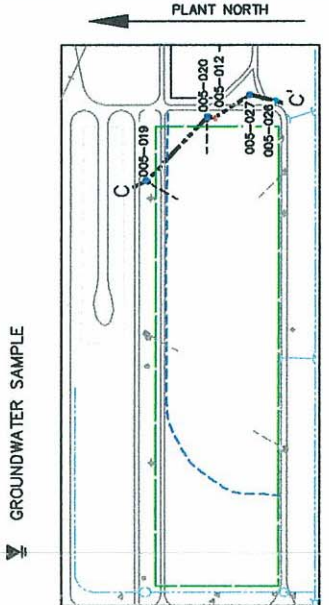


Figure 3.20. WAG 3, SWMU 5 Lithologic Cross-Section C-C'

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UCRS Groundwater Flow and Hydraulic Potential. MW190, screened over the depth interval 17.5 to 22.5 ft bgs (elevation of 348.6 to 353.6 amsl) provides a direct measure of the hydraulic potential in HU2 on the north side of SWMU 5 and an approximation of the elevation of the water table in HU1. With the exclusion of one data outlier (in 59 measurements), the depth of the MW190 water level below ground surface ranges from 1.1 to 6.5 ft bgs. The average elevation of measured water levels in MW190 is 367.3 ft (3.8 ft bgs).

The base of the ditch on the south side of SWMUs 5 and 6, with a local elevation of 358 ft amsl, is a primary control on the elevation of the water table in the area (Figure 3.21). Because the ditch is a linear east-west discharge feature, the area's shallow groundwater flow is likely oriented north-south. The north-south distance between MW190 and the ditch is 350 ft. The difference in elevation of the average MW190 water level and the base of the ditch is 9 ft; thus, the gradient of the water table across SWMU 5 (and similar to that of SWMU 6) is oriented southward with an approximate value of 9/350 ft/ft (0.03 ft/ft). Because HU1 has low transmissivity, the gradient of the water table will tend to be less on the north side of SWMU 5 (although still southward) and significantly greater on the south side of SWMU 5 adjacent to the ditch.

The shallow depth to water in well MW190 (average of 4.1 ft) determines that the vertical hydraulic gradient within the HU1/HU2 hydrogeologic system must be negligible; thus, groundwater flow in HU1 in the area of SWMUs 5 and 6 has a south-oriented vector with a minimal vertical component. The limited shallow groundwater flow beneath SWMU 5 must discharge to the ditch.

Waste was buried to depths of 15 ft (approximate elevation of 355 ft) in SWMU 5; thus, at a minimum, the deepest buried waste cells are saturated over the bottom 3 ft of depth (358 ft amsl/base of ditch–355 ft amsl/base of waste). Assuming a minimal southward gradient of the water table across most of SWMUs 5 and 6, even the shallowest wastes (with top near 365 ft amsl) are likely buried below the water table (at an elevation of approximately 367 ft amsl on the north side of SWMU 5).

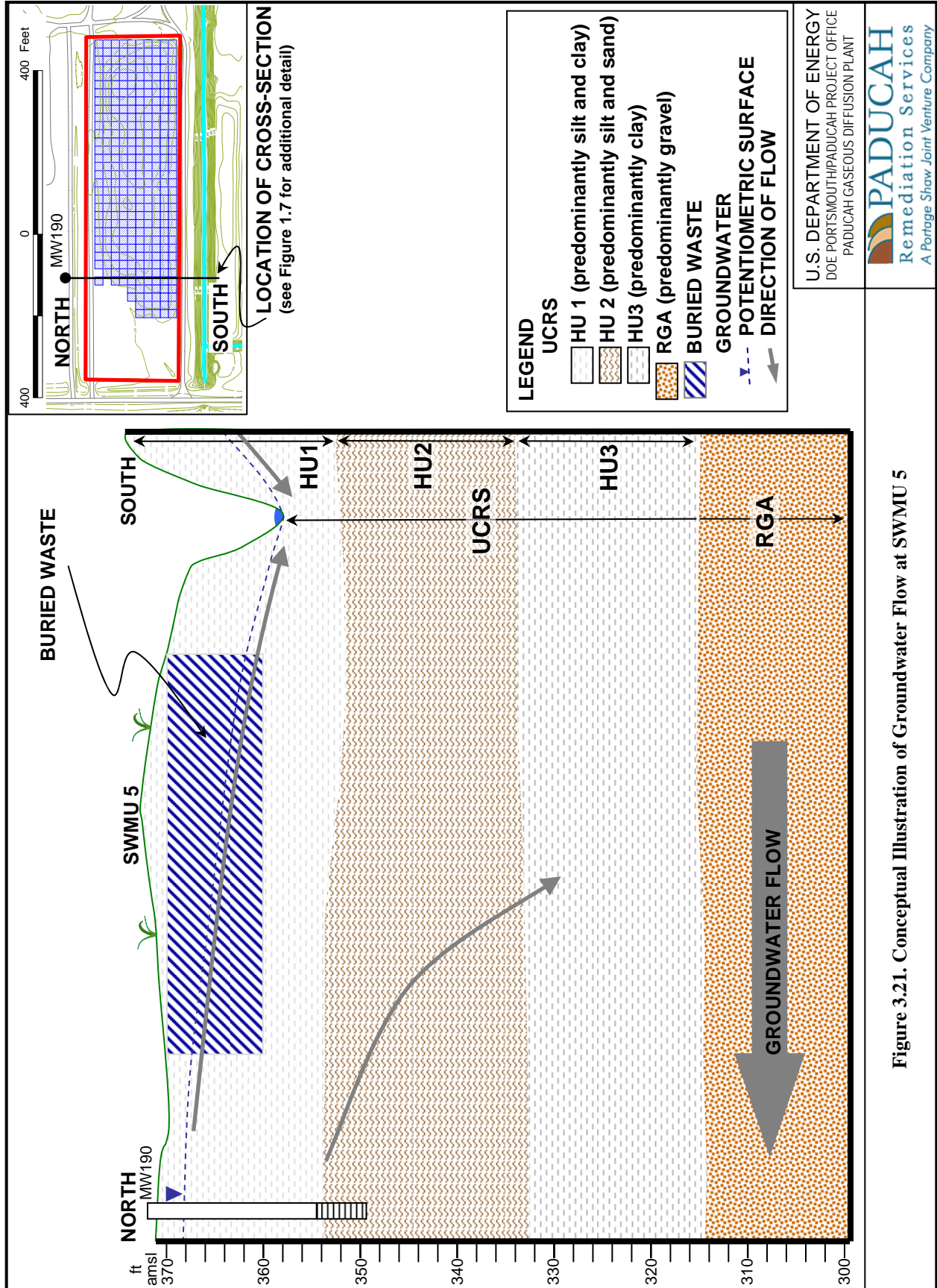
RGA Groundwater Flow and Hydraulic Potential. The high-concentration core of the Northwest Plume passes immediately to the east of SWMU 6 in the RGA. This plume vector defines the direction of RGA groundwater flow below SWMUs 5 and 6. It is anticipated that the hydraulic conductivity of the RGA beneath SWMUs 5 and 6 is similar to that of other on-site areas containing the main contaminant plumes, 1,200 to 1,300 ft/day. Average RGA groundwater flow velocity in the areas of the contaminant plumes is commonly 1 to 3 ft/day.

3.9.3.4 SWMUs 7 and 30 hydrogeologic interpretation

Waste Disposal Background. SWMUs 7 and 30 (C-747-A) are located in the extreme northwest corner of the industrial area of the plant. Both SWMUs are burial grounds. SWMU 7 consists of five distinct burial pit areas that range from 6 to 15 ft deep. (See Section 1.3.6) PGDP buried waste to 12 ft deep at SWMU 30. (See Section 1.3.7)

Stratigraphy. Like all other on-site BGOU SWMUs, the HU1 silt interval contains the burial cells of SWMUs 7 and 30. The base of HU1 is at a depth of 20 ft, approximately 5 ft below the deepest of the burial cells (SWMU 30). A single sand and gravel horizon, in a clay matrix, defines the underlying HU2 interval. The sand and gravel deposits commonly range between 5 and 10 ft thick. Silt and clay members, with a cumulative thickness of 20 to 35 ft, comprise the HU3 interval below SWMUs 7 and 30.

In the area of SWMUs 7 and 30, the RGA consists of an intermittent HU4 sand overlying 20 to 40 ft of the HU5 sand with gravel layers. The top of the RGA commonly occurs at depths of 45 to 60 ft (Figure 3.22, taken from DOE 1998a).



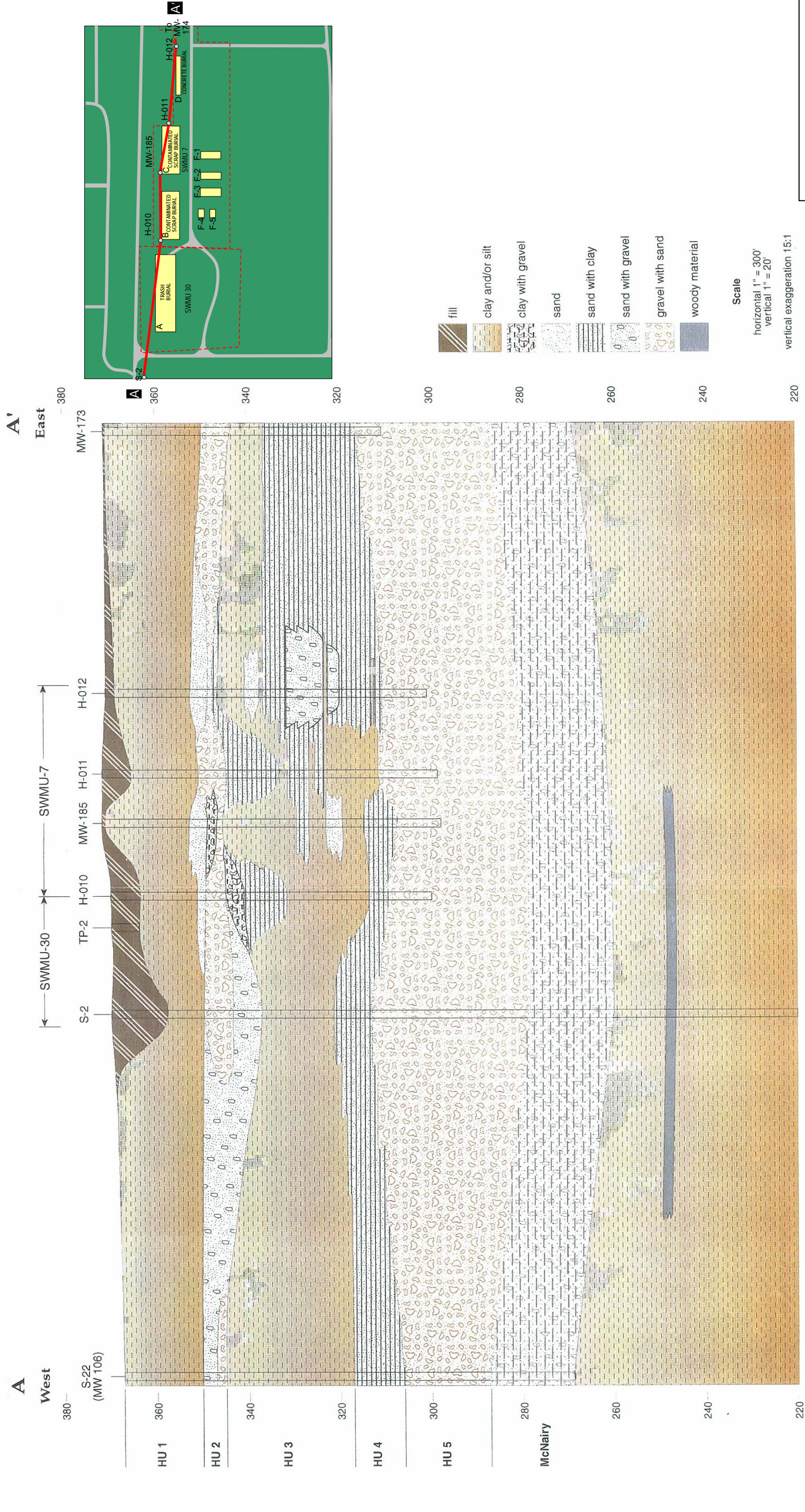


Figure 3.22. SWMUs 7 and 30 Lithologic Cross-Section A-A'

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UCRS Groundwater Flow and Hydraulic Potential. The SWMUs 7 and 30 RI (DOE 1998a) determined that a shallow water table exists approximately 5 ft bgs (Figure 3.23) and within the burial cells. UCRS piezometer and well measurements documented a strong downward gradient within the area UCRS. The vertical downward hydraulic gradient is more than 10 times the lateral hydraulic gradient at SWMUs 7 and 30. This, along with lack of connectivity with shallow sand and gravel strata, leads to predominantly downward groundwater flow through the UCRS. These trends determine that dissolved contaminants from the burial grounds have potential to migrate into the RGA.

The elevation of the water table is above the elevation of the ditches that bound SWMUs 7 and 30 on the north and south sides;⁵ however, although seepage into the bounding ditch just north of SWMUs 7 and 30 was noted following a precipitation event, neither ditch gains significant flow along the reaches adjacent to SWMUs 7 and 30. The SWMU 7 and 30 RI Report (DOE 1998a) concluded that uranium activity concentrations in the ditch sediments suggest SWMUs 7 and 30 are contributing to contamination in the ditch, but the uranium isotope activity ratios in surface water in the ditch argued against waste burial pit waters as contributors to surface water contamination. The increased radiological activity is best explained by surface erosion carrying soil-bound radionuclides to the ditch.

These observations indicate that the UCRS groundwater flow vector must be oriented steeply downward and that the area contributing infiltration to the ditches typically is limited to a thin border along the ditches.

RGA Groundwater Flow and Hydraulic Potential. The high-contamination core of the Northwest Plume passes beneath the west end of SWMU 7 in the RGA. All RGA flow in SWMUs 7 and 30 is to the northwest, as defined by the plume orientation. The south well field of the Northwest Plume containment system is located approximately 650 ft to the northwest of SWMU 7. A pumping test of EW231, an extraction well of the south well field, determines the hydraulic conductivity of the area RGA to be approximately 1,300 ft/day.

The TCE trend in MW66, located near the boundary between SWMUs 7 and 30, exhibits spikes that can be correlated with similar TCE spikes at MW248 in the south well field. The distance between the wells (650 ft) divided by the time lag between TCE “events” in MW66 and MW248 (6 months) defines the local groundwater flow rate to be 3.5 ft/d (Figure 3.24). Typical groundwater flow rates in the Northwest Plume are thought to range from 1 to 3 ft/day. The RGA groundwater flow velocity beneath SWMUs 7 and 30 is accelerated by groundwater extraction in the south well field.

3.9.3.5 SWMU 145 hydrogeologic interpretation

Waste Disposal Background. The waste disposal practices of SWMU 145 (Area P) remain largely undocumented. Anecdotal evidence and historical aerial photographs are sufficient to show that PGDP contractors used the area for disposal of site-related construction debris as early as the construction period of the plant (circa 1952), continuing into the early 1980s. Approximately once a year, plant personnel moved the accumulated scrap into consolidated piles or earth depressions and, wherever practicable, covered them with dirt (Section 1.3.8). By 1973, the disposal area covered approximately 23 acres. Today the area underlies the C-746-S and -T Landfills complex, an area of 44 acres. Area P and the overlying C-746-S and -T Landfills complex form an isolated hill that rises 20 to 40 ft above the surrounding countryside, located 0.25 mile north of the PGDP industrial area.

⁵ The bottom elevation of the ditches on the north and south sides of SWMUs 7 and 30, as well as well and piezometer measurements within SWMUs 7 and 30 provided definitive control of the water table in those areas. The trends of the water table on the east and west ends of SWMUs 7 and 30 were assumed to resemble the land topography.

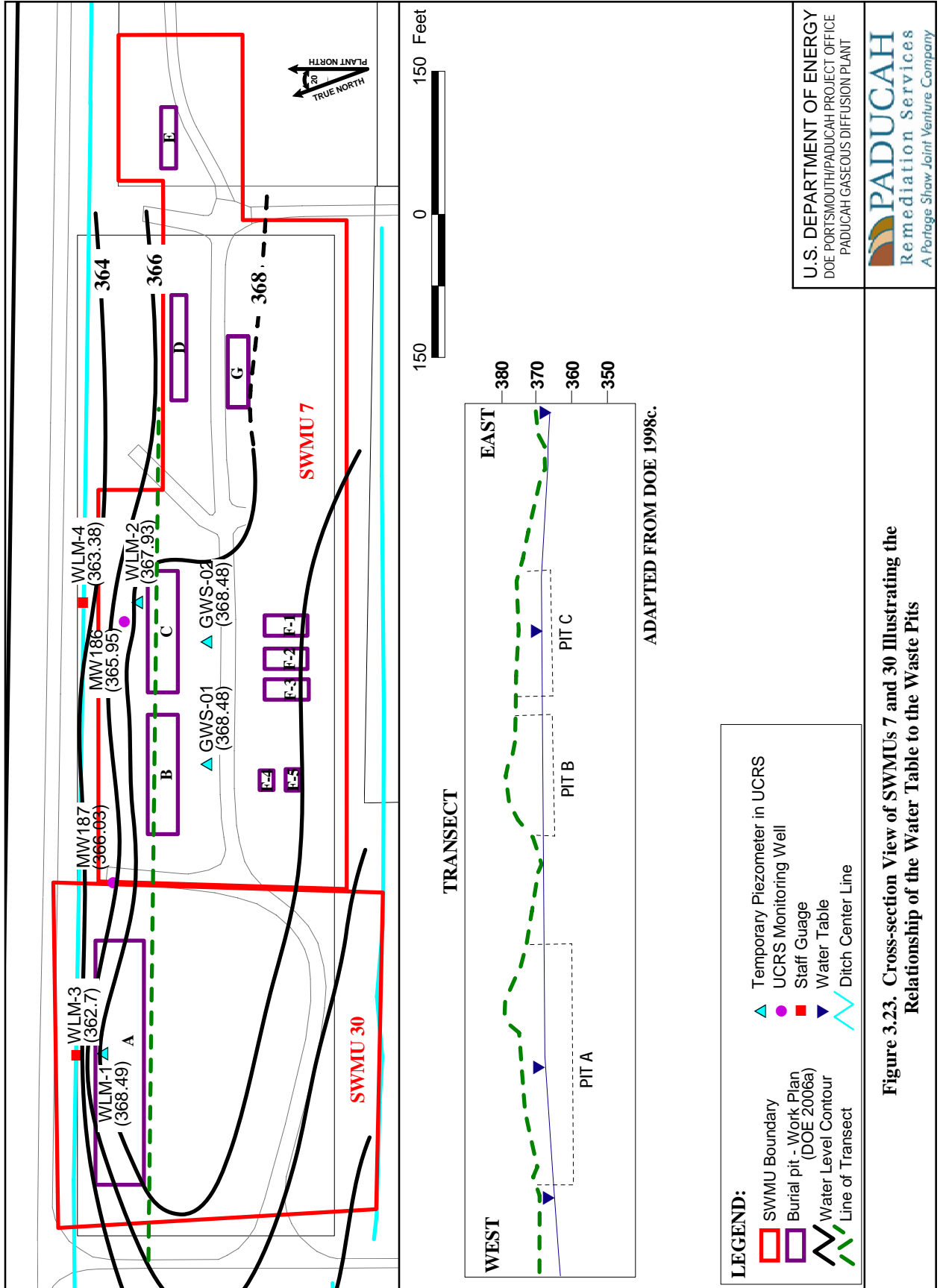


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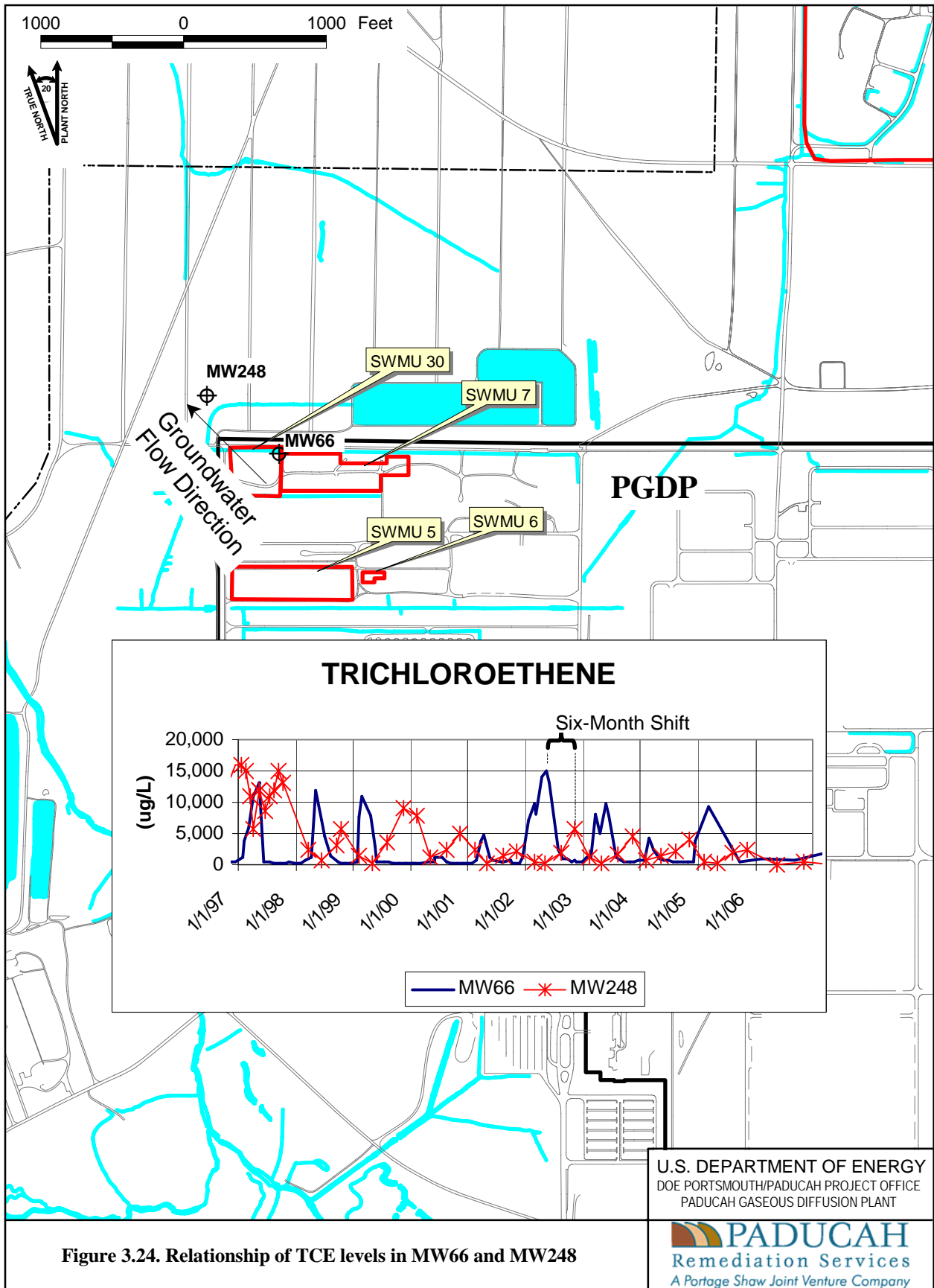


Figure 3.24. Relationship of TCE levels in MW66 and MW248

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Stratigraphy. The UCRS beneath SWMU 145 typically consists of a near continuous sequence of silt members down to the top of the RGA at depths of 40 to 60 ft. A thin (commonly less than 1 ft thick), intermittent, sand horizon at a depth of approximately 20 ft is the only vestige of the HU2 interval. The C-746-S and -T Landfills SI (DOE 2006c) determined that the top of the RGA has approximately 20 ft of relief (elevations of 310 to 330 ft) beneath SWMU 145. Where the RGA is deepest, the UCRS grades downward into a series of fine sand layers with silt interbeds overlying the RGA.

An HU4 sand, averaging 5 ft thick, forms the top of the RGA. This, in turn, overlies 20 to 40 ft of gravelly sand, made up of individual sand and gravel layers that range from 0.2 to 3.4 ft thick (Figure 3.25). The underlying McNairy Formation (top at an elevation of approximately 280 ft) consists of interbedded units of silt and fine sand.

UCRS Groundwater Flow and Hydraulic Potential. Water level elevations of shallow wells at SWMU 145 determine that a vertical hydraulic gradient of approximately 1 ft/ft is characteristic of the local UCRS (Figure 3.26). The area SI developed a conceptual water table map for the SWMU 145 area (Figure 3.27).⁶ This figure presents representative water levels from both now-abandoned and current water wells to incorporate the most data possible. These water levels are not directly comparable because of the strong vertical gradient that is present in the UCRS; the depth of each well's screen (which varies considerably across the area of the map) is an important influence on the water levels in the wells. The conceptual water table map uses the elevation and depth of the well screens and the representative water levels to assess the approximate elevation of the water table across the site. Lateral hydraulic gradients range from 0.03 to 0.12 ft/ft horizontally, as measured from the conceptual water table. The area SI analysis determined that lateral UCRS flow may be important where the horizontal hydraulic gradients are steepest, but that vertical flow predominates in the UCRS under most of SWMU 145.

Table 3.4 summarizes the water level measurements in the SWMU 145 area UCRS wells for the years 2007 and 2008. Because of the strong vertical hydraulic gradient that is prevalent at PGDP, of the SWMU 145 area UCRS wells, MW396 alone provides some measure of the water table. In addition, because all of the UCRS wells are located on the perimeter or exterior to the landfill complex, the wells don't characterize the water table in the fill area.

Table 3.4. UCRS Water Level Measurements at SWMU 145 for 2007 and 2008

Well	Screen Interval		Ground Water Level Measurements				
	(Elevation amsl)		# of Measurements	(Elevation amsl)			
	Top	Bottom		Average	Median	Minimum	Maximum
MW371	341.0	331.0	16	342.9	343.2	341.0	344.9
MW386	343.1	333.1	16	345.6	345.3	343.5	348.2
MW389	341.7	331.7	0 ^a	NA ^a	NA ^a	NA ^a	NA ^a
MW390	330.4	320.4	17	325.8	326.5	322.9	329.3
MW393	336.3	326.3	15 ^b	341.0	340.7	339.6	342.5
MW396	343.2	333.2	17	371.0	371.0	366.7	374.5

^a Either the well was dry or the water level was below the base of the well screen (in the well sump) at each of 16 measurement attempts during the 2007 through 2008 period. "NA" means not available.

^b One measurement of 16 during the 2007 through 2008 period is an outlier and is not included in the summary statistics.

⁶ The elevation of the water table remains poorly documented at SWMU 145. Some buried waste at SWMU 145 is likely saturated.

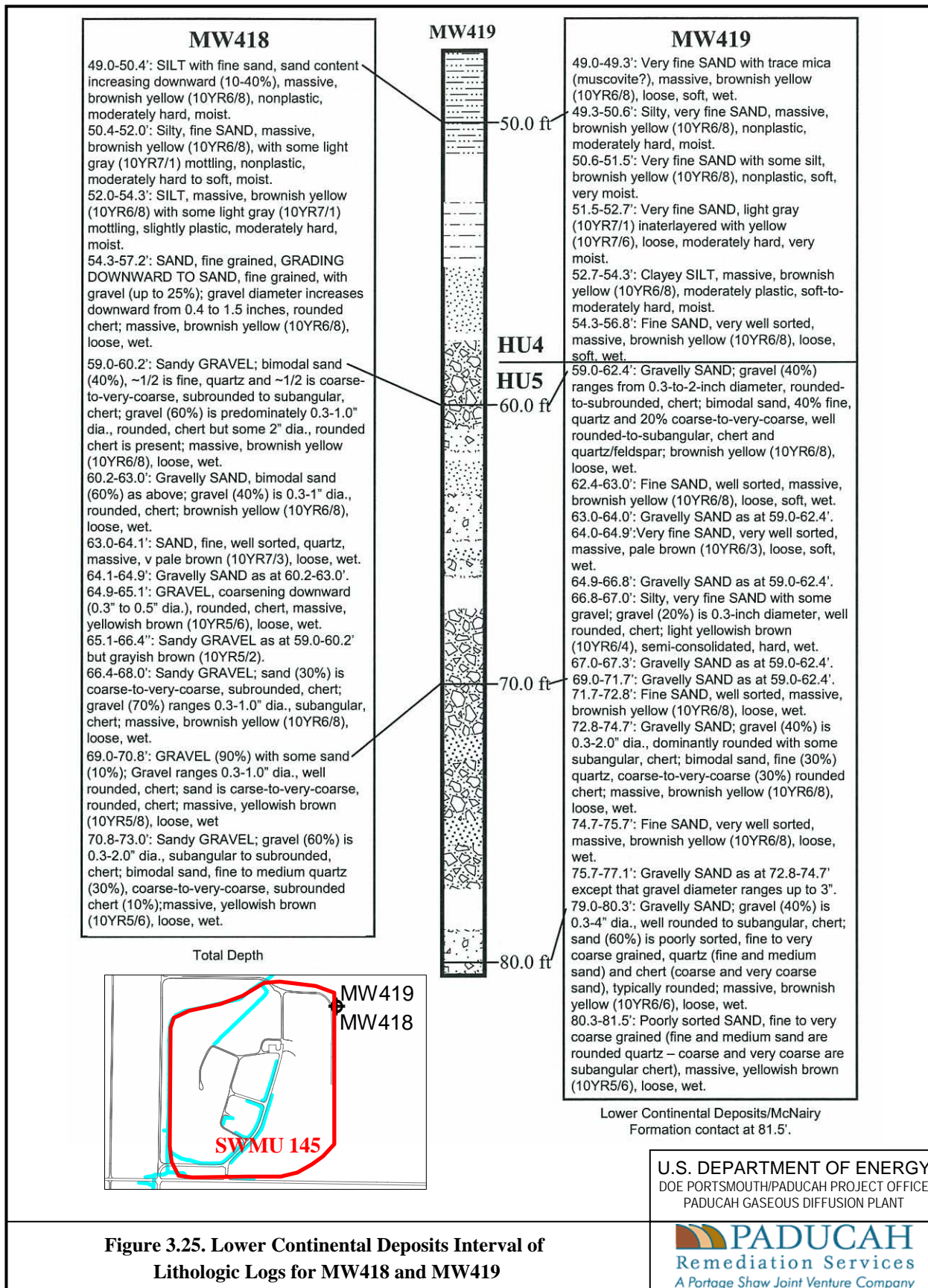


Figure 3.25. Lower Continental Deposits Interval of Lithologic Logs for MW418 and MW419

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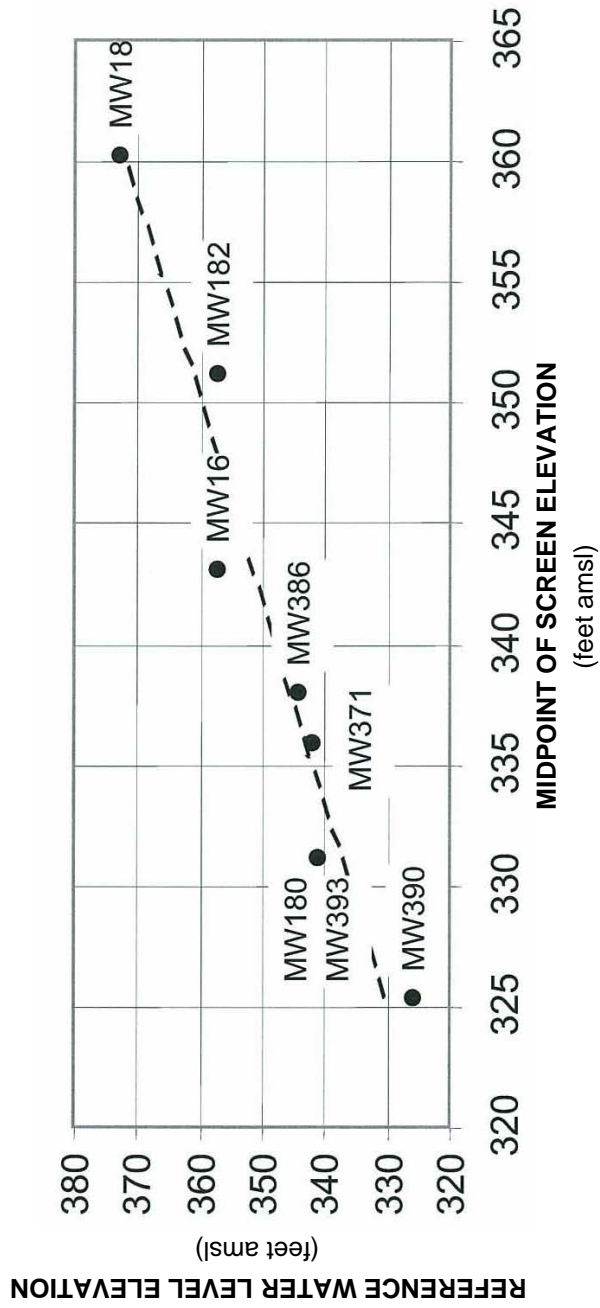


Figure 3.26. UCRS Vertical Gradient

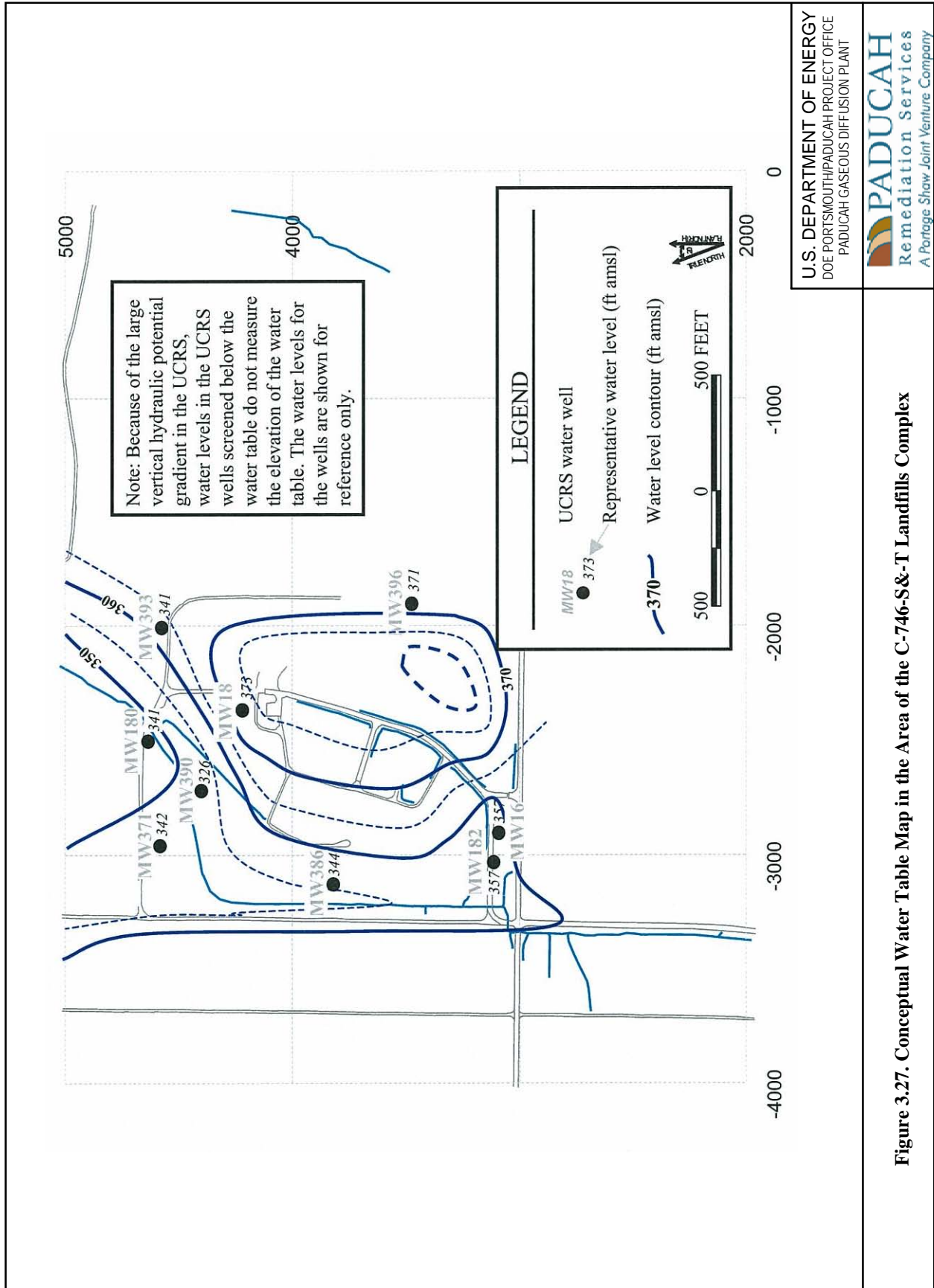


Figure 3.27. Conceptual Water Table Map in the Area of the C-746-S&-T Landfills Complex

Now abandoned MW18 provides water level characterization for the north central area of SWMU 145. The well screen extended over the interval 347.4—367.4 ft amsl. (In this area, the base of the C-746-P Landfill is approximately 375 ft amsl.) Of the 27 measurements of water level in MW18 (for the period February 1988 through July 1994), water level averaged 373.9 ft amsl, with a median of 373.4 ft amsl. The measured water levels ranged from 370.3—373.9 ft amsl. Water level measurements of MW396 and MW18 suggest that the water table occurs near the base of the Area P fill.

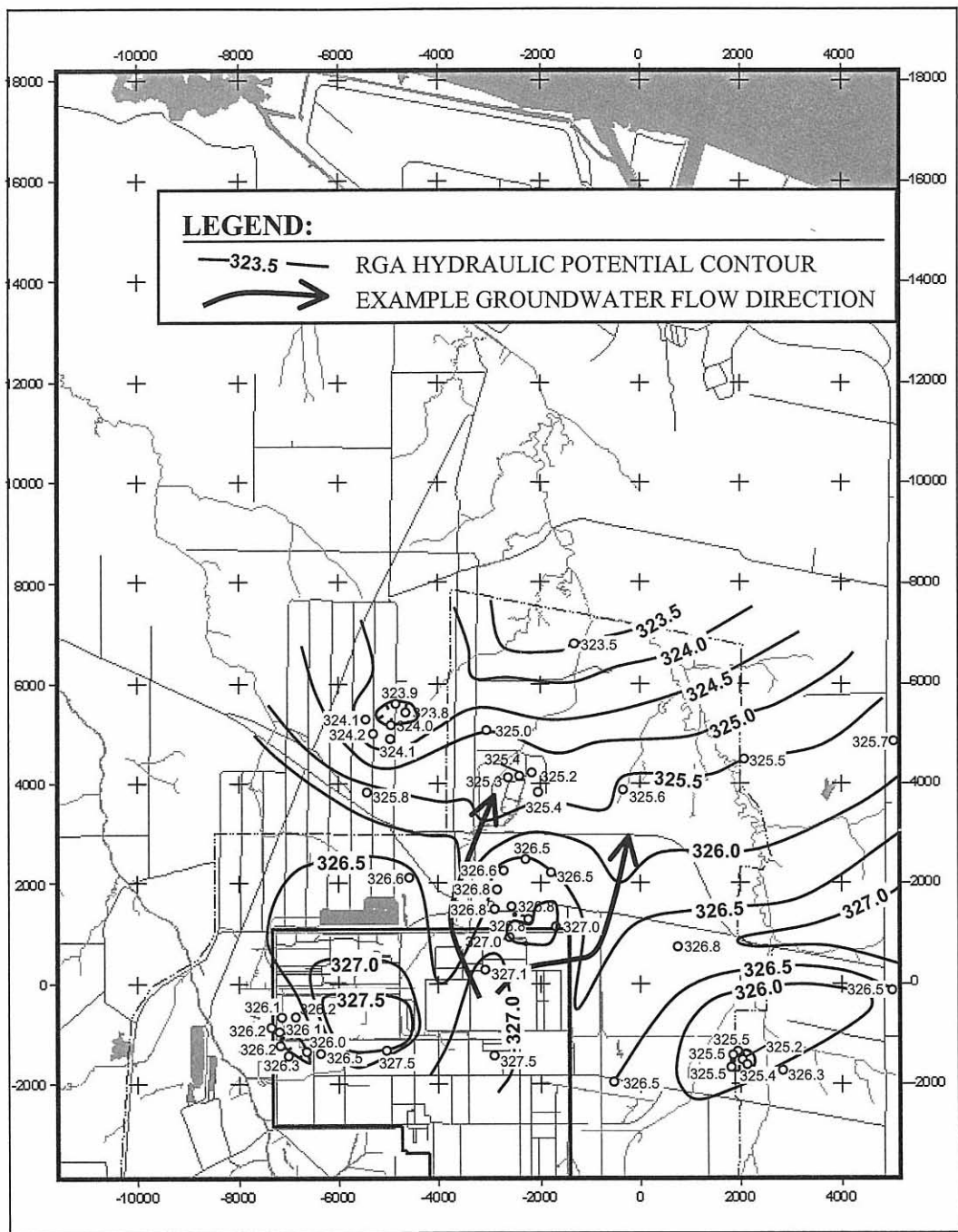
RGA Groundwater Flow and Hydraulic Potential. The regional hydraulic gradient of the RGA in the SWMU 145 area is northward with a typical slope of 10^{-3} ft/ft (Figure 3.28). Water level measurements of RGA wells for the area SI documented the presence of a hydraulic potential mound beneath SWMU 145 (see Figure 4.68). The inferred groundwater flow directions, extending radially from SWMU 145 in the immediate vicinity of the burial ground, were consistent with trends of the direction of dissolved TCE contamination associated with the burial ground.

Groundwater modeling indicates that the hydraulic conductivity of the RGA in the area of SWMU 145 ranges between 200 to 500 ft/d. With the regional hydraulic gradient, average groundwater flow velocity in the RGA should range between 1 and 2 ft/d.

3.9.4 BGOU Hydrogeologic Conceptual Model

Observations from the BGOU RI are consistent with the following conceptual model of the flow system north of the Porters Creek Clay subcrop (Figure 3.29). A shallow water table exists in the area of the on-site BGOU SWMUs. The UCRS is saturated from the water table down. Groundwater flow through the UCRS (HU1, HU2, and HU3) is primarily downward to the top of the RGA (HU4 and HU5). This is due to vertical hydraulic gradients being much greater than lateral hydraulic gradients and also due to a lack of connectivity of the shallow sands and gravel units. Limited lateral dispersion results as groundwater and contaminants migrate vertically through the UCRS. The rate of vertical and horizontal movement (migration) is influenced by the physical properties of a particular contaminant including solubility, specific gravity, and the individual contaminant's affinity to adsorb to the surrounding soils and by the lithology of the individual HUs, most notably the HU1 interval, which contains the burial cells, and the HU3 interval, which serves as the upper semiconfining unit between the UCRS and the RGA. Once groundwater reaches the RGA, then the predominant flow is horizontal. The RGA serves as the primary exit pathway for groundwater from within the PGDP property boundary.

The previously known burial pits and waste cells are the sources of contamination identified in the shallow soils and UCRS groundwater of the BGOU SWMUs. Contaminants are migrating from the waste cells as a result of water infiltrating through the cells into the underlying soils and as a result of water migrating through potential secondary DNAPL sources at SWMU 4 (DOE 2007a) and SWMUs 7 and 30 (DOE 1998a). Once the contaminants reach the RGA, the rate of migration increases as a result of the higher hydraulic conductivity of the RGA sands and gravels. Regional groundwater flow is generally north to northwest in the RGA beneath the BGOU SWMUs.

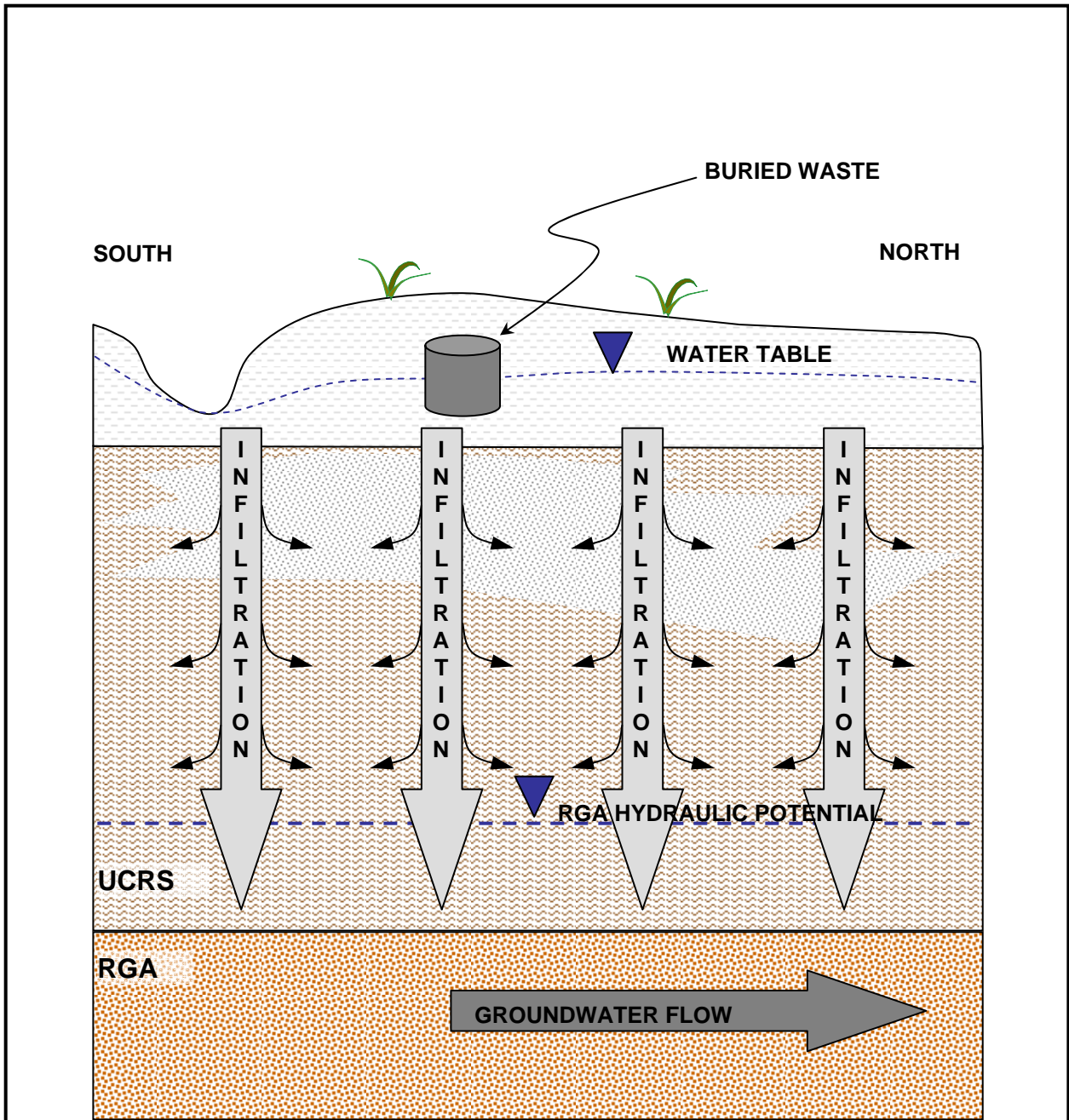


REFERENCE: DOE 2006c





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Figure 3.28. RGA Potentiometric Surface for the C-746-S&T Region, mid-July 2004





LEGEND

-  UCRS (predominantly silt and clay)
-  UCRS sand lens
-  UCRS clay lens
-  RGA (predominantly gravel)

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Figure 3.29. Conceptual Model of the Groundwater Flow System



For SWMU 4, the evidence of DNAPL presence is markedly higher dissolved TCE levels (commonly 1,000 to 4,000 µg/L) in the RGA on the west (downgradient) side of the SWMU. The area of higher TCE levels in the RGA spans the entire western side of SWMU 4, suggestive of a diffuse source of DNAPL contamination in the UCRS soils underlying the burial grounds. Subsurface soil samples also show TCE contamination near the southern burial pit. A discrete area of 10,000 µg/L in the lower RGA also implies the presence of a small pool of DNAPL (zone of higher DNAPL saturation) at the base of the RGA. Figure 3.30, taken from the Southwest Plume SI (DOE 2007a), summarizes the dissolved TCE levels in the RGA on the west side of SWMU 4.

MW66 is an upper RGA well⁷ located near the shared boundary of SWMUs 7 and 30. The analyses of groundwater samples from MW66 reveal abrupt rises of dissolved TCE (Figure 3.31) that commonly occur in the first half of the calendar year, when RGA water levels are highest. In other words, there is a strong correlation of TCE spikes with periods of high hydraulic head. (TCE spikes often exceed 10,000 µg/L.) This suggests that when RGA hydraulic head is high, more TCE in the lower UCRS is dissolved into UCRS groundwater, which then moves downward into the RGA. These high-TCE events typically are limited to years where RGA water level exceeds 324 ft amsl.

In MW66, the contact of the RGA and the overlying UCRS soils occurs at an approximate elevation of 318 ft amsl. The relationship between abrupt rises in TCE levels and high RGA water levels indicates the likely presence of a DNAPL source zone near the boundary of SWMUs 7 and 30 at an elevation of 324 ft amsl, in the silt/clay horizon that overlies the RGA.

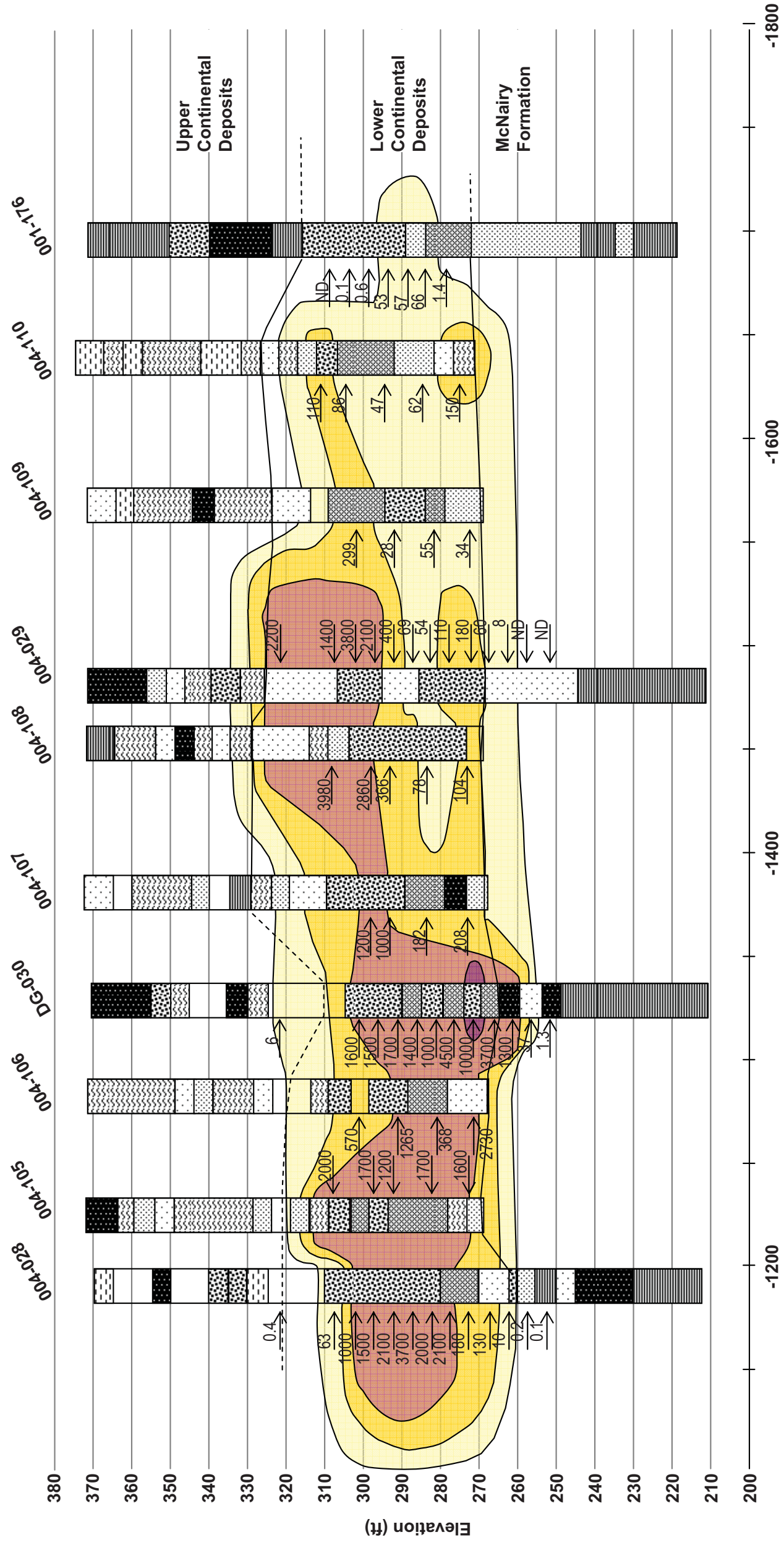
⁷ MW66 is constructed with a 5-ft length well screen installed over the interval 308 to 313 ft amsl.

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S

Gross Lithology and TCE Concentrations (µg/L) along West Side SWMU 4 – Southwest Plume

N



LEGEND

	Sand		Clay
	Fine Sand		Coarse Sand
	Clayey Silt		Gravelly Sand
	Silt		Stiff Clay

Interpreted TCE Depiction

	5- 100 µg/L
	100- 1,000 µg/L
	1,000- 10,000 µg/L
	> 10,000 µg/L

North Plant Coordinate (ft)

REFERENCE: DOE 2007d

001-176 sampled in June 1998;
 004-028, DG-030, 004-029 sampled during WAG 3, Summer 1999;
 All other data from SW Plume SI, Summer 2004

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Figure 3.30. TCE Distribution within the RGA Downgradient of SWMU 4

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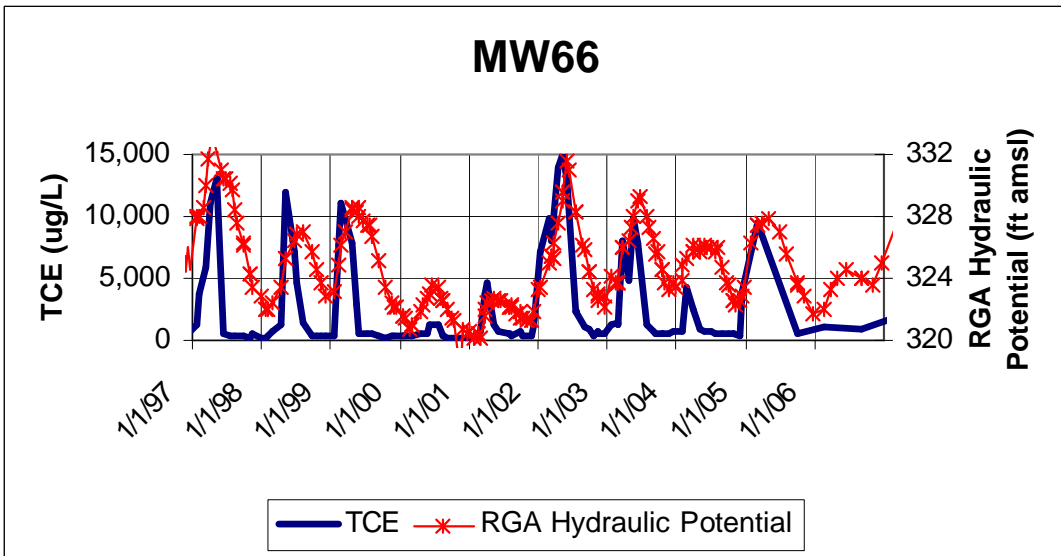


Figure 3.31. TCE Trends in MW66

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4. NATURE AND EXTENT OF CONTAMINATION

The SWMUs comprising the BGOU consist primarily of landfills and below ground burial cells in which various PGDP wastes have been placed. Infiltration descending through the buried waste could mobilize contaminants within the waste. Once mobilized, the most likely pathway of the contaminants would be downward through the UCRS soils, ultimately reaching the RGA (based on the hydrogeologic conceptual model presented in Chapter 3). Some lateral movement of contaminants would occur in the UCRS, but these pathways appear limited since the vertical hydraulic gradient is dominant and connectivity of the shallow sands and gravels also is limited.

Based on this conceptual model, any contamination resulting from buried waste found at the BGOU SWMUs would be expected to be found concentrated in the UCRS soils and groundwater immediately within and under the burial cells and landfills, with a lesser amount of contamination dispersed laterally from the cells. The emphasis of the BGOU RI was the evaluation of samples collected from angled soil borings to characterize the potential contaminants leaking from the bottoms of the cells. This chapter provides an assessment of data from the BGOU RI along with data from historical investigations to evaluate the nature and extent of contamination (vertical and lateral) associated with the BGOU SWMUs.

Environmental data from the BGOU RI field activities were merged with the historical data set used for development of the BGOU RI Work Plan (DOE 2006a). This data set is of sufficient quality to address the data needs identified during the DQO process. These data have been compiled and screened to identify COPCs to be addressed in the evaluation of the nature and extent of contaminants. This chapter presents summary tables containing analytical results for each of the sites (tables that show sample borings and depths only provide results above screening levels) and figures depicting the locations of the samples. Duplicate analyses do not appear in the summary tables. The summary tables for subsurface soil and groundwater in this section may not present all data for all analytes. These tables provide a summary of those analytes that exceeded the screening criteria described in Section 4.1, which were considered for addressing the nature and extent of contamination at each SWMU. Appendix C provides a complete report of analytical results for all samples collected during this investigation and the historical data set in a searchable database on compact disk. Appendix D provides three-dimensional figures for various contaminants. Several of these key figures are presented in this chapter.

This report assesses the extent of contamination based on the presence of contaminants in subsurface soils (below 1 ft bgs) and UCRS and RGA groundwater. Samples from these media were analyzed for suites of constituents and reported as the following analytical groups: VOCs, semivolatile organic compounds (SVOCs), PCBs, metals, and radionuclides.¹ There are minimal data available to assess whether there are releases to surface water (essentially ditches that may occur adjacent to a burial ground) from the burial grounds. A previous report prepared for SWMU 2 indicated a shallow water table and contribution to a surrounding ditch, and it was reported that, after a significant rainfall event, water was observed emanating from the sidewalls of a ditch adjacent to SWMUs 7 and 30. Other than these reported occurrences, there is no other evidence of burial ground contribution to surface water. Existing water level data at the various SWMUs are not of sufficient density or from monitoring points appropriately screened to accurately determine if water from a SWMU will contribute to flow in a surrounding ditch.

¹ This report assesses the extent of contamination based on the presence of contaminants in subsurface soils (below 1 ft bgs) and UCRS and RGA groundwater. Samples from these media were analyzed for suites of constituents and reported as the following analytical groups: VOCs, SVOCs, PCBs, metals, and radionuclides. As noted in Section 2.4, metal and radionuclide analyses of unfiltered groundwater samples from soil borings commonly are biased high by suspended soil particles. The reader should be aware that the assessment of nature and extent of contamination in this section is health-protective in that it reports analyses of unfiltered groundwater samples. This assessment of nature and extent may identify groundwater contaminants that are not mobile in groundwater as dissolved contamination or colloids.

Based on past extreme rainfall events, there is a potential for lateral release of contaminants from burial grounds into adjacent surface water features that would need to be considered in the development and evaluation of remedies for the BGOU SWMUs in the FS. As such, the pathway for surface water contamination will be addressed further, as needed, in the FS. The FS will consider the uncertainties related to releases from the BGOU SWMUs to adjacent surface water features and the need for potential remedial actions to mitigate releases that contribute contamination to surface water.

An understanding of the potential releases is key to the rationale behind the sampling that was performed during the BGOU RI. Section 1.3 presents descriptions of the known processes and possible releases from each site that may have contributed to the nature and extent of contaminants.

4.1 DATA PROCESSING AND SCREENING

One objective of the data processing and screening of this RI was to identify potential site-related contaminants and delineate the vertical and horizontal extent of these potential contaminants. To achieve this goal, the RI compared the analytical results of this RI and historical data to PGDP media-specific background concentrations^{2, 3} and applicable screening values.⁴ Where more than one screening value existed for groundwater analytes, the maximum screening value was used to limit the number of contaminants presented in the nature and extent assessment while remaining inclusive of risk-based criteria. Those analytes with no applicable screening value are not discussed in this section. Seven analytes known to be essential nutrients and known to be toxic only at extremely high concentrations were removed from the selection of contaminants in the groundwater data set. These analytes were calcium, chloride, iodine, magnesium, phosphorus, potassium, and sodium.

Data processing and screening for the BGOU were conducted as a multiphase process. First, data were screened to eliminate those sample results that were less than the minimum detection limit (or, in the case of radionuclides, did not exceed the total propagation error). Data that does not have associated QC criteria, such that detection limits cannot be verified, were considered qualitative only. These data then were compared with screening levels.⁵ Screening levels for subsurface soil consisted of background levels at PGDP and risk-based no action levels (NALs) for the excavation worker as compiled from the Risk Methods Document (DOE 2001). (The COPC screening for groundwater fate and transport modeling, described in Section 5, compared analyte levels in subsurface soil against PGDP-specific Soil Screening Levels.) Groundwater samples with analytes above detection limits were compared to maximum contaminant levels (MCLs), if available, for specific chemicals. Risk-based child resident NALs were used to screen compounds or analytes that did not have an MCL. Additionally, background groundwater values for the RGA and McNairy were used for screening against their respective results. The combined data set of soil and groundwater analyses of the BGOU RI and related historical data is sufficient to address the goals of the BGOU RI (see Table 7.1).

² PGDP background levels for water drawn from the RGA and McNairy Formation are provisional values that are subject to change. Potential concerns regarding the background levels are the data set from which these values were derived and the statistical methods that were used to analyze the data set.

³ Background values were unavailable for UCRS groundwater.

⁴ For UCRS and RGA groundwater, the applicable screening criteria were MCLs, and child residence No Action Levels (and RGA background for RGA samples) (Tables 4.3 and 4.4). For subsurface soils, the applicable screening criteria were background (Table 4.1) and excavation worker No Action Levels (Table 4.2). The identification of analytes for groundwater fate and transport modeling (Section 5) screened the subsurface soils against PGDP-specific Soil Screening Levels.

⁵ The data used to define analytes for the assessment of nature and extent (this section) and the risk assessment (Section 6 and Appendix F) differ. The nature and extent assessment includes all available subsurface soil (>1 ft depth) analyses within a SWMU while the risk assessment addresses soils of 10 ft depth or less (the excavation worker receptor). For groundwater, the nature and extent assessment is based on groundwater analyses within a SWMU while the risk assessment is based on modeled groundwater contaminant levels at the SWMU and at downgradient points of exposure.

Tables 4.1 through 4.4 contain the soil and groundwater standards used to screen BGOU data for the nature and extent assessment. If an analyte was detected at a level higher than these screening values, it was considered a contaminant for evaluation of nature and extent. (Section 6, Baseline Human Health Risk Assessment, continues the assessment of hazard associated with the COPCs. Appendix F explains the more thorough COPC screening process used in the risk assessment.) The following sections in this chapter discuss the contaminants found in the BGOU, using both historical and RI data.

Table 4.1. Background Values^a for Subsurface Soils (DOE 2001)

Analytical Compound	Subsurface Soil Background Data (mg/kg or pCi/g)
Aluminum	12,000
Antimony	0.21
Arsenic	7.9
Barium	170
Beryllium	0.69
Boron	NA
Cadmium	0.21
Calcium	6,100
Chromium	43
Chromium, hexavalent	NA
Cobalt	13
Copper	25
Cyanide	NA
Iron	28,000
Lead	23
Lithium	NA
Magnesium	2,100
Manganese	820
Mercury	0.13
Molybdenum	NA
Nickel	22
Potassium	950
Selenium	0.7
Silica	NA
Silver	2.7
Sodium	340
Strontium	NA
Thallium	0.34
Uranium (metal)	4.6
Vanadium	37
Zinc	60
Americium-241	NA
Cesium-137	0.28
Cobalt-60	NA
Neptunium-237	NA
Protactinium-233*	NA
Plutonium-238	NA
Plutonium-239	NA
Plutonium-239/240*	NA
Potassium-40	16
Protactinium-234m	NA
Radium	NA
Radium-226	1.5
Radon-222	NA
Strontium-90	NA
Technetium-99	2.8
Thorium-230	1.4
Thorium-234	NA

Table 4.1. Background Values^a for Subsurface Soils (DOE 2001) (Continued)

Analytical Compound	Subsurface Soil Background Data (mg/kg or pCi/g)
Uranium (total)	NA
Uranium-233/234*	2.4
Uranium-234	2.4
Uranium-235	0.14
Uranium-235/236*	0.14
Uranium-238	1.2

^a The PGDP studies of background levels for soils are *Background Concentrations and Human Health Risk-based Screening Criteria for Metals in Soil at the Paducah Gaseous Diffusion Plant, Paducah, Kentucky*, DOE/OR/07-1417&D1 (DOE 1995d) and *Background Levels of Selected Radionuclides and Metals in Soils and Geologic Media at the Paducah Gaseous Diffusion Plant, Paducah, Kentucky*, DOE/OR/07-1586&D2 (DOE 1997c).

*NOTE: Data for the undifferentiated isotopes neptunium-237/protactinium-233, plutonium-239/240, uranium-233/234, and uranium-235/236 were compared to the background values for neptunium-237, plutonium-239, uranium-234, and uranium-235, respectively.

NA = not available

Table 4.2. Risk-Based No Action Levels Used in BGOU Soil Screening (DOE 2001)

Chemical	No Action Level	Chemical	No Action Level
<i>Inorganics (mg/kg)</i>		<i>Radionuclides (pCi/g)</i>	
Antimony	0.492	Americium-241	1.74
Arsenic	0.324	Cobalt-60	0.0238
Beryllium	1.26	Cesium-137+Daughters	0.0858
Cadmium (Diet)	15.2	Neptunium-237+Daughters	0.271
Chromium (III) (Insoluble Salts)	476	Plutonium-238	11.7
Chromium (Total)	476	Plutonium -239	11.5
Chromium VI (particulates)	3.69	Plutonium -240	11.5
Copper	427	Radium-226+Daughters	0.0256
Iron	2,170	Radon-222+Daughters	33,900,000
Lead	50	Strontium-90+Daughters	7.44
Manganese	56.6	Technetium-99	362
Mercury	1.17	Thorium-228+Daughters	0.028
Molybdenum	66	Thorium -230	14.9
Nickel Soluble Salts	216	Thorium -232	13.5
Selenium	71.3	Uranium-234	19.8
Silver	41.2	Uranium -235+Daughters	0.395
Thallium (Thallium Chloride)	0.711	Uranium -238+Daughters	1.71
Uranium	11.3	Plutonium-239/240	11.5
Vanadium	4.4	Uranium-235/236	0.395
Zinc	2,660		
<i>Organics (mg/kg)</i>			
Acenaphthene	350	Fluorene	338
Acrylonitrile	0.248	HpCDD, 2,3,7,8-	0.000349
Anthracene	3340	HpCDF, 2,3,7,8-	0.000349
PCB-1016	0.168	HxCDD, 2,3,7,8-	0.0000349
PCB-1221	0.168	HxCDF, 2,3,7,8-	0.0000349
PCB-1232	0.168	Indeno[1,2,3-cd]pyrene	0.232
PCB-1242	0.168	Naphthalene	30.4
PCB-1248	0.168	OCDD	0.00349
PCB-1254	0.168	OCDF	0.00349
PCB-1260	0.168	PeCDD, 2,3,7,8-	0.0000699
Benz(a)anthracene	0.232	PeCDF, 1,2,3,7,8-	0.0000699
Benzene	1.4	PeCDF, 2,3,4,7,8-	0.0000699
Benzo(a)pyrene	0.0232	Phenanthrene	No value
Benzo(b)fluoranthene	0.232	Total PCBs (high risk)	0.168
Benzo(k)fluoranthene	2.32	Total PCBs (lowest risk)	4.81
Carbon Tetrachloride	0.51	Total PAHs	0.0232
Chloroform	0.166	Pyrene	181
Chrysene	23.2	TCDD, 2,3,7,8-	0.0000349
Dibenz(a,h)anthracene	0.0232	TCDF, 2,3,7,8-	0.0000349
1,1-DCE	0.119	Tetrachloroethylene	4.04
1,2-DCE	68.2	TCE	3.25
cis-1,2-DCE	17.1	Vinyl Chloride	0.141
trans-1,2-DCE	28.4	Xylene, Mixture	963
Ethylbenzene	28.7	Xylene, m-	5,560
Fluoranthene	242	Xylene, o-	5,590

Note: No Action Levels are for the excavation worker scenario (from Table A.4 of the 2001 Risk Methods document).

Table 4.3. Background Values for Groundwater Drawing from the RGA and McNairy Formation^a at PGDP (DOE 2001)

Analytical Compound	(mg/L or pCi/L)	
	RGA	McNairy
Aluminum	2.189	0.687
Aluminum, Dissolved	0.311	0.579
Antimony	0.060 ^b	0.060 ^b
Antimony, Dissolved	0.060 ^b	0.060 ^b
Arsenic	0.005 ^b	0.005 ^b
Arsenic, Dissolved	0.005 ^b	0.005 ^b
Barium	0.235	0.296
Barium, Dissolved	0.2	0.268
Beryllium	0.004 ^b	0.017 ^b
Beryllium, Dissolved	0.004 ^b	0.004 ^b
Cadmium	0.010 ^b	0.010 ^b
Cadmium, Dissolved	0.010 ^b	0.010 ^b
Calcium	41.238	38.858
Calcium, Dissolved	38.166	38.829
Chloride	91.021	19.708
Chromium	0.144	0.060 ^b
Chromium, Dissolved	0.050 ^b	0.050 ^b
Cobalt	0.045 ^b	0.096
Cobalt, Dissolved	0.045 ^b	0.045 ^b
Copper	0.036	0.057
Copper, Dissolved	0.02	0.013 ^b
Fluoride	0.27	0.33
Iron	5.03	18.36
Iron, Dissolved	0.267	12.372
Lead	0.129	0.050 ^b
Lead, Dissolved	0.098	0.050 ^b
Magnesium	16.262	13.418
Magnesium, Dissolved	16.215	14.171
Manganese	0.119	0.941
Manganese, Dissolved	0.068	0.894
Mercury	0.0002 ^b	0.0002 ^b
Mercury, Dissolved	0.0002 ^b	0.0002 ^b
Molybdenum	0.050 ^b	0.050 ^b
Molybdenum, Dissolved	0.050 ^b	0.050 ^b
Nickel	0.682	0.109 ^b
Nickel, Dissolved	0.305	0.050 ^b
Nitrate as Nitrogen	15.561	1.474
Potassium	5.195	55.752
Potassium, Dissolved	4.096	51.205
Selenium	0.005 ^b	0.005 ^b
Selenium, Dissolved	0.005 ^b	0.005 ^b
Silica	26.401	36
Silver	0.011 ^b	0.050 ^b
Silver, Dissolved	0.060 ^b	0.050 ^b
Sodium	59.45	29.2
Sodium, Dissolved	60.433	27.98
Sulfate	19.947	28.9
Thallium	0.056 ^b	0.644
Thallium, Dissolved	0.056 ^b	0.056 ^b
Uranium	0.002 ^b	0.001 ^b
Uranium, Dissolved	0.002 ^b	0.001

Table 4.3. Background Values for Groundwater Drawing from the RGA and McNairy Formation^a at PGDP (DOE 2001) (Continued)

Analytical Compound	(mg/L or pCi/L)	
	RGA	McNairy
Vanadium	0.134	0.126
Vanadium, Dissolved	0.134	0.126
Zinc	0.054	0.142
Zinc, Dissolved	0.049	0.116
Gross Alpha	5.8	11.9
Gross Beta	13.8	144.5
Neptunium-237	0.8	0.5
Plutonium-239	0.1	0.2
Radium-226	0.6	1.2
Radon-222	626	295
Technetium-99	22.3	20.6
Thorium-230	1.1	1.5
Total Radium	1.3	0.7
Uranium-234 ^c	0.7	0.3
Uranium-235 ^c	0.3	0.2
Uranium-238 ^c	0.7	0.3

^a Values are for those derived over all observations.

^b Background value was derived qualitatively over all observations because analyte was never detected or was detected infrequently at a concentration near the analyte's detection limit.

^c Uranium isotopic concentrations were derived from the mass concentration of uranium.

Table 4.4. Groundwater MCLs and Child Resident No Action Levels Used in BGOU Screening

Analytical Compound	MCL (mg/L)	Child Resident No Action Level (mg/L) ^a
<i>Inorganics</i>		
Antimony ^b	0.006	0.000564
Arsenic ^b	0.010	0.000035
Barium	2	0.104
Beryllium	0.004	0.00264
Cadmium	0.005	0.000661
Chromium ^b	0.1	1.76
Copper	1.3	0.0557
Cyanide	0.2	0.0284
Fluoride	4.0	No value
Iron	No MCL	0.449
Lead	0.015	0.015
Manganese	No MCL	0.035
Mercury ^b	0.002	0.000444
Molybdenum	No MCL	0.00753
Nickel ^b	No MCL	0.0301
Nitrate	10	2.41
Nitrite	1	0.151
Selenium	0.05	0.00754
Silver	No MCL	0.0075
Thallium ^b	0.002	0.00012

Table 4.4. Groundwater MCLs and Child Resident No Action Levels Used in BGOU Screening (Continued)

Analytical Compound	MCL (mg/L)	Child Resident No Action Level (mg/L)
Uranium	0.03	0.000906
Vanadium	No MCL	0.00925
Zinc	No MCL	0.45
Organics		
Acenaphthene	No MCL	0.0136
Acrylonitrile	No MCL	0.0000426
Anthracene	No MCL	0.0766
Benz(a)anthracene	No MCL	0.0000132
Benzene	0.005	0.000385
Benzo(a)pyrene	0.002	0.000000951
Benzo(b)fluoranthene	No MCL	0.00000951
Benzo(k)fluoranthene	No MCL	0.000168
Carbon tetrachloride	0.005	0.000181
Chloroform	No MCL	0.0000287
Chrysene	No MCL	0.00132
Chlorobenzene	0.1	0.00466
Dibenz(a,h)anthracene	No MCL	0.000000456
Dioxins/Furans (Total)	No MCL	6.09E-11
o-Dichlorobenzene	0.6	0.0166
p-Dichlorobenzene	0.075	0.000578
1,2-Dichloroethane	0.005	0.000147
1,1-DCE	0.007	0.000047
1,2-DCE	No MCL	0.00247
cis-1,2-DCE	0.07	0.00273
trans-1,2-DCE	0.1	0.00548
Methylene chloride	0.005	0.00426
Bis(2-ethylhexyl)phthalate	0.006	0.00312
Ethylbenzene	0.7	0.00468
Fluoranthene	No MCL	0.0226
Fluorene	No MCL	0.00972
Heptachlor	0.0004	0.0000114
HpCDD, 2,3,7,8-	No MCL	5.45E-09
HpCDF, 2,3,7,8-	No MCL	3.51E-08
HxCDD, 2,3,7,8-	No MCL	3.51E-09
HxCDF, 2,3,7,8-	No MCL	3.51E-09
Indeno(1,2,3-cd)pyrene	No MCL	0.00000631
Naphthalene	No MCL	0.000285
OCDD	No MCL	1.91E-08
OCDF	No MCL	2.03E-08
PeCDD, 2,3,7,8-	No MCL	2.5E-10
PeCDF, 1,2,3,7,8-	No MCL	1.48E-10
PeCDF, 2,3,4,7,8-	No MCL	1.29E-09
Total PCBs ^b	0.0005	0.0000793
PCB-1016	Total PCB applies	0.0000468
PCB-1221	Total PCB applies	0.000112
PCB-1232	Total PCB applies	0.000128
PCB-1242	Total PCB applies	0.000123
PCB-1248	Total PCB applies	0.0000775
PCB-1254	Total PCB applies	0.0000194
PCB-1260	Total PCB applies	0.0000428
Total PAHs	No MCL	0.000000951
Pyrene	No MCL	0.0182
TCDD, 2,3,7,8-	No MCL	6.09E-11
TCDF, 2,3,7,8-	No MCL	1.6E-09
Tetrachloroethene	0.005	0.000582

Table 4.4. Groundwater MCLs and Child Resident No Action Levels Used in BGOU Screening (Continued)

Analytical Compound	MCL (mg/L or pCi/L)	Child Resident No Action Level (mg/L or pCi/L)
Toluene	1	0.0338
1,2,4-Trichlorobenzene	0.07	0.00781
1,1,1-Trichloroethane	0.2	0.0335
1,1,2-Trichloroethane	0.005	0.000238
TCE	0.005	0.0016
Vinyl chloride	0.002	0.000035
Total Xylene	10	0.0653
m,p-Xylene	10	0.439
1,3-Dimethylbenzene	10	0.439
Radionuclides		
Americium-241	No MCL	0.371
Cobalt-60	No MCL	2.46
Cesium-137 ^c	200 pCi/L	1.27
Neptunium-237	No MCL	0.573
Plutonium-238	No MCL	0.295
Plutonium-239	No MCL	0.286
Plutonium-240	No MCL	0.286
Radium-226	5	0.1
Radon-222	No MCL	0.866
Strontium-90	No MCL	0.522
Technetium-99 ^c	900 pCi/L	14
Thorium-228	No MCL	0.129
Thorium-230	No MCL	0.424
Thorium-232	No MCL	0.382
Uranium-234 ^d	20 pCi/L	0.546
Uranium-235 ^d	20 pCi/L	0.538
Uranium-238 ^d	20 pCi/L	0.443

^aNALs were taken from Table A.18 of the PGDP Risk Methods Document (2001) for PGDP Primary COPCs.

^bThese NALs were used for the following compounds with more than 1 NAL listed in Table A.18 of the PGDP Risk Methods Document (2001): Antimony (metallic); Arsenic, inorganic; Chromium (total); Mercury, inorganic salts; Nickel soluble salts; Thallium chloride; Total PCBs (low risk)

^cThe activities for cesium-137 and 99Tc are assumed to yield the MCL of 4 mrem/yr for beta emitters (if no other beta or gamma emitters are present).

^dThe MCL for uranium is 0.030 mg/L (activity will vary depending on the specific isotope). The MCL is not based on radiotoxicity, but on kidney toxicity and cancer risk (EPA Directive No. 9283.1-14, (November 6, 2001).

MCL = maximum contaminant level.

4.2 SOURCES OF CONTAMINATION

Previous fate and transport modeling has determined radionuclides such as technetium-99 (⁹⁹Tc), neptunium-237 (²³⁷Np), plutonium-239 (²³⁹Pu), uranium-234 (²³⁴U), uranium-235 (²³⁵U), and uranium-238 (²³⁸U); VOCs such as TCE, 1,1-DCE, vinyl chloride, 1,2-DCE, carbon tetrachloride, and chloroform; metals; and some SVOCs as contributing to groundwater contamination. The COPCs from previous fate and transport modeling for each SWMU are listed in Table 4.5. There was no previous modeling at SWMU 145.

Table 4.5. Summary of Historical Fate and Transport Modeling Results

Location	Chemicals of Potential Concern Determined by Historical Groundwater Modeling
SWMU 2	TCE and other VOCs (<i>cis</i> -1,2-DCE and vinyl chloride) ⁹⁹ Tc
SWMU 3	⁹⁹ Tc and naphthalene
SWMU 4	VOCs (TCE; 1,1-DCE; vinyl chloride; 1,2-DCE; carbon tetrachloride, and chloroform), metals (arsenic, cobalt, copper, iron, and manganese), and radionuclides (²³⁷ Np, ²³⁹ Pu, ⁹⁹ Tc, ²³⁴ U, ²³⁵ U, and ²³⁸ U).
SWMU 5	1,1-DCE; naphthalene; manganese; iron; and ⁹⁹ Tc
SWMU 6	⁹⁹ Tc and iron
SWMUs 7 and 30	⁹⁹ Tc and vinyl chloride

Table 4.6 lists the BGOU RI sampling locations and the historical sampling locations for each SWMU. Figures 4.1 through 4.16 present locations of all historical and RI sampling for soils and groundwater.

Table 4.6. BGOU SWMU Sampling Locations Used in BGOU RI Nature and Extent and Fate and Transport Evaluations

SWMU 2					
Soil Sampling Locations					
RI Sampling Locations			Historical Sampling Locations		
002-001			SWMU2-1	SWMU2-9	
002-002			SWMU2-2	SWMU2-10	
			SWMU2-3	SWMU2-11	
			SWMU2-4	SWMU2-12	
			SWMU2-5	SWMU2-13	
			SWMU2-6	SWMU2-14	
			SWMU2-7	SWMU2-15	
			SWMU2-8	SWMU2-17	
Groundwater Sampling Locations					
RI Sampling Locations			Historical Sampling Locations¹		
None.			UCRS	RGA	McNairy
			MW154 (18.5 ft)	MW333 (79 ft)	SWMU2-10
			PZ334 (19.75 ft)	MW337 (74.33 ft)	SWMU2-17
			PZ335 (19.75 ft)	MW338 (74.33 ft)	SWMU2-3
			PZ336 (19.75 ft)	SWMU2-10	SWMU2-9
			PZ74 (42 ft)	SWMU2-13	
			SWMU2-10	SWMU2-16	
			SWMU2-17	SWMU2-17	
			SWMU2-3	SWMU2-3	
			SWMU2-9	SWMU2-5	
				SWMU2-9	
SWMU 3					
Soil Sampling Locations					
RI Sampling Locations			Historical Sampling Locations		
003-001			003-006		
003-002			003-007		
003-003			003-008		
003-004			003-009		
003-005			003-010		
Groundwater Sampling Locations					
RI Sampling Locations			Historical Sampling Locations¹		
UCRS	RGA	McNairy	UCRS	RGA	McNairy
003-003	None.	None.	MW85 (40.14 ft)	MW226 (88.9 ft)	None.
003-004			MW88 (34.27 ft)	MW227 (74.5 ft)	
			MW91 (33.61 ft)	MW67 (70.3 ft)	
			MW94 (39.06 ft)	MW84 (75.87 ft)	
				MW86 (85.57 ft)	
				MW87 (74.28 ft)	
				MW89 (88.15 ft)	
				MW90 (74.21 ft)	
				MW90A (72 ft)	
				MW92 (89.28 ft)	
				MW93 (79.9 ft)	
				MW95 (88.22 ft)	
				MW95A (88 ft)	

Table 4.6. BGOU SWMU Sampling Locations used in BGOU RI Nature and Extent and Fate and Transport Evaluations (Continued)

<i>SWMU 4</i>						
Soil Sampling Locations						
RI Sampling Locations			Historical Sampling Locations			
None.			004-009	004-032	004-045	
			004-017	004-033	004-046	
			004-019	004-034	004-047	
			004-020	004-035	004-048	
			004-021	004-036	004-049	
			004-022	004-037	004-050	
			004-023	004-038	004-051	
			004-024	004-039	004-052	
			004-025	004-040	004-053	
			004-026	004-041	004-054	
			004-027	004-042	004-055	
			004-030	004-043	004-056	
			004-031	004-044	004-057	
			Groundwater Sampling Locations			
RI Sampling Locations			Historical Sampling Locations ¹			
None.			UCRS		RGA	McNairy
			004-008	004-028	004-028	004-028
			004-009	004-029	004-029	004-029
			004-011	004-032	004-058	004-058
			004-017	004-033	720-026	DG-030
			004-019	004-035	DG-030	
			004-020	004-036		
			004-021	004-037		
			004-022	004-038		
			004-023	004-039		
			004-024	004-040		
			004-024	004-044		
			004-025	004-047		
			004-026	004-049		
004-027						
<i>SWMU 5</i>						
Soil Sampling Locations						
RI Sampling Locations			Historical Sampling Locations			
005-101			005-015	005-020		
005-102			005-016	005-021		
005-103			005-017	005-022		
			005-018	005-027		
			005-019	005-028		
Groundwater Sampling Locations						
RI Sampling Locations			Historical Sampling Locations ¹			
UCRS	RGA	McNairy	UCRS	RGA	McNairy	
005-101	None.	None.	005-015	005-013	005-013	
005-102			005-016	005-026	005-026	
			005-017	DG-002		
			005-018			
			005-019			
			005-020			
			005-021			
			005-022			

Table 4.6. BGOU SWMU Sampling Locations used in BGOU RI Nature and Extent and Fate and Transport Evaluations (Continued)

SWMU 6					
Soil Sampling Locations					
RI Sampling Locations			Historical Sampling Locations		
006-101			006-016	006-022	
006-102			006-017	006-023	
006-103			006-018	006-026	
006-104			006-019	006-027	
			006-020	006-028	
			006-021	006-029	
Groundwater Sampling Locations					
RI Sampling Locations			Historical Sampling Locations¹		
UCRS	RGA	McNairy	UCRS	RGA	McNairy
006-101	None.	None.	006-009	006-024	006-024
006-102			006-011	006-025	006-025
006-103			006-012		
006-104			006-016		
			006-017		
			006-018		
			006-019		
			006-020		
			006-021		
			006-022		
			006-023		
			006-025		
			006-028		
			006-029		
SWMU 7					
Soil Sampling Locations					
RI Sampling Locations			Historical Sampling Locations		
007-001	007-006		TP-3A	WB-7	
007-002	007-007		WB-10	WB-8	
007-003A	007-008		WB-12	WB-9	
007-003B	007-009		WB-13	WBP-12A	
007-004	007-010		WB-14	WBP-13A	
007-005	007-011		WB-6	WBP-9A	
Groundwater Sampling Locations					
RI Sampling Locations			Historical Sampling Locations¹		
UCRS	RGA	McNairy	UCRS	RGA	McNairy
007-001	007-009	None.	DG-005	DG-005	GGW-01
007-002	007-010		GGW-01	GGW-01	GGW-02
007-003B	007-011		GGW-02	GGW-02	GGW-03
007-005			GGW-03	GGW-03	
007-007			MW186 (23 ft)	MW185 (73 ft)	
007-008			MW187 (26.5 ft)	MW339 (95 ft)	
007-009			WB-12 (9 ft)	MW340 (95.3 ft)	
007-010			WB-13 (7 ft)		
007-011			WB-7 (12 ft)		
			WB-8 (11.5 ft)		
			WB-9 (9 ft)		
			WBP-12A (9 ft)		
			WBP-9A (11 ft)		

Table 4.6. BGOU SWMU Sampling Locations used in BGOU RI Nature and Extent and Fate and Transport Evaluations (Continued)

<i>SWMU 30</i>					
Soil Sampling Locations					
RI Sampling Locations			Historical Sampling Locations		
030-001			WB-1	WB-5	
030-002			WB-11	WBP-1A	
030-003			WB-3	WBP-4A	
030-004			WB-4		
Groundwater Sampling Locations					
RI Sampling Locations			Historical Sampling Locations ¹		
UCRS	RGA	McNairy	UCRS	RGA	McNairy
030-003	None.	None.	MW64 (32.8 ft) WB-1 (9.5 ft) WB-4 (8 ft) WB-5 (8 ft) WBP-4A (8 ft)	MW245 (74.58 ft) MW63 (63.5 ft) MW65 (91.2 ft) MW66 (60.2 ft)	None.
<i>SWMU 145</i>					
RI Sampling Locations ^b			Historical Sampling Locations ^c		
145-101			Landfill Locations:	NS-SD-01	
145-102			DG-029	NS-SD-02	
145-103			NSDD Locations:	NS-SD-03	
145-104			A10	NS-SD-04	
145-105			A2	NS-SD-05	
145-106			AIPNSD2000-01	NS-SD-06	
145-107			AIPNSD2000-02	NS-SD-07	
			AIPNSD2000-03	NS-SS-01	
			AIPNSD2000-04	NS-SS-02	
			AIPNSD2000-05	NS-SS-03	
			AIPNSD2000-06	NS-SS-04	
			AIPNSD2000-07	NS-SS-05	
			JP-0100	NS-SS-06	
			NSD030	NS-SS-07	
			NSD2000-01bank	NS-SS-08	
			NSD2000-02bank	NST1S01	
			NSD2000-03bank	NST1S02	
			NSD2000-04bank	NST1S03	
			NSD2000-05bank	NST2S01	
			NSD2000-06bank	NST2S02	
			NSD2000-06sedi	NST2S03	
			NSD2000-06SPT	NST2S04	
			NSD2000-07bank	NST2S05	
			NSD2000-07SPT	SEC3A-5SO	
			NSD2000-08bank	SEC3A-8SO	

Table 4.6. BGOU SWMU Sampling Locations used in BGOU RI Nature and Extent and Fate and Transport Evaluations (Continued)

Groundwater Sampling Locations			
RI Sampling Locations	Historical Sampling Locations^a		
	UCRS	RGA	McNairy
None.	MW180 (27 ft) MW182 (20 ft) MW371 (31.6 ft) MW386 (30 ft) MW390 (37.8 ft) MW393 (38 ft) MW396 (43.6 ft)	DG-014 MW179 (57 ft) MW181 (57 ft) MW220 (69 ft) MW221 (84 ft) MW222 (78.3 ft) MW223 (82.4 ft) MW224 (82 ft) MW225 (78 ft) MW263 (58.96 ft) MW264 (55.2 ft) MW265 (70.3 ft) MW266 (76.6 ft) MW267 (85 ft) MW276 (58.9 ft) MW277 (71.4 ft) MW369 (51.5 ft) MW370 (71 ft) MW384 (60 ft) MW385 (76.2 ft) MW387 (57.4 ft) MW388 (70.2 ft) MW391 (67.2 ft) MW392 (91.2 ft) MW394 (75 ft) MW395 (83 ft) MW397 (95 ft)	DG-014

¹ Bottom screen depths for monitoring wells and piezometers.

^b RI sampling locations all characterized landfill.

^c Surface soil samples not included in fate and transport source term development.

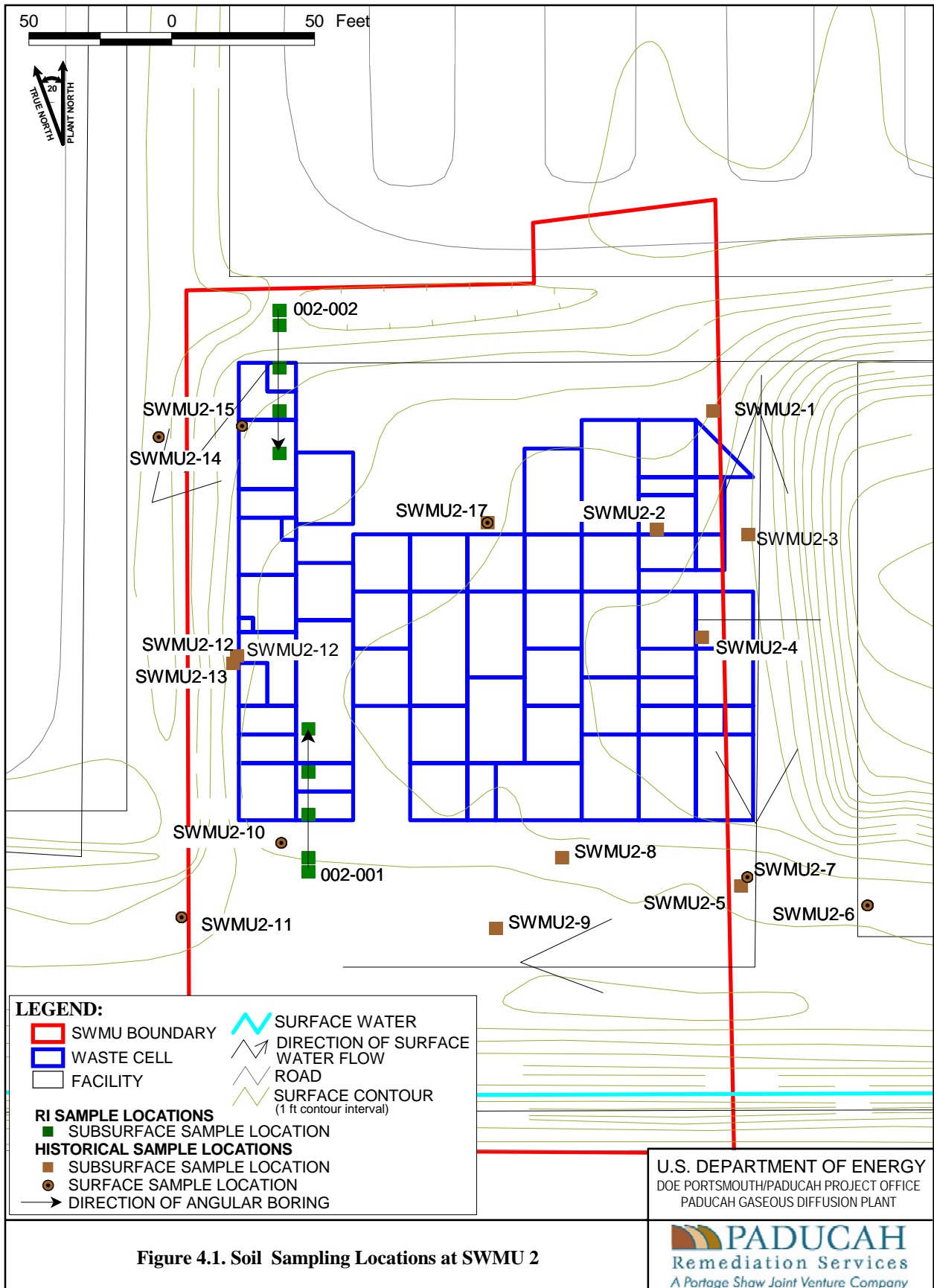


Figure 4.1. Soil Sampling Locations at SWMU 2

Figure No. \BGOU\d2R1-ri_sect1.apr
DATE 08-04-09

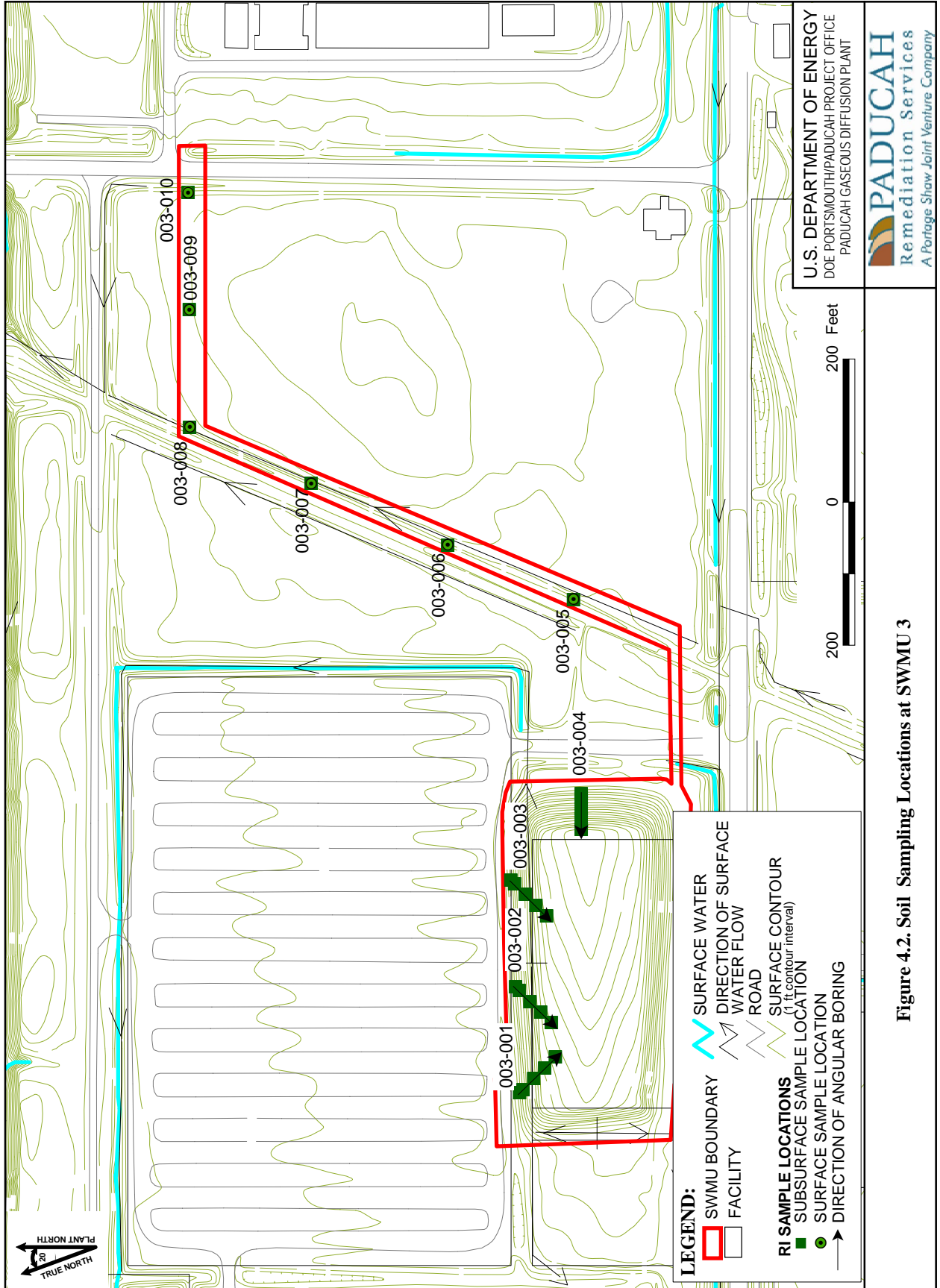


Figure 4.2. Soil Sampling Locations at SWMU 3

Figure No. IBGOUd2R1-ri_sec14.apr
DATE 08-04-09

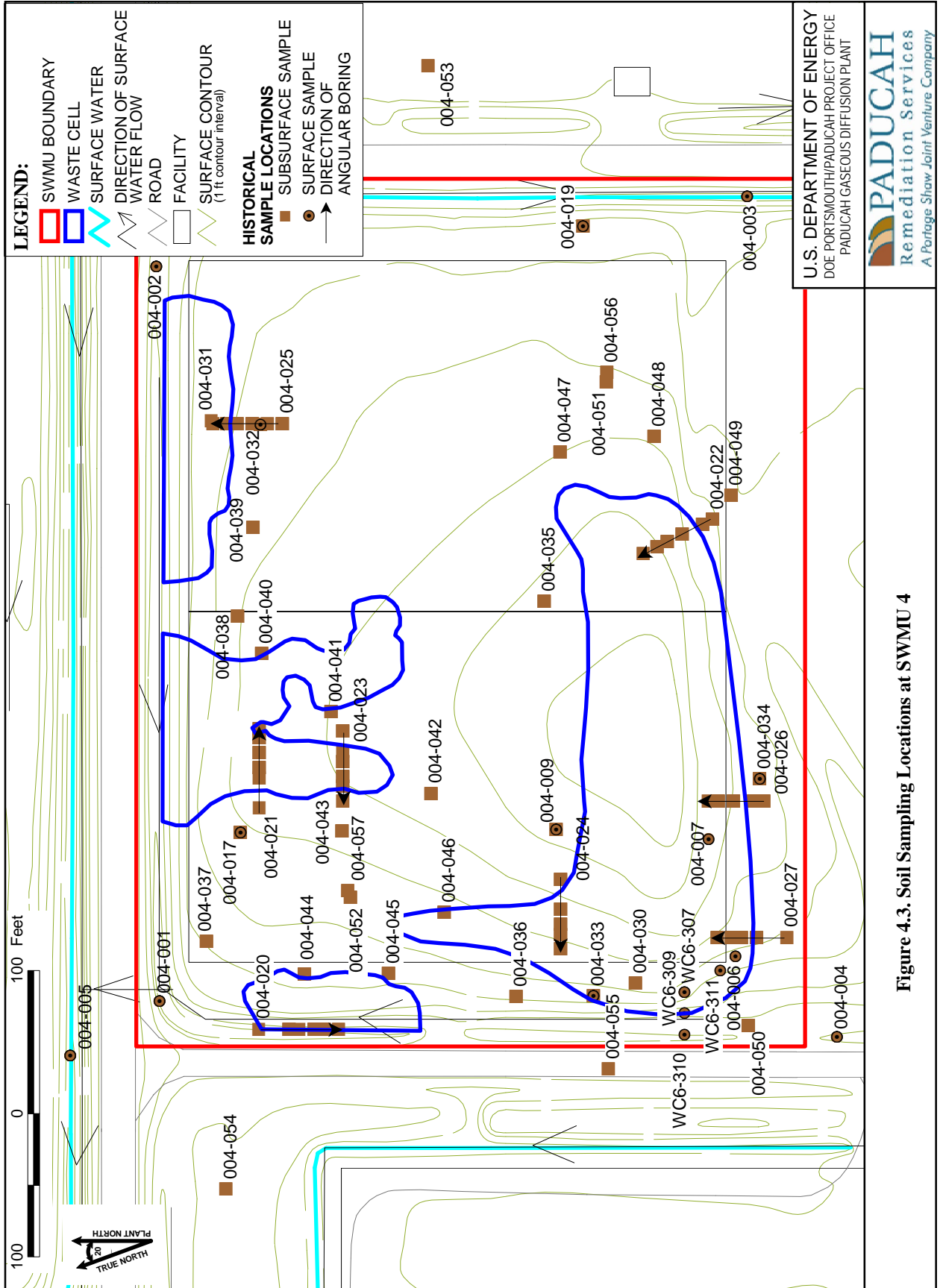
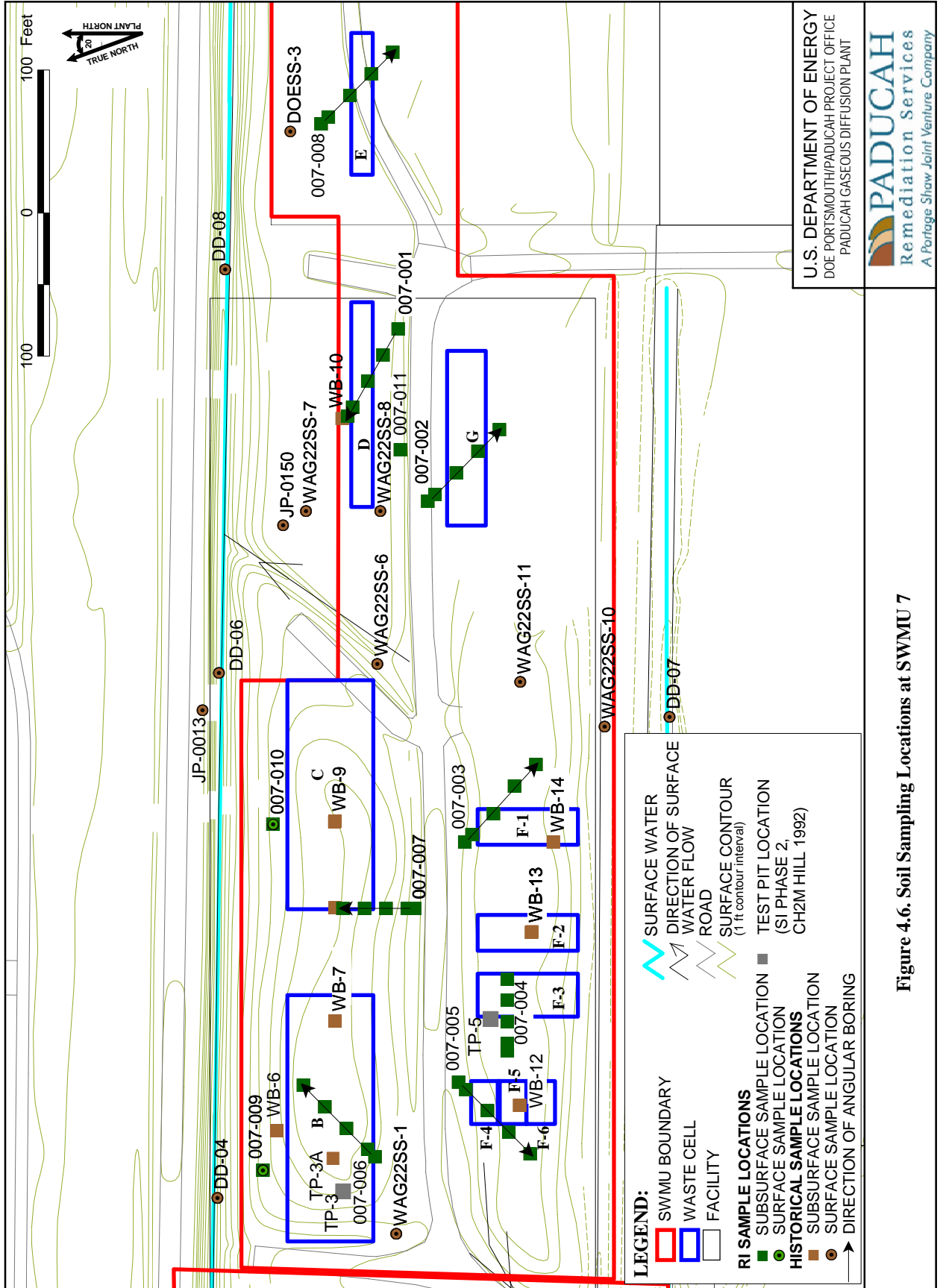


Figure No. IBG0102R1-r1_sec14.apr
DATE 08-04-09

Figure 4.3. Soil Sampling Locations at SWMU 4

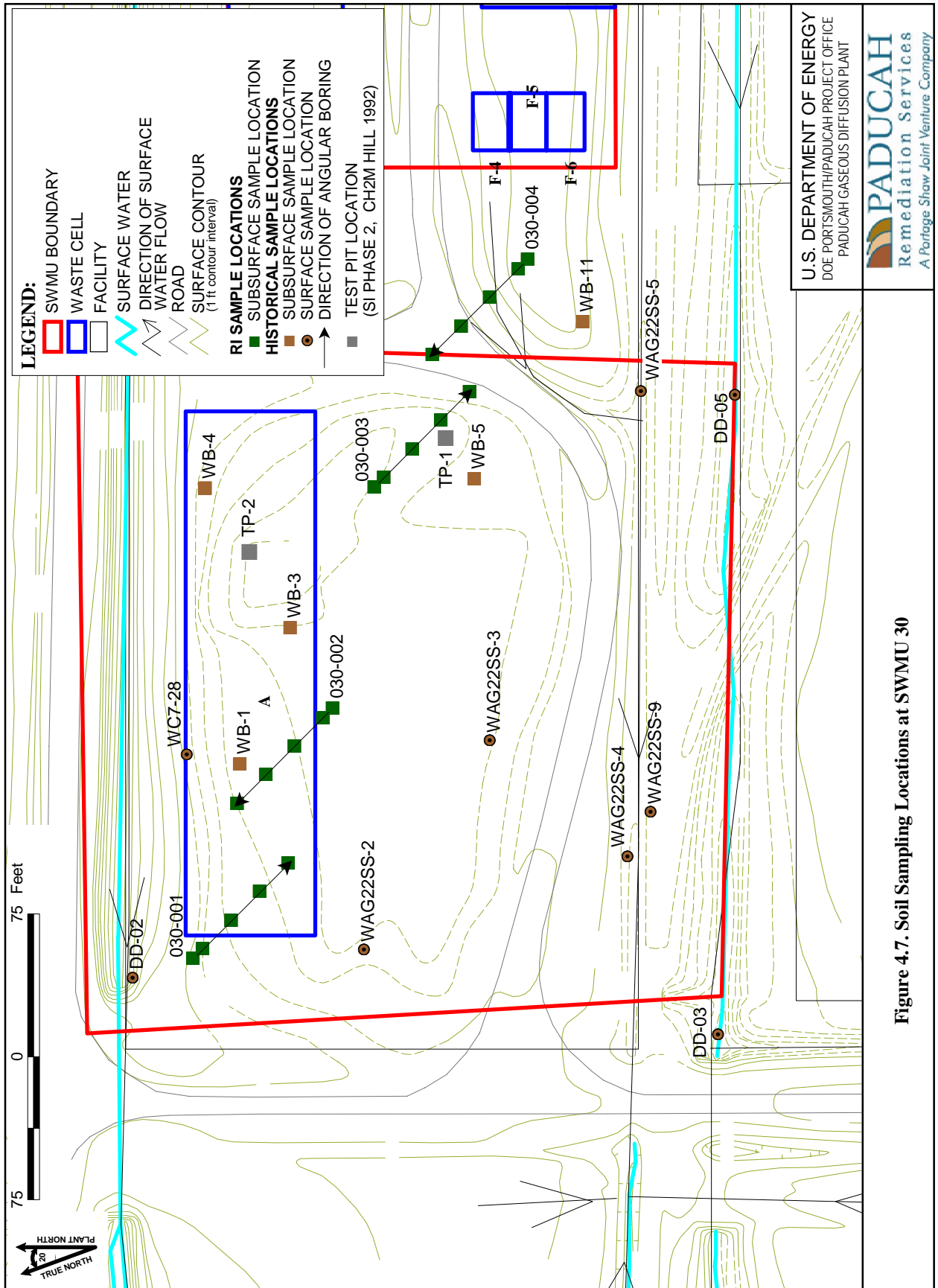


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Remediation Services
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Figure 4-6. Soil Sampling Locations at SWMU 7

Figure No. IBGOU042R1-r1_sec14.apr
DATE 09-29-09



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Figure 4.7. Soil Sampling Locations at SWMU 30

Figure No. IBGOU02R1-r1_sec14.apr
DATE 08-04-09

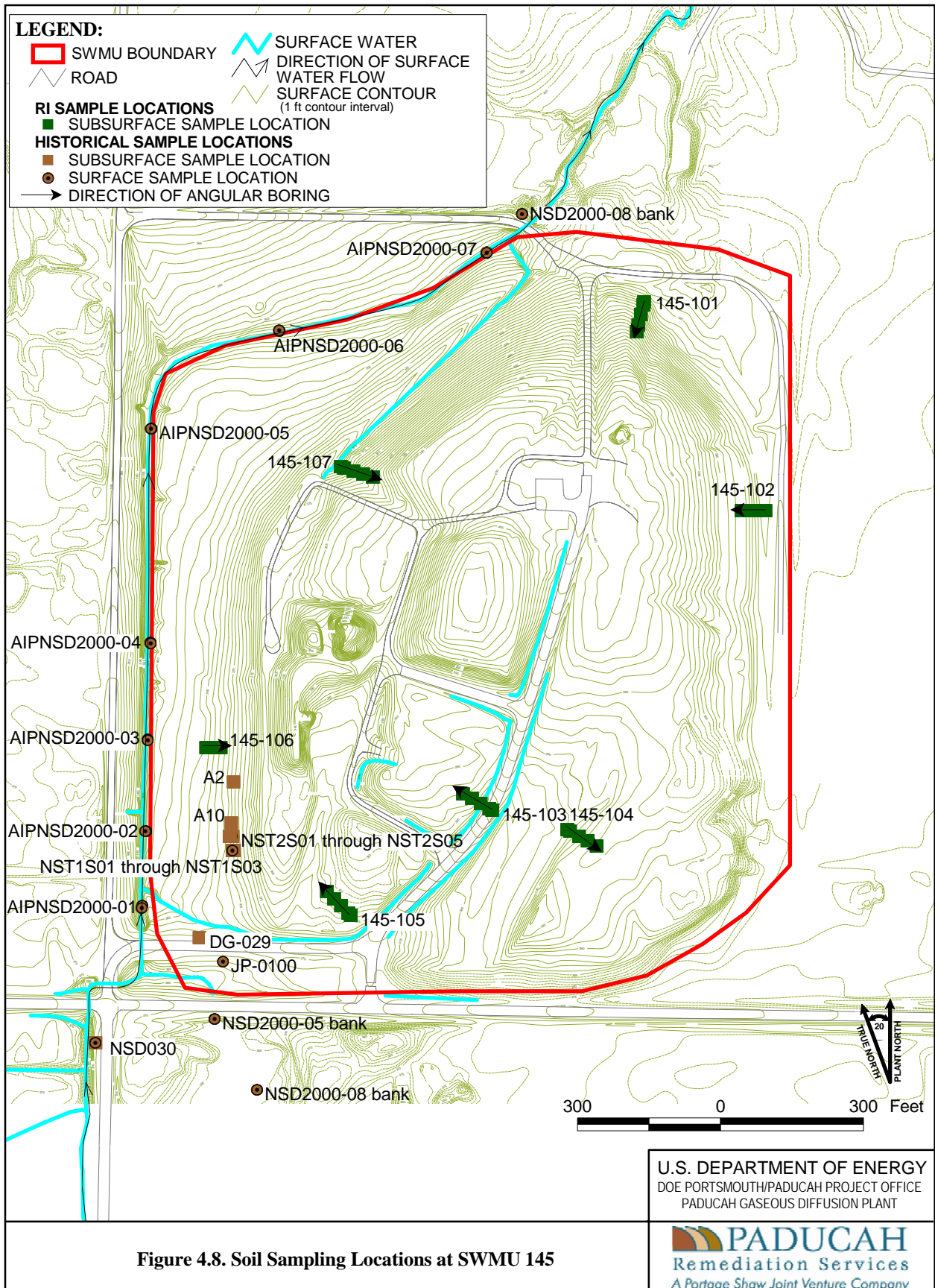


Figure 4.8. Soil Sampling Locations at SWMU 145

Figure No. \BGOU\d2R1-ri_sect4.apr
DATE 08-04-09

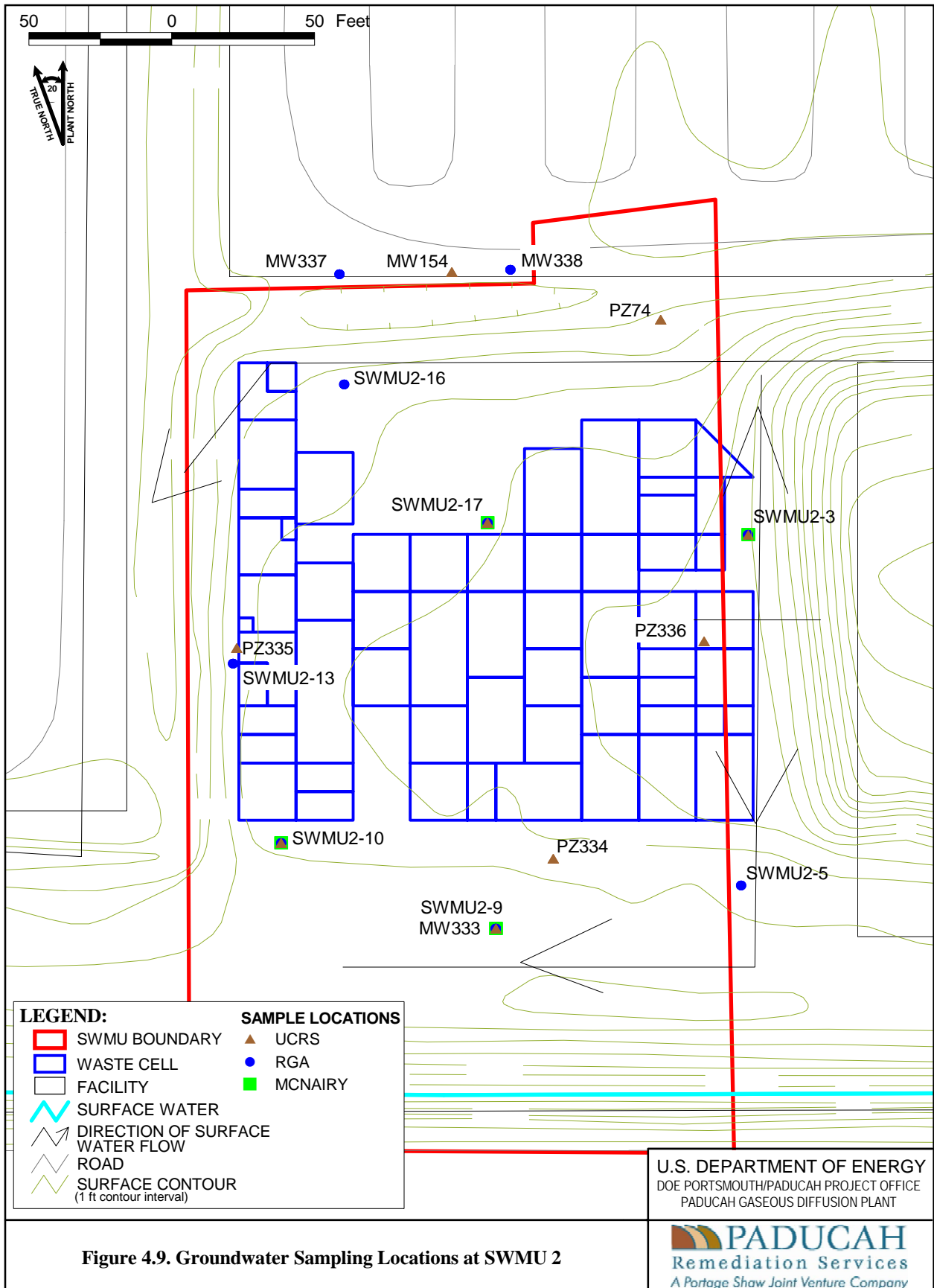
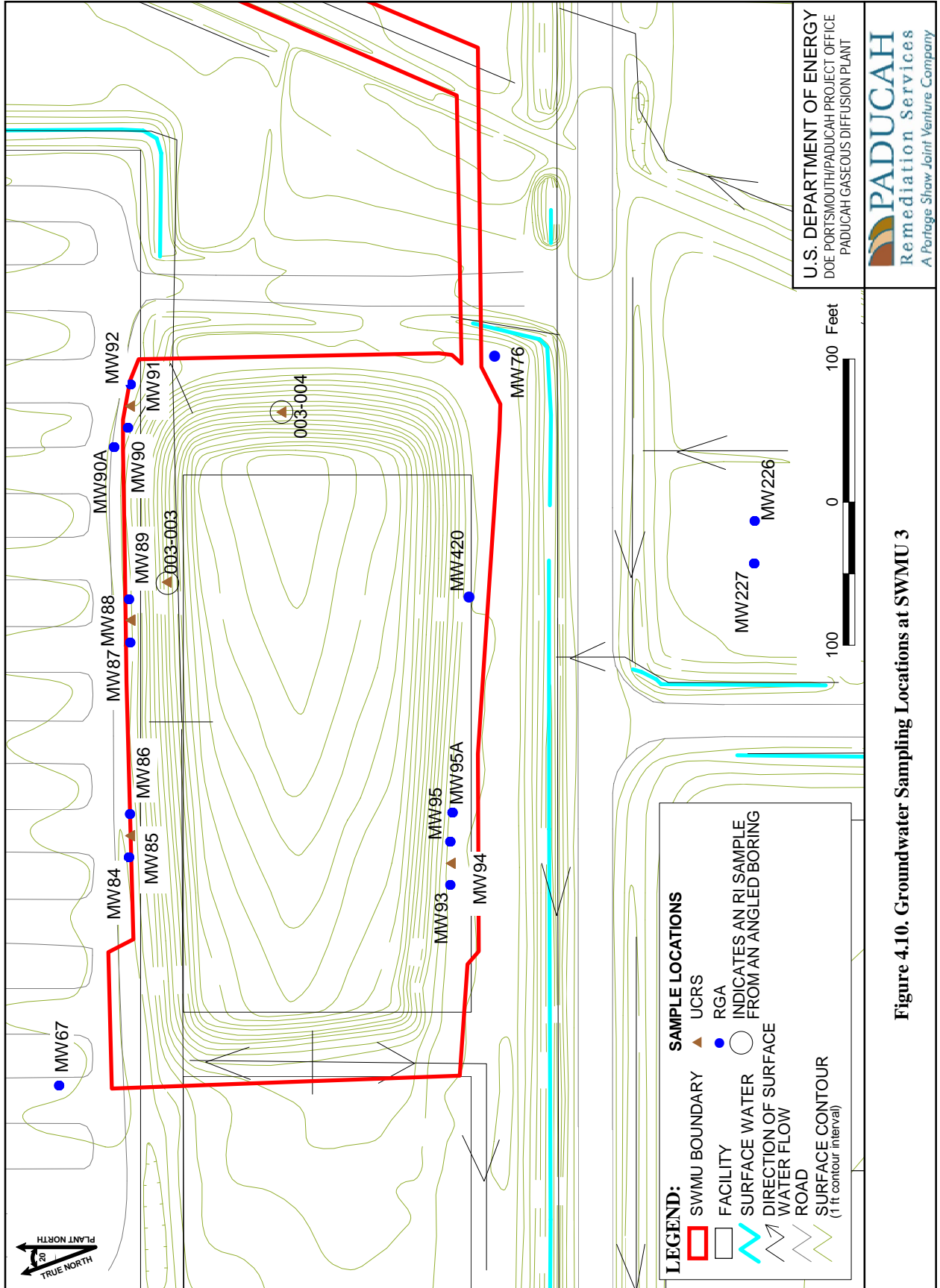


Figure 4.9. Groundwater Sampling Locations at SWMU 2

Figure No. 1BGOU\42R1-ri_sect4.apr
DATE 08-04-09



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Figure 4.10. Groundwater Sampling Locations at SWMU 3

Figure No. IBGOU02R1-rj_sec14.apr
DATE 08-04-09

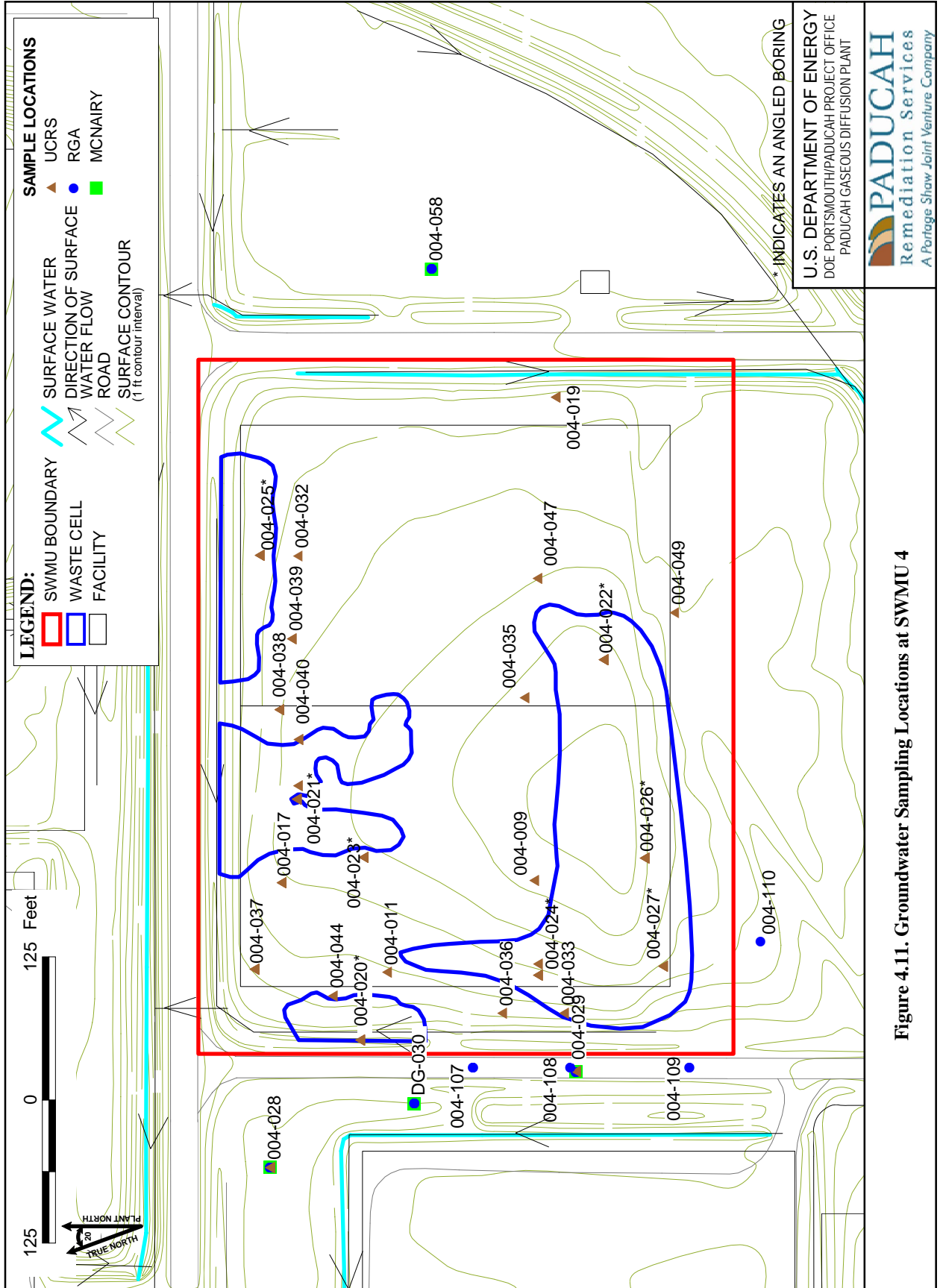


Figure 4.11. Groundwater Sampling Locations at SWMU 4

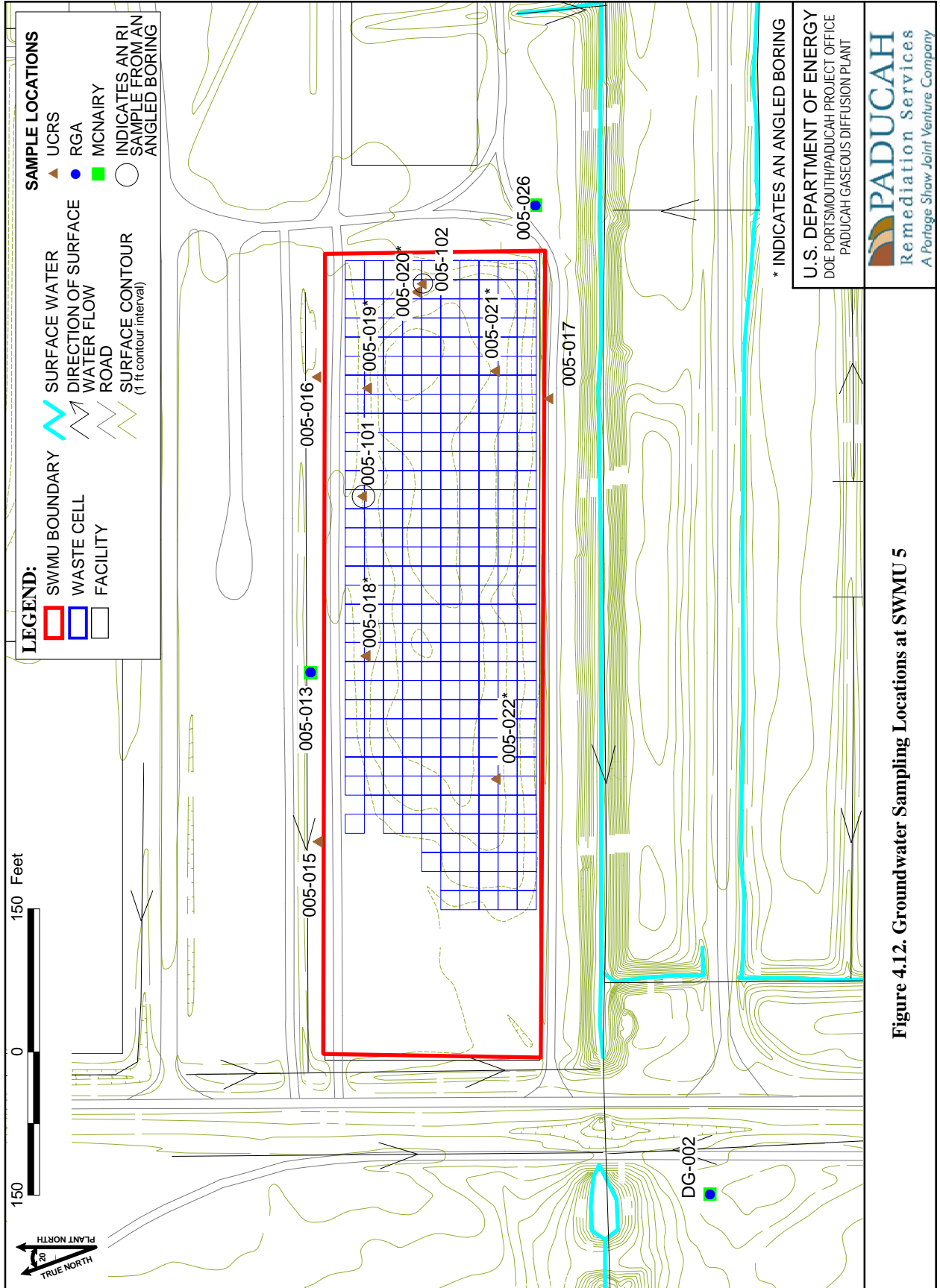


Figure 4.12. Groundwater Sampling Locations at SWMU 5

Figure No. 1BGO01d2R1-r1_sec14.apr
DATE 08-04-09

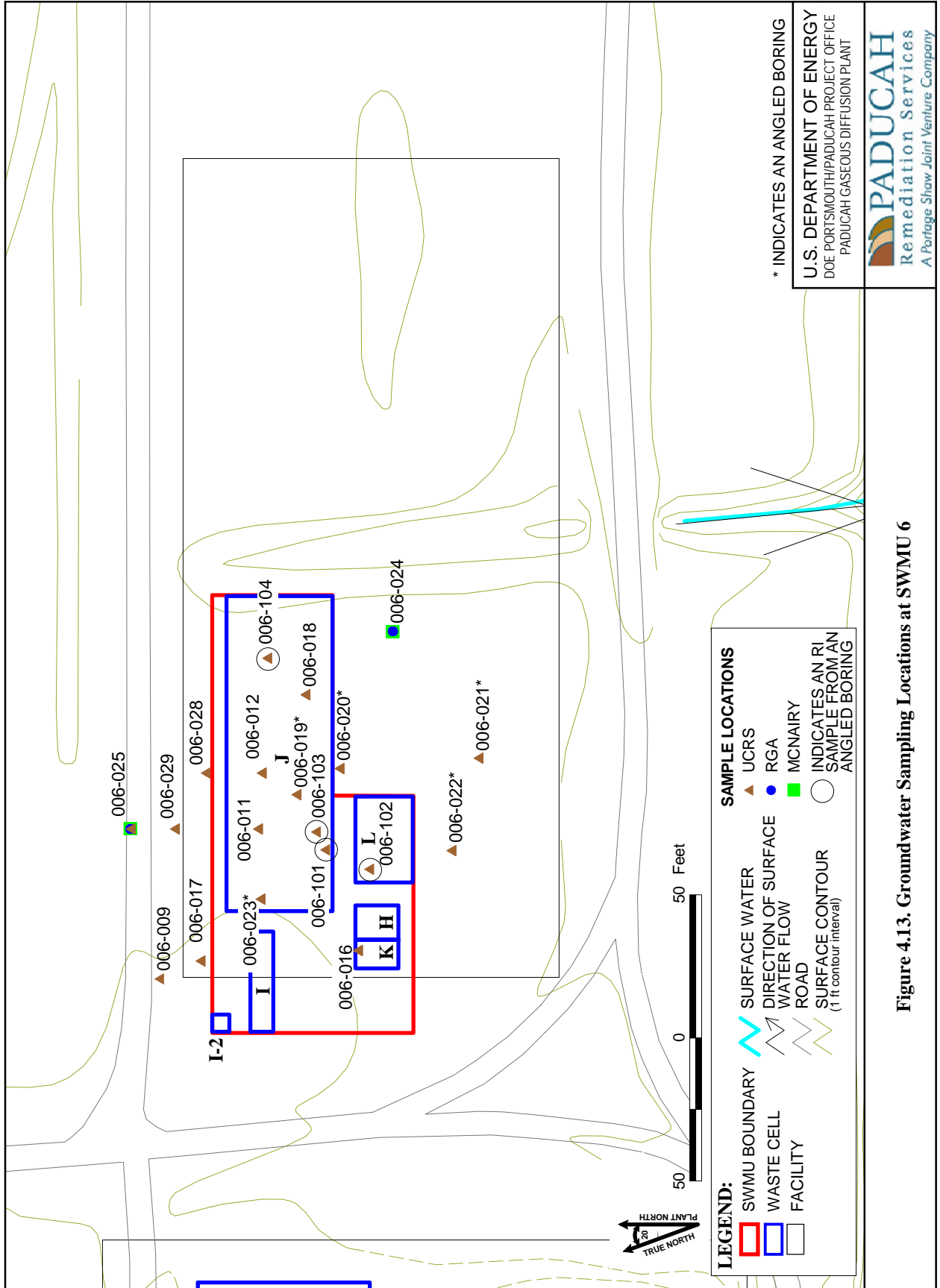
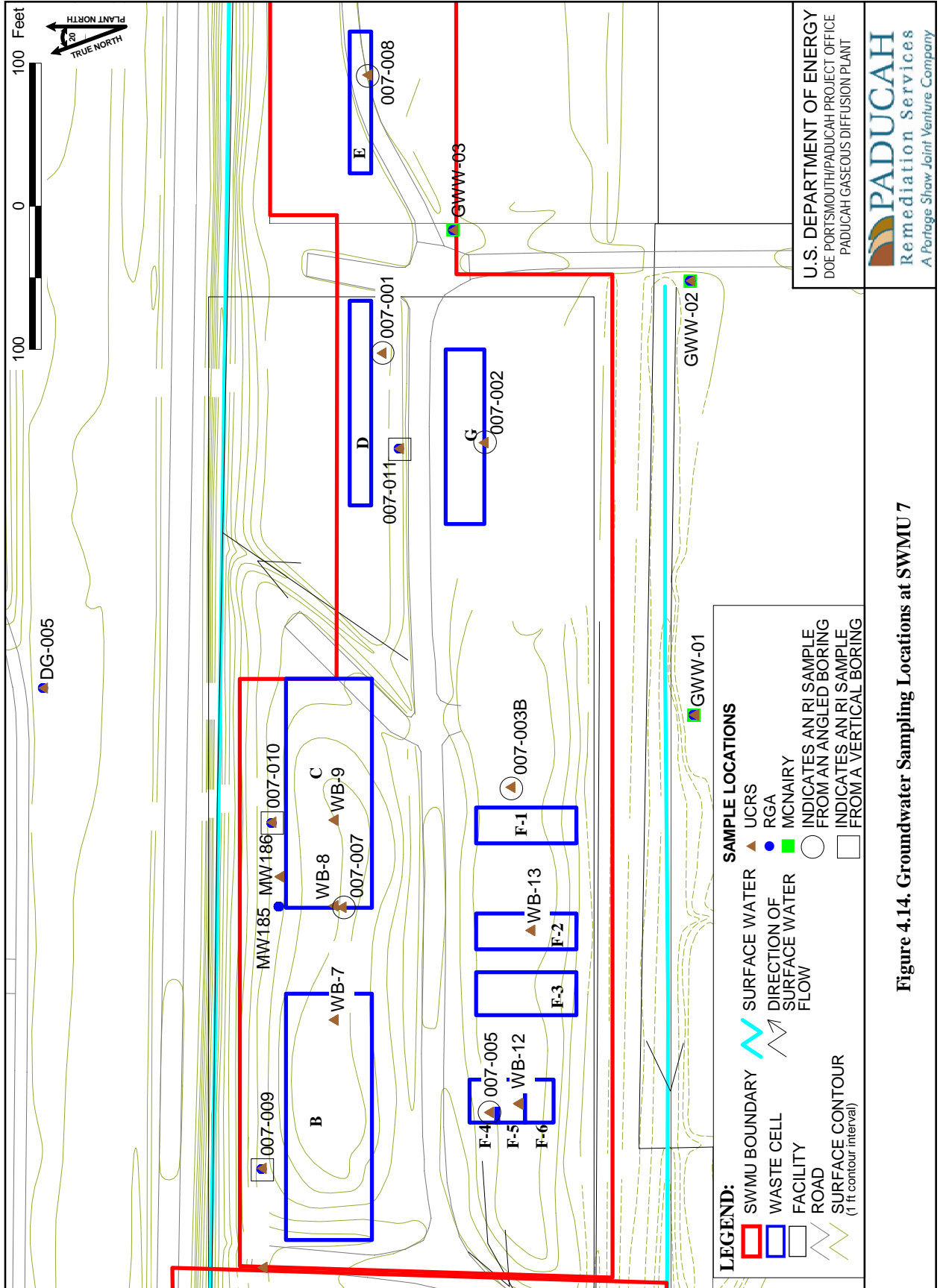


Figure 4.13. Groundwater Sampling Locations at SWMU 6

Figure No. IBG001d2R1-ri_sec14.apr
DATE 09-29-09



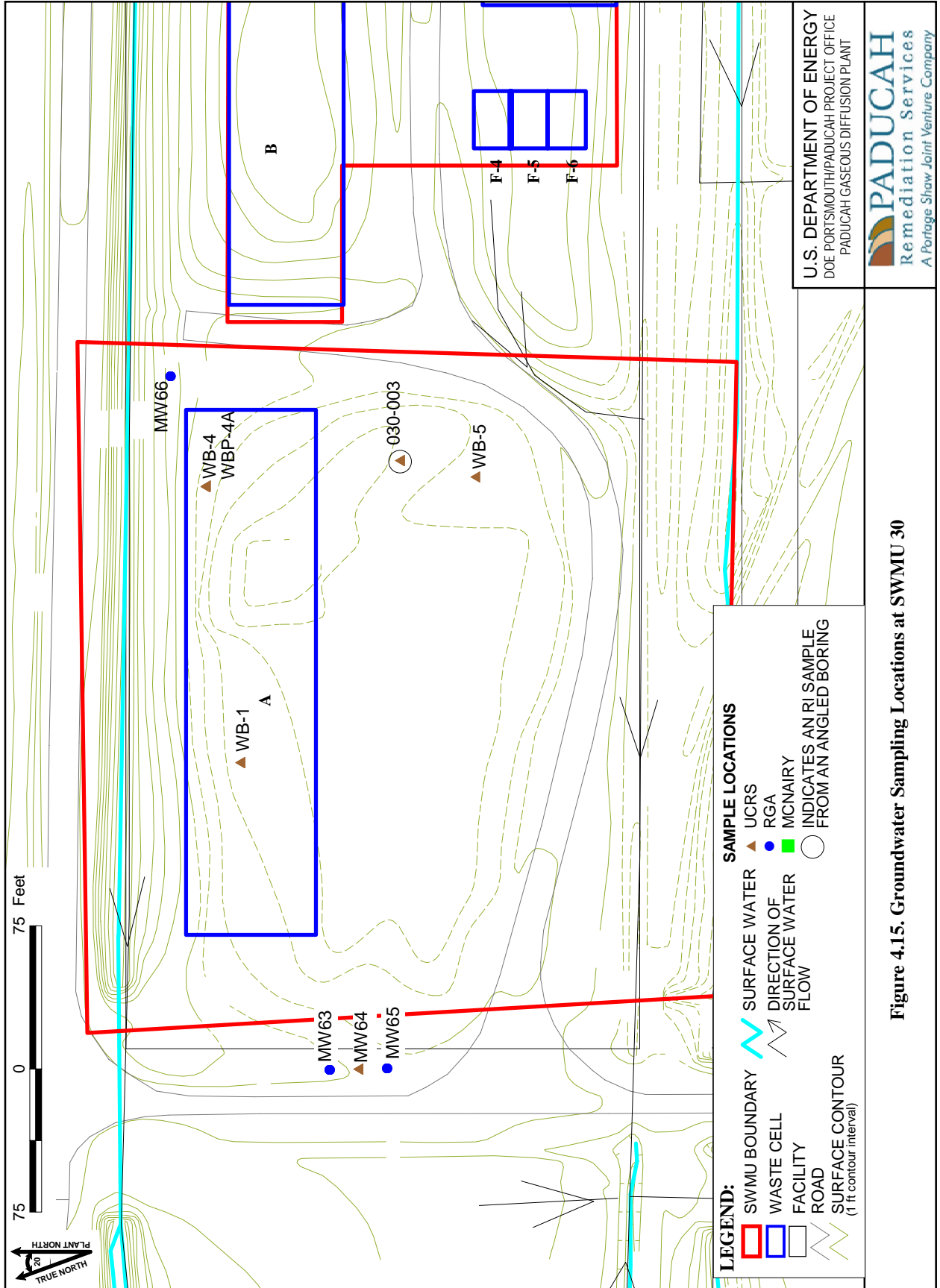


Figure 4.15. Groundwater Sampling Locations at SWMU 30

Figure No. IBGOUd2R1-ri_sec14.apr
DATE 08-04-09

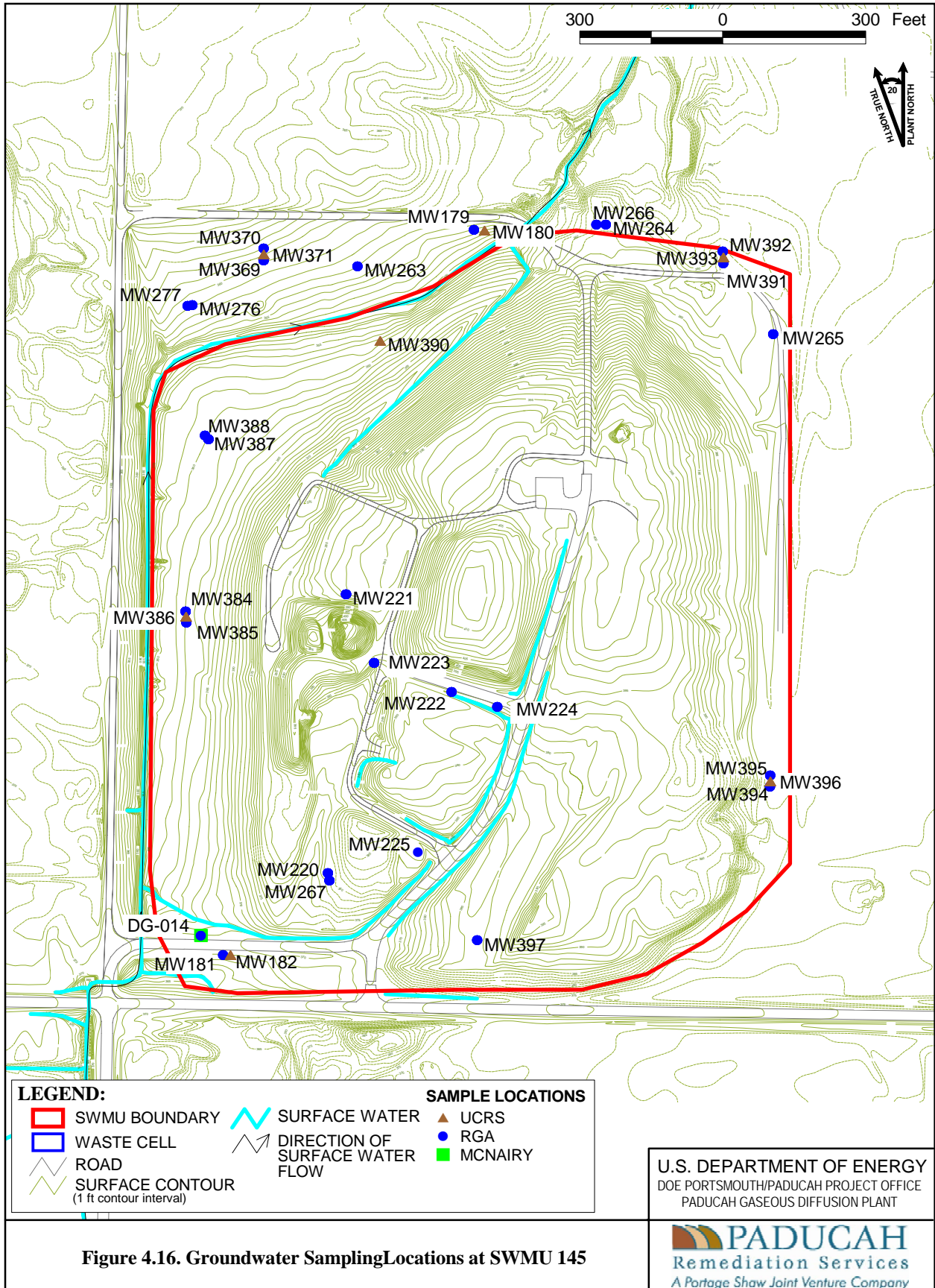


Figure 4.16. Groundwater Sampling Locations at SWMU 145

Figure No. 1BGOU\42R1-ri_sect4.apr
DATE 08-04-09

4.3 SWMU 2

4.3.1 Subsurface Soils

The RI collected subsurface soil samples from two angled borings at SWMU 2. Review of the RI data along with historical data identified the contaminants presented in Table 4.7. Table 4.8 lists the locations of the metals, organics, and radionuclides detected above screening levels.

As stated in Section 1.3.1.2, records for SWMU 2 indicate the waste consisted of uranium, oils (possibly containing PCBs), and TCE. Historical records indicate that a total of 270 tons of uranium and 59,000 gal of oils was disposed of at the burial ground. The maximum result of uranium (1,500 mg/kg) was detected at a depth of 5 ft bgs in boring SWMU 2-12 (subsurface soil locations shown in Figure 4.1). This boring, from the SWMU 2 Interim Remedial Design Investigation, was one of several sample locations intended to characterize soils adjacent to a waste pit.⁶ The next highest uranium result, 33 mg/kg at a depth of 12 ft bgs, was from boring SWMU 2-2 of the SWMU 2 Interim Remedial Design Investigation (also intended to characterize soils adjacent to a waste pit). The waste pits extend to a depth of approximately 17 ft. The most prevalent metals detected above background level in subsurface soil samples at SWMU 2 are arsenic, thallium, and uranium. Arsenic and thallium are commonly associated with uranium. Arsenic was detected above the screening levels throughout the depth of the angled borings (60 ft) installed by the RI (Figure 4.17). Based on Figure 4.17, the areas that exceed the background level are in the shallow soils on the eastern side of the SWMU and an isolated area at 45 ft bgs on the western side (the 60 ft sample at this location was less than background). Because this is a relatively small SWMU, these two zones may be connected spatially. Figure 4.18 shows the distribution of uranium at SWMU 2 with the highest concentrations being found at shallow depths on the western side of the burial ground.

TCE and its degradation products *cis*-1,2-DCE and vinyl chloride were detected at high levels (140 mg/kg, 130 mg/kg, and 1.4 mg/kg, respectively) in the historic sample location SWMU 2-2 at a depth of 12 ft bgs. (This is the only detection of *cis*-1,2-DCE above the excavation worker NAL in all analyses related to the BGOU SWMUs.) This boring was not in a known area of TCE burial in SWMU 2. Figure 4.19 presents the distribution of TCE in soil at SWMU 2. This shows the high level detected on the eastern side of the burial unit at location SWMU 2-2. Because no deeper samples are available from boring SWMU 2-2, the vertical extent of TCE contamination remains unknown. Nearby borings at locations SWMU 2-17 and SWMU 2-3 (Figure 4.1) collected soil samples into the RGA and did not detect TCE above the screening criteria. The RI data, which was used in modeling the source zone but is not shown in Figure 4.19, includes two detections of TCE at a level of 0.428 mg/kg (40 to 45 ft sample) and 0.366 mg/kg (60 ft sample) in borings 002-001 and 002-002, respectively. The degradation products also have a similar limited extent (see Appendix D). All other VOC detections in subsurface soils of SWMU 2 were less than 1 mg/kg. The distribution of *cis*-1,2-DCE is similar to that of TCE, with the exception that concentrations are less. While subsurface soil or groundwater concentrations do not indicate a TCE DNAPL, there is some potential, based on waste disposal records, that a DNAPL might exist. Although PCBs were suspected to be associated with the waste buried in SWMU 2, PCBs were detected above 1 ppm in only one subsurface soil sample below a depth of 6 ft (the approximate depth of the top of buried waste). (The maximum PCB detection in shallower subsurface soils was 0.06 mg/kg.) It remains unclear if PCB-contaminated oil was placed in uranium waste drums at SWMU 2 to prevent spontaneous combustion. The available sample analyses for SWMU 2 do not characterize the PCB levels in soils immediately below the buried waste.

⁶ Boring SWMU2-12 penetrated and sampled a waste drum at a depth of 7 to 8 ft.

The highest activities of the uranium isotopes uranium-234 (155 pCi/g) and uranium-238 (947 pCi/g) were detected at historic sample location SWMU 2-12 at a depth of 5 ft bgs. (The waste sample recovered from the penetrated drum contained 7.6 pCi/g uranium-234 and 43.5 pCi/g uranium-238.) All other detections of uranium isotopes in subsurface soil were less than 10 pCi/g. The distribution of the uranium isotopes are very similar to that of uranium shown in Figure 4.18 (the extent of contamination that exceeds background is mostly in shallow soils on the western side of the unit).

Table 4.7. SWMU 2 Subsurface Soil Contaminants

Analysis	Maximum Result		Frequency of Detection ^a	Frequency of Detection	
	Historical Data	RI Data		Above Background Value	Above Excavation Worker NAL
<i>Inorganics (mg/kg)</i>					
Aluminum	N/A	10,900	11/11	0/11	8/11
Arsenic	22	13.7	28/29	8/29	28/29
Barium	350	127	29/29	1/29	1/29
Beryllium	1.3	1.05	19/29	4/29	1/29
Cadmium	0.26	ND	11/29	5/29	0/29
Calcium	N/A	1,420	11/11	0/11	N/A
Chromium	26	35.7	29/29	0/29	0/29
Cobalt	N/A	20	7/11	1/11	0/11
Copper	N/A	10.9	11/11	0/11	0/11
Iron ^b	N/A	34,900	11/11	1/11	11/11
Lead	N/A	10.3	11/11	0/11	0/11
Magnesium	N/A	1,330	11/11	0/11	0/11
Manganese	1,200	481	29/29	2/29	25/29
Mercury	N/A	0.019	2/11	0/11	0/11
Nickel	23	15.1	25/29	1/29	0/29
Silver	2.5	ND	4/29	0/29	0/29
Sodium	N/A	445	5/11	2/11	0/11
Thallium	1.7	ND	10/29	10/29	7/29
Uranium	1,500	15.3	12/58	10/58	7/58
Vanadium	38	23.2	28/29	1/29	27/29
Zinc	N/A	42.6	4/11	0/11	0/11
<i>Organics--Volatiles (mg/kg)</i>					
<i>cis</i> -1,2-Dichloroethene	130	0.118	6/29	N/A	1/29
Trichloroethene ^c	140	0.428	10/58	N/A	1/58
Vinyl chloride	1.4	ND	1/29	N/A	1/29
<i>Organics--PCBs (mg/kg)</i>					
PCB, Total	4.2	ND	5/28	N/A	1/28
PCB-1248	4.2	ND	5/28	N/A	1/28
<i>Radionuclides(pCi/g)</i>					
Americium-241	0.48	ND	47/58	N/A	0/58
Cesium-137	0.21	ND	7/58	0/58	4/58
Neptunium-237	0.12	ND	43/57	N/A	0/57
Plutonium-239	0.09	N/A	47/47	N/A	0/47
Technetium-99	2.24	ND	46/57	0/57	0/57
Thorium-228	N/A	0.57	11/11	0/11	11/11
Thorium-230	1.55	0.448	52/58	4/58	0/58
Thorium-232	N/A	0.646	10/11	0/11	0/11
Thorium-234	N/A	11	3/11	N/A	N/A
Uranium	N/A	6.77	1/11	N/A	N/A
Uranium-234	155	0.824	52/58	1/58	1/58
Uranium-235	N/A	0.08	1/11	0/11	0/11
Uranium-235/236	25.8	N/A	47/47	3/47	1/47
Uranium-238	947	5.87	52/58	11/58	12/58

^a Frequency of detection is the number of detections of an analyte per number of analyses (includes regular and duplicate samples).

^b One of the samples for iron is a duplicate (10 primary samples shown in Table 4.8).

^c Three of the samples for TCE are duplicates (55 primary samples shown in Tale 4.8).

N/A = not applicable

ND = not detected

Table 4.8. SWMU 2 Locations of Subsurface Soil Contaminants

Analysis <i>Inorganics (mg/kg)</i>	Depth (ft)	Excavation Worker NAL	Background	RI Data		Historical Data													
				002-001	002-002	SWMU2-1	SWMU2-12	SWMU2-13	SWMU2-17	SWMU2-2	SWMU2-3	SWMU2-4	SWMU2-5	SWMU2-8	SWMU2-9				
Aluminum	10-12	5,250	12,000	5,540	5,940														
	15-16	5,250	12,000	6,660	7,290														
	30-35	5,250	12,000	6,670	5,350														
	40-45	5,250	12,000	5,110	10,900														
	60-70	5,250	12,000	2,620	7,960														
	01-05	0.324	7.9								12								
	05-10	0.324	7.9																
Arsenic	10-12	0.324	7.9	2.47	1.22	1.8	6.8	4.5											
	15-16	0.324	7.9	2.06	1.21	1.9	4.6												
	20-25	0.324	7.9			1.1	1.7												
	30-35	0.324	7.9	3.98	2.52														
	40-45	0.324	7.9	1.26	13.7														
	60-70	0.324	7.9	2.38	2.02														
	01-05	272	170								160								
Barium	05-10	272	170																
	10-12	272	170	61.9	54.7	89	350												
	15-16	272	170	62.7	55.8	63	86												
	20-25	272	170			49	86												
	30-35	272	170	20.1	17.4														
	40-45	272	170	16.9	127														
	60-70	272	170	43.4	38.8														
Beryllium	01-05	1.26	0.69																
	05-10	1.26	0.69			1.3													
	10-12	1.26	0.69	0.487U	0.427U	0.52	0.55												
	15-16	1.26	0.69	0.479U	0.45U	0.38	0.75												
	20-25	1.26	0.69			0.29	0.39												
	30-35	1.26	0.69	0.453U	0.498U														
	40-45	1.26	0.69	0.469U	1.05														
Cadmium	60-70	1.26	0.69	0.475U	0.432U														
	01-05	15.2	0.21																
	05-10	15.2	0.21			0.091													
	10-12	15.2	0.21	1.95U	1.71U	0.033U	0.25												
	15-16	15.2	0.21	1.92U	1.8U	0.033U	0.25												
	20-25	15.2	0.21			0.033U	0.033U												
	30-35	15.2	0.21	1.81U	1.99U														
40-45	15.2	0.21	1.88U	1.84U															
60-70	15.2	0.21	1.9U	1.73U															

Table 4.8. SWMU 2 Locations of Subsurface Soil Contaminants (Continued)

Analysis	Depth (ft)	Excavation Worker NAL	Background	Historical Data																
				RI Data	SWMU2-1	SWMU2-12	SWMU2-13	SWMU2-17	SWMU2-2	SWMU2-3	SWMU2-4	SWMU2-5	SWMU2-8	SWMU2-9						
Calcium	10-12	n/a	6,100	002-001	1,420	858														
	15-16	n/a	6,100		1,100	965														
	30-35	n/a	6,100		867	657														
	40-45	n/a	6,100		445	915														
	60-70	n/a	6,100		281	577														
				43				15			15									
Chromium	05-10	476	43																	
	10-12	476	43		10.7	8.62				21			13							9.4
	15-16	476	43		9.51	9.12				21			13							19
	20-25	476	43										11							11
	30-35	476	43										17							11
	40-45	476	43		35.7	4.11														
Cobalt	40-45	476	43		4.94	12.2														
	60-70	476	43		5.6	5.89														
	10-12	1,110	13		2.43U	2.8														
	15-16	1,110	13		3.17	5.3														
	30-35	1,110	13		4.43	2.49U														
	40-45	1,110	13		2.59	20														
Copper	60-70	1,110	13		4.18	2.16U														
	10-12	427	25		6.92	5.26														
	15-16	427	25		4.82	5.51														
	30-35	427	25		7.34	3.74														
	40-45	427	25		3.34	10.9														
	60-70	427	25		2.9	5.91														
Iron	10-12	2,170	28,000		8,250	5,950														
	15-16	2,170	28,000		7,900	6,830														
	30-35	2,170	28,000		13,600	10,600														
	40-45	2,170	28,000		7,110	34,900														
	60-70	2,170	28,000		7,190	12,800														
				23		4.7	5.4													
Lead	10-12	50	23		6.07	6.58														
	15-16	50	23		6.07	6.58														
	30-35	50	23		6.9	4.31														
	40-45	50	23		4.19	10.3														
	60-70	50	23		1.85	4.63														
				23		4.7	5.4													
Magnesium	10-12	n/a	2,100		1,330	750														
	15-16	n/a	2,100		857	812														
	30-35	n/a	2,100		442	325														
	40-45	n/a	2,100		238	882														
	60-70	n/a	2,100		238	814														
				2,100		238	814													

Table 4.8. SWMU 2 Locations of Subsurface Soil Contaminants (Continued)

Analysis	Depth (ft)	Excavation Worker NAL	Background	RI Data		Historical Data																		
				002-001	002-002	SWMU2-1	SWMU2-12	SWMU2-13	SWMU2-17	SWMU2-2	SWMU2-3	SWMU2-4	SWMU2-5	SWMU2-8	SWMU2-9									
PCB-1248	01-05	0.168	n/a				0.058				0.00074U													
	05-10	0.168	n/a			0.00074U					0.00074U					0.00074U								
	10-12	0.168	n/a	0.1U	0.1U	0.00074U	0.031				4.2					0.00074U								
	15-16	0.168	n/a	0.1U	0.09U	0.00074U	0.015									0.00074U								
	20-25	0.168	n/a				0.041									0.00074U								
	30-35	0.168	n/a	0.1U	0.09U																			
	40-45	0.168	n/a	0.09U	0.09U																			
	60-70	0.168	n/a	0.1U	0.09U																			
Radionuclides (pCi/g) Americium-241	01-05	1.74	n/a								0.13										0.37			
	05-10	1.74	n/a			0.17					0.18											0.11		
	10-12	1.74	n/a	0.0297U	0.0295U	0.12	0.24	0.31			0.15					0.28						0.13		
	15-16	1.74	n/a	0.0294U	0.0293U	0.16	0.18				0.14					0.15						0.13		
	20-25	1.74	n/a			0.12	0.14	0.14								0.13						0.15		
	30-35	1.74	n/a	0.0296U	0.0293U						0.12					0.08						0.1		
	40-45	1.74	n/a	0.0294U	0.0298U			0.1	0.48		0.09					0.08						0.11		
	50-55	1.74	n/a					0.31	0.1		0.39					0.08						0.08		
	60-70	1.74	n/a	0.0298U	0.0292U						0.08					0.08							0.08	
	75	1.74	n/a					0.04															0.04	
Cesium-137	85	1.74	n/a																					
	95	1.74	n/a																					
	01-05	0.115	0.28				0.00009U				0.05												0.14	
	05-10	0.115	0.28			0.0001U					0.00004U					0.00008U							0.00008U	
	10-12	0.115	0.28	0.0431U	0.0496U	0.00009U	0.00004U	0.04			0.00004U					0.00009U							0.00007U	
	15-16	0.115	0.28	0.0444U	0.0404U	0.00007U	0.00005U				0.00005U					0.00008U							0.00008U	
	20-25	0.115	0.28			0.00009U	0.00004U				0.21					0.00009U							0.00007U	
	30-35	0.115	0.28	0.0524U	0.039U						0.00004U					0.00009U							0.00003U	
	40-45	0.115	0.28	0.0393U	0.0567U						0.15					0.00009U								0.00005U
	50-55	0.115	0.28								0.13					0.00004U								0.00004U
60-70	0.115	0.28	0.0374U	0.0478U						0.00005U					0.00004U								0.00004U	
75	0.115	0.28								0.00003U					0.00004U								0.00003U	
85	0.115	0.28								0.0002U														
95	0.115	0.28								0.00003U														

Table 4.8. SWMU 2 Locations of Subsurface Soil Contaminants (Continued)

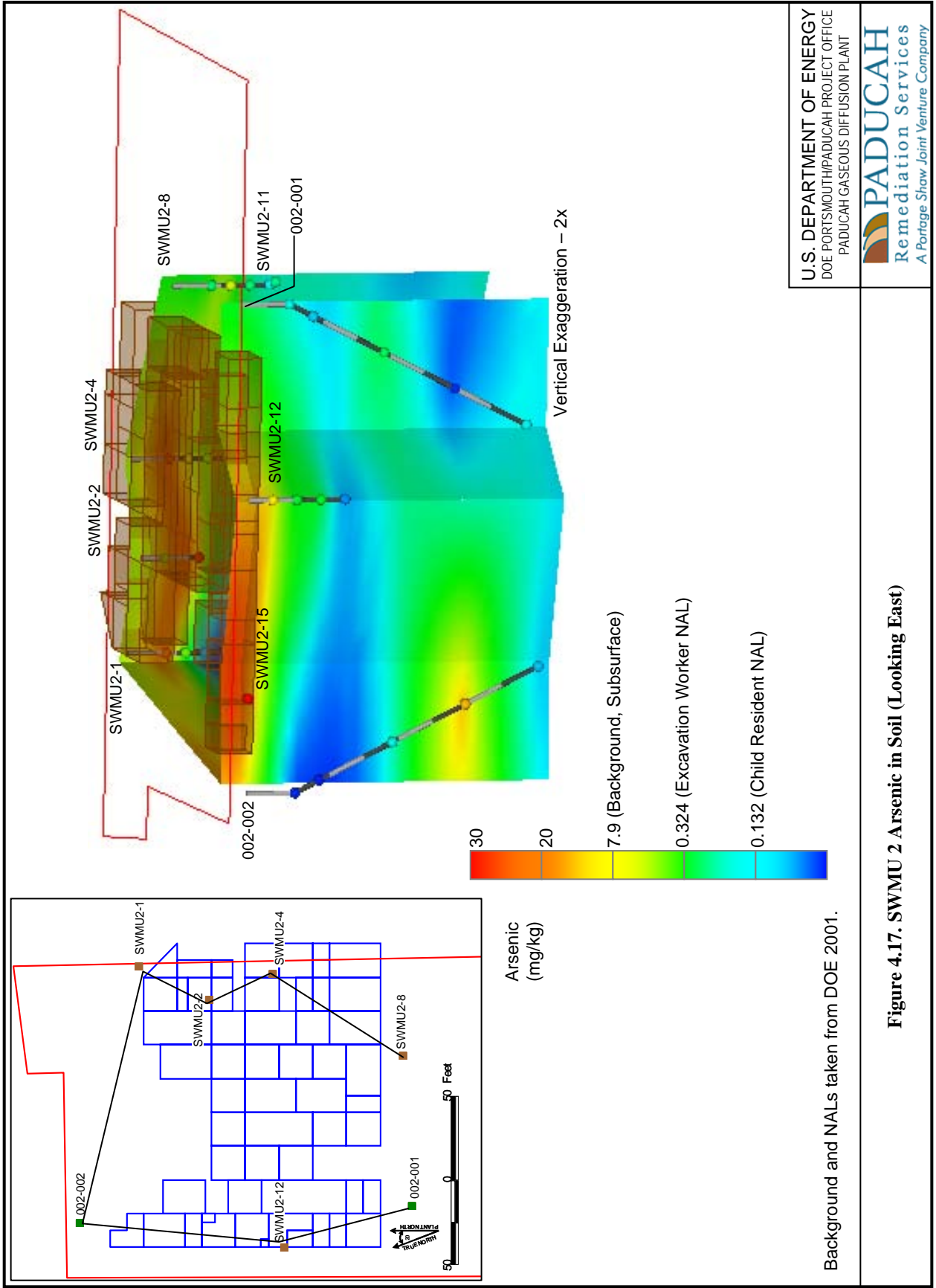
Analysis	Depth (ft)	Excavation Worker NAL	Background	RI Data		Historical Data															
				002-001	002-002	SWMU2-1	SWMU2-12	SWMU2-13	SWMU2-17	SWMU2-2	SWMU2-3	SWMU2-4	SWMU2-5	SWMU2-8	SWMU2-9						
Uranium-235	10-12	0.455	0.14	0.0374U	0.08																
	15-16	0.455	0.14	0.0385U	0.0376U																
	30-35	0.455	0.14	0.0383U	0.0383U																
	40-45	0.455	0.14	0.0382U	0.0379U																
	60-70	0.455	0.14	0.0377U	0.0383U																
	01-05	0.455	0.14			0.19															0.1
	05-10	0.455	0.14																		
	10-12	0.455	0.14			0.07															
	15-16	0.455	0.14			0.08															
	20-25	0.455	0.14			0.04															
30-35	0.455	0.14																			
40-45	0.455	0.14																			
50-55	0.455	0.14																			
60-70	0.455	0.14																			
75	0.455	0.14																			
85	0.455	0.14																			
95	0.455	0.14																			
Uranium-238	01-05	1.17	1.2																		
	05-10	1.17	1.2																		
	10-12	1.17	1.2	0.13U	5.87	6.25															
	15-16	1.17	1.2	0.13U	0.13U	1.02															
	20-25	1.17	1.2			0.84															
	30-35	1.17	1.2	0.319	0.132																
	40-45	1.17	1.2	0.131U	0.241																
	50-55	1.17	1.2																		
	60-70	1.17	1.2	0.13U	0.206																
	75	1.17	1.2																		
85	1.17	1.2																			
95	1.17	1.2																			

Blank cells indicate interval was not sampled for the specified analysis. Maximum value shown for each depth interval.

Bold indicates result is greater than NAL value.

Italics indicates result is greater than background value.

Bold + Italics indicate result is greater than both NAL and background values.

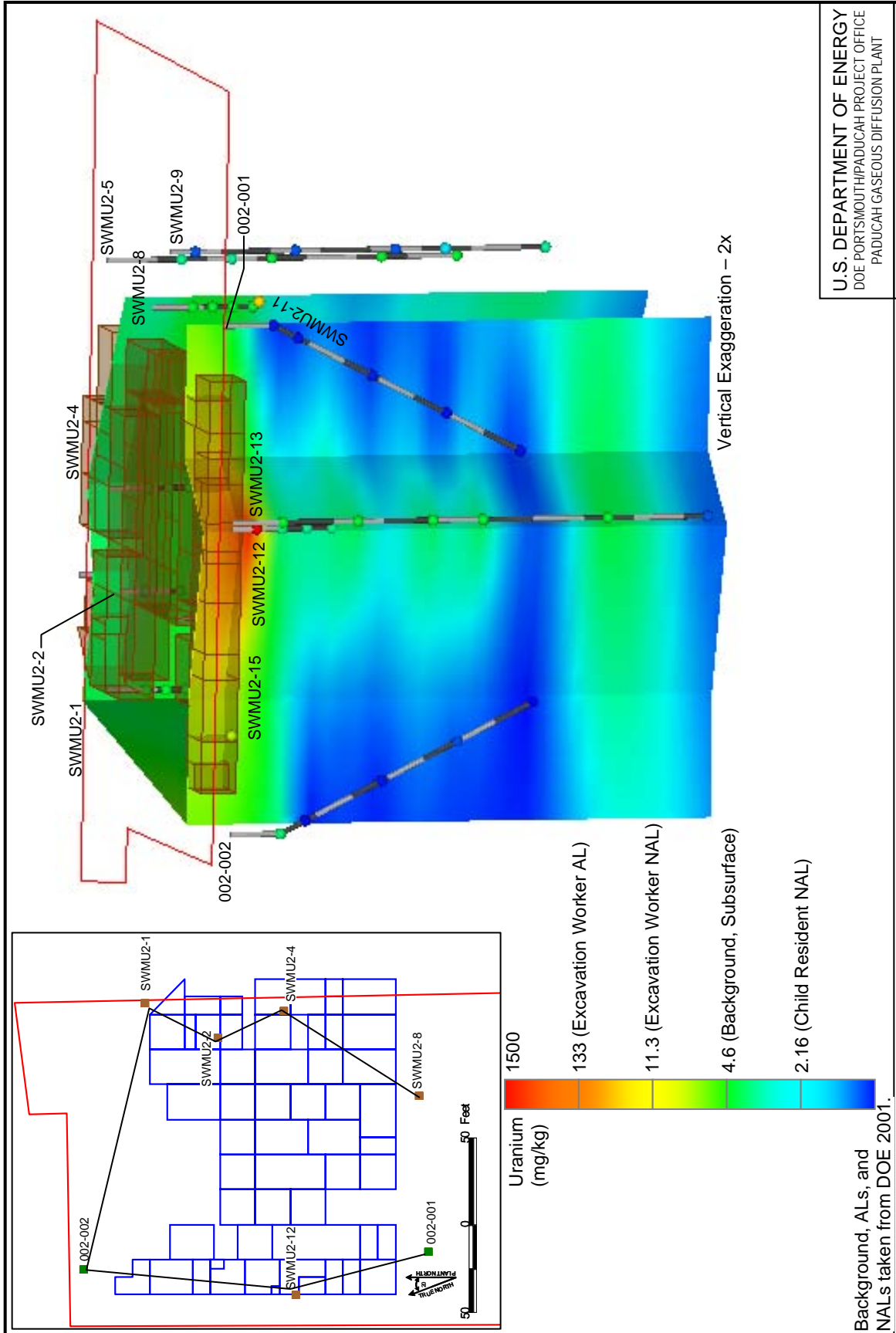


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Figure 4.17. SWMU 2 Arsenic in Soil (Looking East)

Background and NALs taken from DOE 2001.



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Figure 4.18. SWMU 2 Uranium in Soil (Looking East)

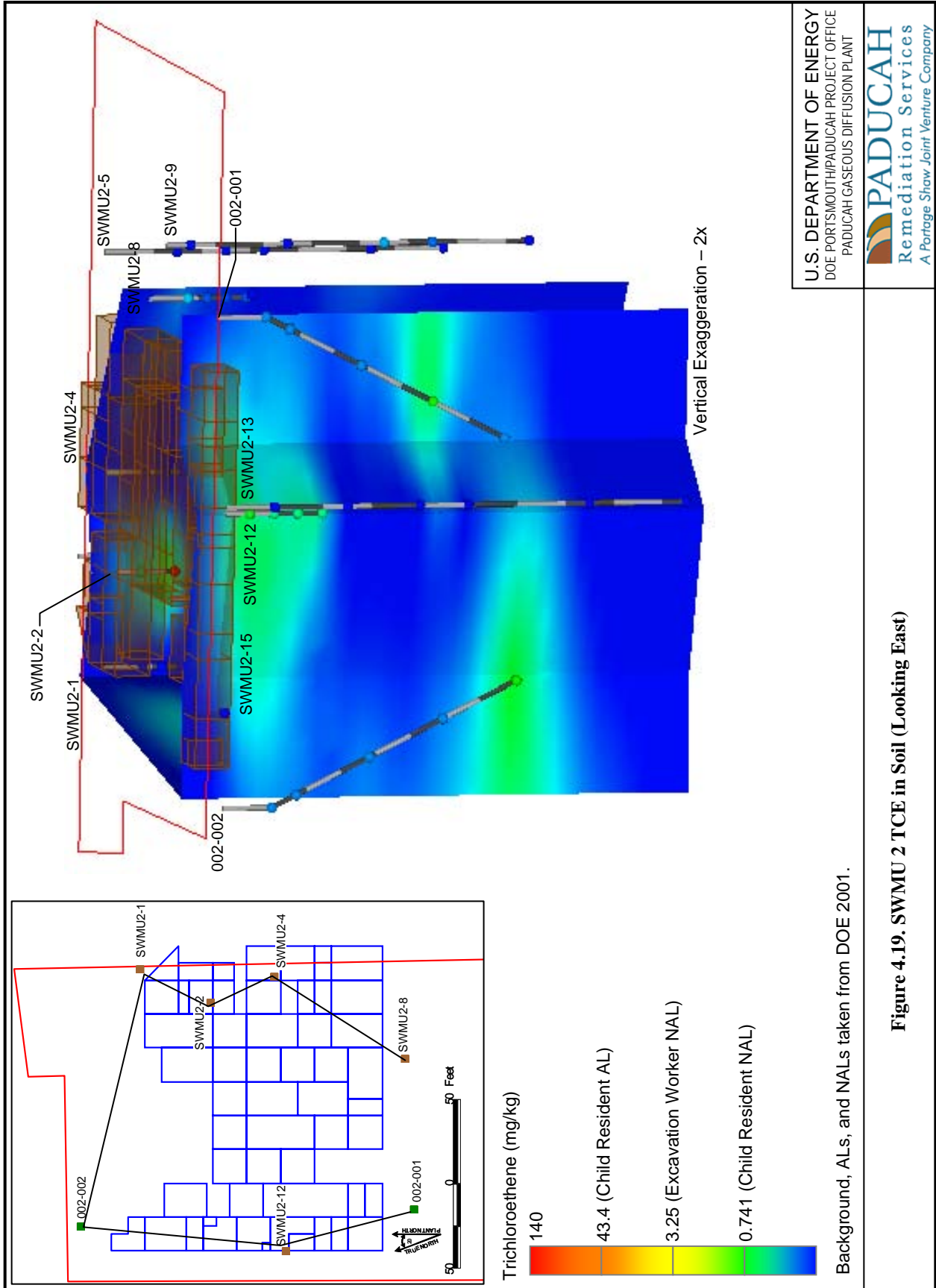


Figure 4.19. SWMU 2 TCE in Soil (Looking East)

4.3.2 SWMU 2 Groundwater

Groundwater samples were attempted at the two angled borings installed at SWMU 2 as part of this RI; however, none was collected (even where the UCRS is saturated, the low hydraulic conductivity of the unit restricts groundwater yield). A review of historical data, identified the contaminants listed in Tables 4.9, 4.10, and 4.11 for UCRS, RGA, and McNairy groundwater, respectively. Table 4.12 provides detail (depth, sample location, and analytical results) for SWMU 2 groundwater samples, including nondetects and detections above screening levels, and Figure 4.9 shows groundwater sampling locations.

UCRS characterization data are derived from three sources (all of these wells, piezometers, and borings are located on the perimeter of the burial cells):

- 1) Samples from PZ74 (1996 through 1998) and MW154 (1990 through 1996);
- 2) Samples from temporary borings (SWMU2-3, SWMU2-9, SWMU2-10, and SWMU 2-17) drilled during the SWMU 2 Interim Remedial Design Investigation of 1996 (DOE 1997a); and
- 3) Samples from piezometers PZ334, PZ335, and PZ336 in 1998 to assess the potential mobility of dissolved uranium.

Locations MW154, PZ334, PZ335, and PZ336 (Figure 4.9) directly monitor the horizon of the buried waste around the perimeter of the burial ground. Samples from all of the temporary borings at depths of 22 to 26 ft (within the HU2 interval of the UCRS) characterize groundwater immediately below the depth of the waste pits (excavated at depths of 7 to 17 ft) but on the perimeter of the burial ground. Locations PZ74 and SWMU2-9 (42-43 ft sample) sample the deeper HU3 interval within the UCRS. Because these wells, piezometers, and soil borings are located on the perimeter of the burial cells, the contaminant loading to the RGA from SWMU 2 is uncertain.

The screen of the SWMU 2 analyses identified the metals beryllium, iron, manganese, uranium, and vanadium and the organics TCE; *cis*-1,2-DCE; and vinyl chloride (TCE and its reductive dechlorination products) and 1,1-DCE as UCRS contaminants. In addition, uranium-234 and uranium-238 levels frequently exceeded background and child resident NALs.

The only metal and radionuclides that exceeded screening criteria in the horizon of the burial cells were uranium and the uranium isotopes (similar to soil contamination shown in Figure 4.18). Beryllium, manganese, and vanadium, the uranium isotopes, and TCE and its degradation products occurred at levels that exceed screening criteria throughout the UCRS interval below but peripheral to the waste pits.

Characterization data for the RGA come from seven temporary borings of the SWMU 2 Interim Remedial Design Investigation of 1996 (DOE 1997a) and MW333, MW337, and MW338 (for the period 1996 to present) (Figure 4.9). The metals that exceeded screening criteria include beryllium, iron, manganese, uranium, vanadium (also identified as UCRS contaminants) and arsenic and cadmium.

TCE, with a maximum value of 5.35 mg/L (5,350 µg/L), was the most widely detected organic contaminant in RGA groundwater at SWMU 2. Another VOC, 1,1-DCE, showed high levels in the RGA (47.9 mg/L) from historical boring SWMU 2-5. Because TCE is believed to be buried in SWMU 2, it is likely that higher levels of TCE and other VOCs may have occurred directly beneath the buried wastes. The hydrogeological assessment of the SWMUs 2 and 3 area (PRS 2007a) determined that an upgradient source is responsible for the high TCE levels in the area. Figure 4.20 shows SWMUs 2, 3, and 4 in

relation to the underlying TCE plume in the RGA.⁷ The C-400 TCE DNAPL zone, located central to the plant, appears to be a major contributor. It is difficult to separate any potential impacts of VOC migration to the RGA from SWMU 2 due to the migration of contamination from upgradient areas. Figures 4.21, 4.22, and 4.23 present trend graphs of TCE, ⁹⁹Tc, and *cis*-1,2-DCE, respectively, in RGA MWs. The increases in TCE and *cis*-1,2-DCE in MW333 and the increase in technetium-99 in MW337 are caused by upgradient sources. Similar patterns are seen upgradient in SWMU 3 MWs. The elevated contaminant levels in the RGA due to upgradient sources potentially may be masking the migration of the same contaminants from SWMU 2.

RGA groundwater samples from the location SWMU2-17 contained both uranium-234 and uranium-238 above screening criteria at 50.6 and 55.1 pCi/L, respectively. The analysis of a sample from location SWMU2-16 detected uranium-238 at 91.7 pCi/L.

Four of the temporary borings of the SWMU 2 Interim Remedial Design Investigation of 1996 (DOE 1997a), SWMU2-3, SWMU2-9, SWMU2-10, and SWMU2-17 (Figure 4.9), characterized groundwater in the McNairy Formation immediately below the RGA. TCE and 1,1-DCE were the only groundwater contaminants identified by comparison against the RI screening criteria.

⁷ Because of its age (circa 1996), the SWMU 2 TCE analysis of 5.35 mg/L was not used in preparation of the TCE plume map of the Southwest Plume Site Investigation, representing calendar year 2004. Thus, Figure 4.20, taken from the Southwest Plume Site Investigation, does not depict the majority of the SWMU 2 area within the 1,000-to-10,000 µg/L area of the TCE plume.

Table 4.9. SWMU 2 UCRS Groundwater Contaminants

Analysis	Historical Data	RI	Frequency of Detection ^a	Frequency of Detection		
				Above Background	Above NAL	Above MCL
Metals (mg/L)						
Beryllium	0.078	N/A	5/6	N/A	3/6	3/6
Calcium	130	N/A	11/11	N/A	N/A	N/A
Iron	14	N/A	1/1	N/A	1/1	N/A
Magnesium	56	N/A	11/11	N/A	0/11	N/A
Manganese	37	N/A	6/6	N/A	5/6	N/A
Potassium	36	N/A	6/11	N/A	N/A	N/A
Sodium	193	N/A	11/11	N/A	0/11	N/A
Uranium	0.075	N/A	7/15	N/A	7/15	3/15
Vanadium	4.1	N/A	5/6	N/A	5/6	N/A
Radionuclides (pCi/L)						
Americium-241	0.34	N/A	6/6	N/A	0/6	N/A
Neptunium-237	0.02	N/A	4/6	N/A	0/6	N/A
Plutonium-239	0.17	N/A	5/6	N/A	0/6	N/A
Technetium-99	170	N/A	6/6	N/A	5/6	0/6
Thorium-230	0.77	N/A	6/6	N/A	6/6	N/A
Uranium-234	10.3	N/A	10/10	N/A	9/10	N/A
Uranium-235	0.48	N/A	4/4	N/A	0/4	N/A
Uranium-235/236	1.1	N/A	4/6	N/A	N/A	N/A
Uranium-238	55.8	N/A	10/10	N/A	9/10	N/A
Volatiles (mg/L)						
1,1-Dichloroethene	8.33	N/A	4/4	N/A	4/4	4/4
cis-1,2-Dichloroethene	0.28	N/A	4/12	N/A	2/12	2/12
Trichloroethene	0.04	N/A	7/12	N/A	6/12	6/12
Vinyl chloride	0.005	N/A	2/7	N/A	2/7	1/7

^a Frequency of detection is the number of detections of an analyte per number of analyses (includes regular and duplicate samples).

^b N/A = not applicable

Table 4.10. SWMU 2 RGA Groundwater Contaminants

Analysis	Historical Data	RI	Frequency of Detection ^a	Frequency of Detection		
				Above Background	Above NAL	Above MCL
Metals (mg/L)						
Arsenic	0.081	N/A	3/4	2/4	3/4	2/4
Barium	2.4	N/A	4/4	1/4	2/4	1/4
Beryllium	0.092	N/A	24/28	21/28	22/28	21/28
Cadmium	0.012	N/A	1/4	1/4	1/4	1/4
Calcium	960	N/A	29/29	14/29	N/A	N/A
Chromium	0.35	N/A	2/4	2/4	0/4	2/4
Cobalt	0.0019	N/A	1/3	0/3	0/3	N/A
Iron	23,000	N/A	12/13	10/13	11/13	N/A
Magnesium	110	N/A	29/29	17/29	0/29	N/A
Manganese	96	N/A	28/28	25/28	26/28	N/A
Nickel	0.33	N/A	3/4	0/4	2/4	N/A
Potassium	86	N/A	26/29	20/29	N/A	N/A
Sodium	120	N/A	29/29	2/29	0/29	N/A
Uranium	0.41	N/A	5/145	4/145	5/145	4/145
Vanadium	1.9	N/A	25/28	19/28	24/28	N/A
Radionuclides (pCi/L)						
Americium-241	1.39	N/A	23/32	N/A	6/32	N/A
Neptunium-237	0.6	N/A	21/36	0/36	1/36	N/A
Plutonium-239	4.29	N/A	22/24	10/24	4/24	N/A
Technetium-99	229	N/A	54/115	32/115	38/115	0/115
Thorium-230	1.05	N/A	24/42	0/42	12/42	N/A
Uranium-234	50.6	N/A	24/33	19/33	22/33	N/A
Uranium-235/236	6.47	N/A	23/24	7/24	N/A	N/A
Uranium-238	91.7	N/A	23/34	18/34	21/34	N/A
Volatiles (mg/L)						
1,1-Dichloroethane	0.0022	N/A	1/92	N/A	0/92	N/A
1,1-Dichloroethene	47.9	N/A	20/39	N/A	20/39	20/39
Chloroform	0.0029	N/A	2/19	N/A	2/19	N/A
cis-1,2-Dichloroethene	0.75	N/A	31/137	N/A	24/137	10/137
trans-1,2-Dichloroethene	0.0043	N/A	3/137	N/A	0/137	0/137
Trichloroethene	5.35	N/A	113/137	N/A	109/137	97/137
Vinyl chloride	0.0014	N/A	1/57	N/A	1/57	0/57
Wetchem Parameters (mg/L)						
Cyanide	0.0075	N/A	1/1	N/A	0/1	0/1

^a Frequency of detection is the number of detections of an analyte per number of analyses (includes regular and duplicate samples).

^b N/A = not applicable

Table 4.11. SWMU 2 McNairy Groundwater Contaminants

Analysis	Historical Data	RI	Frequency of Detection ^a	Frequency of Detection		
				Above Background	Above PRG	Above MCL
Metals (mg/L)						
Beryllium	0.067	N/A	4/4	2/4	2/4	2/4
Calcium	120	N/A	4/4	2/4	N/A	N/A
Magnesium	86	N/A	4/4	2/4	0/4	N/A
Manganese	80	N/A	4/4	4/4	4/4	N/A
Potassium	83	N/A	4/4	1/4	N/A	N/A
Sodium	17	N/A	4/4	0/4	0/4	N/A
Vanadium	1.5	N/A	4/4	2/4	4/4	N/A
Radionuclides (pCi/L)						
Americium-241	0.32	N/A	4/4	N/A	0/4	N/A
Neptunium-237	0.15	N/A	4/4	0/4	0/4	N/A
Plutonium-239	0.08	N/A	4/4	0/4	0/4	N/A
Technetium-99	20	N/A	3/3	0/3	1/3	0/3
Thorium-230	0.94	N/A	4/4	0/4	3/4	N/A
Uranium-234	16.3	N/A	4/4	3/4	3/4	N/A
Uranium-235/236	1.7	N/A	4/4	2/4	N/A	N/A
Uranium-238	19.6	N/A	4/4	3/4	3/4	N/A
Volatiles (mg/L)						
1,1-Dichloroethene	7.5	N/A	4/4	N/A	4/4	4/4
cis-1,2-Dichloroethene	0.0232	N/A	2/9	N/A	1/9	0/9
Trichloroethene	0.055	N/A	7/9	N/A	5/9	4/9
Wetchem Parameters (mg/L)						
Cyanide	0.013	N/A	1/1	N/A	0/1	0/1

^a Frequency of detection is the number of detections of an analyte per number of analyses (includes regular and duplicate samples).

^b N/A = not applicable

Table 4.12. SWMU 2 Locations of Groundwater Contaminants

Hydro Unit	Depth Interval	Analysis	Child Resident NAL	Bkgd	MCL	MW154	MW333	MW337	MW338	PZ334	PZ335	PZ336	PZ74	SWMU2-10	SWMU2-13	SWMU2-16	SWMU2-17	SWMU2-3	SWMU2-5	SWMU2-9	
UCRS	18-20	Metals (mg/L)																			
		Beryllium	0.00264	N/A	0.004	0.0002U															
		Calcium	N/A	N/A	N/A	29					70.6	28.8	44.5								
		Magnesium	N/A	N/A	N/A	12					23.2	12.9	25.5								
		Manganese	0.035	N/A	N/A	0.019															
		Potassium	N/A	N/A	N/A	5U					5U	5U	9.34								
		Sodium	N/A	N/A	N/A	78					182	173	193								
		Uranium	0.000906	N/A	0.03	0.002					0.061	0.001U	0.023								
		Vanadium	0.00925	N/A	N/A	0.0017U															
		Radionuclides (pCi/L)																			
		Americium-241	0.371	N/A	N/A	0.19															
	Neptunium-237	0.573	N/A	N/A	0.37U																
	Plutonium-239	0.286	N/A	N/A	0.1U																
	Technetium-99	14	N/A	900	170																
	Thorium-230	0.424	N/A	N/A	0.5																
	Uranium-234	0.546	N/A	N/A	0.73						5.62		2.55								
	Uranium-235	0.538	N/A	N/A							0.48		0.17								
	Uranium-235/236	N/A	N/A	N/A	0.39U																
	Uranium-238	0.443	N/A	N/A	1.55						26.8		6.7								
	Volatiles (mg/L)																				
	cis-1,2-Dichloroethene	0.00273	N/A	0.07	0.00047U																
	Trichloroethene	0.0016	N/A	0.005	0.04																
	Vinyl chloride	0.000035	N/A	0.002	0.0013U																
Metals (mg/L)																					
Beryllium	0.00264	N/A	0.004											0.0069			0.0006			0.078	
Calcium	N/A	N/A	N/A											29			43			89	
Magnesium	N/A	N/A	N/A											24			17			48	
Manganese	0.035	N/A	N/A											6.3			3			37	
Potassium	N/A	N/A	N/A											13			3.4			19	
Sodium	N/A	N/A	N/A											110			55			87	
Uranium	0.000906	N/A	0.03											0.009U			0.0092U			0.075	
Vanadium	0.00925	N/A	N/A											0.42			0.012			4.1	
Radionuclides (pCi/L)																					
Americium-241	0.371	N/A	N/A											0.26			0.19			0.1	
Neptunium-237	0.573	N/A	N/A											0.02			-0.0125			-0.0124	
Plutonium-239	0.286	N/A	N/A											0.04			0.03			-0.0001	
Technetium-99	14	N/A	900											5.4			65.7			16.5	
Thorium-230	0.424	N/A	N/A											0.56			0.63			0.77	
Uranium-234	0.546	N/A	N/A											1.76			10.3			4.11	
Uranium-235/236	N/A	N/A	N/A											0.06			1.1			0.31	
Uranium-238	0.443	N/A	N/A											2.57			55.8			3.96	

Table 4.12. SWMU 2 Locations of Groundwater Contaminants (Continued)

Hydro Unit	Depth Interval	Analysis	Child Resident NAL	Bkgd	MCL	MW154	MW333	MW337	MW338	PZ334	PZ335	PZ336	PZ74	SWMU2-10	SWMU2-13	SWMU2-16	SWMU2-17	SWMU2-3	SWMU2-5	SWMU2-9		
UCRS	22-26	Volatiles (mg/L)																				
		I,1-Dichloroethene	0.000047	N/A	0.007													0.153	0.0143		8.33	
		cis-1,2-Dichloroethene	0.00273	N/A	0.07										0.1U			0.01U	0.28		0.00212	
		Trichloroethene	0.0016	N/A	0.005										0.1U			0.01U	0.039		0.00073	
		Vinyl chloride	0.000035	N/A	0.002										0.0013U			0.0013U	0.005			
		Trichloroethene	0.0016	N/A	0.005														0.039		0.00073	
		Vinyl chloride	0.000035	N/A	0.002														0.005			
		Metals (mg/L)																				
		Beryllium	0.00264	N/A	0.004										0.0004							0.023
		Calcium	N/A	N/A	N/A										11							130
	Iron	0.449	N/A	N/A										14							56	
	Magnesium	N/A	N/A	N/A										4.9							7.1	
	Manganese	0.035	N/A	N/A										0.071							36	
	Potassium	N/A	N/A	N/A										2.5							47	
	Sodium	N/A	N/A	N/A										23.3								
	Uranium	0.000906	N/A	0.03										0.0092U							0.0092U	
	Vanadium	0.00925	N/A	N/A										0.029							0.93	
	Radionuclides (pCi/L)																					
	Americium-241	0.371	N/A	N/A										0.34							0.21	
	Neptunium-237	0.573	N/A	N/A										0.11U							-0.0122	
Plutonium-239	0.286	N/A	N/A										0.17							0.05		
Technetium-99	14	N/A	900										170							24.2		
Thorium-230	0.424	N/A	N/A										0.67							0.61		
Uranium-234	0.546	N/A	N/A										0.34							4.14		
Uranium-235/236	N/A	N/A	N/A										0.04U							0.29		
Uranium-238	0.443	N/A	N/A										0.35							3.72		
Volatiles (mg/L)																						
I,1-Dichloroethene	0.000047	N/A	0.007										0.0013							0.667		
cis-1,2-Dichloroethene	0.00273	N/A	0.07										0.013							0.001U		
Trichloroethene	0.0016	N/A	0.005										0.013							0.01		
Vinyl chloride	0.000035	N/A	0.002										0.0013U							0.0013U		
Metals (mg/L)																						
Beryllium	0.00264	0.004	0.004																	0.0031		
Calcium	N/A	41.238	N/A																	16		
Iron	0.449	5.03	N/A																	93		
Magnesium	N/A	16.262	N/A																	7.5		
Manganese	0.035	0.119	N/A																	2.1		
Potassium	N/A	5.195	N/A																	13		
Sodium	N/A	59.45	N/A																	12		
Uranium	0.000906	0.002	0.03																	0.0092U		
RGA	58	Metals (mg/L)																				
		Beryllium	0.00264	0.004	0.004																0.0031	
		Calcium	N/A	41.238	N/A																16	
		Iron	0.449	5.03	N/A																93	
		Magnesium	N/A	16.262	N/A																7.5	
		Manganese	0.035	0.119	N/A																2.1	
		Potassium	N/A	5.195	N/A																13	
		Sodium	N/A	59.45	N/A																12	
		Uranium	0.000906	0.002	0.03																0.0092U	

Table 4.12. SWMU 2 Locations of Groundwater Contaminants (Continued)

Hydro Unit	Depth Interval	Analysis	Child Resident NAL	Bkgd	MCL	MW154	MW333	MW337	MW338	PZ334	PZ335	PZ336	PZ74	SWMU2-10	SWMU2-13	SWMU2-16	SWMU2-17	SWMU2-3	SWMU2-5	SWMU2-9	
	58	Vanadium	0.00925	0.134	N/A									1.9					0.065		
		Radionuclides (pCi/L)																			
		Americium-241	0.371	N/A	N/A									1.39					0.19		
		Neptunium-237	0.573	0.8	N/A									-0.0194					0.04		
		Plutonium-239	0.286	0.1	N/A									0.04					0.04		
		Technetium-99	14	22.3	900									36.3					0.89		
		Thorium-230	0.424	1.1	N/A									0.19					0.47		
		Uranium-234	0.546	0.7	N/A									14.2					0.75		
		Uranium-235/236	N/A	0.3	N/A									0.88					0.05		
		Uranium-238	0.443	0.7	N/A									17.6					0.55		
		Volatiles (mg/L)																			
		1,1-Dichloroethene	0.000047	N/A	0.007									1.79					2.16		
		cis-1,2-Dichloroethene	0.00273	N/A	0.07									0.75					0.001U		
		trans-1,2-Dichloroethene	0.00548	N/A	0.1									0.0043					0.001U		
		Trichloroethene	0.0016	N/A	0.005									5.35					0.001U		
		Vinyl chloride	0.000035	N/A	0.002									0.0014					0.0013U		
	61-63	Metals (mg/L)																			
		Beryllium	0.00264	0.004	0.004									0.047					0.012		
		Calcium	N/A	41.238	N/A									960					21		
		Magnesium	N/A	16.262	N/A									98					17		
		Manganese	0.035	0.119	N/A									20					3.5		
		Potassium	N/A	5.195	N/A									62					19		
		Sodium	N/A	59.45	N/A									27					8.7		
		Uranium	0.000906	0.002	0.03									0.009U					0.009U		
		Vanadium	0.00925	0.134	N/A									0.97					0.24		
		Radionuclides (pCi/L)																			
		Americium-241	0.371	N/A	N/A									0.23					0.06		
		Neptunium-237	0.573	0.8	N/A									-0.0792					0.68		
		Plutonium-239	0.286	0.1	N/A									0.02					3.98		
		Technetium-99	14	22.3	900									6.46					0.2		
		Thorium-230	0.424	1.1	N/A									0.28					3.84		
		Uranium-234	0.546	0.7	N/A									1.01					0.51		
		Uranium-235/236	N/A	0.3	N/A									0.05					0.07U		
		Uranium-238	0.443	0.7	N/A									0.62					9.07		
		Volatiles (mg/L)																			
		1,1-Dichloroethene	0.000047	N/A	0.007									0.295					3.24		
		cis-1,2-Dichloroethene	0.00273	N/A	0.07									0.0017					0.00139		
		trans-1,2-Dichloroethene	0.00548	N/A	0.1									0.077U					0.005U		
		Trichloroethene	0.0016	N/A	0.005									0.12					0.005U		
		Vinyl chloride	0.000035	N/A	0.002									0.0013U					0.0013U		

Table 4.12. SWMU 2 Locations of Groundwater Contaminants (Continued)

Hydro Unit	Depth Interval	Analysis	Child Resident NAL	Bkgd	MCL	MW154	MW333	MW337	MW338	PZ334	PZ335	PZ336	PZ74	SWMU2-10	SWMU2-13	SWMU2-16	SWMU2-17	SWMU2-3	SWMU2-5	SWMU2-9		
66-68		Metals (mg/L)																				
		Beryllium	0.00264	0.004	0.004											0.087	0.04	0.04		0.0061		
		Calcium	N/A	41.238	N/A											200	220	78		21		
		Iron	0.449	5.03	N/A											2000	690			110		
		Magnesium	N/A	16.262	N/A											110	67	52		15		
		Manganese	0.035	0.119	N/A											39	32	26		1.5		
		Potassium	N/A	5.195	N/A											84	50	69		18		
		Sodium	N/A	59.45	N/A											72	15	16		14		
		Uranium	0.000906	0.002	0.03											0.0092U	0.15	0.0092U		0.0092U		
		Vanadium	0.00925	0.134	N/A											1.7	0.6	0.83		0.11		
		Radionuclides (pCi/L)																				
		Americium-241	0.371	N/A	N/A											0.08	0.28	0.11		0.17		
		Neptunium-237	0.573	0.8	N/A											0.16	0.6	0.21		0.09		
		Plutonium-239	0.286	0.1	N/A											0.11	0.79	0.05		-0.0153		
		Technetium-99	14	22.3	900											14.6	-3.43	77.1		0.65		
		Thorium-230	0.424	1.1	N/A											0.84	0.21	0.49		0.46		
		Uranium-234	0.546	0.7	N/A											0.91	0.61	50.6		1.22		
		Uranium-235/236	N/A	0.3	N/A											0.03	0.16	6.47		0.06		
		Uranium-238	0.443	0.7	N/A											0.76	2.12	55.1		0.94		
		Volatiles (mg/L)																				
1,1-Dichloroethene	0.00047	N/A	0.007											0.356	4.21	0.0165		4.12				
cis-1,2-Dichloroethene	0.00273	N/A	0.07											0.221	0.01U	0.001U		0.077U				
trans-1,2-Dichloroethene	0.00548	N/A	0.1											0.001	0.01U	0.001U		0.077U				
Trichloroethene	0.0016	N/A	0.005											1.6	0.0043	0.00049		0.077U				
Vinyl chloride	0.00035	N/A	0.002											0.0013U	0.0013U	0.0013U		0.0013U				
Metals (mg/L)																						
Beryllium	0.00264	0.004	0.004											0.092				0.038	0.0059	0.068		
Calcium	N/A	41.238	N/A											390				43	30	190		
Magnesium	N/A	16.262	N/A											50				36	13	72		
Manganese	0.035	0.119	N/A											51				96	14	71		
Potassium	N/A	5.195	N/A											42				47	11	67		
Sodium	N/A	59.45	N/A											34				14	14	17		
Uranium	0.000906	0.002	0.03											0.009U				0.009U	0.049	0.41		
Vanadium	0.00925	0.134	N/A											1.5				0.6	0.12	1.2		
Radionuclides (pCi/L)																						
Americium-241	0.371	N/A	N/A											0.26				0.4	0.13	0.19		
Neptunium-237	0.573	0.8	N/A											-0.0375				-0.0192	-0.0377	0.02		
Plutonium-239	0.286	0.1	N/A											0.23				0.08	0	0.49		
Technetium-99	14	22.3	900											6.85				-0.395	1.82	36.3		
Thorium-230	0.424	1.1	N/A											0.71				0.45	0.45	0.87		
Uranium-234	0.546	0.7	N/A											0.43				2.56	3	6.68		
71-72		Metals (mg/L)																				
		Beryllium	0.00264	0.004	0.004																	
		Calcium	N/A	41.238	N/A																	
		Magnesium	N/A	16.262	N/A																	
		Manganese	0.035	0.119	N/A																	
		Potassium	N/A	5.195	N/A																	
		Sodium	N/A	59.45	N/A																	
		Uranium	0.000906	0.002	0.03																	
		Vanadium	0.00925	0.134	N/A																	
		Radionuclides (pCi/L)																				

Table 4.12. SWMU 2 Locations of Groundwater Contaminants (Continued)

Hydro Unit	Depth Interval	Analysis	Child Resident NAL	Bkgd	MCL	MW154	MW333	MW337	MW338	PZ334	PZ335	PZ336	PZ74	SWMU-10	SWMU-13	SWMU-16	SWMU-17	SWMU-2-3	SWMU-2-5	SWMU-2-9
	71-72	Uranium-235/236	N/A	0.3	N/A									-0.0148				0.22	0.11	0.27
		Uranium-238	0.443	0.7	N/A									0.18				7.55	2.92	6.53
		Volatiles (mg/L)																		
		1,1-Dichloroethene	0.000047	N/A	0.007									0.375					47.9	2.14
		cis-1,2-Dichloroethene	0.00273	N/A	0.07									0.0014				0.02U	0.099U	0.1U
		trans-1,2-Dichloroethene	0.00548	N/A	0.1									0.001U				0.02U	0.099U	0.1U
		Trichloroethene	0.0016	N/A	0.005									0.1				0.02U	0.099U	0.015
		Vinyl chloride	0.000035	N/A	0.002									0.0013U				0.0013U	0.0013U	0.0013U
	74-77	Metals (mg/L)																		
		Arsenic	0.000035	0.005	0.01			0.0175	0.001U							0.081				
		Barium	0.104	0.235	2			0.078	0.097							2.4				
		Beryllium	0.00264	0.004	0.004			0.0014	0.0003							0.068			0.03	
		Cadmium	0.000661	0.01	0.005			0.001U	0.001U							0.012				
		Calcium	N/A	41.238	N/A			16	15							96			52	
		Chromium	1.76	0.144	0.1			0.02U	0.269							0.35				
		Cobalt	0.0906	0.045	N/A			0.001U	0.0019											
		Iron	0.449	5.03	N/A			56	5.7											
		Magnesium	N/A	16.262	N/A			7.3	6.07										36	
		Manganese	0.035	0.119	N/A			2.1	1.1							65		65	19	41
		Nickel	0.0301	0.682	N/A			0.005U	0.0908							0.33				
		Potassium	N/A	5.195	N/A			3.9	0.471										48	
		Sodium	N/A	59.45	N/A			12.2	14										15	
		Uranium	0.000906	0.002	0.03			0.05U	0.35							0.009U			0.0092U	
		Vanadium	0.00925	0.134	N/A			0.052	0.0078							1.3			0.28	0.61
		Radionuclides (pCi/L)																		
		Americium-241	0.371	N/A	N/A			0.19	0.1							0.18			0.32	0.08
		Neptunium-237	0.573	0.8	N/A			1.33U	1.35U							0.06			0.46	0.04
		Plutonium-239	0.286	0.1	N/A			0.14U	0.13							0.15			0.22	0.03
		Technetium-99	14	22.3	900			229	19.04							5.32			-1.74	17.1
		Thorium-230	0.424	1.1	N/A			0.52	0.44							1.05			0.38	0.61
		Uranium-234	0.546	0.7	N/A			0.38	0.56							3.35			13.9	12.3
		Uranium-235/236	N/A	0.3	N/A			0.11	0.13U							0.28			2.43	1.41
		Uranium-238	0.443	0.7	N/A			0.27	0.67							3.94			91.7	14.9
		Volatiles (mg/L)																		
		1,1-Dichloroethane	0.0363	N/A	N/A			0.0022	0.01U											
		1,1-Dichloroethene	0.000047	N/A	0.007			0.05U	0.005U							3.55			1.09	0.159
		Chloroform	0.0000287	N/A	N/A			0.0029	0.0015							0.0605			0.01U	0.00041
		cis-1,2-Dichloroethene	0.00273	N/A	0.07			0.062	0.0073							0.0714U			0.01U	0.001U
		trans-1,2-Dichloroethene	0.00548	N/A	0.1			0.05U	0.01U							0.422			0.0072	0.00026
		Trichloroethene	0.0016	N/A	0.005			0.78	0.14							0.0013U			0.0013U	0.0013U
		Vinyl chloride	0.000035	N/A	0.002			0.02U	0.002U											

RGA

Table 4.12. SWMU 2 Locations of Groundwater Contaminants (Continued)

Hydro Unit	Depth Interval	Analysis	Child Resident NAL	Bkgd	MCL	MW154	MW333	MW337	MW338	PZ334	PZ335	PZ336	PZ74	SWMU2-10	SWMU2-13	SWMU2-16	SWMU2-17	SWMU2-3	SWMU2-5	SWMU2-9		
RGA	74-77	Wetchem (mg/L)	0.0284	N/A	0.2												0.0075					
	79-82	Metals (mg/L)																				
		Arsenic	0.000035	0.005	0.01		0.0029															
		Barium	0.104	0.235	2		0.13															
		Beryllium	0.00264	0.004	0.004		0.001U								0.015				0.021		0.016	
		Cadmium	0.000661	0.01	0.005		0.001U															
		Calcium	N/A	41.238	N/A		24								87				35		50	
		Chromium	1.76	0.144	0.1		0.02U															
		Cobalt	0.0906	0.045	N/A		0.001U															
		Iron	0.449	5.03	N/A		6.2															
		Magnesium	N/A	16.262	N/A		9.2								27				19		21	
		Manganese	0.035	0.119	N/A		2.6								15				38		38	
		Nickel	0.0301	0.682	N/A		0.0182															
		Potassium	N/A	5.195	N/A		1.2								22				15		21	
		Sodium	N/A	59.45	N/A		16								17				14		14	
		Uranium	0.000906	0.002	0.03		0.05U								0.009U				0.009U		0.0092U	
		Vanadium	0.00925	0.134	N/A		0.0097								0.37				0.3		0.3	
		Radionuclides (pCi/L)																				
			Americium-241	0.371	N/A	N/A		0.19							0.22				0.52		0.24	
			Neptunium-237	0.573	0.8	N/A		1.33U							-0.13				0.12		0.04	
			Plutonium-239	0.286	0.1	N/A		0.1U							4.29				0.07		0	
			Technetium-99	14	22.3	900		19.27							2.33				2.21			
			Thorium-230	0.424	1.1	N/A		0.25							0.4				0.42		0.31	
			Uranium-234	0.546	0.7	N/A		9.66							1.14				3.55		0.56	
			Uranium-235/236	N/A	0.3	N/A		0.35							0.08				0.21		0.03	
			Uranium-238	0.443	0.7	N/A		0.65U							1.08				9.91		0.78	
		Volatiles (mg/L)																				
		1,1-Dichloroethane	0.0363	N/A	N/A		0.12U															
		1,1-Dichloroethene	0.00047	N/A	0.007		0.12U							1.45				4.1		22.8		
		Chloroform	0.000287	N/A	N/A		0.12U															
		cis-1,2-Dichloroethene	0.00273	N/A	0.07		0.2							0.00029				0.01U		0.02U		
		trans-1,2-Dichloroethene	0.00548	N/A	0.1		0.12U							0.001U				0.01U		0.02U		
		Trichloroethene	0.0016	N/A	0.005		1.6							0.052				0.0022		0.0089		
		Vinyl chloride	0.000035	N/A	0.002		0.05U							0.0013U				0.0013U		0.0013U		
	Metals (mg/L)																					
	87	Beryllium	0.00264	0.004	0.004										0.035			0.029				
		Calcium	N/A	41.238	N/A										60			95				
		Iron	0.449	5.03	N/A									1500								
		Magnesium	N/A	16.262	N/A										35			41				
		Manganese	0.035	0.119	N/A										43			23				

Table 4.12. SWMU 2 Locations of Groundwater Contaminants (Continued)

Hydro Unit	Depth Interval	Analysis	Child Resident NAL	Bkgd	MCL	MW154	MW333	MW337	MW338	PZ334	PZ335	PZ336	PZ74	SWMU2-10	SWMU2-13	SWMU2-16	SWMU2-17	SWMU2-3	SWMU2-5	SWMU2-9		
RGA	87	Potassium	N/A	5.195	N/A										26	31						
		Sodium	N/A	59.45	N/A											24	14					
		Uranium	0.000906	0.002	0.03											0.009U	0.0092U					
		Vanadium	0.00925	0.134	N/A											0.65	0.48					
		Radionuclides (pCi/L)																				
		Americium-241	0.371	N/A	N/A											0.43	1.13					
		Neptunium-237	0.573	0.8	N/A											-0.0565	0.24					
		Plutonium-239	0.286	0.1	N/A											0.05	0.15					
		Technetium-99	14	22.3	900											-3.55	5.03					
		Thorium-230	0.424	1.1	N/A											0.33	0.41					
		Uranium-234	0.546	0.7	N/A											2.28	9.97					
		Uranium-235/236	N/A	0.3	N/A											0.22	1.32					
		Uranium-238	0.443	0.7	N/A											2.56	57.8					
		Volatiles (mg/L)																				
		1,1-Dichloroethene	0.00047	N/A	0.007											0.0651	0.008					
		cis-1,2-Dichloroethene	0.00273	N/A	0.07											0.034	0.001U					
		trans-1,2-Dichloroethene	0.00548	N/A	0.1											0.0668U	0.001U					
		Trichloroethene	0.0016	N/A	0.005											0.37	0.011					
		Vinyl chloride	0.00035	N/A	0.002												0.0013U					
		Metals (mg/L)																				
Beryllium	0.00264	0.017	0.004														0.044					
Calcium	N/A	38.858	N/A														62					
Magnesium	N/A	13.418	N/A														35					
Manganese	0.035	0.941	N/A														80					
Potassium	N/A	55.752	N/A														32					
Sodium	N/A	29.2	N/A														15					
Vanadium	0.00925	0.126	N/A														0.73					
Radionuclides (pCi/L)																						
Americium-241	0.371	N/A	N/A														0.15					
Neptunium-237	0.573	0.5	N/A														0.15					
Plutonium-239	0.286	0.2	N/A														0.05					
Technetium-99	14	20.6	900														20					
Thorium-230	0.424	1.5	N/A														0.51					
Uranium-234	0.546	0.3	N/A														16.3					
Uranium-235/236	N/A	0.2	N/A														1.7					
Uranium-238	0.443	0.3	N/A														19.6					
Volatiles (mg/L)																						
1,1-Dichloroethene	0.00047	N/A	0.007														3.3					
cis-1,2-Dichloroethene	0.00273	N/A	0.07														0.0232					
Trichloroethene	0.0016	N/A	0.005														0.0017					
Wetchem Parameters (mg/L)																						
Cyanide	0.0284	N/A	0.2														0.013					
McNairy	87																					

Table 4.12. SWMU 2 Locations of Groundwater Contaminants (Continued)

Hydro Unit	Depth Interval	Analysis	Child Resident NAL	Bkgd	MCL	MW154	MW333	MW337	MW338	PZ334	PZ335	PZ336	PZ74	SWMU2-10	SWMU2-13	SWMU2-16	SWMU2-17	SWMU2-3	SWMU2-5	SWMU2-9		
McNairy	92-93	Metals (mg/L)																				
		Beryllium	0.00264	0.017	0.004										0.0026				0.0005		0.067	
		Calcium	N/A	38,858	N/A											24				14		120
		Magnesium	N/A	13,418	N/A											8.8				6.2		86
		Manganese	0.035	0.941	N/A											3				1.7		70
		Potassium	N/A	55,752	N/A											3.6				2.6		83
		Sodium	N/A	29.2	N/A											15				1.3		17
		Vanadium	0.00925	0.126	N/A											0.074				0.029		1.5
		Radionuclides (pCi/L)																				
		Americium-241	0.371	N/A	N/A											0.29				0.29		0.32
		Neptunium-237	0.573	0.5	N/A											0.11				0.08		0.05
		Plutonium-239	0.286	0.2	N/A											0.06				0.08		-0.0132
		Technetium-99	14	20.6	900											7.72				1.26		
		Thorium-230	0.424	1.5	N/A											0.54				0.36		0.94
		Uranium-234	0.546	0.3	N/A											0.57				0.23		5.09
		Uranium-235/236	N/A	0.2	N/A											0.05				0.03		0.29
		Uranium-238	0.443	0.3	N/A											0.48				0.28		5.1
		Volatiles (mg/L)																				
		1,1-Dichloroethene	0.000047	N/A	0.007											0.0715				7.5		
		cis-1,2-Dichloroethene	0.00273	N/A	0.07											0.00038				0.005U		0.1U
		Trichloroethene	0.0016	N/A	0.005											0.0464				0.00126		0.055

Blank cells indicate interval was not sampled for the specified analysis. Maximum value shown for each depth interval.

N/A = not applicable

Bold indicates result is greater than NAL value.

Italics indicates result is greater than MCL value.

Bold + Italics indicate result is greater than both NAL and MCL values.

Shading indicates result is greater than the background value.

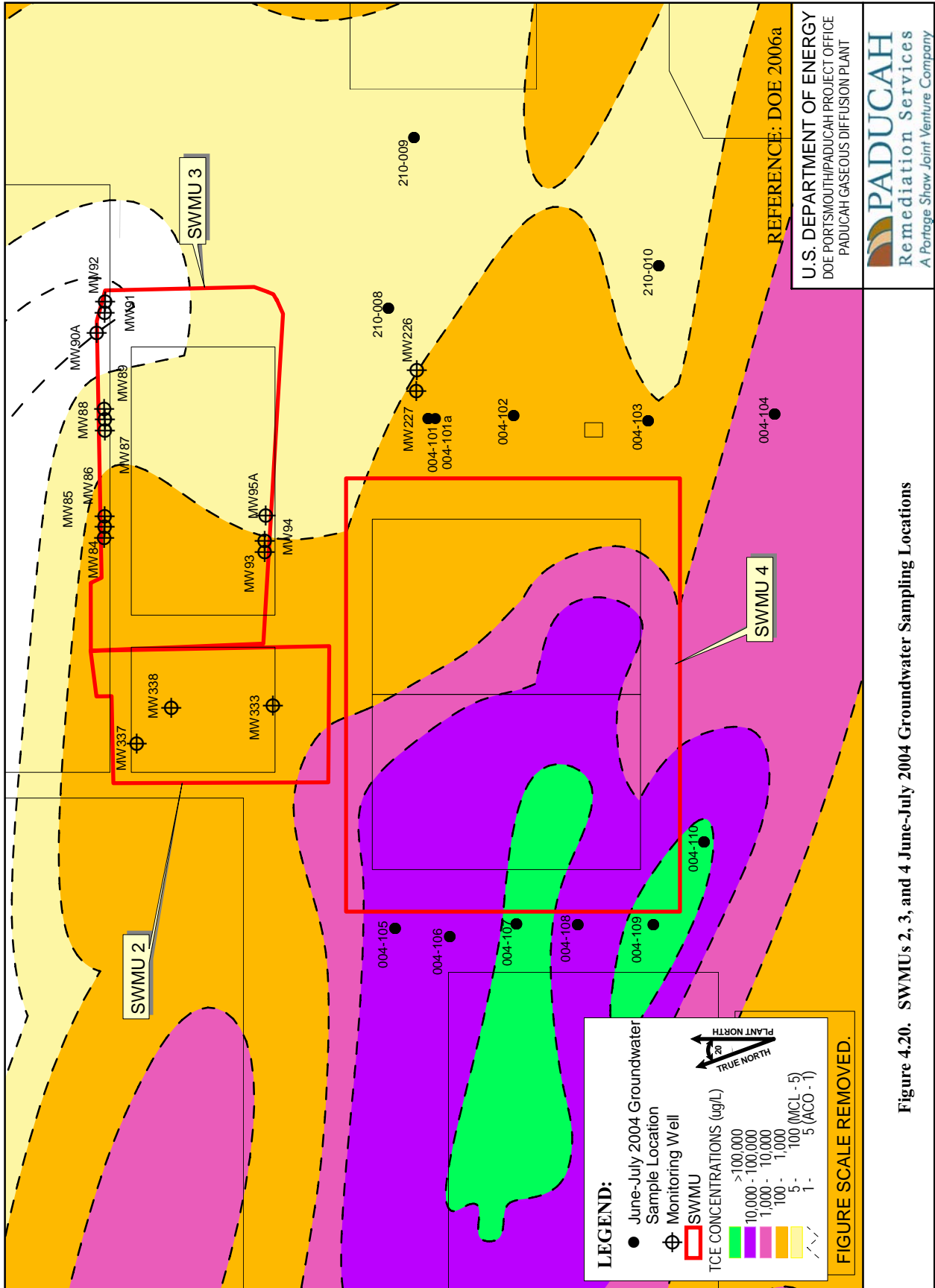


Figure No. \BGO\bgwmap.apr
DATE 08-08-06

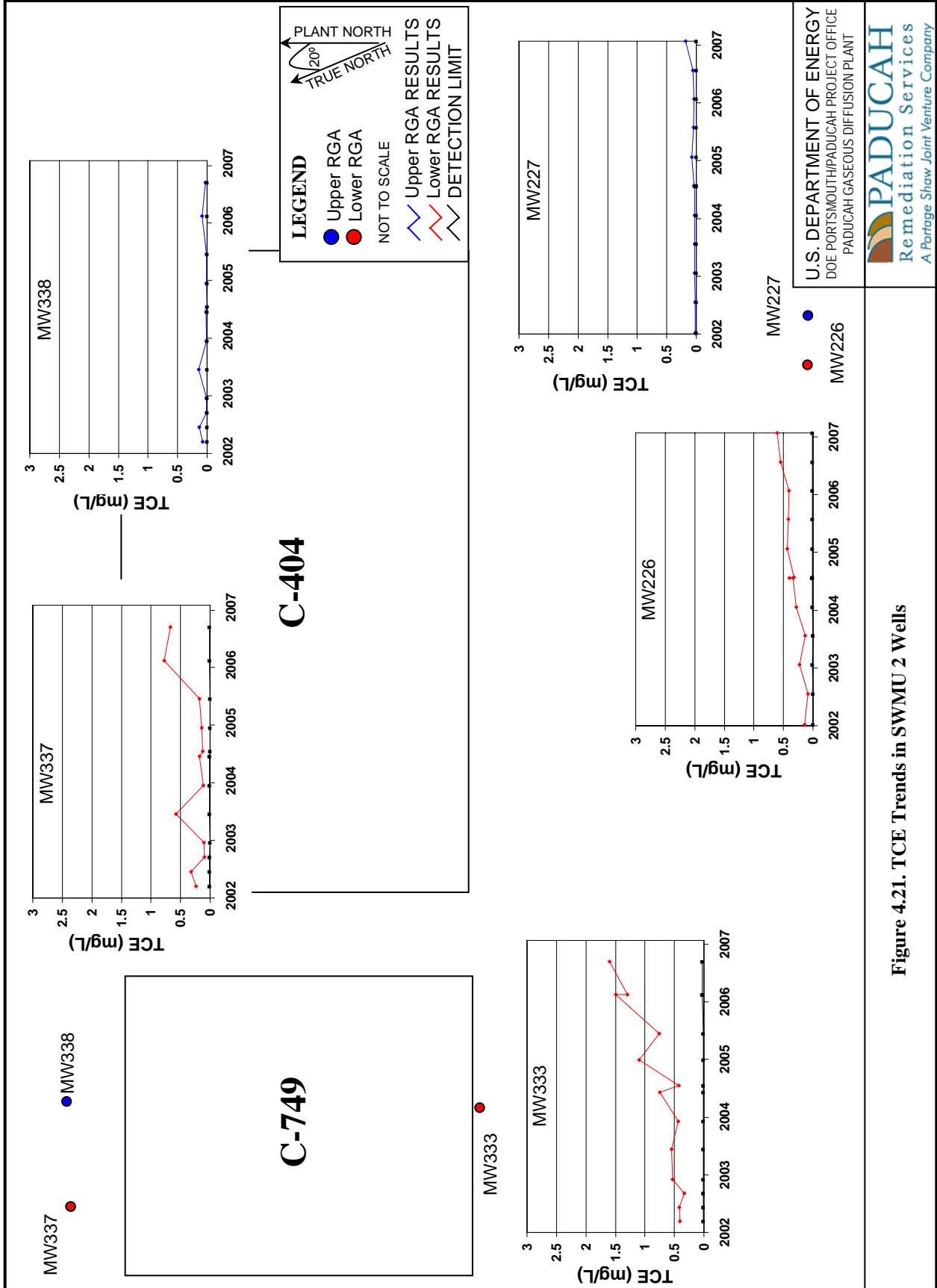
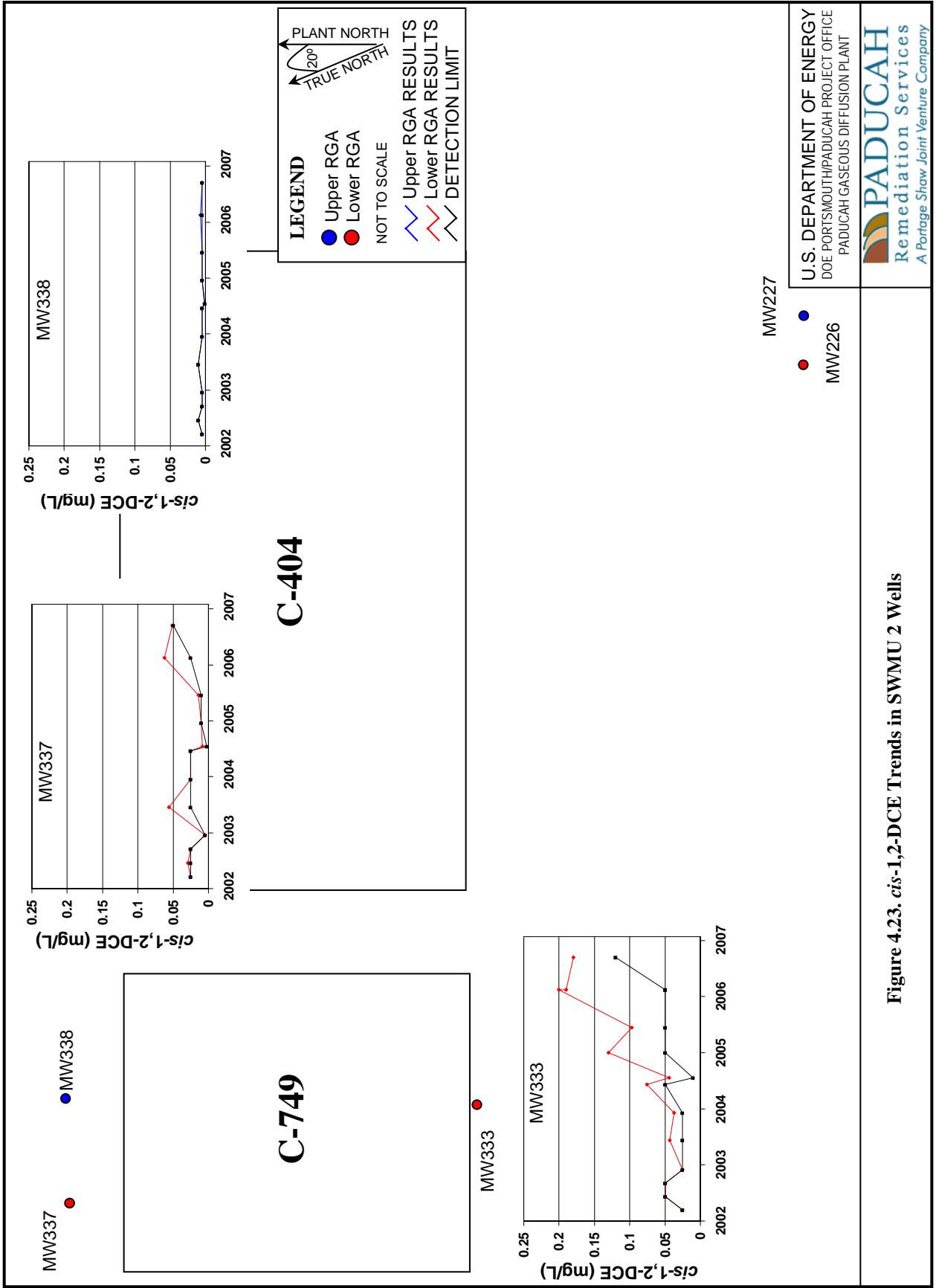


Figure 4.21. TCE Trends in SWMU 2 Wells



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DOE PORTSMOUTH/PADUCAH PROJECT OFFICE
PADUCAH GASEOUS DIFFUSION PLANT

PADUCAH
Remediation Services
A Portage Shaw Joint Venture Company

4.4 SWMU 3

4.4.1 Subsurface Soils

No historical subsurface soil data were available for SWMU 3; however, subsurface soil samples were collected from four angled borings at C-404 as part of this RI (Figure 4.2). Six shallow borings also were drilled and sampled along the former discharge ditch associated with SWMU 3 as part of this RI. A review of RI data identified the contaminants listed in Table 4.13.

Table 4.14 shows the locations of the samples with metals and radionuclides detected above screening levels.

Wastes disposed of in SWMU 3 include all liquid effluents from C-400 operations from 1952 through 1957. C-404 continued to receive solid uranium-contaminated and radioactively contaminated wastes from 1957 until 1986.

The most prevalent metal detected above its background values in subsurface soil at SWMU 3 is uranium, followed by antimony. Uranium contamination has migrated to a depth of 10 to 15 ft under C-404 and as much as 10 ft under the former discharge ditch (as a metal). Figure 4.24 provides the distribution of uranium based on the RI data. The higher concentrations are found in shallow soils on the western side of the unit. The 10 ft sample from boring 003-001 had a uranium concentration of 83.6 mg/kg, but this decreased to approximately 1 mg/kg or less in deeper samples indicating a limited extent to uranium contamination. Uranium was not detected above screening levels in the 15 ft samples along the former discharge ditch (Appendix D, page D-25). Cesium-137 was detected above screening in one sample at a depth of 5 ft in boring 003-005 along the former discharge ditch (Figure 4.2).

Antimony contamination is limited to a depth of 5 to 10 ft along the former discharge ditch. (SWMUs 3 and 145 are the only BGOU SWMUs to have antimony concentrations that exceed the contaminant screening criteria.) Arsenic was detected frequently in subsurface soil samples, but exceeded the PGDP background level in only 1 of 40 samples.

Table 4.13. SWMU 3 Subsurface Soil Contaminants

Analysis	Maximum Result		Frequency of Detection ^a	Frequency of Detection	
	Historical Data	RI Data		Above Background Value	Above Excavation Worker NAL
<i>Inorganics (mg/kg)</i>					
Aluminum	N/A	11,800	40/40	0/40	35/40
Antimony	N/A	11	3/40	3/40	3/40
Arsenic	N/A	8.25	36/40	1/40	36/40
Barium	N/A	122	40/40	0/40	0/40
Beryllium	N/A	1.06	4/40	1/40	0/40
Calcium	N/A	35,600	40/40	1/40	N/A
Chromium	N/A	25.8	40/40	0/40	0/40
Cobalt	N/A	14.9	28/40	1/40	0/40
Copper	N/A	15.9	40/40	0/40	0/40
Iron	N/A	25,700	40/40	0/40	40/40
Lead	N/A	23.9	40/40	1/40	0/40
Magnesium	N/A	1,770	40/40	0/40	0/40
Manganese	N/A	644	40/40	0/40	32/40
Mercury	N/A	0.024	5/40	0/40	0/40
Molybdenum	N/A	3.78	1/40	N/A	0/40
Nickel	N/A	16.1	33/40	0/40	0/40
Sodium	N/A	249	18/40	0/40	0/40
Uranium	N/A	83.6	11/40	7/40	4/40
Vanadium	N/A	32.1	40/40	0/40	40/40
Zinc	N/A	50	22/40	0/40	0/40
<i>Organics--Volatiles (mg/kg)</i>					
Acetone	N/A	0.04	2/40	N/A	0/40
Trichloroethene	N/A	0.02	1/40	N/A	0/40
<i>Radionuclides(pCi/g)</i>					
Cesium-137	N/A	0.456	1/40	1/40	1/40
Plutonium-239/240	N/A	0.0399	1/40	N/A	0/40
Technetium-99	N/A	56.9	4/40	3/40	0/40
Thorium-228	N/A	0.566	40/40	0/40	40/40
Thorium-230	N/A	0.573	27/40	0/40	0/40
Thorium-232	N/A	0.603	40/40	0/40	0/40
Thorium-234	N/A	14.1	8/40	N/A	N/A
Uranium	N/A	25.8	9/40	N/A	N/A
Uranium-234	N/A	3.02	14/40	1/40	1/40
Uranium-235	N/A	0.362	5/40	1/40	0/40
Uranium-238	N/A	22.4	18/40	6/40	6/40

^a Frequency of detection is the number of detections of an analyte per number of analyses (includes regular and duplicate samples).
N/A = not applicable ND = not detected

Table 4.14. SWMU 3 Locations of Subsurface Soil Contaminants

Analysis	Depth (ft)	Excavation Worker NAL	Background	RI Data										
				003-001	003-002	003-003	003-004	003-005	003-006	003-007	003-008	003-009	003-010	
Inorganics (mg/kg)	Aluminum	5	12,000						8,370	6,540	7,190	8,700	9,340	4,920
		10	12,000	11,800	9,990	6,450	11,300	7,710	7,910	5,100	6,670	7,990	6,300	6,300
		15	12,000	6,520	7,270	6,480	8,470	7,140	6,200	5,760	6,760	6,520	4,620	
		30	12,000	5,800	5,420	5,490	5,670							
		45	12,000	7,590	4,370	2,820	6,690							
		60	12,000	7,230	5,640	8,060	5,770							
Antimony	5	0.492	0.21					9.64U	8.08U	9.89	11	9.96U	9.22U	
	10	0.492	0.21	8.36U	9.18U	7.73U	9.8U	9.89U	9.58U	7.31U	9.57U	9.85U	10.1	
	15	0.492	0.21	9.39U	6.88U	9.95U	9.72U	9.47U	9.15U	8.92U	8.99U	8.94U	9.34U	
	30	0.492	0.21	9.46U	9.65U	9.74U	9.89U							
	45	0.492	0.21	7.83U	9.62U	9.84U	10U							
	60	0.492	0.21	7.52U	9.25U	9.84U	9.52U							
Arsenic	5	0.324	7.9					3.76	3.9	7.03	1.25	2.93	2.68	
	10	0.324	7.9	5.69	3.09	2.47	3.81	4.67	2.96	2.57	1.4	2.02	2.61	
	15	0.324	7.9	0.956	2.12	5.19	1.46	0.956U	1.68	2.3	1.18	1.16	1.27	
	30	0.324	7.9	0.951U	3.32	0.831U	3.02							
	45	0.324	7.9	8.25	2.39	2.61	3.01							
	60	0.324	7.9	1.65	0.89U	1.28	2.36							
Barium	5	272	170					79.1	33.6	41.9	111	116	35.3	
	10	272	170	99.2	71.3	78.8	50.9	87.1	53.8	81.3	83.1	92.6	122	
	15	272	170	61.7	62.5	53.7	80.7	86.4	60.3	44.7	54.4	53.2	57.5	
	30	272	170	13.4	14.8	12.5	21.3							
	45	272	170	30.9	19.4	19.9	17.8							
	60	272	170	78.1	38.5	66.1	58.9							
Beryllium	5	1.26	0.69					0.488U	0.479U	0.465U	0.462U	0.448U	0.468U	
	10	1.26	0.69	0.468U	0.46U	0.442U	0.494U	0.458U	0.494U	0.468U	0.44U	0.484U	0.462U	
	15	1.26	0.69	0.428U	0.488U	0.512	0.486U	0.478U	0.454U	0.494U	0.497U	0.476U	0.453U	
	30	1.26	0.69	0.476U	0.475U	0.416U	0.484U							
	45	1.26	0.69	0.44U	0.456U	0.414U	0.489U							
	60	1.26	0.69	0.495U	0.445U	0.619	1.06							
Calcium	5	n/a	6,100					5,080	35,600	557	673	2,160	643	
	10	n/a	6,100	1,550	2,320	1,190	1,080	952	1,410	976	1,110	651	1,190	
	15	n/a	6,100	947	1,160	1,000	1,120	1,500	850	838	927	1,050	873	
	30	n/a	6,100	625	665	546	451							
	45	n/a	6,100	888	653	584	706							
	60	n/a	6,100	697	941	1,350	1,090							
Chromium	5	476	43					10	17.9	11.5	9.6	11.5	5.7	
	10	476	43	14.1	12.7	10.3	13	10.7	9.11	8.58	10.2	8.77	9.9	
	15	476	43	7.37	10	25.8	12.6	10.5	10.2	12.8	10	9.48	6.93	
	30	476	43	6.39	4.41	6.33	6.09							

Table 4.14. SWMU 3 Locations of Subsurface Soil Contaminants (Continued)

Analysis	Depth (ft)	Excavation Worker NAL	Background	RI Data																
				003-001	003-002	003-003	003-004	003-005	003-006	003-007	003-008	003-009	003-010							
Chromium	45	476	43	8.96	4.95	6.04	12.3													
	60	476	43	6.14	9.85	18.1	8.95													
Cobalt	5	1,110	13					3.57	9.2	4.58										
	10	1,110	13	5.22	3.6	5.78	4.25	4.09	4.51	3.11	2.76	2.48U	2.38U	4.87	4.95	4.74				
Copper	15	1,110	13	10.1	5.18	7.88	3.49	2.39U	2.29	2.72										
	30	1,110	13	2.38U	2.37U	2.08U	2.67													
Iron	45	1,110	13	2.76	2.28U	10.7	2.45U													
	60	1,110	13	12.8	2.23U	2.43U	74.9													
Lead	5	427	25					6.19	4.33	13.6	6.56	9.55	4.17							
	10	427	25	12.8	11.3	6.19	15.9	7.25	11.3	7.08	6.5	8.06	7.79							
Magnesium	15	427	25	4.73	7.28	8.69	6.31	4.93	4.88	5.18	4.71	4.98	4.68							
	30	427	25	3.8	3.42	2.49	3.01													
Mercury	45	427	25	8.05	3.69	2.34	4.8													
	60	427	25	6.85	2.9	8.8	8.39													
Manganese	5	2,170	28,000					9,730	14,900	16,000	6,820	11,500	6,690							
	10	2,170	28,000	13,900	15,100	8,600	25,700	13,500	10,800	9,360	8,630	9,130	9,930							
Zinc	15	2,170	28,000	6,620	7,770	16,800	8,410	7,180	8,300	10,300	7,250	6,590	6,300							
	30	2,170	28,000	9,940	14,800	5,620	8,860													
Cadmium	45	2,170	28,000	23,000	9,840	5,310	18,000													
	60	2,170	28,000	12,500	5,000	15,600	15,600													
Chloride	5	50	23					8.9	9.37	11	7.67	7.64	6.67							
	10	50	23	23.9	8.85	6.5	7.95	7.13	5.89	7.34	5.62	5.35	6.48							
Sulfate	15	50	23	6.14	7.07	14.2	6.21	3.87	5.17	7.68	5.51	4.95	5.73							
	30	50	23	5.58	5.39	4.75	7.74													
Nitrate	45	50	23	10.4	4.66	6.11	5.42													
	60	50	23	5.1	2.94	7.7	6.07													
Ammonia	5	n/a	2,100					720	1,510	1,770	704	1,330	534							
	10	n/a	2,100	1,600	1,500	1,260	1,240	1,110	1,140	1,200	1,120	1,110	1,290							
Nitrogen	15	n/a	2,100	840	1,060	716	960	1,350	807	902	806	995	778							
	30	n/a	2,100	312	374	319	271													
Fluoride	45	n/a	2,100	640	322	266	354													
	60	n/a	2,100	926	638	1,520	1,190													
Selenium	5	56.6	820					400	182	282	201	323	233							
	10	56.6	820	507	339	342	279	242	203	227	154	298	438							
Vanadium	15	56.6	820	644	288	638	80.5	63.5	87.3	109	104	107	96.7							
	30	56.6	820	46.8	41.6	26	156													
Molybdenum	45	56.6	820	45.9	35.2	59	95.6													
	60	56.6	820	45.1	9.12	34.8	175													
Cyanide	5	1.17	0.13					0.019U	0.018U	0.02U	0.02U	0.02U	0.017U							
	10	1.17	0.13	0.019U	0.02	0.02U	0.017U	0.02U	0.015U	0.019U	0.019U	0.018U	0.017U							
Arsenic	15	1.17	0.13	0.019U	0.018U	0.017U	0.018	0.02U	0.016U	0.018U	0.019U	0.018U	0.018U							
	30	1.17	0.13	0.016U	0.015U	0.019U	0.019U	0.015U	0.016U	0.018U	0.019U	0.018U	0.018U							

Table 4.14. SWMU 3 Locations of Subsurface Soil Contaminants (Continued)

Analysis	Depth (ft)	Excavation Worker NAL	Background	RI Data																	
				003-001	003-002	003-003	003-004	003-005	003-006	003-007	003-008	003-009	003-010								
Acetone	30	421	n/a	0.00502U	0.00497U	0.00503U	0.00504U														
	45	421	n/a	0.00498U	0.00496U	0.00502U	0.005U														
	60	421	n/a	0.00504U	0.00502U	0.00501U	0.00501U														
Trichloroethene	5	3.25	n/a					0.00499U	0.00503U	0.00498U	0.00501U	0.00504U	0.005U								
	10	3.25	n/a	0.00504U	0.00502U	0.00503U	0.00503U	0.00502U	0.005U	0.00497U	0.00499U	0.00499U	0.00498U	0.00498U	0.00499U	0.00499U	0.00498U	0.00498U	0.00498U	0.00498U	0.00498U
	15	3.25	n/a	0.00497U	0.00502U	0.00498U	0.00498U	0.00495U	0.00495U	0.00504U	0.00496U	0.00501U	0.00504U	0.00502U	0.00504U	0.00504U	0.00504U	0.00504U	0.00502U	0.00502U	0.00502U
	30	3.25	n/a	0.00502U	0.00497U	0.00503U	0.00504U														
	45	3.25	n/a	0.00498U	0.00496U	0.0152	0.005U														
	60	3.25	n/a	0.00504U	0.00502U	0.00501U	0.00501U														
Radionuclides (pCi/g)																					
Cesium-137	5	0.115	0.28																		
	10	0.115	0.28	0.0628U	0.0629U	0.0505U	0.0496U	0.0518U	0.0507U	0.0801U	0.0512U	0.0795U	0.0795U	0.0866U	0.0866U	0.0866U	0.0866U	0.0866U	0.0866U	0.0866U	0.0866U
	15	0.115	0.28	0.0468U	0.0498U	0.0481U	0.045U	0.0469U	0.0417U	0.0502U	0.0421U	0.0445U	0.0445U	0.0479U	0.0479U	0.0479U	0.0479U	0.0479U	0.0479U	0.0479U	0.0479U
	30	0.115	0.28	0.0473U	0.0596U	0.0437U	0.042U														
	45	0.115	0.28	0.0533U	0.0475U	0.0371U	0.0503U														
	60	0.115	0.28	0.046U	0.0472U	0.0591U	0.0543U														
Plutonium-239/240	5	1.63	n/a																		
	10	1.63	n/a	0.0173U	0.0181U	0.0177U	0.018U	0.0197U	0.0223U	0.0194U	0.0211U	0.0203U	0.0204U	0.0204U	0.0204U	0.0204U	0.0204U	0.0204U	0.0204U	0.0204U	0.0204U
	15	1.63	n/a	0.0187U	0.019U	0.0169U	0.0184U	0.0196U	0.0197U	0.0202U	0.0198U	0.0196U	0.0196U	0.0208U	0.0216U	0.0216U	0.0216U	0.0216U	0.0216U	0.0216U	0.0216U
	30	1.63	n/a	0.0178U	0.0193U	0.0179U	0.0179U														
	45	1.63	n/a	0.0166U	0.0182U	0.0181U	0.0183U														
	60	1.63	n/a	0.0188U	0.0176U	0.018U	0.0174U														
Technetium-99	5	57.9	2.8					7.35	56.9	3.04	1.85U	1.85U	1.85U	1.85U	1.85U	1.85U	1.85U	1.85U	1.85U	1.85U	
	10	57.9	2.8	1.8U	1.8U	1.85U	1.85U	1.81U	1.81U	1.81U	1.81U	1.85U	1.85U	1.85U	1.85U	1.85U	1.85U	1.85U	1.85U	1.85U	
	15	57.9	2.8	1.8U	1.8U	2.4	1.85U	1.81U	1.81U	1.81U	1.81U	1.85U	1.85U	1.85U	1.85U	1.85U	1.85U	1.85U	1.85U	1.85U	
	30	57.9	2.8	1.8U	1.8U	1.85U	1.85U														
	45	57.9	2.8	1.8U	1.8U	1.85U	1.85U														
	60	57.9	2.8	1.8U	1.8U	1.85U	1.85U														
Thorium-228	5	0.0357	1.6																		
	10	0.0357	1.6	0.365	0.454	0.258	0.331	0.443	0.284	0.275	0.341	0.405	0.307	0.307	0.307	0.307	0.307	0.307	0.307	0.307	
	15	0.0357	1.6	0.202	0.388	0.282	0.356	0.302	0.323	0.372	0.322	0.395	0.335	0.335	0.335	0.335	0.335	0.335	0.335	0.335	
	30	0.0357	1.6	0.341	0.201	0.213	0.18														
	45	0.0357	1.6	0.566	0.211	0.162	0.234														
	60	0.0357	1.6	0.344	0.39	0.496	0.257														
Thorium-230	5	2.22	1.4																		
	10	2.22	1.4	0.245	0.573	0.237U	0.371	0.505	0.238	0.191	0.278	0.338	0.194	0.194	0.194	0.194	0.194	0.194	0.194	0.194	
	15	2.22	1.4	0.241	0.323	0.238U	0.295	0.135U	0.224	0.152	0.134	0.344	0.205	0.205	0.205	0.205	0.205	0.205	0.205	0.205	
	30	2.22	1.4	0.237U	0.237U	0.237U	0.238U														
	45	2.22	1.4	0.382	0.241U	0.238U	0.238U														
	60	2.22	1.4	0.237U	0.237U	0.407	0.238U														

Table 4.14. SWMU 3 Locations of Subsurface Soil Contaminants (Continued)

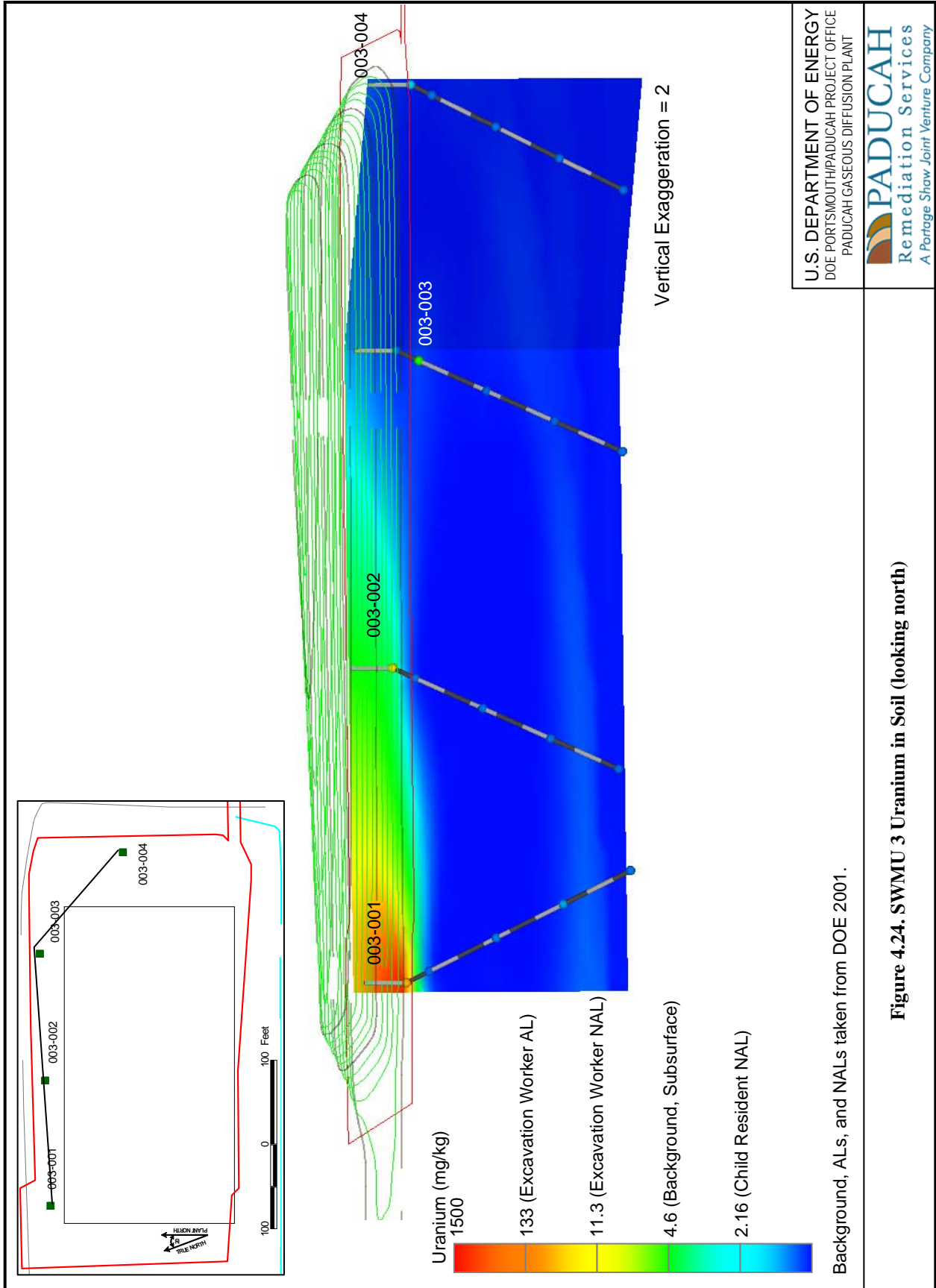
Analysis	Depth (ft)	Excavation Worker NAL	Background	RI Data									
				003-001	003-002	003-003	003-004	003-005	003-006	003-007	003-008	003-009	003-010
Thorium-232	5	1.95	1.5					0.274	0.223	0.554	0.414	0.445	0.375
	10	1.95	1.5	0.328	0.432	0.25	0.364	0.512	0.288	0.304	0.376	0.377	0.324
	15	1.95	1.5	0.265	0.451	0.277	0.389	0.266	0.289	0.331	0.306	0.431	0.338
	30	1.95	1.5	0.471	0.218	0.292	0.234						
	45	1.95	1.5	0.603	0.27	0.197	0.27						
	60	1.95	1.5	0.34	0.405	0.472	0.333						
Thorium-234	5	n/a	n/a					11.6	3.03	9.81	1.59	1.06U	1.36U
	10	n/a	n/a	14.1	14.1	0.877U	0.87U	0.951U	0.851U	11	0.854U	0.91U	0.916U
	15	n/a	n/a	0.918U	0.864U	0.867U	0.847U	0.912U	1.82	0.909U	0.84U	0.881U	0.867U
	30	n/a	n/a	0.894U	0.975U	0.91U	0.796U						
	45	n/a	n/a	1.1U	0.882U	0.834U	0.958U						
	60	n/a	n/a	0.882U	0.946U	1U	1.02U						
Uranium	5	n/a	n/a					7.46	3.48	6.55	0.28U	0.282U	0.281U
	10	n/a	n/a	8.53	25.8	0.3U	0.299U	0.283U	0.283U	2.7	0.282U	0.283U	0.281U
	15	n/a	n/a	0.464	0.297U	0.302U	0.301U	0.285U	0.286U	0.284U	0.282U	0.281U	0.283U
	30	n/a	n/a	0.297U	0.299U	0.303U	0.303U						
	45	n/a	n/a	0.602	0.301U	0.304U	0.304U						
	60	n/a	n/a	0.299U	0.297U	0.371	0.303U						
Uranium-234	5	2.84	2.4					1.08	0.533	0.913	0.126U	0.142	0.127U
	10	2.84	2.4	0.927	3.02	0.139U	0.138U	0.127U	0.127U	0.392	0.127U	0.127U	0.126U
	15	2.84	2.4	0.133U	0.132U	0.139U	0.138U	0.127U	0.128U	0.127U	0.127U	0.126U	0.127U
	30	2.84	2.4	0.18	0.133U	0.139U	0.139U						
	45	2.84	2.4	0.305	0.144	0.139U	0.139U						
	60	2.84	2.4	0.178	0.211	0.249	0.139U						
Uranium-235	5	0.455	0.14					0.102	0.0397	0.0799	0.036U	0.0371U	0.0365U
	10	0.455	0.14	0.14	0.362	0.0374U	0.0373U	0.0364U	0.0364U	0.037U	0.0366U	0.0371U	0.0362U
	15	0.455	0.14	0.0387U	0.0383U	0.0383U	0.0379U	0.037U	0.0374U	0.0372U	0.0372U	0.0366U	0.0372U
	30	0.455	0.14	0.038U	0.039U	0.0389U	0.0385U						
	45	0.455	0.14	0.0382U	0.04U	0.0402U	0.0401U						
	60	0.455	0.14	0.0385U	0.0381U	0.0388U	0.0395U						
Uranium-238	5	1.17	1.2					6.29	2.9J	5.55	0.127	0.201	0.118U
	10	1.17	1.2	7.47	22.4	0.124U	0.2	0.142	0.12U	2.28	0.118U	0.119U	0.118U
	15	1.17	1.2	0.354	0.325	0.125U	0.124U	0.12U	0.121U	0.12U	0.118U	0.118U	0.118U
	30	1.17	1.2	0.192	0.127U	0.125U	0.125U						
	45	1.17	1.2	0.271	0.129	0.125U	0.125U						
	60	1.17	1.2	0.147	0.127U	0.19	0.125U						

Blank cells indicate interval was not sampled for the specified analysis. Maximum value shown for each depth interval.

Bold indicates result is greater than NAL value.

Italics indicates result is greater than background value.

Bold + Italics indicate result is greater than both NAL and background values.



Background, ALs, and NALs taken from DOE 2001.

Figure 4.24. SWMU 3 Uranium in Soil (looking north)

4.4.2 SWMU 3 Groundwater

A large amount of historical UCRS and RGA groundwater data were available for SWMU 3. MW67 and MW76 (Figure 4.10) were evaluated for inclusion in the SWMU 3 monitoring system, but only one new well installed during the RI (MW420) will be added to the compliance monitoring system. Results from those three wells are discussed below. Additionally, UCRS groundwater samples were collected from two of the four angled borings at SWMU 3 as part of this RI. The UCRS data sources included well samples collected from MW85, MW88, MW91, and MW94 for the period 1995 through present (these wells are screened between 29 and 40 ft bgs around the perimeter of C-404) and samples from the BGOU RI temporary borings 003-003 (at 28 ft) and 003-004 (at 30 ft) (shown on Figure 4.10). All of these samples represent the HU2 interval within the UCRS.

A review of RI and historical data identified the UCRS contaminants listed in Table 4.15. All sample locations documented levels of TCE, technetium-99, and at least one metal (arsenic, iron, lead, manganese, molybdenum, and uranium) that exceed screening criteria. Arsenic and uranium are the only contaminants in this table that also are found in the SWMU 3 subsurface soil contaminant list exceeding the subsurface soil screening criteria.

Leachate from the SWMU 3 sump is periodically removed, sampled, and treated prior to release. Some of the constituents found in this leachate from CY 2003 through 2008 and the maximum concentrations are shown in Table 4.16).

Table 4.15. Contaminants Found in SWMU 3 Leachate

Analyte	Maximum	Frequency of Detection
Arsenic (mg/L)	0.0011	2/18
Barium (mg/L)	0.0871	18/18
Cadmium (mg/L)	NA	0/18
Fluoride (mg/L)	10	17/18
Lead (mg/L)	NA	0/18
Neptunium-237 (pCi/L)	1.62	12/18
PCB (µg/L)	1.18	17/18
Selenium (mg/L)	NA	0/18
Technetium-99 (pCi/L)	365	18/18
TCE (µg/L)	22	5/18
Uranium (mg/L)	115	18/18
Uranium-234 (pCi/L)	3,390	18/18
Uranium-235 (pCi/L)	1,050	15/15
Uranium-238 (pCi/L)	37,900	18/18

Wells characterize the upper and lower RGA around the perimeter of C-404 to monitor for potential contamination derived from C-404. Upper RGA wells include MW67, MW76, MW84, MW87, MW90/90A, MW93, and MW227. The lower RGA wells are MW86, MW89, MW92, MW95/95A, and MW226. The data base for screening RGA contaminants is the monitoring data for the period 1995 to present. RGA groundwater contaminants for SWMU 3 are listed in Table 4.17.

Table 4.16. SWMU 3 UCRS Groundwater Contaminants

Analysis	Historical Data	RI	Frequency of Detection ^a	Frequency of Detection		
				Above Background	Above NAL	Above MCL
Metals (mg/L)						
Aluminum	N/A	59	2/2	N/A	2/2	N/A
Arsenic	0.012	ND	34/46	N/A	34/46	9/46
Arsenic, Dissolved	0.012	0.00159	36/39	N/A	36/39	8/39
Barium	N/A	0.167	2/2	N/A	2/2	0/2
Barium, Dissolved	N/A	0.102	2/2	N/A	0/2	0/2
Beryllium	N/A	0.00135	2/2	N/A	0/2	0/2
Cadmium	ND	0.000711	1/126	N/A	1/126	0/126
Calcium	N/A	37.8	2/2	N/A	N/A	N/A
Calcium, Dissolved	N/A	35.2	2/2	N/A	N/A	N/A
Chromium	1.16	ND	45/126	N/A	0/126	23/126
Chromium, Dissolved	0.043	ND	2/72	N/A	0/72	0/72
Cobalt	N/A	0.0118	1/2	N/A	0/2	N/A
Cobalt, Dissolved	N/A	0.00691	2/2	N/A	0/2	N/A
Iron	N/A	43.5	2/2	N/A	2/2	N/A
Iron, Dissolved	N/A	14	2/2	N/A	2/2	N/A
Lead	0.00539	0.0172	4/46	N/A	2/46	2/46
Magnesium	N/A	20.2	2/2	N/A	0/2	N/A
Magnesium, Dissolved	N/A	16.3	2/2	N/A	0/2	N/A
Manganese	N/A	1.45	2/2	N/A	2/2	N/A
Manganese, Dissolved	N/A	1.14	2/2	N/A	2/2	N/A
Mercury	ND	0.000026	2/121	N/A	0/121	0/121
Mercury, Dissolved	0.0004	ND	1/41	N/A	0/41	0/41
Molybdenum	N/A	0.0184	2/2	N/A	1/2	N/A
Molybdenum, Dissolved	N/A	0.0191	1/2	N/A	1/2	N/A
Nickel, Dissolved	N/A	0.0122	1/2	N/A	0/2	N/A
Selenium	0.0148	ND	10/112	N/A	6/112	0/112
Selenium, Dissolved	0.00676	ND	4/41	N/A	0/41	0/41
Sodium	N/A	76	2/2	N/A	0/2	N/A
Sodium, Dissolved	N/A	77	2/2	N/A	0/2	N/A
Uranium	0.0518	0.00193	14/150	N/A	14/150	2/150
Uranium, Dissolved	0.0168	ND	14/52	N/A	14/52	0/52
Zinc, Dissolved	N/A	0.0227	1/2	N/A	0/2	N/A
Radionuclides (pCi/L)						
Plutonium-238	N/A	0.0508	1/2	N/A	0/2	N/A
Technetium-99	998	8.72	161/166	N/A	159/166	1/166
Thorium-228	N/A	0.632	2/2	N/A	2/2	N/A
Thorium-230	N/A	0.405	2/2	N/A	0/2	N/A
Thorium-232	N/A	0.478	2/2	N/A	1/2	N/A
Uranium	N/A	3.3	2/2	N/A	N/A	N/A
Uranium-234	14.39	2.33	11/27	N/A	10/27	N/A
Uranium-235	0.91	0.0613	5/27	N/A	1/27	N/A
Uranium-238	34.81	0.912	13/27	N/A	13/27	N/A
Uranium-238, Dissolved	N/A	0.158	1/2	N/A	0/2	N/A
Volatiles (mg/L)						
cis-1,2-Dichloroethene	N/A	0.014	1/2	N/A	1/2	0/2
Trichloroethene	1.8	0.046	105/126	N/A	104/126	82/126

^aFrequency of detection is the number of detections of an analyte per number of analyses (includes regular and duplicate samples).

N/A = not applicable ND = not detected

Table 4.17. SWMU 3 RGA Groundwater Contaminants

Analysis	Historical Data	RI	Frequency of Detection ^a	Frequency of Detection		
				Above Background	Above NAL	Above MCL
Metals (mg/L)						
Aluminum	0.547	N/A	6/18	0/18	0/18	N/A
Arsenic	0.12	N/A	28/79	2/79	28/79	2/79
Arsenic, Dissolved	0.00265	N/A	3/53	0/53	3/53	0/53
Barium	0.314	N/A	18/18	1/18	13/18	0/18
Beryllium	0.00078	N/A	1/19	0/19	0/19	0/19
Boron	0.0195	N/A	10/10	N/A	0/10	N/A
Cadmium	0.0037	N/A	1/329	0/329	1/329	0/329
Calcium	30.8	N/A	20/20	0/20	N/A	N/A
Chromium	0.949	N/A	45/329	18/329	0/329	29/329
Chromium, Dissolved	0.037	N/A	2/121	0/121	0/121	0/121
Cobalt	0.00295	N/A	3/18	0/18	0/18	N/A
Copper	0.0099	N/A	2/18	0/18	0/18	0/18
Iron	6.02	N/A	14/19	1/19	6/19	N/A
Lead	0.09	N/A	5/124	0/124	2/124	2/124
Magnesium	12	N/A	20/20	0/20	0/20	N/A
Manganese	1.4	N/A	19/19	10/19	14/19	N/A
Mercury	0.000046	N/A	1/319	0/319	0/319	0/319
Nickel	0.232	N/A	5/18	0/18	2/18	N/A
Potassium	3.27	N/A	11/20	0/20	N/A	N/A
Selenium	0.00774	N/A	12/298	9/298	1/298	0/298
Silicon	13.9	N/A	10/10	N/A	N/A	N/A
Sodium	19.2	N/A	20/20	0/20	0/20	N/A
Thallium	0.0028	N/A	1/18	0/18	0/18	1/18
Uranium	0.09	N/A	15/395	6/395	15/395	2/395
Vanadium	0.0008	N/A	2/19	0/19	0/19	N/A
Zinc	0.01	N/A	5/18	0/18	0/18	N/A
Radionuclides (pCi/L)						
Americium-241	0.35	N/A	1/1	N/A	0/1	N/A
Technetium-99	385	N/A	179/439	80/439	108/439	0/439
Thorium-230	0.74	N/A	1/1	0/1	1/1	N/A
Uranium-234	199.68	N/A	10/67	8/67	8/67	N/A
Uranium-235	9.96	N/A	7/65	3/65	3/65	N/A
Uranium-238	210.83	N/A	11/67	8/67	10/67	N/A
Volatiles (mg/L)						
1,1-Dichloroethane	0.0013	N/A	1/19	N/A	0/19	N/A
1,1-Dichloroethene	0.012	N/A	7/19	N/A	7/19	1/19
2-Butanone	0.035	N/A	1/19	N/A	0/19	N/A
Carbon tetrachloride	0.00049	N/A	3/19	N/A	3/19	0/19
Chloroform	0.0005	N/A	8/20	N/A	8/20	N/A
cis-1,2-Dichloroethene	0.014	N/A	9/20	N/A	5/20	0/20
Tetrachloroethene	0.00052	N/A	5/20	N/A	0/20	0/20
Trichloroethene	0.61	N/A	212/334	N/A	190/334	147/334

^a Frequency of detection is the number of detections of an analyte per number of analyses (includes regular and duplicate samples).

N/A = not applicable ND = not detected

Arsenic, iron, manganese, and uranium concentrations in RGA groundwater samples (also contaminants in the UCRS) exceeded screening levels for metals at SWMU 3. All of these metals were found in both the upper and lower RGA with no significant differentiation with depth. 1,1-DCE and TCE exceeded screening levels for organics in RGA groundwater at SWMU 3. The hydrogeological assessment of SWMUs 2 and 3 that was completed as part of this RI (PRS 2007a) documents that an upgradient source accounts for the high TCE levels (refer to Figure 4.20). It remains possible that the upgradient contamination may mask some TCE loading to the RGA from SWMU 3. Because the 1,1-DCE detects occurred only in upgradient wells, it also appears to be related to an upgradient source.

The only radionuclides in the SWMU 3 RGA groundwater samples to exceed background were uranium-234 and uranium-238, occurring in both the upper and lower RGA. Uranium isotope activities were markedly higher in well cluster MW93/MW95A. In both MW93 and MW95A, the markedly higher analyses are singular events [1 of 14 analyses available for MW93 (March 1991 through February 2009) and 1 of 9 analyses available for MW95A (July 2004 through July 2008)] that may not be representative of groundwater quality in the wells. (Note: the mass concentration of uranium metal represented by the markedly higher uranium isotope analyses for MW93 and MW95A would exceed the MCL for uranium.)

Two existing monitoring wells (MW67 and MW76) and a new well (MW420) were evaluated for inclusion in the SWMU 3 compliance monitoring system. MW67 and MW76 were rehabilitated and purged prior to sampling since they had not been used in a long time. After analytical results were available and discussions were held with KEEC, it was determined that MW67 and MW76 would not be included in the compliance monitoring system and only MW420 would be added. Table 4.18 shows the contaminants that exceed the screening levels in these wells. All three wells monitor the upper part of the RGA.

Groundwater monitoring under the RCRA permit for the unit, however, has shown statistically significant increases of TCE above background in one of three downgradient compliance wells in the upper RGA (MW84). C-404 Landfill Source Demonstration, Paducah Gaseous Diffusion Plant, Paducah, Kentucky (PRS 2007b) stated that, while C-404 may be a source of groundwater contamination, the statistically significant increase of TCE in MW84 was due to an upgradient source migrating through the C-404 area. Additionally, trends of arsenic, chromium, selenium, technetium-99, uranium-234, and uranium-238 have required statistical analysis for contaminant determination. Figures 4.25 through 4.45 present trend graphs of these contaminants in UCRS and upper and lower RGA MWs. These graphs show both the result and the laboratory detection limit. In some cases, particularly with radiological constituents, the reported result is less than the detection limit (this would be qualified as a “nondetect” in the database). With these charts, it’s best to focus on “long-term” trends rather than individual fluctuations. Some of the notable trends in UCRS MWs include increasing levels of arsenic in MW85 and increasing levels of chromium, TCE, and technetium-99 in MW91. TCE was decreasing in UCRS wells MW88 and MW94, and technetium-99 was decreasing in UCRS wells MW85, MW88, and MW94. While MW88 historically has had a maximum TCE value of 1.8 mg/L, as indicated in Table 4.18, the recent contaminant levels are less than 0.005 mg/L (Figure 4.34). While SWMU 3 may have contributed TCE to groundwater in the past, it should be noted that any TCE source at SWMU 3 appears diminished due mostly to decreasing trends in UCRS MWs and no detections of TCE in SWMU 3 leachate since 2004 (Figure 4.46). The high levels of TCE in the RGA are related to groundwater flow and contaminant transport from upgradient sources such as the C-400 source (DOE 2007b). Table 4.19 provides detail (depth, sample location, and analytical results) for SWMU 3 groundwater samples, including nondetects and detections above screening levels.

Table 4.18. RGA Groundwater Contaminants in MW67, MW76, and MW420

Analysis	Depth (ft)	MW67	MW76	MW420
<i>Inorganics (mg/L)</i>				
Arsenic	67-77	0.0156	ND	ND
Chromium		0.0155	ND	ND
Selenium		ND	0.00874	ND
Arsenic, Dissolved		0.0158	ND	ND
Chromium, Dissolved		0.0124	ND	ND
<i>Organics-Volatiles (mg/L)</i>				
1,1,1-Trichloroethane	67-77	ND	0.00019	ND
1,1,2-Trichloroethane		0.0011	ND	ND
1,1-Dichloroethane		0.0002	0.00025	ND
1,1-Dichloroethene		ND	ND	ND
Carbon disulfide		ND	ND	0.0012
Carbon tetrachloride		ND	ND	ND
Chloroform		0.034	0.0012	0.00067
<i>cis</i> -1,2-Dichloroethene		0.12	ND	ND
Methylene chloride		0.00043	ND	ND
Tetrachloroethene		ND	0.0017	ND
Toluene		ND	ND	0.001
<i>trans</i> -1,2-Dichloroethene		0.0011	ND	ND
Trichloroethene		1.1	0.46	0.21
Trichlorofluoromethane		ND	0.0005	ND
Vinyl chloride		0.00047	ND	ND
<i>Radionuclides (pCi/L)</i>				
Technetium-99	67-77	43.6	103	22.83
Uranium-238		0.31	0.305	ND

ND = not detected or not detected above screening levels

Table 4.19. SWMU 3 Locations of Groundwater Contaminants (Continued)

Hydro Unit	Depth Interval	Analysis	Child Resident NAL	Bkgd	MCL	003-003	003-004	MW226	MW227	MW67	MW84	MW85	MW86	MW87	MW88	MW89	MW90	MW90A	MW91	MW92	MW93	MW94	MW95	MW95A			
UCRS	30-40	Metals (mg/L)																									
		Arsenic	0.000035	N/A	0.01								0.012			0.011				0.00389			0.01				
		Arsenic, Dissolved	0.000035	N/A	0.01								0.012			0.0081				0.00225			0.0025				
		Cadmium	0.000661	N/A	0.005								0.005U			0.005U				0.005U			0.005U				
		Chromium	1.76	N/A	0.1								0.05U			0.2U				0.947			1.16				
		Chromium, Dissolved	1.76	N/A	0.1								0.02U			0.02U				0.043			0.05U				
		Lead	0.015	N/A	0.015								0.05U			0.05U				0.05U			0.00539				
		Mercury	0.000444	N/A	0.002								0.0002U			0.0002U				0.0002U			0.0002U				
		Mercury, Dissolved	0.000444	N/A	0.002								0.0002U			0.0002U				0.0004			0.0002U				
		Selenium	0.00754	N/A	0.05								0.01U			0.01U				0.0148			0.00797				
		Selenium, Dissolved	0.00754	N/A	0.05								0.005U			0.005U				0.00676			0.005U				
		Uranium	0.000906	N/A	0.03								0.0518			1U				1U			0.00433				
		Uranium, Dissolved	0.000906	N/A	0.03								0.0168			0.001U				0.001U			0.0038				
		Radionuclides (pCi/L)																									
		Technetium-99	14	N/A	900								406			524				998			660				
		Uranium-234	0.546	N/A	N/A								3.9			3.2U				3.22U			14.39				
		Uranium-235	0.538	N/A	N/A								0.515			0.45U				0.45U			0.91				
		Uranium-238	0.443	N/A	N/A								23.3			1.09U				1.12U			34.81				
		Volatiles (mg/L)																									
		Trichloroethene	0.0016	N/A	0.005								0.045			1.8				0.079			0.11				
		RGA	67-77	Metals (mg/L)																							
				Aluminum	1.49	2.189	N/A					0.547		0.2U			0.2U				0.0335			0.0432			
				Arsenic	0.000035	0.005	0.01					0.12		0.00319			0.00116				0.01U			0.01U			
Arsenic, Dissolved	0.000035			0.005	0.01					0.12		0.00265			0.001U				0.001U			0.01U					
Barium	0.104			0.235	2					0.12		0.0744			0.0635				0.12			0.179					
Beryllium	0.00264			0.004	0.004					0.001U		0.0002U			0.001U				0.005U			0.005U					
Boron	0.136			N/A	N/A					0.001U		0.01			0.0106				0.0195			0.0195					
Cadmium	0.000661			0.01	0.005					0.0037		0.005U			0.005U				0.005U			0.005U					
Calcium	N/A			41.238	N/A					25.8		14			30.1				30.8			30.8					
Chromium	1.76			0.144	0.1					0.949		19.7			0.05U				0.05U			0.025U					
Chromium, Dissolved	1.76			0.05	0.1					0.037		0.02U			0.02U				0.01U			0.02U					
Cobalt	0.0906			0.045	N/A					0.00295		0.05U			0.001U				0.05U			0.05U					
Copper	0.0557			0.036	1.3					0.02U		0.02U			0.02U				0.0015			0.025U					
Iron	0.449			5.03	N/A					3.14		0.98			0.1U				0.0652			2.46					
Lead	0.015			0.129	0.015					0.09		0.05U			0.05U				0.003U			0.05U					
Magnesium	N/A			16.262	N/A					9.38		5.9			7.6				7.75			9.01					
Manganese	0.035			0.119	N/A					0.144		0.027			0.0396				0.491			1.06					
Mercury	0.000444			0.0002	0.002					0.0002U		0.0002U			0.0002U				0.0002U			0.00046					
Nickel	0.0301			0.682	N/A					0.232		0.04U			0.005U				0.0019			0.04U					
Potassium	N/A			5.195	N/A					0.706		0.83			0.478				2.93			2.38					
Selenium	0.00754			0.005	0.05					0.00551		0.01U			0.01U				0.0017			0.0019					
Silicon	N/A			N/A	N/A					7.22		7.22			13.9				13.9			13.5					
Sodium	N/A			59.45	N/A					15.6		13			19.2				19.2			18.4					
Thallium	N/A			0.056	0.002					0.002U		0.002U			0.01U				0.01U			0.0028					
Uranium	0.000906			0.002	0.03					0.02		0.0092U			0.5U				0.0008			0.05U					
Vanadium	0.00925			0.134	N/A					0.02U		0.0017U			0.002U				0.0068			0.0038					
Zinc	0.45			0.054	N/A					0.02U		0.01			0.02U				0.0068			0.0038					
Radionuclides (pCi/L)																											
Americium-241	0.371			N/A	N/A					0.35		0.35							20			25.6					
Technetium-99	14			22.3	900					367		8U			385				20			25.6					
Thorium-230	0.424			1.1	N/A					0.74		0.74			3.21U				3.21U			3.21U					
Uranium-234	0.546			0.7	N/A					2.9		0.2			0.43U				0.47U			0.47U					
Uranium-235	0.538	0.3	N/A					0.18		0.18			1.1U				1.1U			1.1U							
Uranium-238	0.443	0.7	N/A					6.69		0.16U			1.1U				1.1U			1.1U							

Table 4.19. SWMU 3 Locations of Groundwater Contaminants (Continued)

Hydro Unit	Depth Interval	Analysis	Child Resident NAL	Bkgd	MCL	003-003	003-004	MW226	MW227	MW67	MW84	MW85	MW86	MW87	MW88	MW89	MW90	MW90A	MW91	MW92	MW93	MW94	MW95	MW95A		
RGA	67-77	Volatiles (mg/L)																								
		1,1-Dichloroethane	0.0363	N/A	N/A							0.0013						0.001U	0.001U							
		1,1-Dichloroethene	0.000047	N/A	0.007							0.001U						0.001U	0.001U							
		2-Butanone	0.0868	N/A	N/A							0.005U						0.005U	0.005U							
		Carbon tetrachloride	0.000181	N/A	0.005							0.001U						0.001U	0.001U							
		Chloroform	0.0000287	N/A	N/A							0.001U						0.00019	0.00017							
		cis-1,2-Dichloroethene	0.00273	N/A	0.07							0.00047U						0.0005U	0.0005U							
		Tetrachloroethene	0.000582	N/A	N/A							0.001U						0.001U	0.001U							
		Trichloroethene	0.0016	N/A	0.005							0.0022	0.48					0.005	0.0024							
				Metals (mg/L)																						
			79-89	Aluminum	1.49	2.189	N/A			0.2U					0.204			0.2U					0.2U		0.0273	0.0264
				Arsenic	0.000035	0.005	0.01			0.00195					0.00399			0.005U					0.00105		0.01U	0.00151
				Arsenic, Dissolved	0.000035	0.005	0.01			0.001U					0.001U			0.005U					0.00124		0.001U	0.001U
				Barium	0.104	0.235	2			0.314					0.155			0.087					0.122		0.131	0.132
				Beryllium	0.00264	0.004	0.004			0.001U					0.001U			0.001U					0.005U		0.005U	0.005U
				Boron	0.136	N/A	N/A											0.01					0.01		0.0132	0.016
				Cadmium	0.000661	0.01	0.005			0.005U					0.005U			0.005U					0.005U		0.005U	0.005U
				Calcium	N/A	41.238	N/A			29.6					20.2			14.2					20.3		21	22.4
				Chromium	1.76	0.144	0.1			0.428					0.0147			0.017					0.05U		0.05U	0.025U
				Chromium, Dissolved	1.76	0.05	0.1			0.05U					0.011			0.02U					0.02U		0.01U	0.02U
				Cobalt	0.0906	0.045	N/A			0.00172					0.001U			0.001U					0.05U		0.0017	0.05U
				Copper	0.0557	0.036	1.3			0.02U					0.02U			0.02U					0.0099		0.025U	0.025U
				Iron	0.449	5.03	N/A			6.02					0.139			0.1U					0.19		0.226	0.0434
				Lead	0.015	0.129	0.015			0.05U					0.05U			0.05U					0.00146		0.003U	0.05U
				Magnesium	N/A	16.262	N/A			12					7.92			5.96					8.22		8.45	8.88
		Manganese	0.035	0.119	N/A			1.4					0.408			0.591					0.0151		0.342	0.199		
		Mercury	0.000444	0.0002	0.002			0.0002U					0.0002U			0.0002U					0.0002U		0.0002U	0.0002U		
		Nickel	0.0301	0.682	N/A			0.0345					0.00522			0.005U					0.04U		0.0017	0.04U		
		Potassium	N/A	5.195	N/A			3.27					0.487			1.56					0.41		5U	0.378		
		Selenium	0.00754	0.005	0.05			0.00774					0.01U			0.01U					0.00531		0.005U	0.00607		
		Silicon	N/A	N/A	N/A								0.01U			0.01U					6.9		7.57	7.29		
		Sodium	N/A	59.45	N/A			17.6					13.4			11.8					13.6		14.4	13.8		
		Thallium	N/A	0.056	0.002			0.002U					0.002U			0.002U					0.01U		0.01U	0.01U		
		Uranium	0.000906	0.002	0.03			1U					0.05			1U					1U		0.09	1U		
		Vanadium	0.00925	0.134	N/A			0.02U					0.02U			0.02U					0.05U		0.00062	0.05U		
		Zinc	0.45	0.054	N/A			0.02U					0.02U			0.02U					0.01		0.02U	0.0053		
		Radionuclides (pCi/L)																								
		Technetium-99	14	22.3	900			166					29			25					22		24.2	30.1		
		Uranium-234	0.546	0.7	N/A			3.29U					1.37			3.35					3.19U		1.54	55.4		
		Uranium-235	0.538	0.3	N/A			0.43U					0.1			0.2					0.43U		0.09	3.21		
		Uranium-238	0.443	0.7	N/A			1.1U					5.02			0.522					1.1U		3.35	7.85		
		Volatiles (mg/L)																								
		1,1-Dichloroethane	0.0363	N/A	N/A			0.005U					0.001U			0.001U					0.001U		0.001U	0.001U		
		1,1-Dichloroethene	0.000047	N/A	0.007			0.012					0.001U			0.001U					0.0018		0.00071	0.0022		
		2-Butanone	0.0868	N/A	N/A			0.035					0.005U			0.005U					0.005U		0.005U	0.005U		
		Carbon tetrachloride	0.000181	N/A	0.005			0.005U					0.001U			0.001U					0.005U		0.005U	0.005U		
		Chloroform	0.0000287	N/A	N/A			0.005U					0.001U			0.001U					0.005U		0.005U	0.005U		
		cis-1,2-Dichloroethene	0.000287	N/A	N/A			0.005U					0.001U			0.001U					0.001U		0.00021	0.0005		
		Tetrachloroethene	0.00273	N/A	0.07			0.005U					0.0027			0.001U					0.0045		0.00089	0.014		
		Trichloroethene	0.000582	N/A	0.005			0.005U					0.001U			0.001U					0.001U		0.00045	0.00052		
		Trichloroethene	0.0016	N/A	0.005			0.61					0.38			0.0026					0.001		0.047	0.35		

Blank cells indicate interval was not sampled for the specified analysis. Maximum value shown for each depth interval.

N/A = not applicable

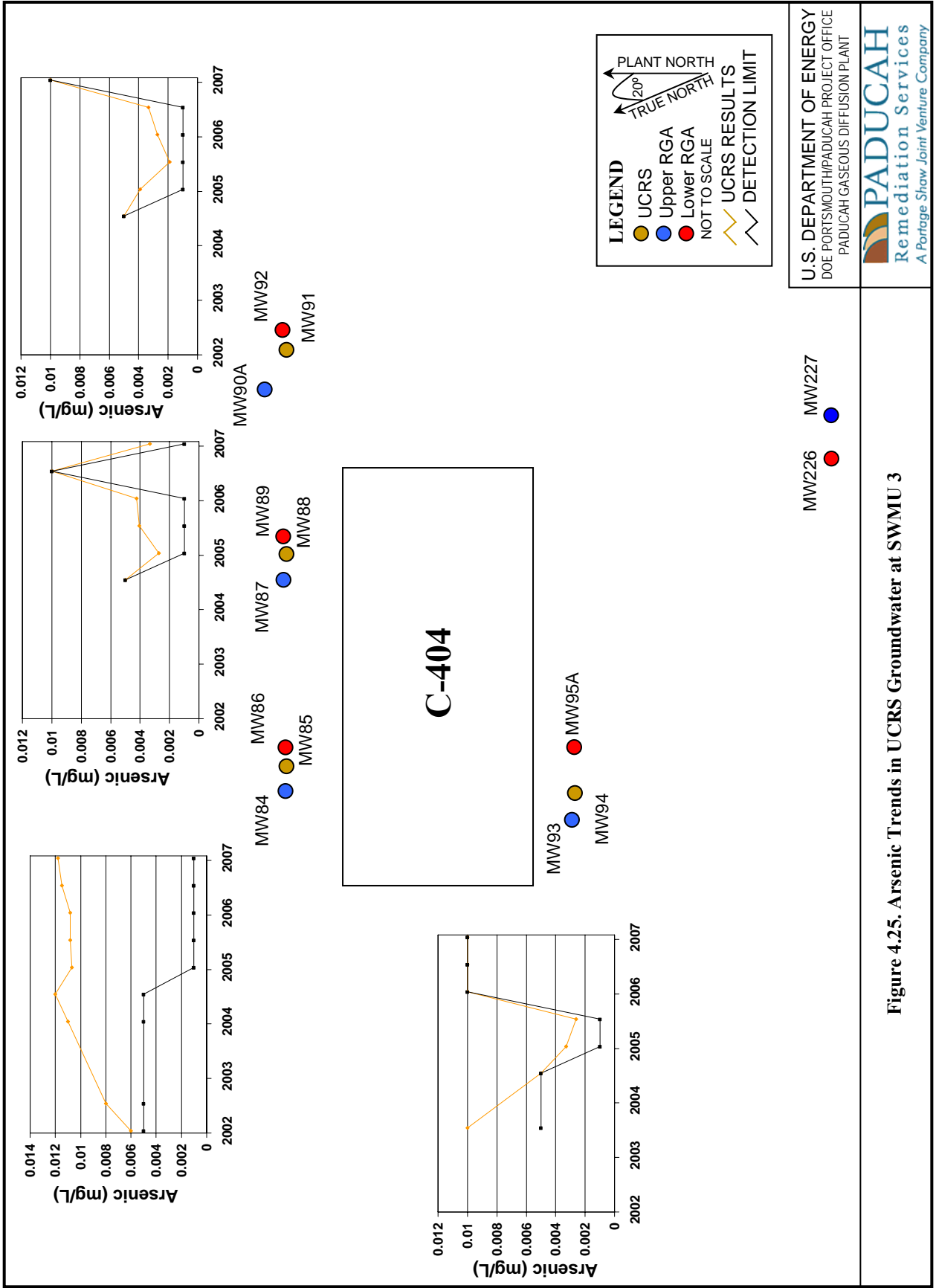
Bold indicates result is greater than NAL value.

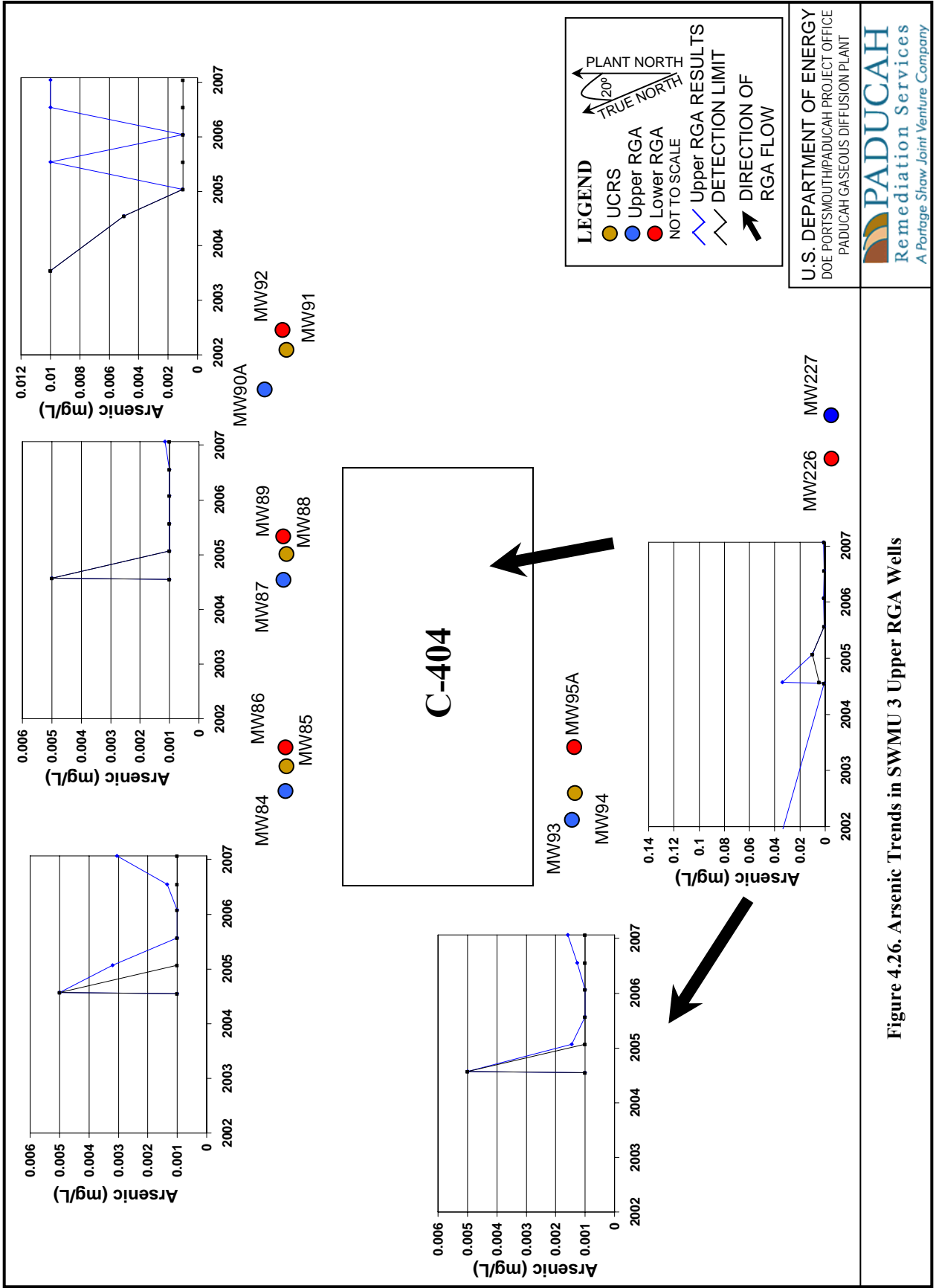
Italics indicates result is greater than MCL value.

Bold + Italics indicate result is greater than both NAL and MCL values.

Shading indicates result is greater than the background value.

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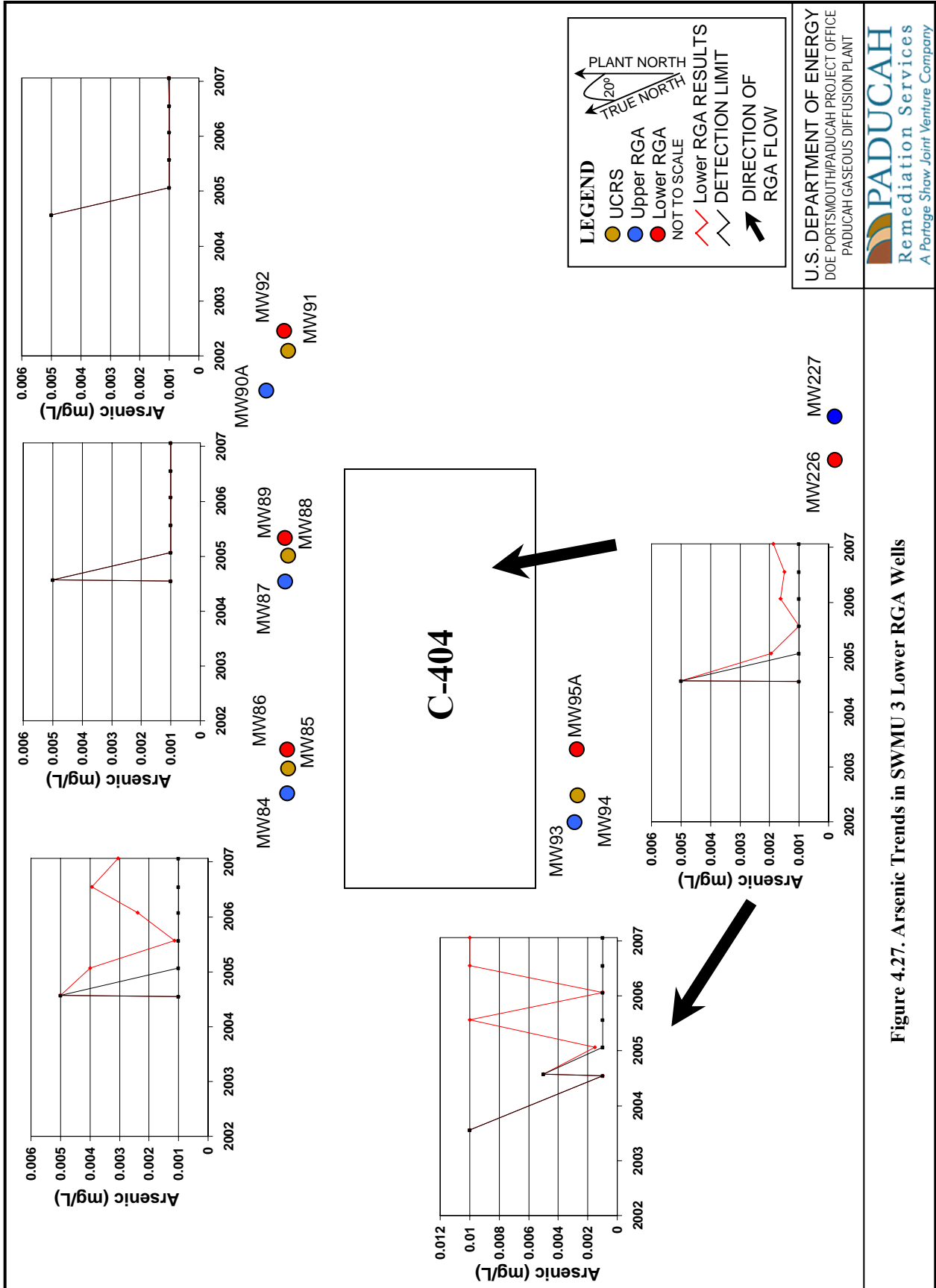


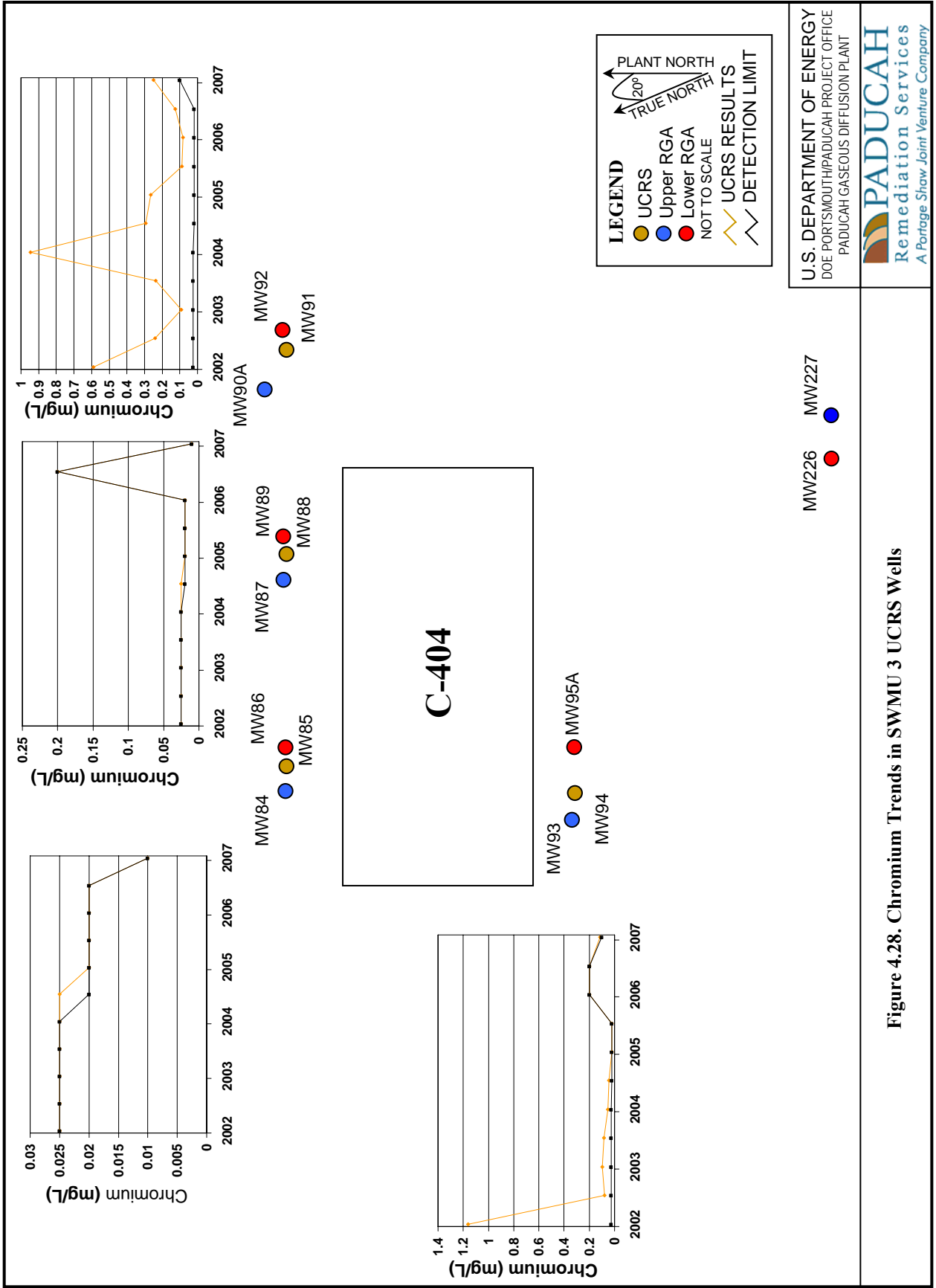


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Figure 4.26. Arsenic Trends in SWMU 3 Upper RGA Wells





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Figure 4.28. Chromium Trends in SWMU 3 UCERS Wells

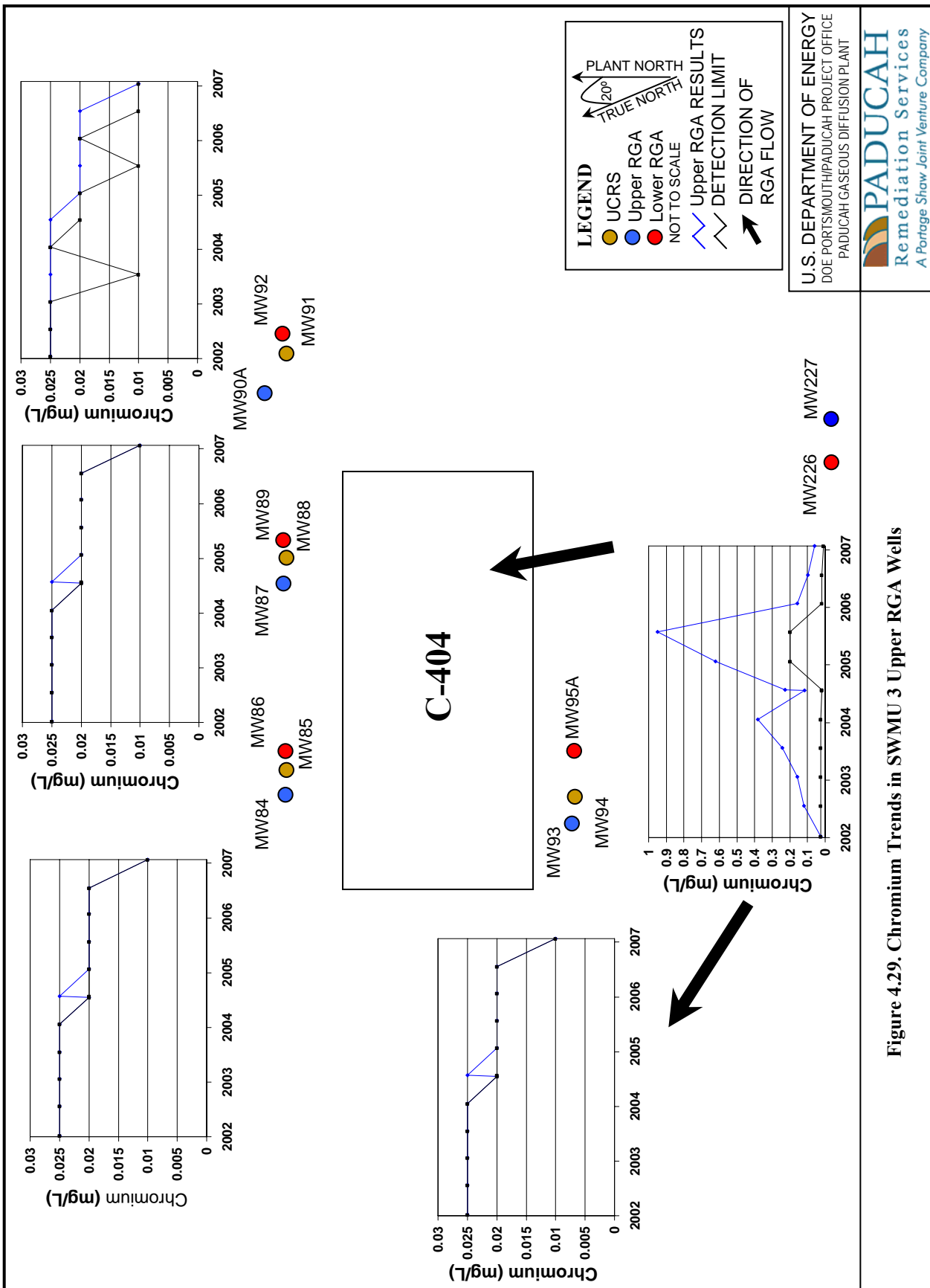


Figure 4.29. Chromium Trends in SWMU 3 Upper RGA Wells

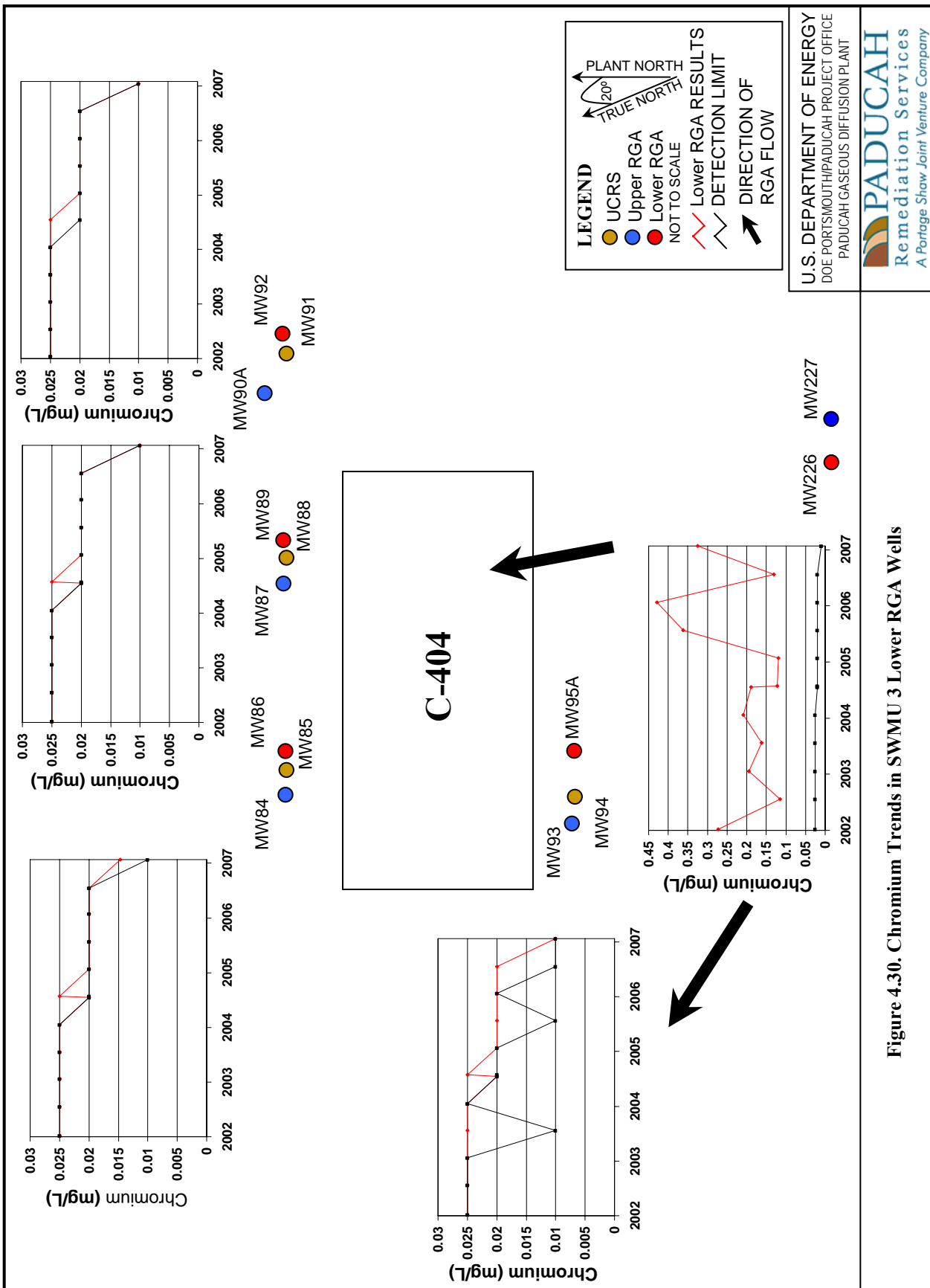
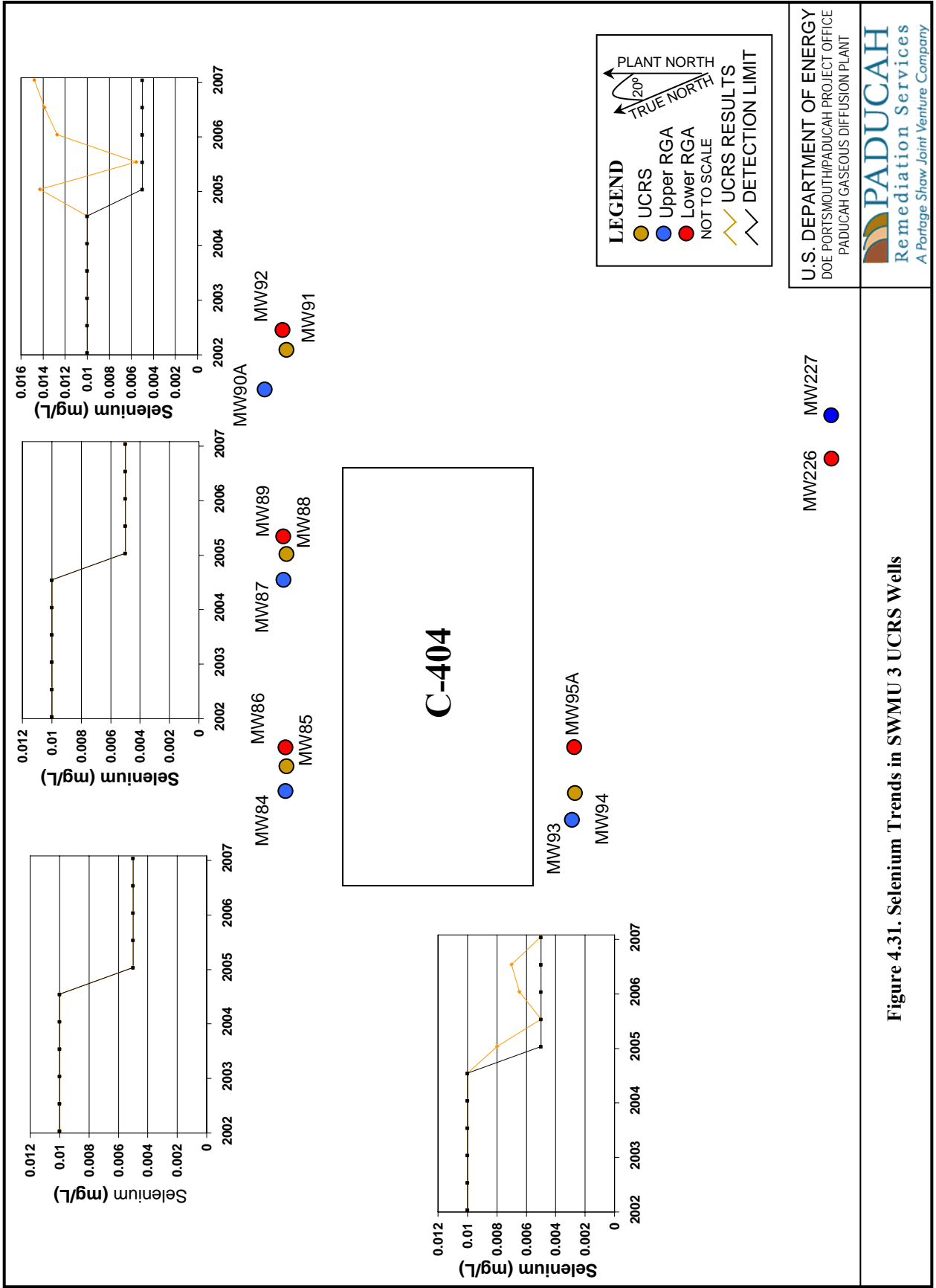


Figure 4.30. Chromium Trends in SWMU 3 Lower RGA Wells

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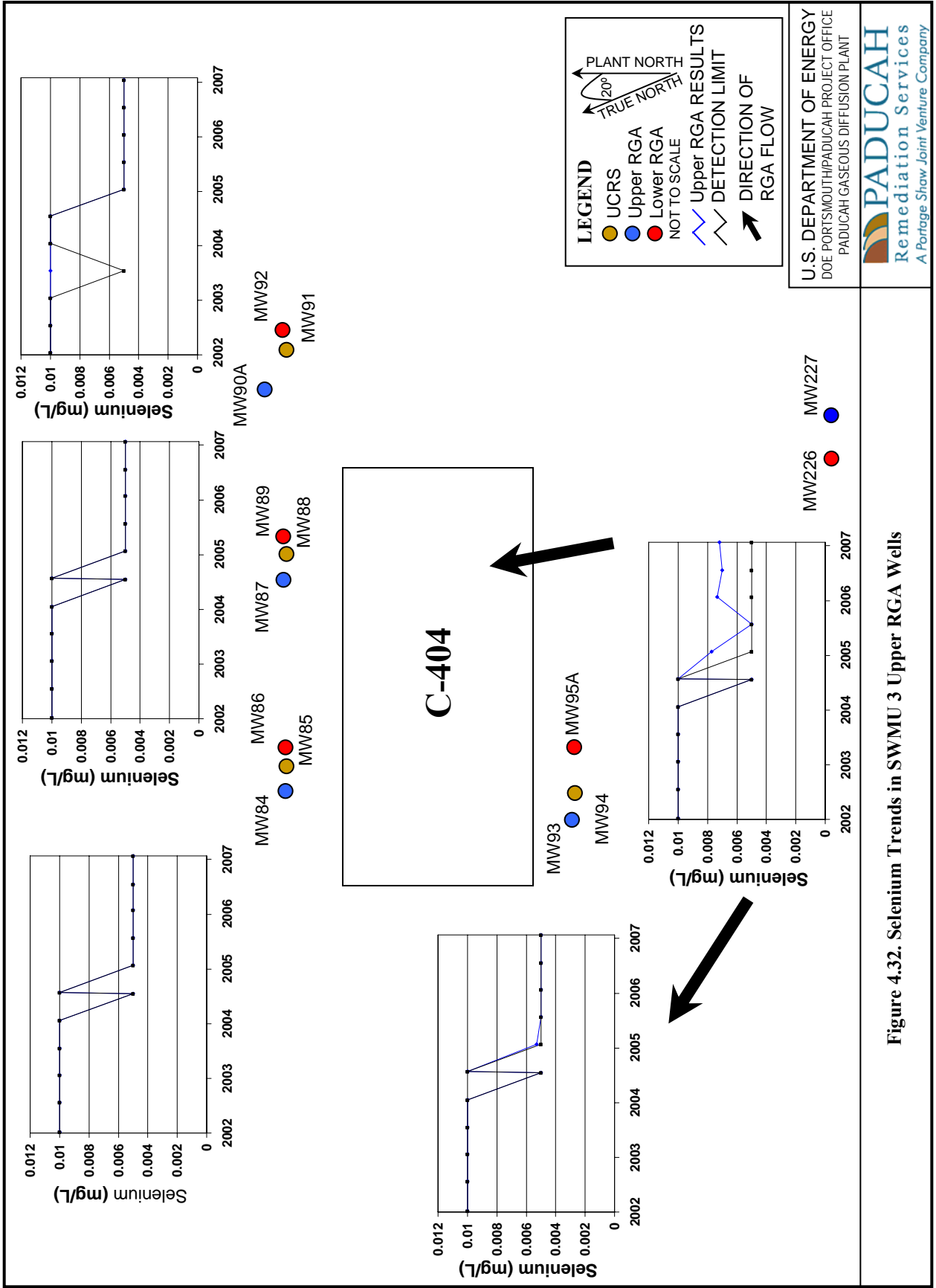
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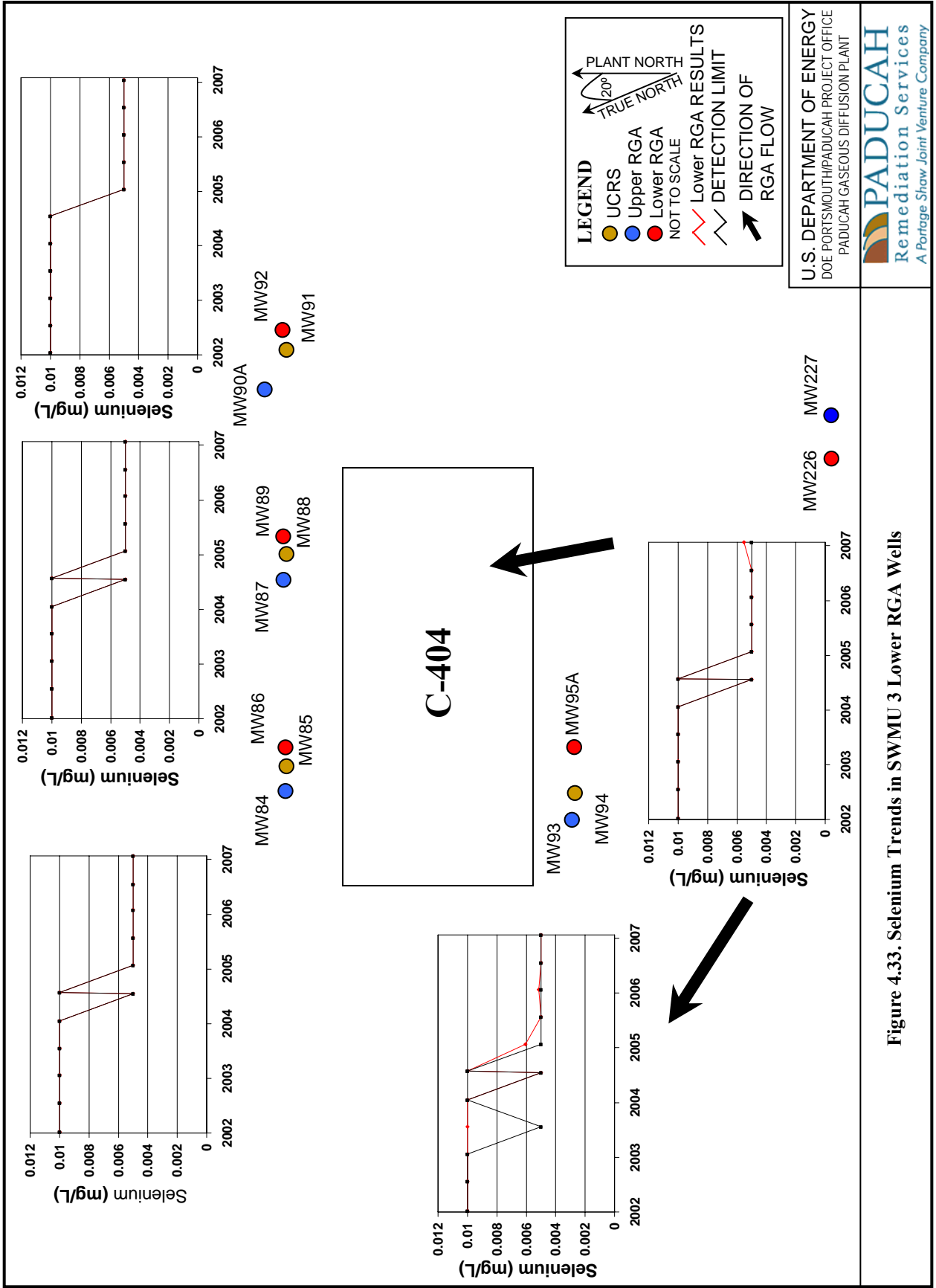


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Figure 4.31. Selenium Trends in SWMU 3 UCRS Wells





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Figure 4.33. Selenium Trends in SWMU 3 Lower RGA Wells

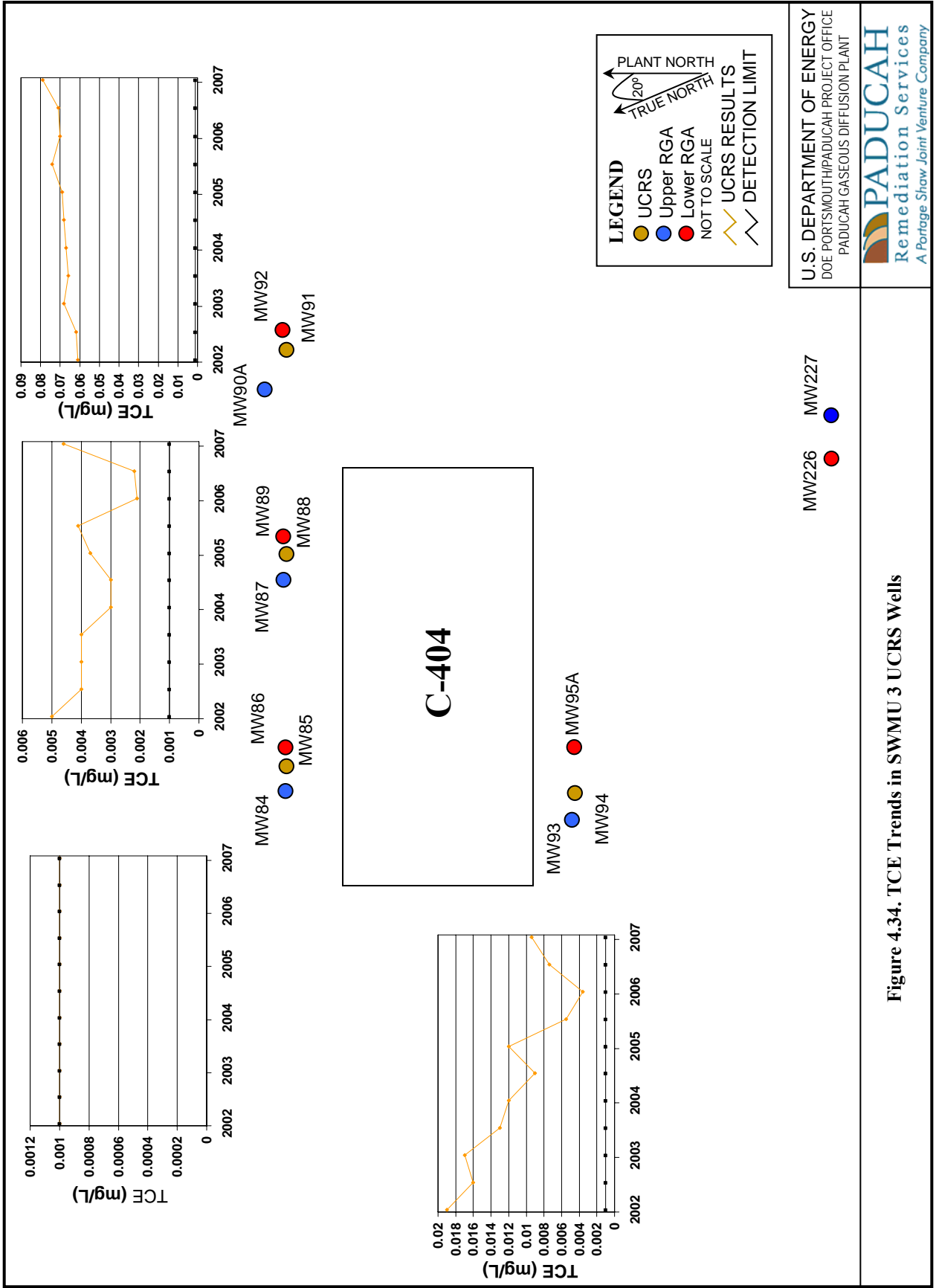
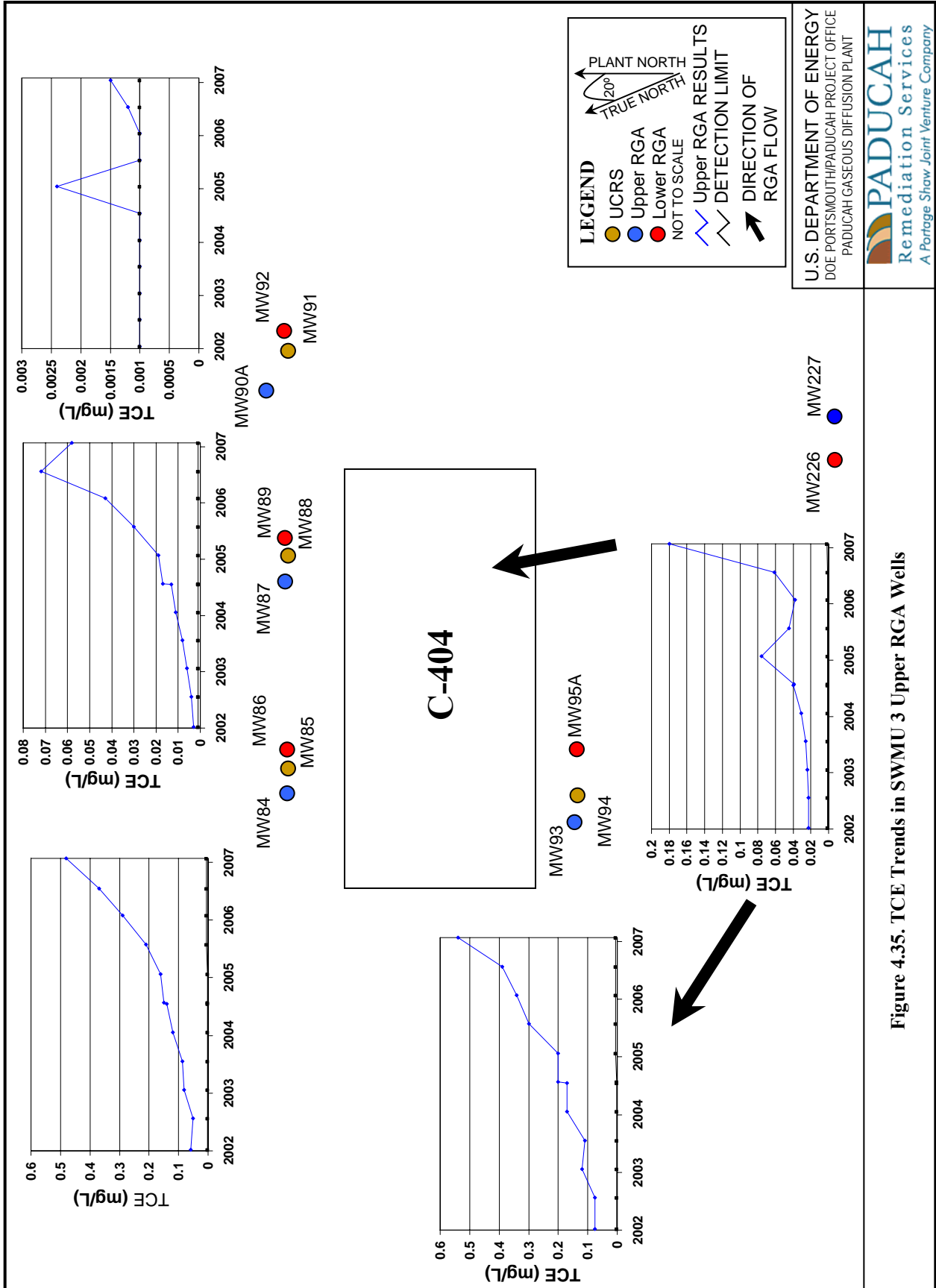


Figure 4.34. TCE Trends in SWMU 3 UCRS Wells

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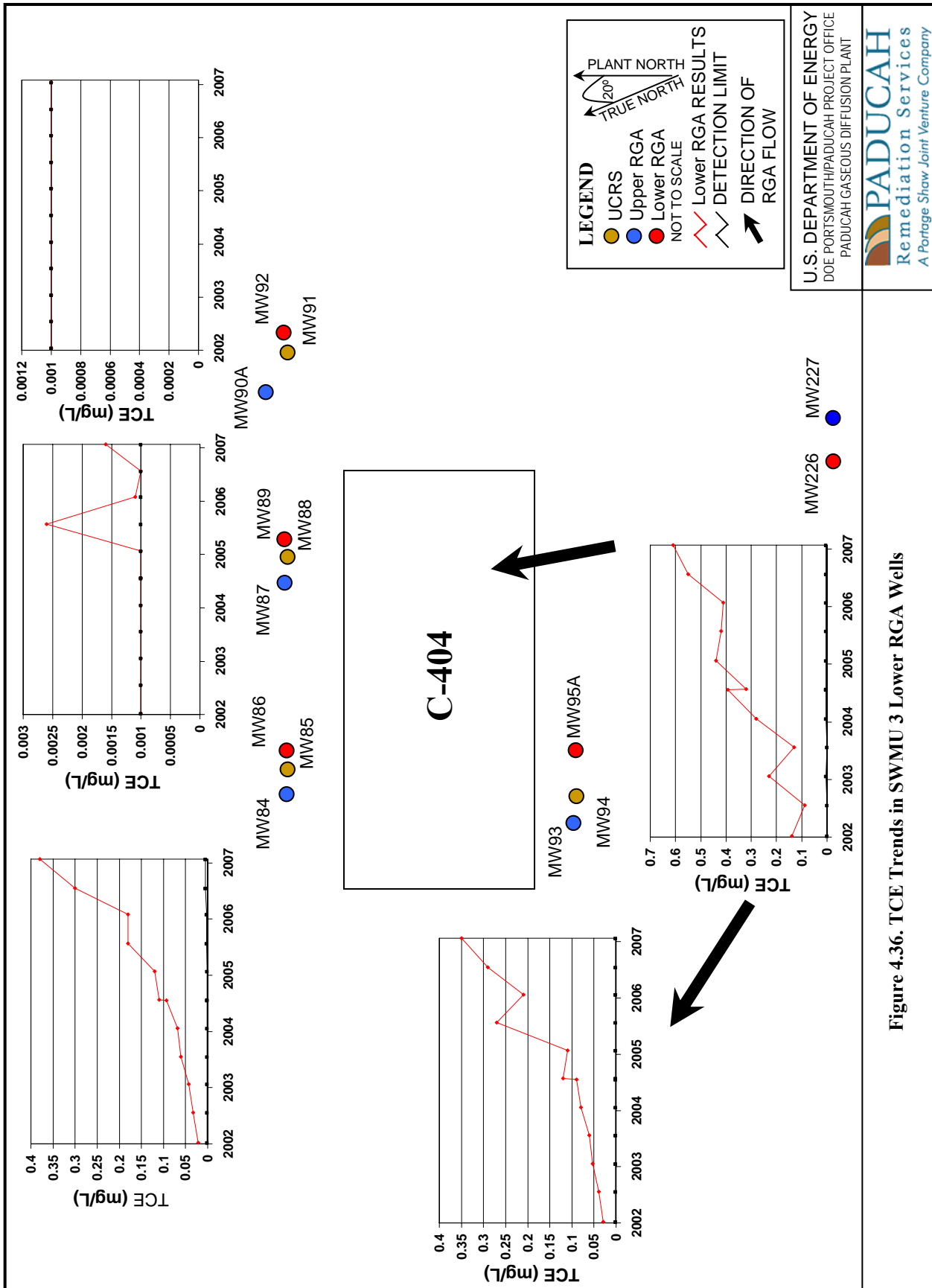


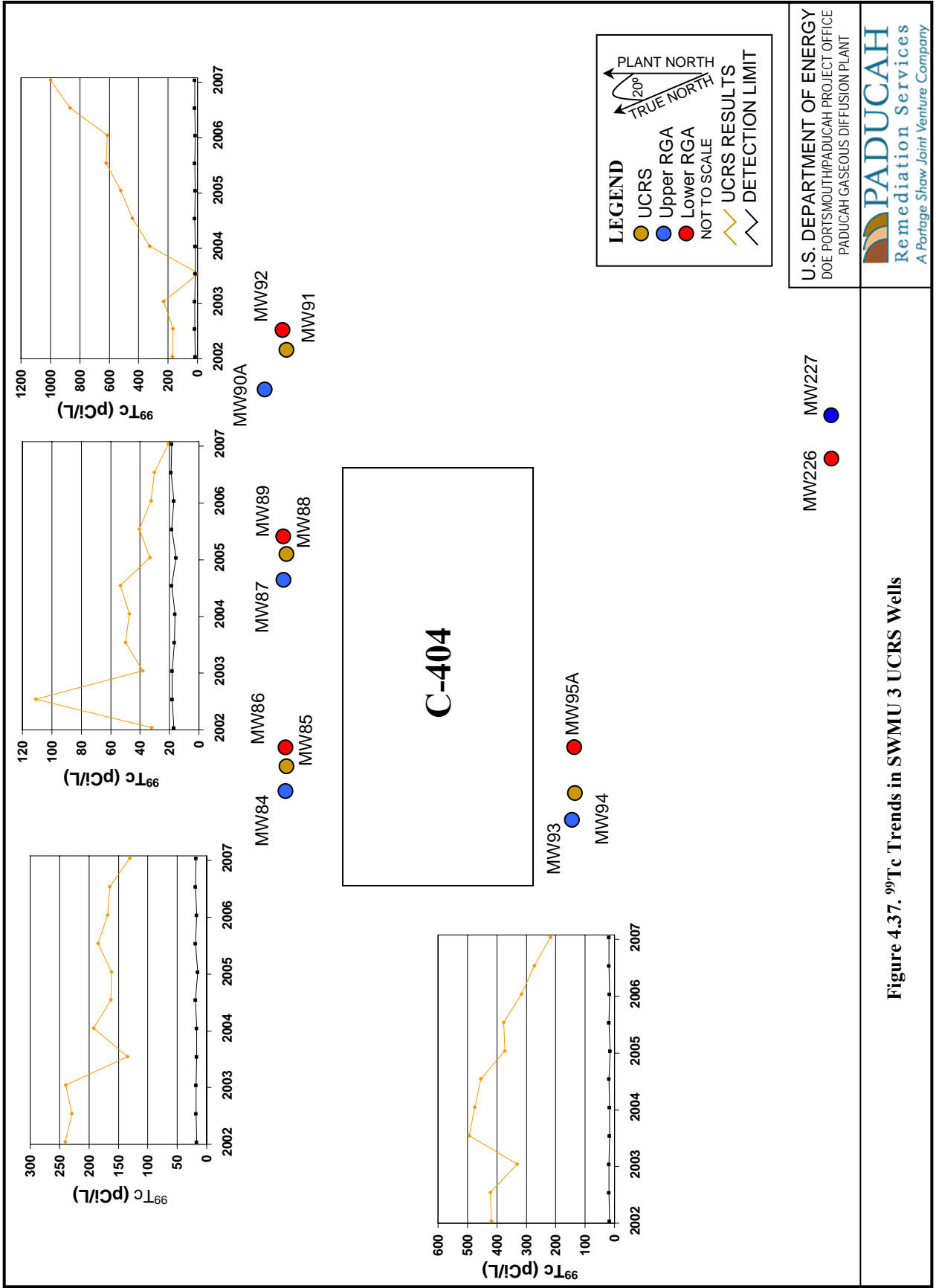


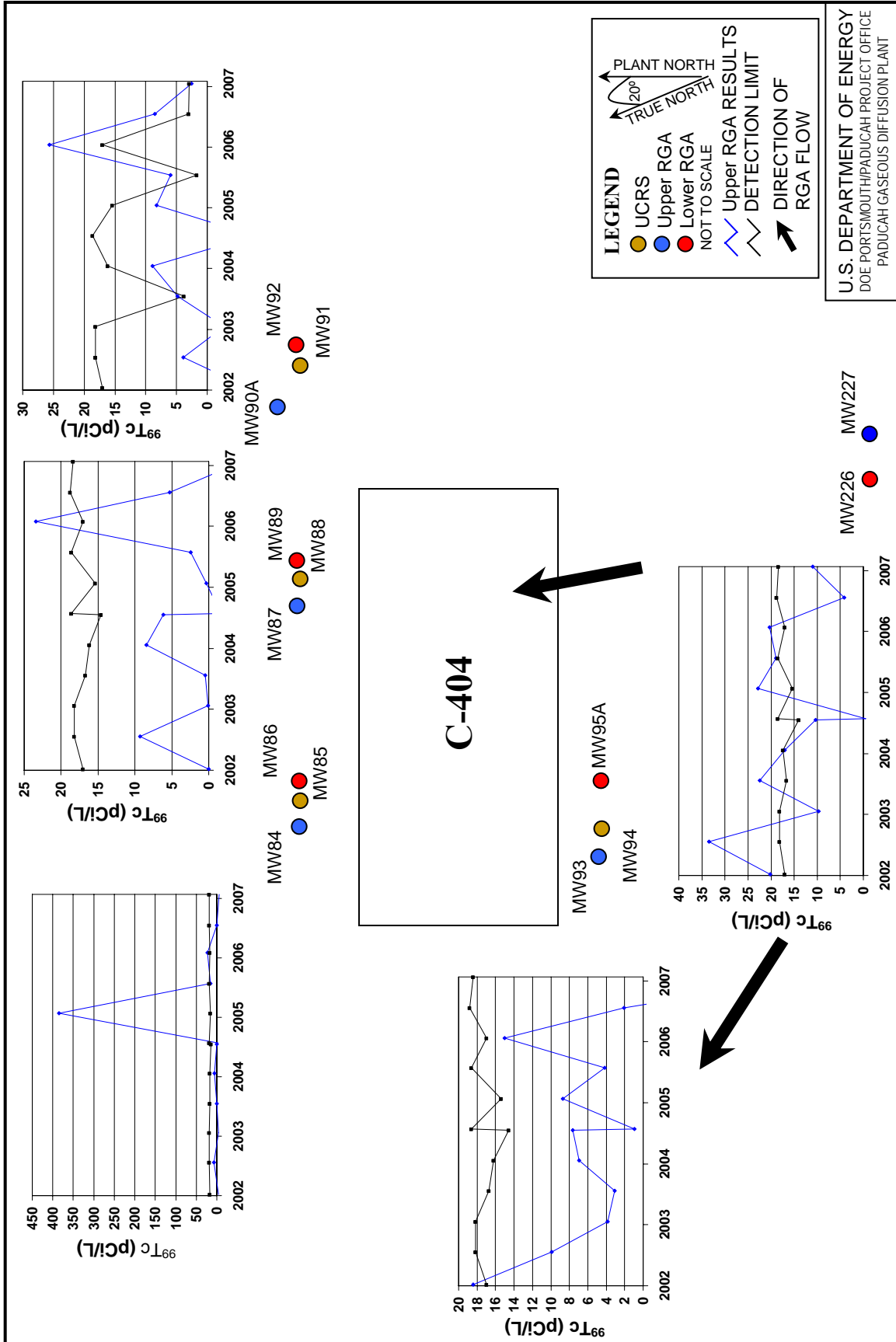
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Figure 4.35. TCE Trends in SWMU 3 Upper RGA Wells







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Figure 4.38. ⁹⁹Tc Trends in SWMU 3 Upper RGA Wells

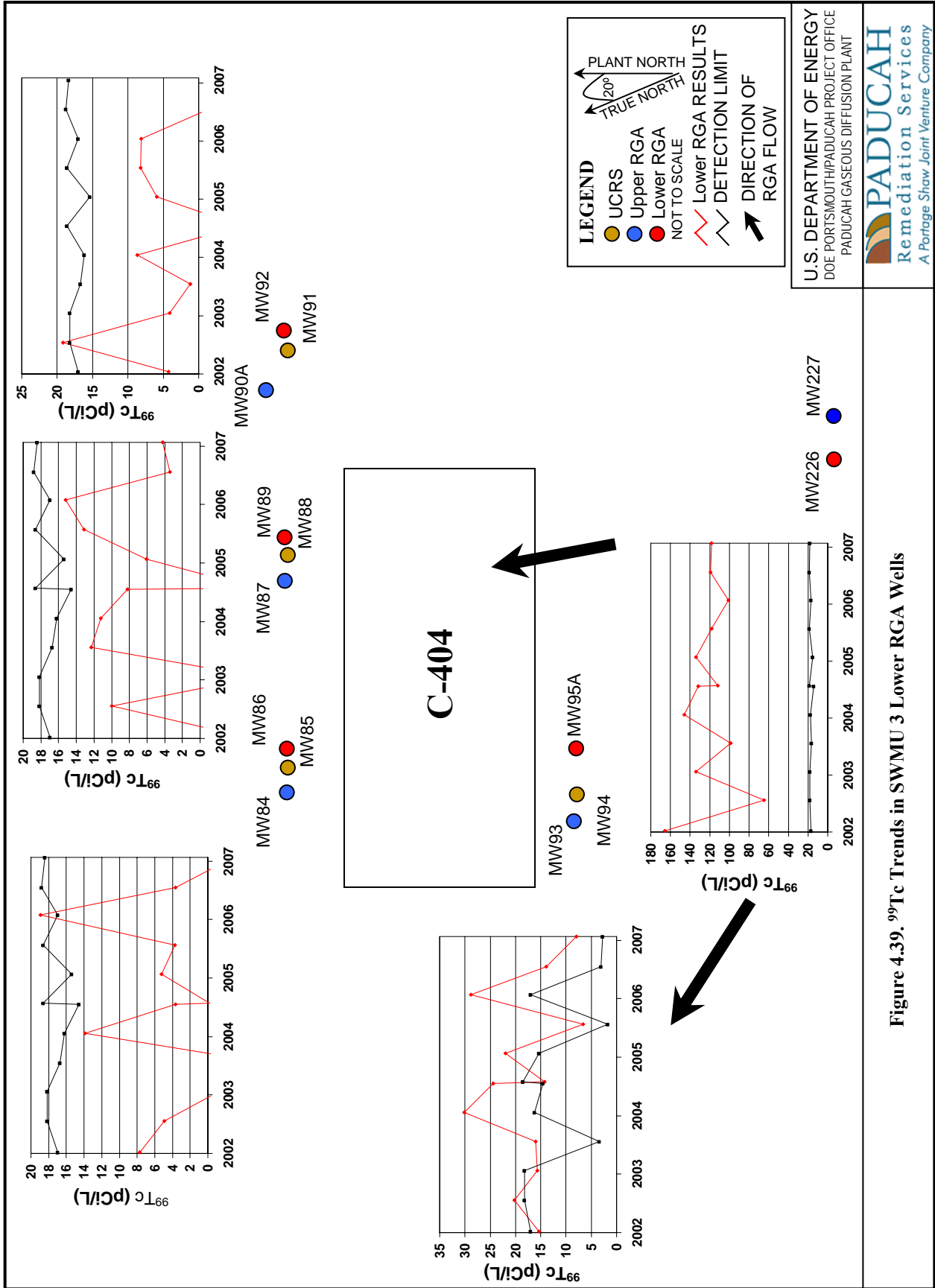
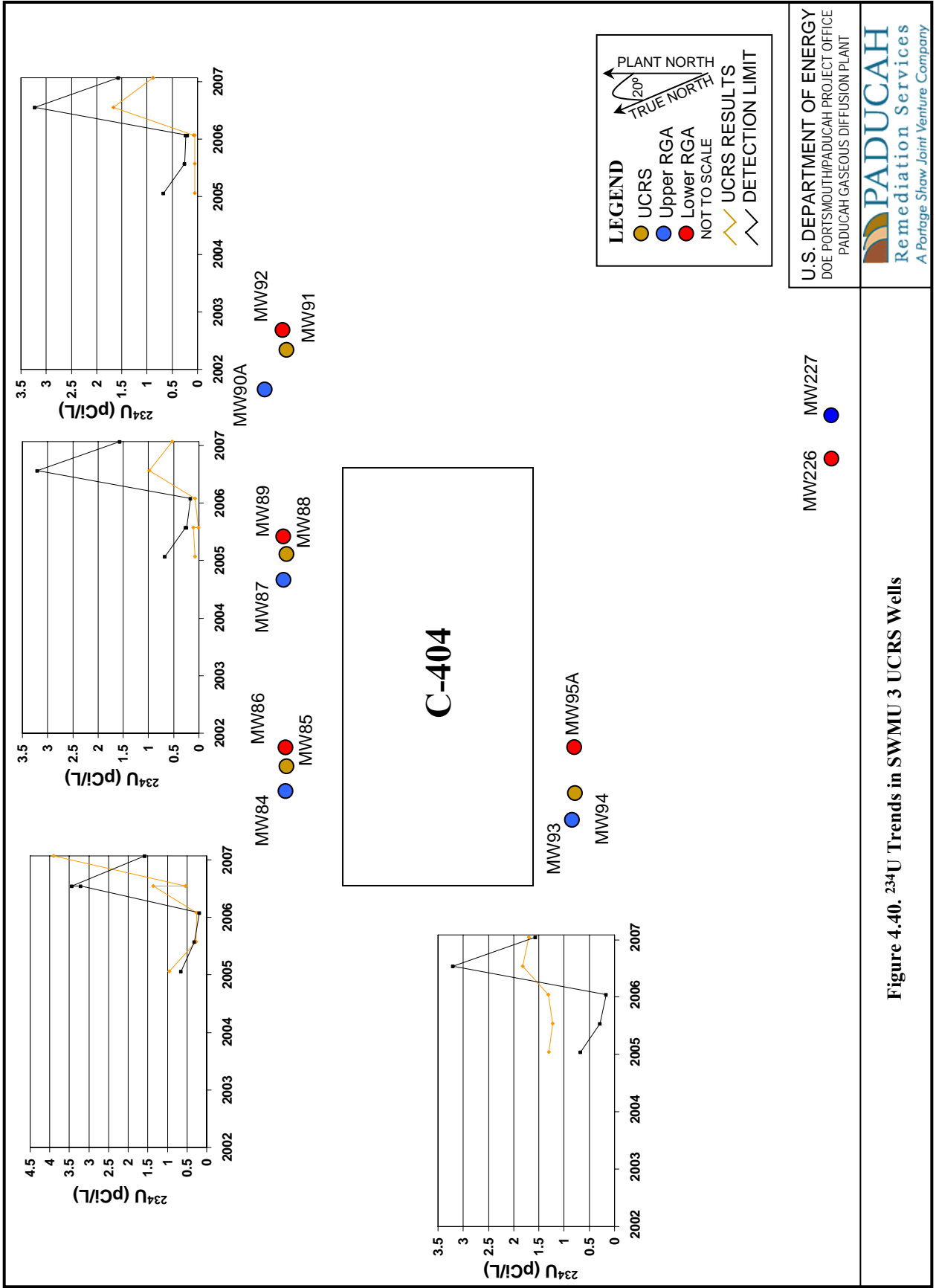


Figure 4.39. ⁹⁹Tc Trends in SWMU 3 Lower RGA Wells



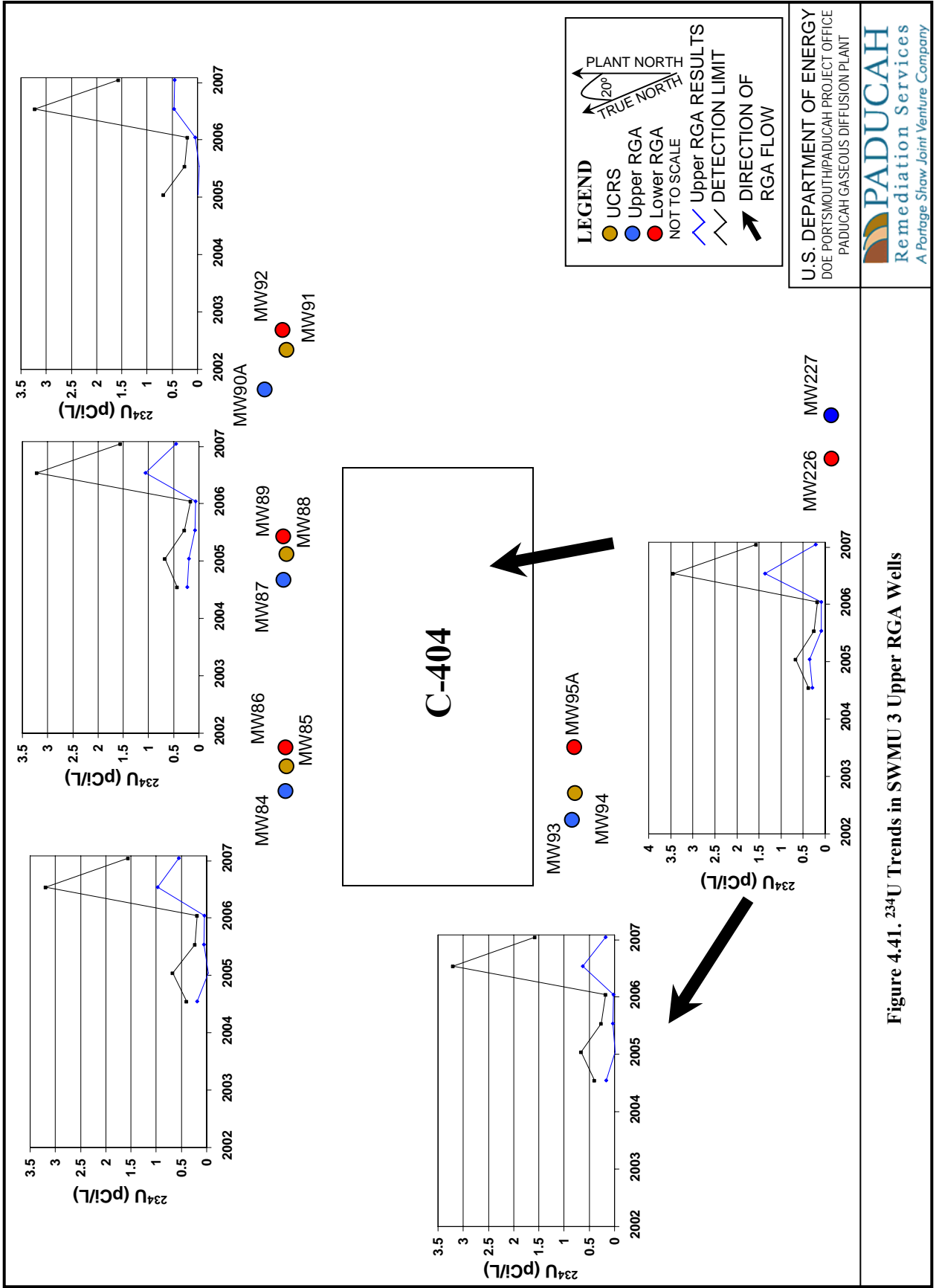


Figure 4.41. ²³⁴U Trends in SWMU 3 Upper RGA Wells

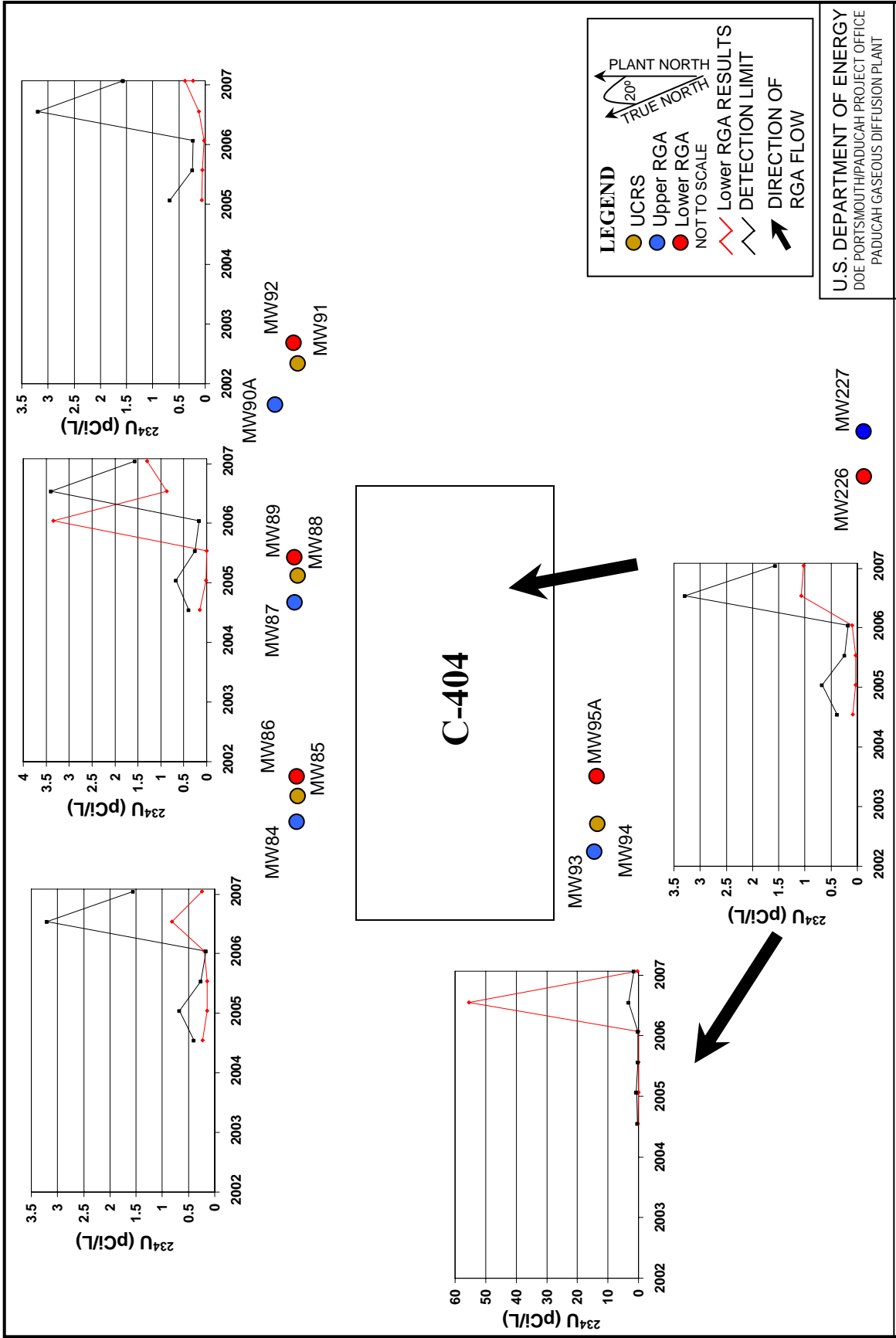
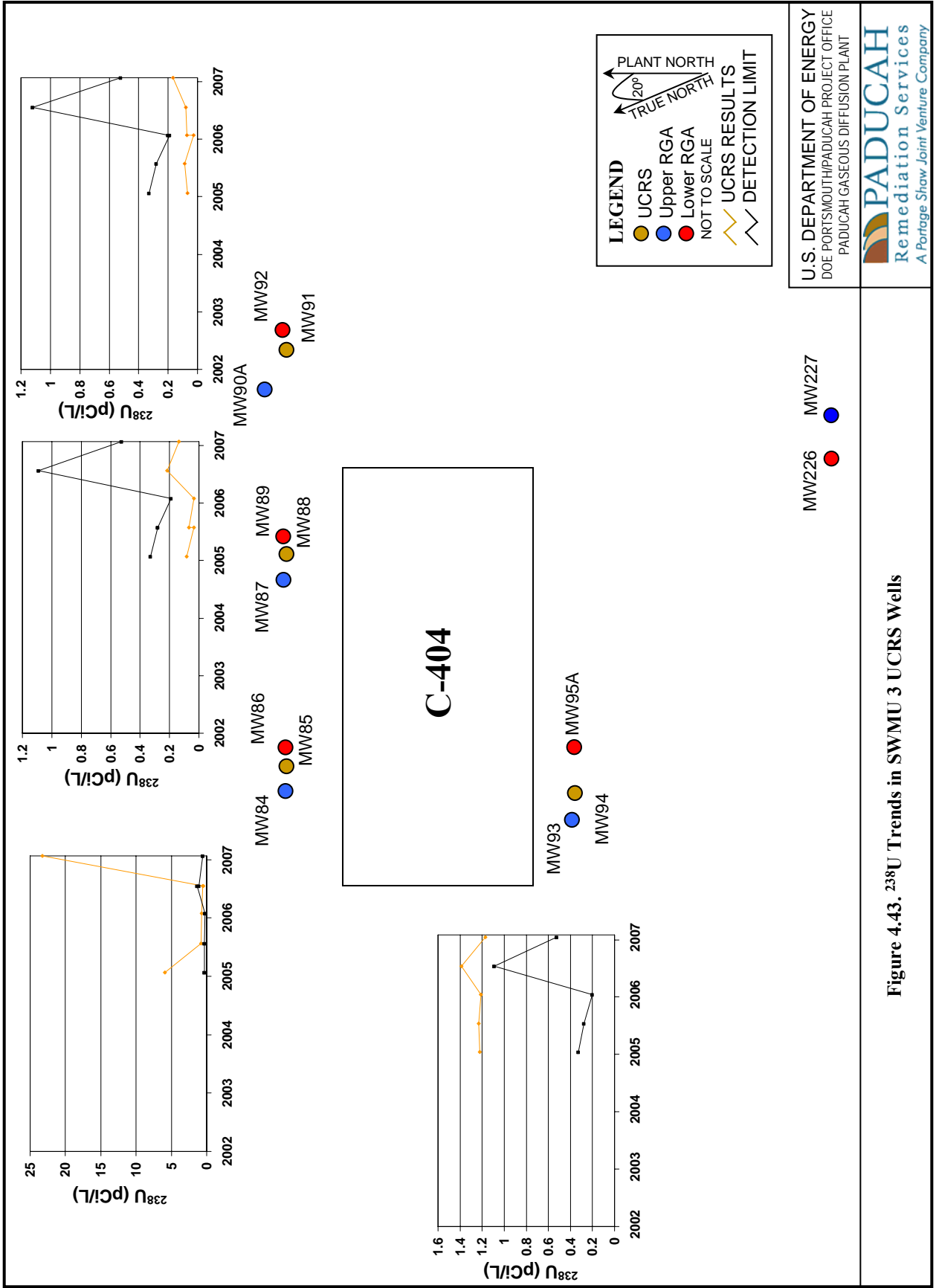


Figure 4.42. ²³⁴U Trends in SWMU 3 Lower RGA Wells



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Figure 4.43. ²³⁸U Trends in SWMU 3 UCRS Wells

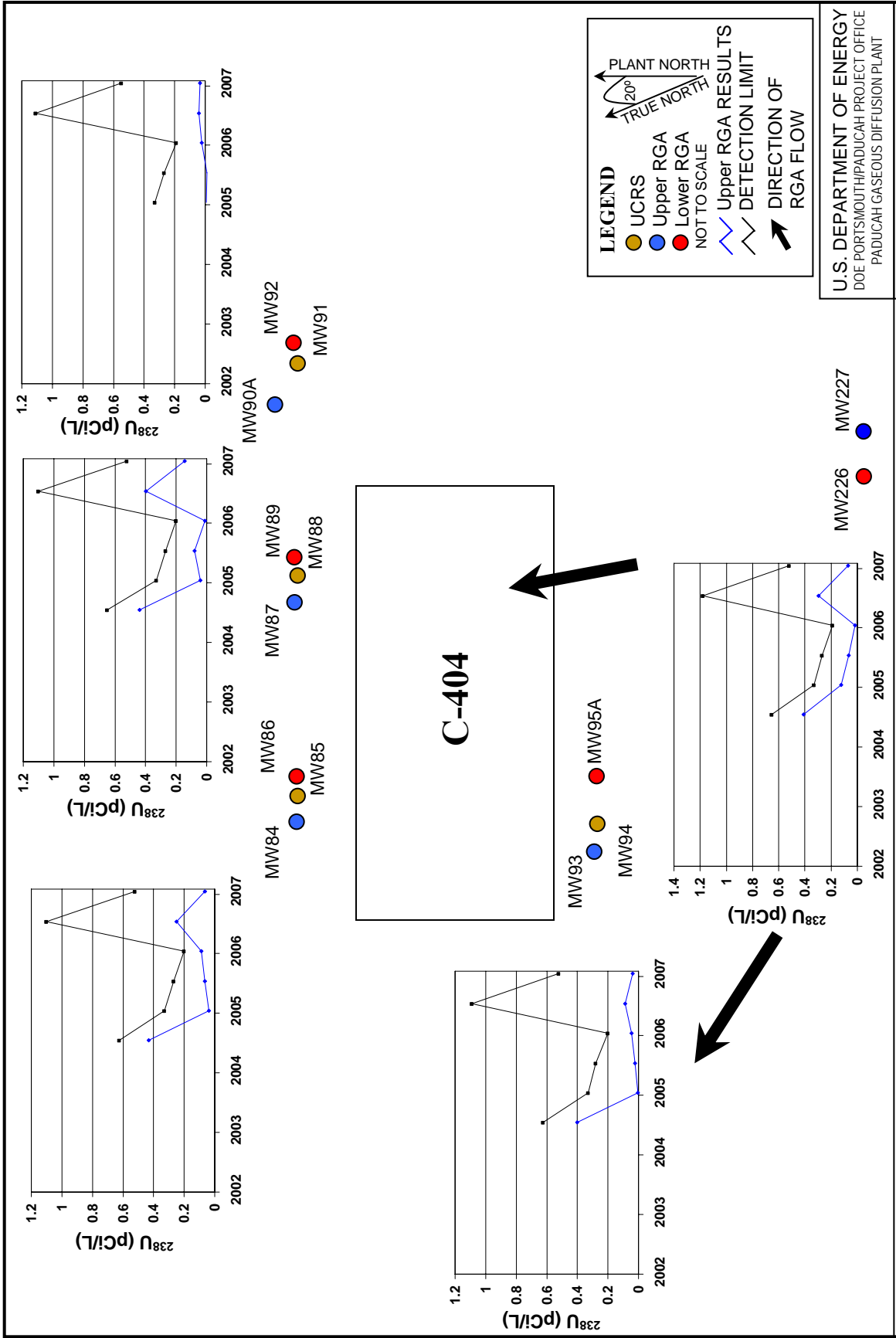


Figure 4.44. ²³⁸U Trends in SWMU 3 Upper RGA Wells

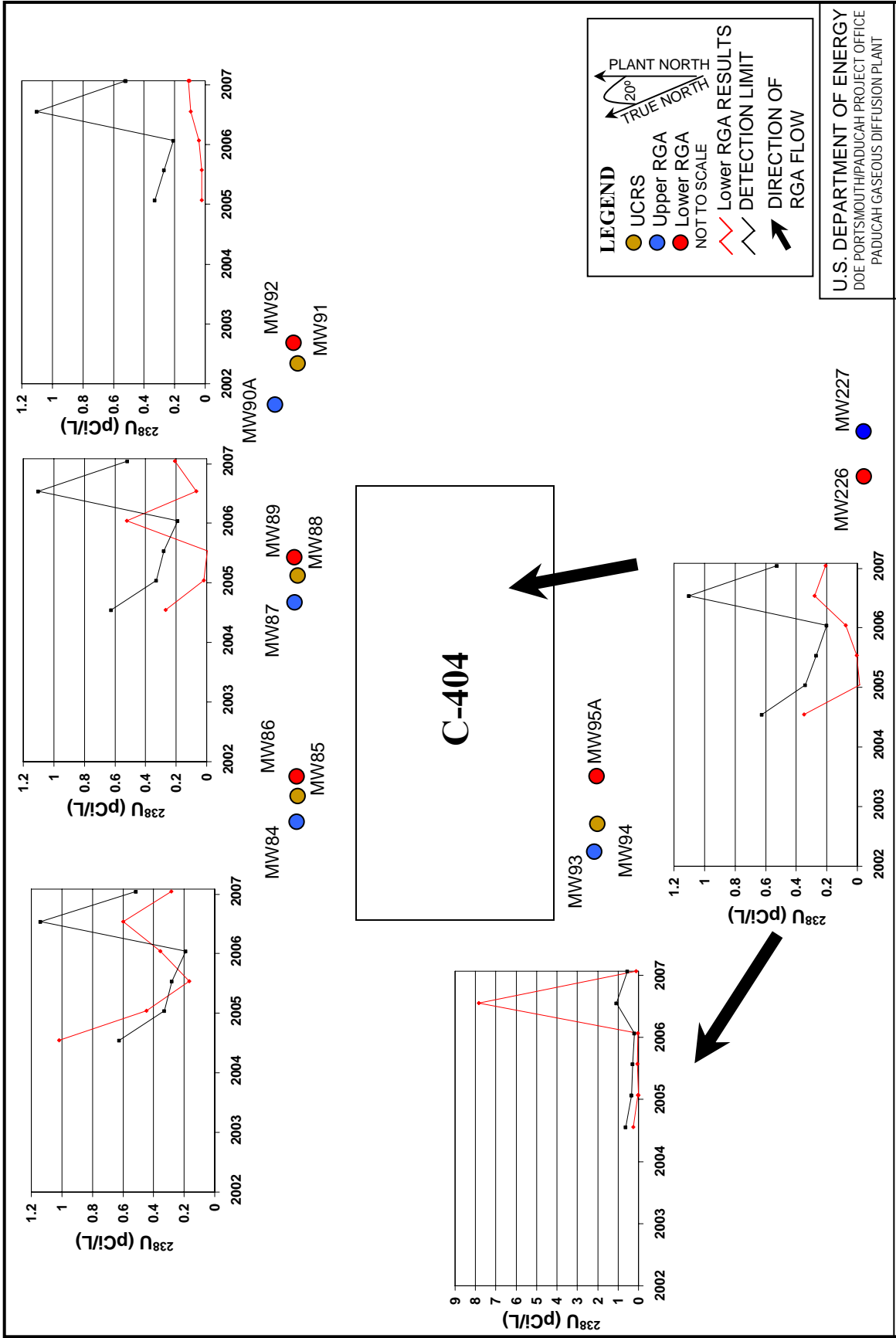
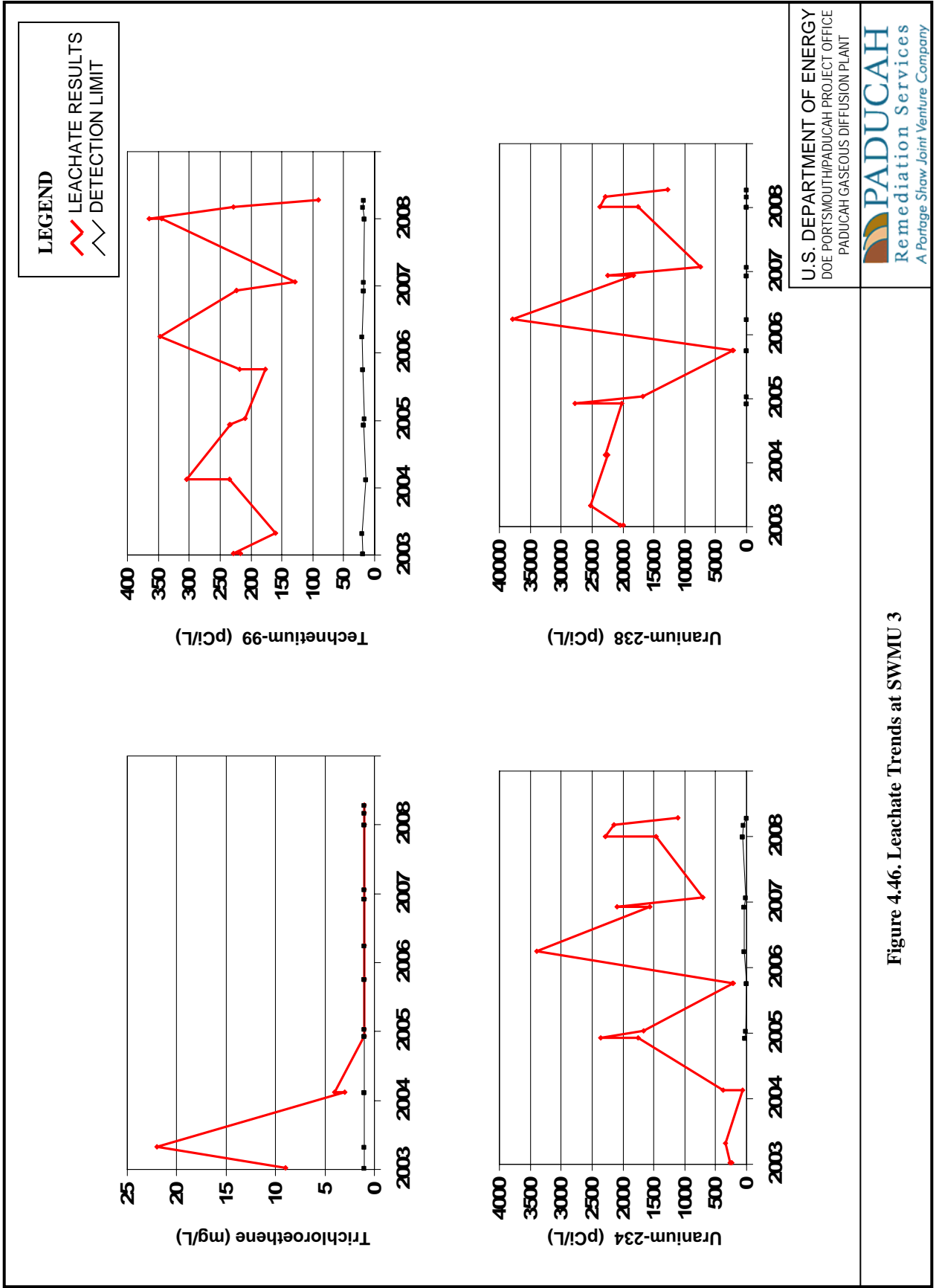


Figure 4.45. ²³⁸U Trends in SWMU 3 Lower RGA Wells



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Figure 4.46. Leachate Trends at SWMU 3

4.5 SWMU 4

4.5.1 Subsurface Soils

Table 4.20 summarizes the review of SWMU 4 subsurface soil data to identify site-related contaminants. Table 4.21 shows the locations and depths of the contaminants that were detected above screening levels (sampling locations are shown in Figure 4.3). During scoping for the BGOU RI/FS Work Plan, it was determined that sufficient data existed to move forward to the FS, so no additional data from SWMU 4 were collected during this RI.

SWMU 4 served as a disposal repository of radiologically contaminated and uncontaminated debris originating from the C-410 UF₆ feed plant. Beryllium is the most widely detected metal in subsurface soils above background (52 of 126 analyses), but exceeds the NAL in only 6 of 126 analyses. Figure 4.47 shows the distribution of beryllium in the subsurface soil at SWMU 4. Most of the higher concentrations (> 1 mg/kg) occur in a horizon at 40 to 55 ft bgs. Iron and vanadium are the most common metals to exceed both PGDP background (in 7 of 126 analyses for both) and the NAL (in 126 of 126 analyses of iron and 125 of 126 analyses of vanadium). Manganese exceeds PGDP background in 6 of 126 analyses and exceeds the NAL in 92 of 126 analyses. The iron and vanadium exceedances are well distributed across SWMU 4. Most of the exceedances occur at depths of 20 to 55 ft. Figure 4.48 provides the vanadium distribution in soil at the SWMU.

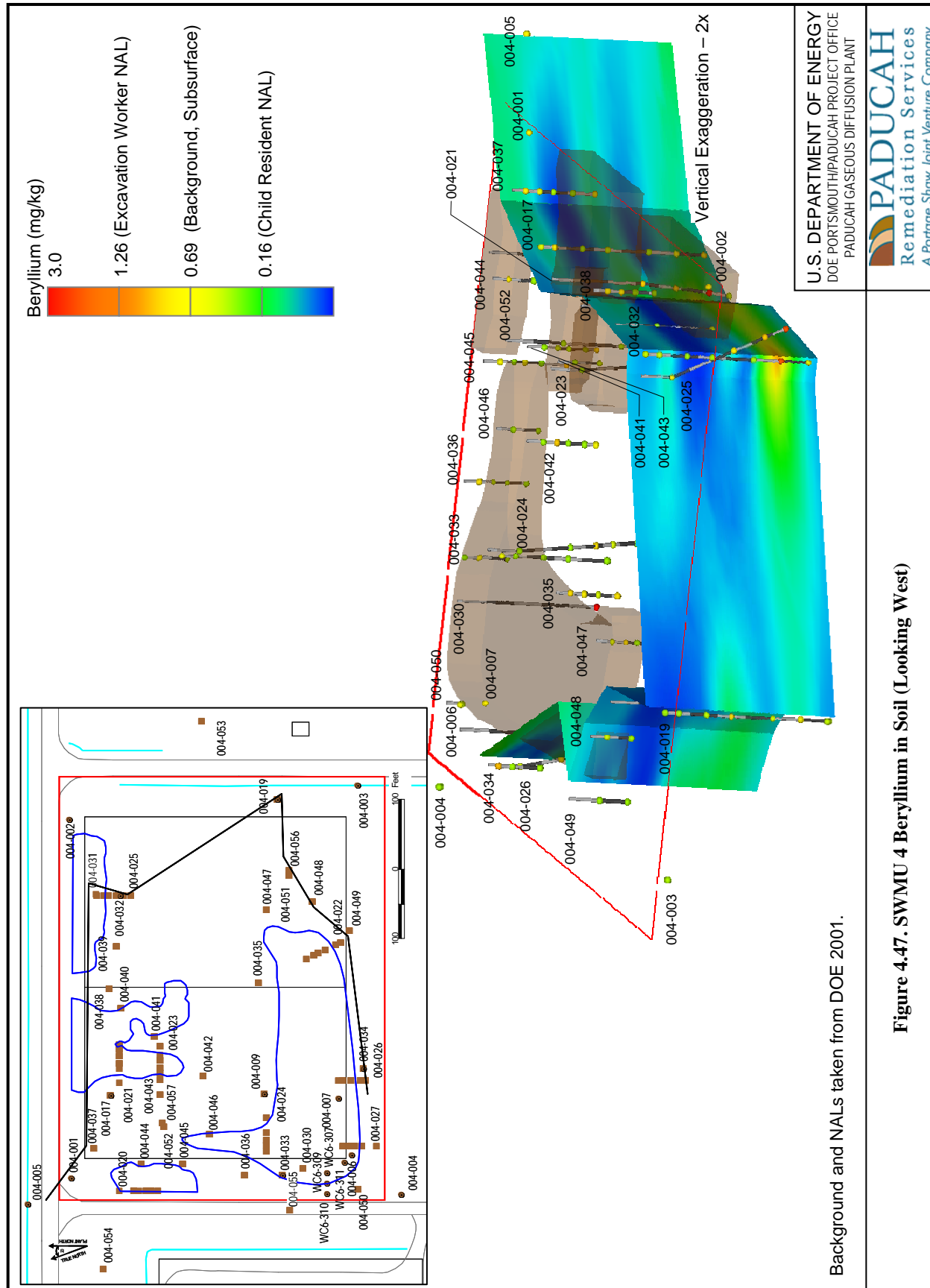
TCE is widely present (47 of 314 analyses) in subsurface samples from borings located within burial pits. Highest levels (up to 41 mg/kg) are commonly found in the soils below the large southern burial pit, with levels as high as 25 mg/kg at the maximum depth of the soil samples (61 ft). Figure 4.49 illustrates the distribution of TCE in soil at the unit and shows the predominant detections in the southern half of the SWMU. A potential DNAPL source is suspected in the UCRS at SWMU 4 near the southern burial pit. Subsurface soil analyses also document the TCE degradation product vinyl chloride above screening levels in 3 of 318 subsurface samples from borings within the area of the large southern burial pit (Figure 4.50). The vinyl chloride may be the result of anaerobic degradation occurring in the UCRS; however, dissolved oxygen data are not available for this SWMU and remains an uncertainty. The highest levels of PCBs cluster around the east end of the southern burial pit (in soils of 6 ft depth or less).

The most common radionuclides with activities that exceed background and the excavation worker NAL are the uranium isotopes uranium-234 and uranium-238. These detections are commonly limited to soils less than 10 ft deep and occur across the site. Figure 4.51 shows the widespread shallow uranium contamination associated with SWMU 4. Uranium levels decrease quickly below a depth of 10 ft.

Table 4.20. SWMU 4 Subsurface Soil Contaminants

Analysis	Maximum Result		Frequency of Detection ^a	Frequency of Detection	
	Historical Data	RI Data		Above Background Value	Above Excavation Worker NAL
<i>Inorganics (mg/kg)</i>					
Aluminum	19,000	N/A	126/126	21/126	119/126
Arsenic	17.1	N/A	12/125	5/125	12/125
Barium	313	N/A	126/126	4/126	1/126
Beryllium	2.02	N/A	85/126	52/126	6/126
Calcium	131,000	N/A	126/126	2/126	N/A
Chromium	60.8	N/A	126/126	1/126	0/126
Cobalt	31.6	N/A	124/126	5/126	0/126
Copper	46.4	N/A	121/126	2/126	0/126
Iron	34,500	N/A	126/126	7/126	126/126
Lead	30.2	N/A	2/126	2/126	0/126
Magnesium	2,650	N/A	125/126	6/126	0/126
Manganese	2,700	N/A	125/126	6/126	92/126
Nickel	53	N/A	84/126	14/126	0/126
Potassium	2,390	N/A	125/126	8/126	N/A
Sodium	3,150	N/A	99/126	49/126	0/126
Vanadium	75.5	N/A	126/126	7/126	125/126
Zinc	79.7	N/A	104/126	9/126	0/126
<i>Organics--Volatiles (mg/kg)</i>					
1,1,2-Trichloroethane	0.021	N/A	1/105	N/A	0/105
1,1-Dichloroethene	0.014	N/A	1/318	N/A	0/318
Acetone	0.012	N/A	1/83	N/A	0/83
Carbon tetrachloride	0.074	N/A	1/105	N/A	0/105
Chloroform	0.012	N/A	1/104	N/A	0/104
<i>cis</i> -1,2-Dichloroethene	9.8	N/A	23/317	N/A	0/317
Methylene chloride	0.054	N/A	18/105	N/A	0/105
<i>trans</i> -1,2-Dichloroethene	0.45	N/A	1/318	N/A	0/318
Trichloroethene	41	N/A	47/314	N/A	9/314
Vinyl chloride	0.29	N/A	7/318	N/A	3/318
<i>Organics--PCBs (mg/kg)</i>					
PCB, Total	27	N/A	10/153	N/A	11/153
PCB-1016	2.5	N/A	1/172	N/A	1/172
PCB-1248	0.8	N/A	2/172	N/A	2/172
PCB-1254	27	N/A	8/172	N/A	7/172
PCB-1260	0.5	N/A	2/172	N/A	2/172
<i>Radionuclides(pCi/g)</i>					
Americium-241	2.8	N/A	1/193	N/A	1/193
Cesium-137	1.48	N/A	2/160	2/160	2/160
Neptunium-237	5.78	N/A	10/35	N/A	7/35
Plutonium-239/240	4.17	N/A	8/36	N/A	1/36
Protactinium-234m	380	N/A	2/158	N/A	N/A
Radium-226	2.51	N/A	17/35	8/35	17/35
Technetium-99	269	N/A	13/182	13/182	2/182
Thorium-230	68.7	N/A	2/2	2/2	1/2
Thorium-234	158	N/A	27/193	N/A	N/A
Uranium	6,260	N/A	16/24	N/A	N/A
Uranium-234	69	N/A	15/23	14/23	13/23
Uranium-235	4.2	N/A	1/158	1/158	1/158

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Figure 4.47. SWMU 4 Beryllium in Soil (Looking West)

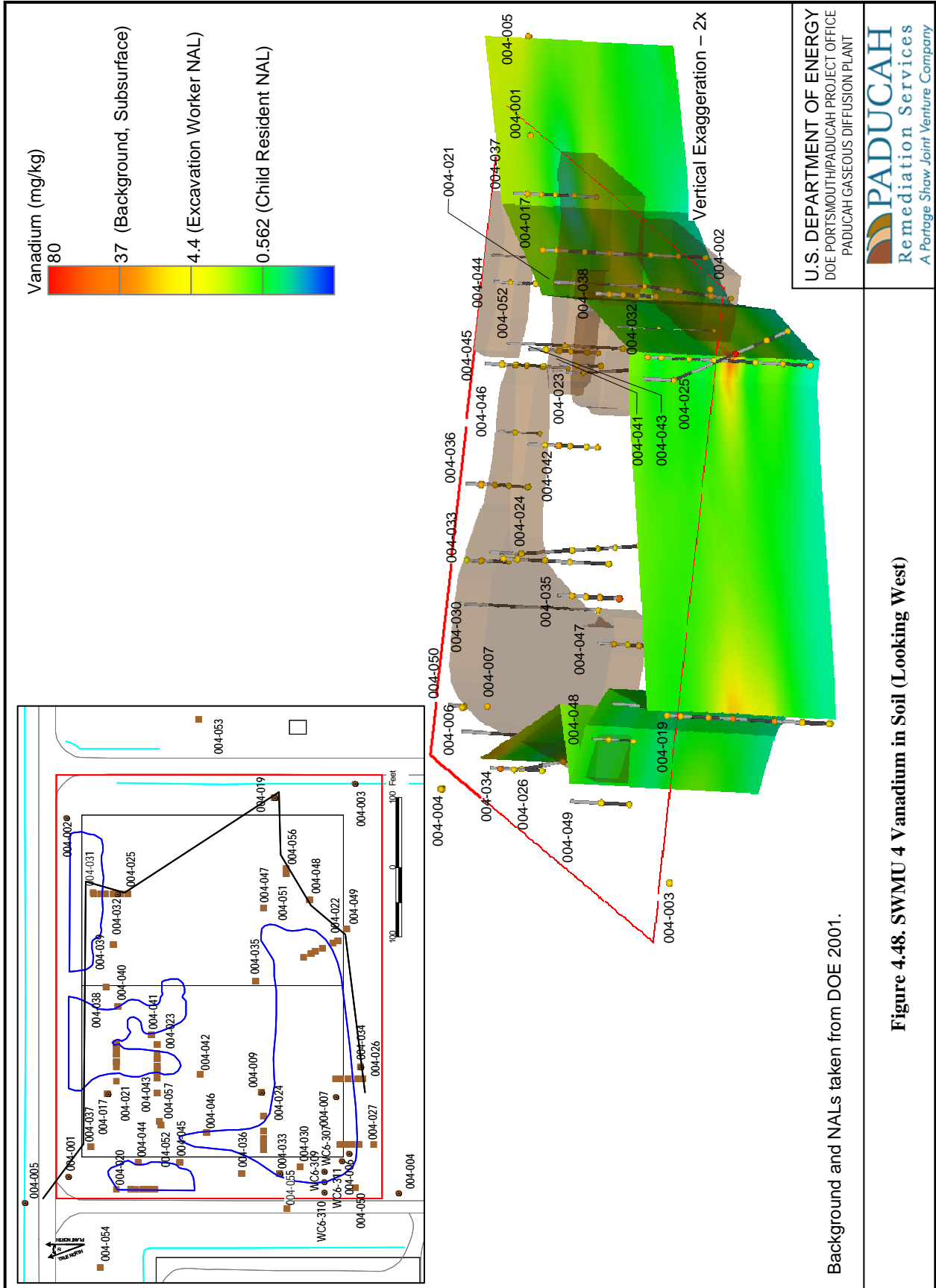
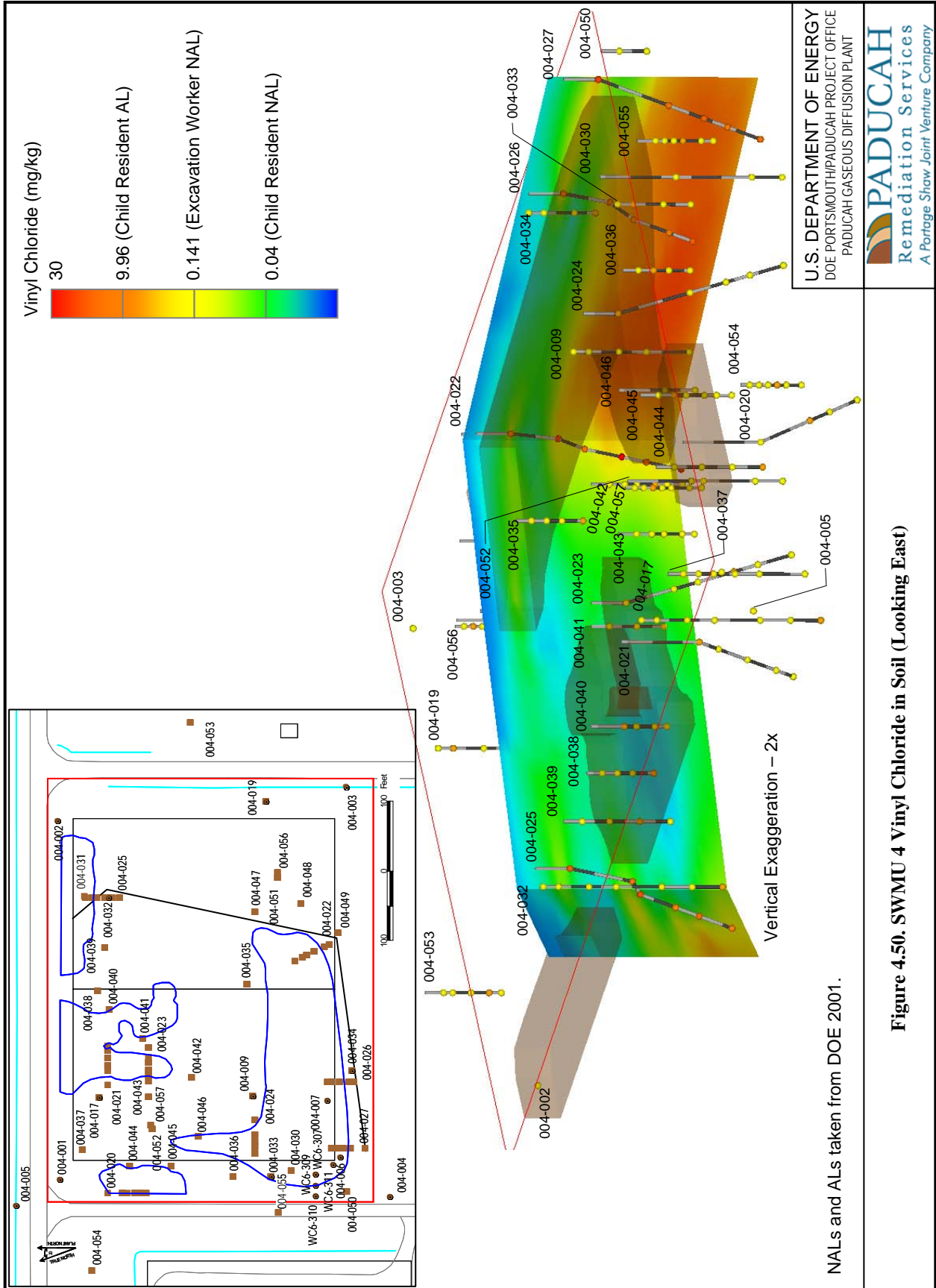


Figure 4.48. SWMU 4 Vanadium in Soil (Looking West)

Background and NALs taken from DOE 2001.



NALs and ALs taken from DOE 2001.

Figure 4.50. SWMU 4 Vinyl Chloride in Soil (Looking East)

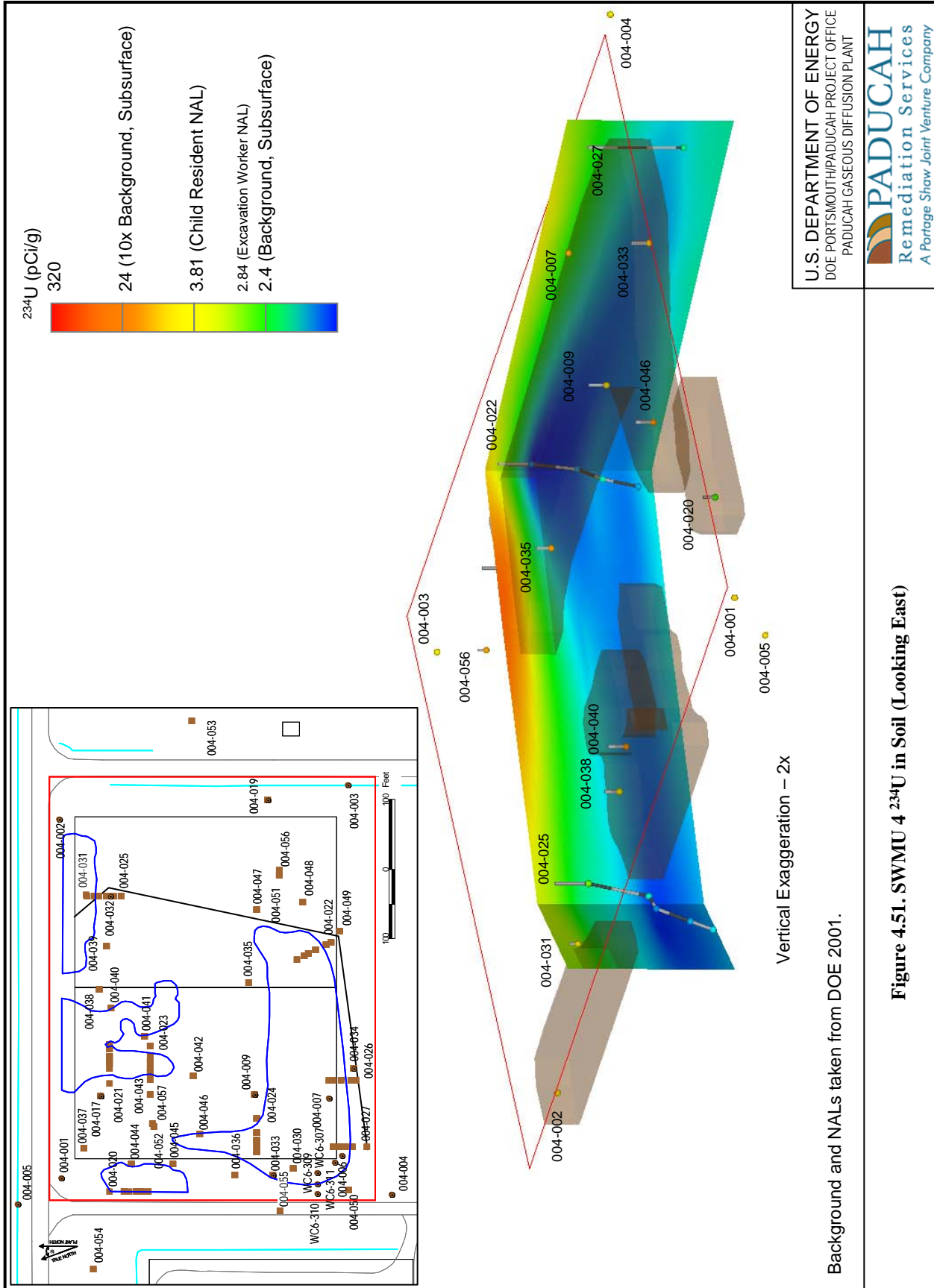


Figure 4.51. SWMU 4 ^{234}U in Soil (Looking East)

4.5.2 SWMU 4 Groundwater

No additional sampling was conducted at SWMU 4 as part of this RI. The WAG 3 RI (DOE 2000a) provided the majority of data to characterize SWMU 4. Single temporary borings of the WAG 27 RI (DOE 1999d) and a sitewide remedial evaluation for source areas (DOE 2000b) supplied additional RGA data for the perimeter of the SWMU 4 area. The WAG 3 RI (DOE 2000a) provided analyses of UCRS groundwater from 26 temporary borings (Figure 4.11), both angled below and located along the perimeter of the burial cells, shown in Figure 4.3.

Three temporary soil borings of the WAG 3 RI (004-028, 004-029, and 004-058) and single temporary soil borings from the WAG 27 RI, 720-026 (DOE 1999d) and from a sitewide remedial evaluation for source areas, DG-030 (DOE 2000b) provided groundwater analyses to characterize the RGA. Tables 4.22 and 4.23 summarize the review of SWMU 4 UCRS and RGA groundwater data (primarily derived from the WAG 3 RI) to identify site-related contaminants. All RGA soil borings at SWMU 4, with the exception of 720-026, also sampled the McNairy. Groundwater samples at SWMU 4 characterized groundwater down to 50 ft below the base of the RGA. The screening steps determined that the only contaminant among the McNairy groundwater samples from SWMU 4 was TCE. Table 4.24 provides detail (depth, sample location, and analytical results) for SWMU 4 groundwater samples, including nondetects and detections above screening levels.

The metals arsenic, iron, lead, and manganese frequently exceeded screening levels in both the UCRS and RGA. The WAG 3 investigation did not collect pH for UCRS groundwater samples so the potential for acidic leachate is an uncertainty. VOCs also were common contaminants of the UCRS and RGA associated with SWMU 4. TCE levels exceeded the MCL in 43 of 45 analyses in the RGA. TCE degradation products, notably 1,1-DCE and *cis*-1,2-DCE, also frequently exceeded MCLs. Other VOCs present in the RGA at SWMU 4 include carbon tetrachloride and chloroform.

Dissolved TCE trends indicate that a potential TCE DNAPL source is present in the UCRS at SWMU 4, related to the elevated soil concentrations found in the southern burial area. A 2004 SI of Southwest Plume sources identified markedly higher dissolved TCE and ⁹⁹Tc levels in the RGA on the west (downgradient) side of SWMU 4. (See Figure 4.20.) Highest groundwater contaminant levels in the 4 temporary soil borings on the east side of SWMU 4 (all at the base of the RGA) (borings 004-101 to 004-104) ranged from 140 to 680 µg/L TCE and below the method detection limit to 93.2 pCi/L ⁹⁹Tc. The SI sampled 5 temporary soil borings on the west side of SWMU 4 (borings 004-105 through 004-109). Groundwater samples from the 4 northernmost borings exhibited a pronounced increase in contaminant levels over those observed in samples from the east side. Highest groundwater contaminant levels (all in the upper RGA) ranged from 1,200 to 3,980 µg/L TCE and 81 to 663 pCi/L ⁹⁹Tc (see Figure 3.29). The evidence of the potential UCRS DNAPL presence is markedly higher dissolved TCE levels in the RGA on the west (downgradient) side of the SWMU. The area of higher TCE levels spans the entire west side of SWMU 4, suggestive of a diffuse source of DNAPL contamination in the UCRS soils underlying the burial grounds (DOE 2007a). The volume of soil potentially contaminated with TCE DNAPL at this SWMU is estimated to be approximately 31,480 yd³. This estimate (based on the cross sectional width of the south burial pit of SWMU 4 and fate and transport modeling results) assumes a source area that is 100 ft by 100 ft with a thickness of 85 ft (depth to base of RGA, which is 100 ft minus the estimated depth to base of the waste cell of 15 ft). Note that the location and extent of the DNAPL zone is poorly constrained. The volumetric extent of this DNAPL zone will be estimated for alternatives evaluation in the FS.

Table 4.22. SWMU 4 UCRS Groundwater Contaminants

Analysis	Historical Data	RI	Frequency of Detection ^a	Frequency of Detection		
				Above Background	Above NAL	Above MCL
Metals (mg/L)						
Aluminum	1,270	N/A	23/38	N/A	16/38	N/A
Arsenic	0.311	N/A	23/27	N/A	23/27	21/27
Barium	13.6	N/A	37/38	N/A	29/38	9/38
Beryllium	0.13	N/A	12/34	N/A	12/34	12/34
Cadmium	0.031	N/A	7/23	N/A	7/23	7/23
Calcium	226	N/A	38/38	N/A	N/A	N/A
Chromium	5.11	N/A	15/38	N/A	6/38	15/38
Cobalt	3.58	N/A	37/38	N/A	11/38	N/A
Copper	1.55	N/A	14/36	N/A	14/36	1/36
Iron	2,560	N/A	33/38	N/A	31/38	N/A
Lead	1	N/A	9/9	N/A	9/9	9/9
Magnesium	102	N/A	38/38	N/A	0/38	N/A
Manganese	118	N/A	38/38	N/A	38/38	N/A
Mercury	0.004	N/A	7/31	N/A	7/31	1/31
Nickel	1.26	N/A	14/34	N/A	14/34	N/A
Potassium	63.1	N/A	23/38	N/A	N/A	N/A
Selenium	0.015	N/A	3/19	N/A	3/19	0/19
Sodium	312	N/A	38/38	N/A	0/38	N/A
Vanadium	4.01	N/A	13/36	N/A	13/36	N/A
Zinc	8.2	N/A	24/38	N/A	15/38	N/A
PCBs (mg/L)						
PCB, Total	0.00091	N/A	2/5	N/A	3/5	1/5
PCB-1248	0.00019	N/A	1/5	N/A	1/5	0/5
PCB-1254	0.00091	N/A	2/5	N/A	2/5	1/5
Radionuclides (pCi/L)						
Technetium-99	1,640	N/A	17/26	N/A	17/26	2/26
Volatiles (mg/L)						
Bis(2-ethylhexyl)phthalate	0.012	N/A	2/18	N/A	2/18	2/18
Diethyl phthalate	0.03	N/A	2/17	N/A	0/17	N/A
Naphthalene	0.007	N/A	1/17	N/A	1/17	N/A
1,1-Dichloroethene	0.34	N/A	11/23	N/A	11/23	4/23
2-Butanone	0.031	N/A	1/11	N/A	0/11	N/A
Acetone	10	N/A	5/9	N/A	3/9	N/A
Chloroethane	0.019	N/A	2/7	N/A	2/7	N/A
cis -1,2-Dichloroethene	12	N/A	22/33	N/A	20/33	14/33
trans -1,2-Dichloroethene	0.11	N/A	13/31	N/A	5/31	1/31
Trichloroethene	56	N/A	29/34	N/A	25/34	24/34
Vinyl chloride	0.44	N/A	12/26	N/A	12/26	9/26

^a Frequency of detection is the number of detections of an analyte per number of analyses (includes regular and duplicate samples).

N/A = not applicable

Table 4.23. SWMU 4 RGA Groundwater Contaminants

Analysis	Historical Data	RI	Frequency of Detection ^a	Frequency of Detection		
				Above Background	Above NAL	Above MCL
Metals (mg/L)						
Aluminum	854	N/A	49/108	32/108	39/108	N/A
Aluminum, Dissolved	0.0265	N/A	2/4	0/4	0/4	N/A
Antimony	0.00582	N/A	1/4	0/4	1/4	0/4
Arsenic	0.045	N/A	51/57	50/57	51/57	26/57
Barium	10.3	N/A	103/107	31/107	81/107	2/107
Barium, Dissolved	0.409	N/A	3/3	3/3	3/3	0/3
Beryllium	0.15	N/A	15/62	11/62	11/62	11/62
Cadmium	0.007	N/A	6/42	0/42	4/42	1/42
Cadmium, Dissolved	0.00043	N/A	4/4	0/4	0/4	0/4
Calcium	154	N/A	107/107	3/107	N/A	N/A
Calcium, Dissolved	32.6	N/A	3/3	0/3	N/A	N/A
Chromium	1.52	N/A	10/51	4/51	0/51	4/51
Chromium, Dissolved	0.0058	N/A	3/3	0/3	0/3	0/3
Cobalt	2.86	N/A	84/102	22/102	9/102	N/A
Cobalt, Dissolved	0.0065	N/A	4/4	0/4	0/4	N/A
Copper	0.59	N/A	8/46	4/46	4/46	0/46
Copper, Dissolved	0.0025	N/A	4/4	0/4	0/4	0/4
Iron	1830	N/A	86/108	41/108	72/108	N/A
Iron, Dissolved	1.22	N/A	4/4	4/4	4/4	N/A
Lead	0.328	N/A	7/8	3/8	4/8	4/8
Lithium	0.148	N/A	2/15	N/A	2/15	N/A
Magnesium	149	N/A	107/107	6/107	0/107	N/A
Magnesium, Dissolved	13.1	N/A	3/3	0/3	0/3	N/A
Manganese	56.2	N/A	107/107	106/107	107/107	N/A
Manganese, Dissolved	1.56	N/A	3/3	3/3	3/3	N/A
Mercury	0.0064	N/A	4/46	3/46	2/46	2/46
Nickel	0.9	N/A	15/61	1/61	14/61	N/A
Nickel, Dissolved	0.0264	N/A	4/4	0/4	0/4	N/A
Potassium	90.8	N/A	20/70	7/70	N/A	N/A
Sodium	28.5	N/A	101/102	0/102	0/102	N/A
Strontium	0.639	N/A	19/29	N/A	0/29	N/A
Vanadium	4.01	N/A	12/56	8/56	8/56	N/A
Vanadium, Dissolved	0.00276	N/A	3/4	0/4	0/4	N/A
Zinc	3.54	N/A	22/72	21/72	7/72	N/A
Zinc, Dissolved	0.0145	N/A	4/4	0/4	0/4	N/A
Radionuclides (pCi/L)						
Technetium-99	663	N/A	43/60	35/60	42/60	0/60
Volatiles (mg/L)						
1,1-Dichloroethane	0.017	N/A	2/24	N/A	0/24	N/A
1,1-Dichloroethene	0.042	N/A	34/60	N/A	34/60	19/60
1,2-Dichloroethane	0.2	N/A	1/24	N/A	1/24	1/24
2-Butanone	0.031	N/A	1/24	N/A	0/24	N/A
2-Propanol	0.54	N/A	1/2	N/A	0/2	N/A
Bromomethane	0.0041	N/A	1/21	N/A	1/21	N/A
Carbon tetrachloride	0.12	N/A	10/24	N/A	10/24	8/24
Chloroform	0.13	N/A	12/23	N/A	12/23	N/A
cis-1,2-Dichloroethene	0.2	N/A	45/61	N/A	37/61	18/61
Methylene chloride	0.59	N/A	2/24	N/A	2/24	2/24
Tetrachloroethene	0.004	N/A	1/24	N/A	1/24	0/24
trans-1,2-Dichloroethene	0.018	N/A	25/64	N/A	15/64	0/64
Trichloroethene	10	N/A	78/78	N/A	76/78	76/78
Vinyl chloride	0.017	N/A	20/57	N/A	20/57	3/57
Wetchem Parameters (mg/L)						
Cyanide	0.04	N/A	3/39	N/A	3/39	0/39

^a Frequency of detection is the number of detections of an analyte per number of analyses (includes regular and duplicate samples).

N/A = not applicable

Table 4.24. SWMU 4 Locations of Groundwater Contaminants (Continued)

Hydro Unit	Depth Interval	Analysis	Child Resident NAL	Bkgd	MCL	004-008	004-009	004-011	004-017	004-019	004-020	004-021	004-022	004-023	004-024	004-025	004-026	004-027	004-028	004-029	004-032	004-033	004-035	004-036	004-037	004-038	004-039	004-040	004-044	004-047	004-049	004-058	720-026	DG-030	004-107	004-108	004-109	004-110												
	92-93	trans-1,2-Dichloroethene	0.00548	N/A	0.1																																													
		Trichloroethene	0.0016	N/A	0.005																																													
		Vinyl chloride	0.000035	N/A	0.002																																													
		Wetchem (mg/L)																																																
		Cyanide	0.0284	N/A	0.2																																													
	98	Metals (mg/L)																																																
		Aluminum	1.49	2.189	N/A																																													
		Arsenic	0.000035	0.005	0.01																																													
		Barium	0.104	0.235	2																																													
		Beryllium	0.00264	0.004	0.004																																													
		Cadmium	0.000661	0.01	0.005																																													
		Calcium	N/A	41.238	N/A																																													
		Chromium	1.76	0.144	0.1																																													
		Cobalt	0.0906	0.045	N/A																																													
		Copper	0.0557	0.036	1.3																																													
		Iron	0.449	5.03	N/A																																													
		Lead	0.015	0.129	0.015																																													
		Lithium	0.0302	N/A	N/A																																													
		Magnesium	N/A	16.262	N/A																																													
		Manganese	0.035	0.119	N/A																																													
		Mercury	0.000444	0.002	0.002																																													
		Nickel	0.0301	0.682	N/A																																													
		Potassium	N/A	5.195	N/A																																													
		Sodium	N/A	59.45	N/A																																													
		Strontium	0.901	N/A	N/A																																													
		Vanadium	0.00925	0.134	N/A																																													
		Zinc	0.45	0.054	N/A																																													
		Radionuclides (pCi/L)																																																
		Technetium-99	14	22.3	900																																													
		Volatiles (mg/L)																																																
		1,1-Dichloroethane	0.0363	N/A	N/A																																													
		1,1-Dichloroethene	0.00047	N/A	0.007																																													
		1,2-Dichloroethane	0.000147	N/A	0.005																																													
		2-Butanone	0.0868	N/A	N/A																																													
		Bromomethane	0.000391	N/A	N/A																																													
		Carbon tetrachloride	0.000181	N/A	0.005																																													
		Chloroform	2.87E-05	N/A	N/A																																													
		cis-1,2-Dichloroethene	0.00273	N/A	0.07																																													
		Methylene chloride	0.00426	N/A	0.005																																													
		Tetrachloroethene	0.000582	N/A	0.005																																													
		trans-1,2-Dichloroethene	0.00548	N/A	0.1																																													
		Trichloroethene	0.0016	N/A	0.005																																													
		Vinyl chloride	0.000035	N/A	0.002																																													
		Wetchem (mg/L)																																																
		Cyanide	0.0284	N/A	0.2			</																																										

Table 4.24. SWMU 4 Locations of Groundwater Contaminants (Continued)

Blank cells indicate interval was not sampled for the specified analysis. Maximum value shown for each depth interval.

N/A = not applicable

Bold indicates result is greater than NAL value.

Italics indicates result is greater than MCL value.

Bold + Italics indicate result is greater than both NAL and MCL values.

Shading indicates result is greater than the background value.

Figure 4.20 (from the Southwest Plume SI report) interprets a discrete area with TCE concentrations greater than 10,000 µg/L in the RGA immediately downgradient of SWMU 4, derived from the DNAPL zone. Identification of this area of elevated dissolved TCE is primarily based on a conceptual model of dissolved TCE levels derived from a DNAPL zone. The only confirmatory analysis represents a single groundwater sample from historic temporary boring DG-030.

4.6 SWMU 5

4.6.1 Subsurface Soils

This RI collected subsurface soil samples from three angled borings at SWMU 5 (Figure 4.4). Review of RI and historical data for SWMU 5 identified the contaminants listed in Table 4.25.

The SWMU 5 burial pits were not excavated below 10 to 15 ft due to the shallow water table. Metals and radionuclides are the primary potential contaminants of interest at SWMU 5, since the majority of items believed to be buried there include some radionuclide-contaminated scrap metal and slag from PGDP nickel and aluminum smelters. The most prevalent metal detected in subsurface soils above background levels is beryllium (26 of 59 analyses), followed by iron and vanadium (4 of 59 analyses). The metals exceedances are well distributed across SWMU 5. High levels of vanadium tended to occur at moderate depths (15 to 30 ft), while beryllium exceedances mostly are at depths of 40 ft or greater. Figure 4.52 provides the distribution of beryllium at SWMU 5. The highest beryllium concentration was 2.59 mg/kg from the 40 to 45 ft sample at historical soil boring 005-022, with the next sample at 50 to 55 ft showing only 0.57 mg/kg. Figure 4.53 shows the distribution of vanadium at SWMU 5 and illustrates the horizon at 15 to 30 ft where most exceedances occur. High levels of iron range across depths of 20 to 55 ft. Table 4.26 shows the locations of detections above screening levels. Uranium-238 is the only radionuclide identified during screening exceeding both the NAL and background. The screening process did not identify any organic compounds as potential contaminants for SWMU 5. Based on review of the available inventories for SWMU 5, scoping for the BGOU RI/FS Work Plan (DOE 2006a; DOE 2006b) determined that organic contaminants likely were not associated with the buried materials. As such, soils collected from angled borings at SWMU 5 were not evaluated for VOCs or SVOCs.

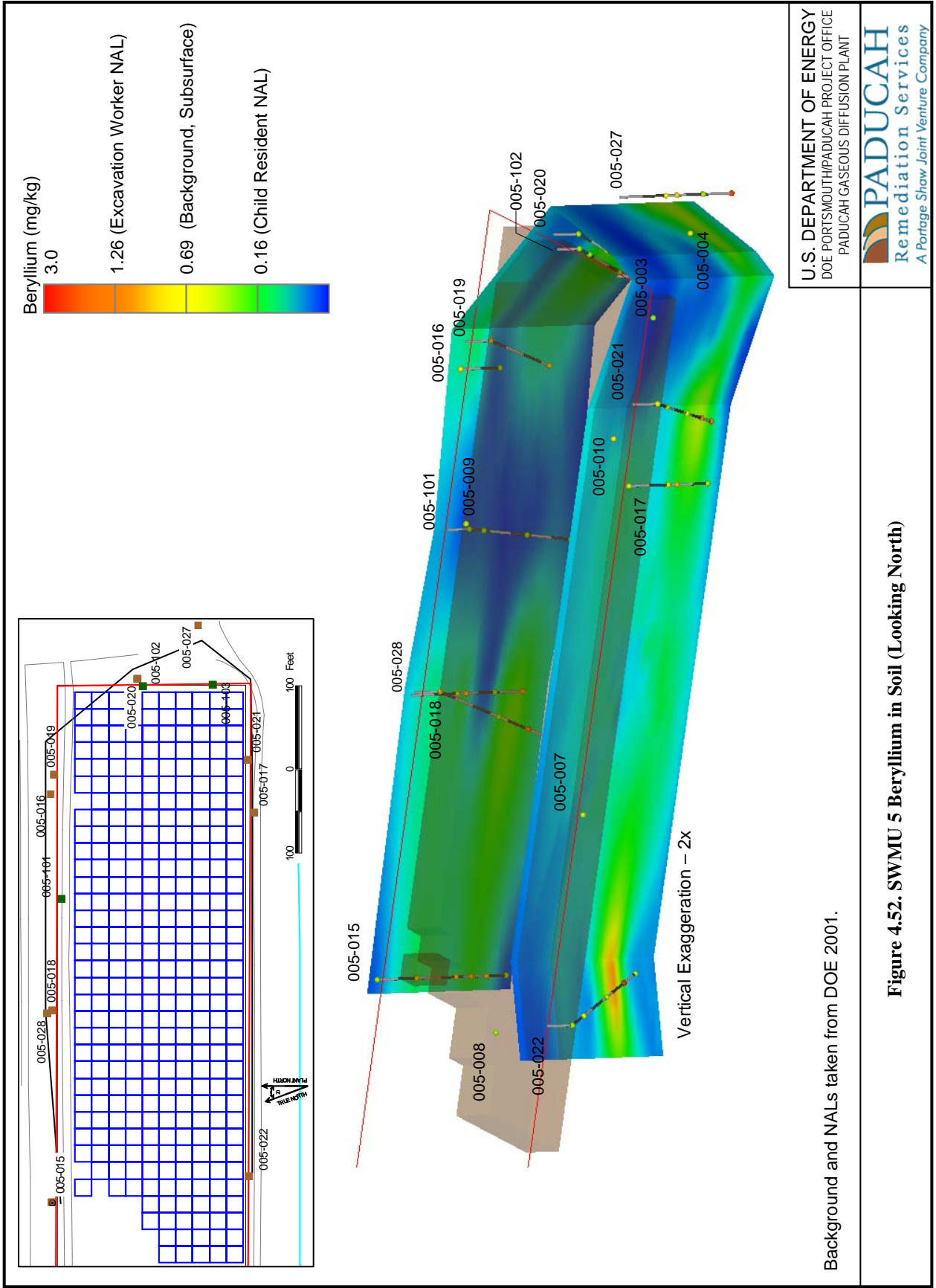
Table 4.25. SWMU 5 Subsurface Soil Contaminants

Analysis	Maximum Result		Frequency of Detection ^a	Frequency of Detection	
	Historical Data	RI Data		Above Background Value	Above Excavation Worker NAL
<i>Inorganics (mg/kg)</i>					
Aluminum	16,400	9,950	59/59	5/59	49/59
Arsenic	6.77	4.21	17/59	0/59	17/59
Barium	343	141	59/59	2/59	1/59
Beryllium	2.59	ND	31/59	26/59	8/59
Calcium	2,050	1,330	59/59	0/59	N/A
Chromium	296	29.3	58/59	2/59	0/59
Cobalt	19.4	28.5	45/59	5/59	0/59
Copper	14.4	8.09	57/59	0/59	0/59
Iron	32,900	21,800	59/59	4/59	57/59
Lead	ND	11.3	17/59	0/59	0/59
Magnesium	1,930	1,180	59/59	0/59	0/59
Manganese	1,750	690	59/59	2/59	44/59
Mercury	ND	0.036	1/59	0/59	0/59
Nickel	38	13.7	44/59	3/59	0/59
Potassium	1,890	N/A	42/42	6/42	N/A
Silver	5.14	ND	1/47	1/47	0/47
Sodium	389	183	26/59	2/59	0/59
Uranium	N/A	1.24	4/17	0/17	0/17
Vanadium	56.9	34.3	59/59	4/59	59/59
Zinc	79.9	39	35/59	8/59	0/59
<i>Organics--Semivolatiles (mg/kg)</i>					
Bis(2-ethylhexyl)phthalate	0.64	N/A	5/40	N/A	0/40
Butyl benzyl phthalate	0.73	N/A	1/29	N/A	0/29
Di-n-butyl phthalate	7.3	N/A	24/40	N/A	0/40
Fluoranthene	0.202	N/A	1/40	N/A	0/40
<i>Organics--Volatiles (mg/kg)</i>					
Trichloroethene	0.0051	N/A	12/85	N/A	0/85
<i>Radionuclides(pCi/g)</i>					
Radium-226	2.22	N/A	14/27	4/27	14/27
Technetium-99	3.89	ND	1/64	1/64	0/64
Thorium-228	N/A	0.383	16/17	0/17	16/17
Thorium-230	N/A	0.377	7/17	0/17	0/17
Thorium-232	N/A	0.494	17/17	0/17	0/17
Thorium-234	1.86	1.36	13/62	N/A	N/A
Uranium	3.3	0.658	7/28	N/A	N/A
Uranium-234	1.47	0.322	10/28	0/28	0/28
Uranium-238	2	0.297	9/28	6/28	6/28

^a A Frequency of detection is the number of detections of an analyte per number of analyses (includes regular and duplicate samples).

N/A = not applicable

ND = not detected



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DOE PORTSMOUTH/PADUCAH PROJECT OFFICE
PADUCAH GASEOUS DIFFUSION PLANT



Background and NALs taken from DOE 2001.

Figure 4.52. SWMU 5 Beryllium in Soil (Looking North)

Table 4.26. SWMU 5 Locations of Subsurface Soil Contaminants

Analysis	Depth (ft)	Excavation Worker NAL	Background	RI Data			Historical Data									
				005-101	005-102	005-103	005-015	005-016	005-017	005-018	005-019	005-020	005-021	005-022	005-027	005-028
Inorganics (mg/kg)	Aluminum	10-15	12,000	6,150	5,370	7,820	5,440	11,800	7,760	8,930	10,200	8,020	10,000	9,510		
		15-20	12,000										6,760	7,980		
		20-25	12,000						6,420						9,260	3,720
		25-30	12,000									7,430			5,970	5,950
		30-35	12,000	5,630	4,950	9,000	6,950				5,630	8,110	7,370	6,940	5,880	4,540
		35-40	12,000				6,950			3,940						
		40-45	12,000	992	2,820	9,950	11,700				11,300		13,600	16,400	15,600	
		50-55	12,000				8,250				9,260		12,600	11,100	4,340	7,750
		60-65	12,000	4,950	6,240	7,500	4,730				8,650					
	Arsenic	10-15	0.324	7.9	2.58	3.07	2.55	5U	5U	5U	5U	5U	5U	5U	5U	
15-20		0.324	7.9				5U	5U	5U	6.77				5U	5.64	
20-25		0.324	7.9											5U	5.04	
25-30		0.324	7.9											5U		
30-35		0.324	7.9	3.16	3.45	4.21	5U	5U	5U		5U	5U	5U	5U	5U	
35-40		0.324	7.9				5U		5U					5U	5U	
40-45		0.324	7.9	0.913U	0.935U	1.57	5U		5U		5U	5U	5U	5U	5U	
50-55		0.324	7.9				5U		5U		5U	5U	5U	5U	5U	
60-65		0.324	7.9	1.04	0.923U	1.53	5U		5U		5U					
Barium		10-15	272	170	141	97.5	101	23.8	61	31.2	71	76	66.1	58.6	54.8	
	15-20	272	170						16.2				28.5	33.9		
	20-25	272	170												38.5	18.4
	25-30	272	170												12.7	19.7
	30-35	272	170	18.4	11.1	15.8				19.7	16.5	19.5	16.7	18.3		
	35-40	272	170				39.2		14.4						10.9	14.1
	40-45	272	170	6.16	32.9	36.7	170			104		70.4	69.6	95		
	50-55	272	170				343			91.7		56.5	136	37.8	62.6	73.4
	60-65	272	170	32.1	65.1	56.4	31.6			216						
	Beryllium	10-15	1.26	0.69	0.48U	0.451U	0.459U	0.92	0.5U	0.87	0.83	0.93	0.5U	0.67	0.5U	
15-20		1.26	0.69					0.5U	1.23				0.5U	0.64		
20-25		1.26	0.69												0.69	0.74
25-30		1.26	0.69												0.91	0.96
30-35		1.26	0.69	0.463U	0.454U	0.494U				0.5U	0.71	0.72	0.64	0.5U		
35-40		1.26	0.69				0.75		0.5U						0.5U	0.5U
40-45		1.26	0.69	0.457U	0.468U	0.499U	1.02			1.47		1.64	1.47	2.59		
50-55		1.26	0.69				0.87			1.16		1.39	1.68	0.57	2.27	1.26
60-65		1.26	0.69	0.493U	0.462U	0.466U	0.5U			0.84						
Calcium		10-15	n/a	6,100	911	967	1,080				1,230	956	1,240	1,880	960	
	15-20	n/a	6,100										534	892		
	20-25	n/a	6,100				1,170	1,120	1,120						941	1,040
	25-30	n/a	6,100					630							541	953
	30-35	n/a	6,100	659	638	765				638	624	650	553	640		

Table 4.26. SWMU 5 Locations of Subsurface Soil Contaminants (Continued)

Analysis	Depth (ft)	Excavation Worker NAL	Background	RI Data			Historical Data									
				005-101	005-102	005-103	005-015	005-016	005-017	005-018	005-019	005-020	005-021	005-022	005-027	005-028
Nickel	40-45	216	22	4.57U	4.68U	7.86	13			15.8	19.3	19.8	27			
	50-55	216	22				9.68		13.6	14.4	17.3	5.1	15.9	12.8		
	60-65	216	22	5.61	7.46	9.55	5U		12.6							
	10-15	n/a	950						269	363	312	307				
	15-20	n/a	950				161	246	187		186	203				
	20-25	n/a	950						136					279	128	
Potassium	25-30	n/a	950								233			158	119	
	30-35	n/a	950						217	247	228	217				
	35-40	n/a	950				339	181		1,080	1,480	1,890		131	164	
	40-45	n/a	950				1,070		934	748	906	462	1,090		849	
	50-55	n/a	950				647		781							
	60-65	n/a	950				446									
	10-15	41.2	2.7	2.4U						4U	4U	4U	4U			
	15-20	41.2	2.7				4U	4U			4U	4U				
	20-25	41.2	2.7						4U					4U	4U	
	25-30	41.2	2.7											4U	4U	
Sodium	30-35	41.2	2.7	2.32U						4U	4U	4U	4U			
	35-40	41.2	2.7				4U	4U						4U	4U	
	40-45	41.2	2.7	2.28U			4U			4U	4U	4U	4U			
	50-55	41.2	2.7				4U			4U	5.74	4U	4U	4U	4U	
	60-65	41.2	2.7	2.47U			4U			4U						
	10-15	n/a	340	164	150	179			218	221	228	288	206			
	15-20	n/a	340				278	389	272			200U	200U			
	20-25	n/a	340						243					343	251	
	25-30	n/a	340											293	200U	
	30-35	n/a	340	92.6U	90.9U	98.7U	274			200U	200U	200U	200U			
Uranium	35-40	n/a	340				281							207	200U	
	40-45	n/a	340	91.3U	93.5U	99.8U			200U		200U	200U	200U			
	50-55	n/a	340				272		200U		200U	200U	200U			
	60-65	n/a	340	98.7U	92.3U	93.3U	225		200U							
	10-15	11.3	4.6	0.959U	0.901U	0.918U										
	30-35	11.3	4.6	0.926U	0.921	1.07										
	40-45	11.3	4.6	0.913U	0.935U	1.24										
	60-65	11.3	4.6	0.987U	0.923U	0.933U										
	10-15	4.4	37	20.3	20.5	15.6				27.2	33.3	18.9	19.1			
	15-20	4.4	37				41.4	16.5	36.1			17.3	27.8			
Vanadium	20-25	4.4	37					56.9						33	42.4	
	25-30	4.4	37											46.6	36.6	
	30-35	4.4	37	27.7	31.7	34.3			13.8	22.4	23	9.65				
	35-40	4.4	37				29.9		14.5		29.6			9.45	21.6	
	40-45	4.4	37	6.28	9.28	8.63	20.4		20.1	26.7	36.8					

Table 4.26. SWMU 5 Locations of Subsurface Soil Contaminants (Continued)

Analysis	Depth (ft)	Excavation Worker NAL	Background	RI Data			Historical Data													
				005-101	005-102	005-103	005-015	005-016	005-017	005-018	005-019	005-020	005-021	005-022	005-027	005-028				
Vanadium	50-55	4.4	37				12.4					18.8			17.2	31.1	11.2	34	15.7	
	60-65	4.4	37	15.1	18.4	14.6	7.23					15.5			26.4	24.3	20U			
	10-15	2,660	60	19.4	26.9	25.2						31.4	32.8							
	15-20	2,660	60				17.3	15.8	17.3											
	20-25	2,660	60						19.5									20.9	18	
	25-30	2,660	60												20U			15U	15U	
	30-35	2,660	60	18.5U	18.2U	19.7U					20U	20U	20U	20U						
	35-40	2,660	60				19.2		15U										15U	15U
	40-45	2,660	60	18.3U	18.7U	27.5	50.5					67.8			79.9	73.6	78.6			
	50-55	2,660	60				32.9					54.7			65.7	63.5	25	73.4		42.3
60-65	2,660	60	24.5	28.8	39	18.1					65.2									
Organics--Semi-volatiles (mg/kg)																				
Bis(2-ethylhexyl)phthalate	10-15	10.1	n/a									0.48U	0.64	0.47U	0.49U	0.48U	0.48U			
	15-20	10.1	n/a						0.5U	0.5U						0.48U	0.59			
	20-25	10.1	n/a						0.5U									0.5U	0.5U	0.5U
	25-30	10.1	n/a											0.47U				0.5U	0.5U	0.5U
	30-35	10.1	n/a									0.46U	0.52	0.48U	0.49U	0.48U	0.48U			
	35-40	10.1	n/a				0.5U					0.47U			0.5	0.45U	0.58		0.5U	0.5U
	40-45	10.1	n/a				0.5U					0.44U			0.48U	0.47U	0.46U	0.5U	0.5U	0.5U
	50-55	10.1	n/a				0.5U					0.47U			0.48U	0.47U	0.48U			
	60-65	10.1	n/a									0.48U	0.46U	0.47U	0.49U	0.48U	0.47U			
	10-15	2,330	n/a													0.48U	0.47U			
Butyl benzyl phthalate	15-20	2,330	n/a															0.5U		0.5U
	20-25	2,330	n/a																	
	25-30	2,330	n/a																	
	30-35	2,330	n/a																	
	40-45	2,330	n/a																	
	50-55	2,330	n/a																	
	60-65	2,330	n/a																	
	10-15	1,520	n/a																	
	15-20	1,520	n/a						0.5U											
	Di-n-butyl phthalate	20-25	1,520	n/a																
25-30		1,520	n/a																	
30-35		1,520	n/a																	
35-40		1,520	n/a				0.5U													
40-45		1,520	n/a				0.5U													
50-55		1,520	n/a				0.5U													
60-65		1,520	n/a				0.5U													
10-15		242	n/a																	
15-20		242	n/a																	
Fluoranthene		20-25	242	n/a						0.5U	0.5U									
	25-30	242	n/a																	

4.6.2 SWMU 5 Groundwater

UCRS groundwater samples were collected from two of three angled borings installed at SWMU 5 as part of this RI. Samples of 10 temporary borings of the WAG 3 RI (DOE 2000a), both angled below the burial cells and peripheral to the burial ground, provided historical data for the UCRS at SWMU 5 (Figure 4.12). RI data were reviewed with historical data to determine the UCRS contaminants listed in Table 4.27.

The SWMU 5 disposal pits extend to a depth of 6 to 15 ft. These are underlain by the HU2 horizon of the UCRS at depths of 20 to 40 ft. The shallowest groundwater samples were from two WAG 3 RI soil borings at depths of 20-30 ft.

The remainder of the UCRS groundwater samples were from depths of 40 to 61 ft. Screening identified many metals in these UCRS groundwater samples from SWMU 5 with concentrations that exceed screening criteria. Of these, iron, lead, manganese, and molybdenum analyses had the highest frequency of exceedances. (Lead exceeded its MCL at three locations.) Locations with metals that exceed screening criteria were well distributed across the SWMU. Because it was determined during scoping meetings for the BGOU RI/FS Work Plan that SWMU 5 should not be a source for VOC contamination, organics were not analyzed during this RI; however, analyses of historical samples of UCRS groundwater documented single detections of pyrene and TCE at concentrations that exceed screening levels. TCE was detected in UCRS groundwater at a concentration of 29 µg/L in boring 005-022. This occurrence, at a depth of 61 ft, is likely related to dissolved-phase contamination in the RGA, located immediately below the sample depth. No radionuclide analyses exceeded screening criteria in the UCRS groundwater samples.

This RI did not collect RGA and McNairy groundwater at SWMU 5; however, historical data were reviewed to determine the RGA groundwater contaminants listed in Table 4.28. (The screening determined that there were no McNairy groundwater contaminants.) Only locations 005-013 and 005-026 of the WAG 3 RI were sampled for metals, organics, and radionuclides in RGA and McNairy groundwater. Additionally, the location DG-002 was sampled for VOCs and radionuclides (DOE 2000b). Manganese exceeded screening criteria in all 51 RGA groundwater samples from SWMU 5. Iron was the only other metal that commonly was present at levels exceeding screening criteria. TCE concentrations exceeded screening criteria throughout the depth of the RGA. These occurrences likely are related to the Northwest Plume, which passes to the east of the SWMU 5 area. Figure 4.54 shows the relationship of the Northwest Plume in the RGA with SWMUs 5 and 6.

Table 4.29 provides detail (depth, sample location, and analytical results) for SWMU 5 groundwater samples, including nondetects and detections above screening levels.

Table 4.27. SWMU 5 UCRS Groundwater Contaminants

Analysis	Historical Data	RI	Frequency of Detection ^a	Frequency of Detection		
				Above Background	Above NAL	Above MCL
Metals (mg/L)						
Aluminum	591	9.59	13/20	N/A	10/20	N/A
Arsenic	0.014	0.00182	3/4	N/A	3/4	1/4
Arsenic, Dissolved	N/A	0.00192	2/2	N/A	2/2	0/2
Barium	10.5	0.164	18/20	N/A	17/20	4/20
Barium, Dissolved	N/A	0.151	2/2	N/A	2/2	0/2
Beryllium	0.144	ND	5/18	N/A	5/18	5/18
Calcium	202	38.7	20/20	N/A	N/A	N/A
Calcium, Dissolved	N/A	39.1	2/2	N/A	N/A	N/A
Chromium	6.47	ND	6/20	N/A	3/20	6/20
Cobalt	1.82	0.00548	17/20	N/A	4/20	N/A
Cobalt, Dissolved	N/A	0.00837	2/2	N/A	0/2	N/A
Copper	1.81	ND	6/20	N/A	6/20	1/20
Iron	2,090	37.9	16/20	N/A	16/20	N/A
Iron, Dissolved	N/A	29.4	2/2	N/A	2/2	N/A
Lead	0.816	0.00208	6/6	N/A	4/6	4/6
Magnesium	88.6	15.6	20/20	N/A	0/20	N/A
Magnesium, Dissolved	N/A	15.3	2/2	N/A	0/2	N/A
Manganese	54.8	0.712	20/20	N/A	20/20	N/A
Manganese, Dissolved	N/A	0.626	2/2	N/A	2/2	N/A
Mercury	0.0025	ND	4/16	N/A	3/16	1/16
Molybdenum	N/A	0.0386	2/2	N/A	2/2	N/A
Molybdenum, Dissolved	N/A	0.0308	2/2	N/A	2/2	N/A
Nickel	1.37	0.0311	7/20	N/A	7/20	N/A
Nickel, Dissolved	N/A	0.0372	2/2	N/A	2/2	N/A
Potassium	28	N/A	7/18	N/A	N/A	N/A
Selenium	0.005	0.0102	3/9	N/A	1/9	0/9
Selenium, Dissolved	N/A	0.0099	2/2	N/A	1/2	0/2
Sodium	56.9	43.5	20/20	N/A	0/20	N/A
Sodium, Dissolved	N/A	44	2/2	N/A	0/2	N/A
Vanadium	2.62	ND	5/18	N/A	5/18	N/A
Zinc	6.77	0.064	10/18	N/A	5/18	N/A
Zinc, Dissolved	N/A	0.0556	2/2	N/A	0/2	N/A
Radionuclides (pCi/L)						
Radon	737	N/A	5/7	N/A	N/A	N/A
Technetium-99	31	78.4	3/10	N/A	2/10	0/10
Thorium-228	N/A	0.142	1/2	N/A	1/2	N/A
Uranium-238	N/A	0.354	3/4	N/A	0/4	N/A
Semivolatiles (mg/L)						
Bis(2-ethylhexyl)phthalate	0.007	N/A	2/6	N/A	2/6	1/6
Diethyl phthalate	0.019	N/A	1/6	N/A	0/6	N/A
Pyrene	0.023	N/A	1/6	N/A	1/6	N/A
Volatiles (mg/L)						
Acetone	0.038	N/A	1/7	N/A	1/7	N/A
cis-1,2-Dichloroethene	0.011	N/A	2/10	N/A	2/10	0/10
Trichloroethene	0.029	N/A	1/10	N/A	1/10	1/10

^a Frequency of detection is the number of detections of an analyte per number of analyses (includes regular and duplicate samples).

N/A = not applicable

ND = not detected

Table 4.28. SWMU 5 RGA Groundwater Contaminants

Analysis	Historical Data	RI	Frequency of Detection ^a	Frequency of Detection		
				Above Background	Above PRG	Above MCL
Metals (mg/L)						
Aluminum	151	N/A	21/51	17/51	20/51	N/A
Barium	9.18	N/A	48/51	16/51	33/51	1/51
Beryllium	0.032	N/A	6/29	6/29	6/29	6/29
Cadmium	0.044	N/A	1/19	1/19	1/19	1/19
Calcium	44.9	N/A	51/51	1/51	N/A	N/A
Chromium	0.81	N/A	6/29	3/29	0/29	4/29
Cobalt	0.248	N/A	29/49	5/49	2/49	N/A
Iron	2160	N/A	41/51	19/51	35/51	N/A
Lead	0.655	N/A	1/1	1/1	1/1	1/1
Magnesium	28.2	N/A	51/51	2/51	0/51	N/A
Manganese	51.3	N/A	51/51	51/51	51/51	N/A
Mercury	0.0002	N/A	1/19	0/19	0/19	0/19
Nickel	0.24	N/A	10/37	0/37	10/37	N/A
Potassium	22.8	N/A	13/31	5/31	N/A	N/A
Sodium	20	N/A	51/51	0/51	0/51	N/A
Vanadium	1.21	N/A	6/29	2/29	6/29	N/A
Zinc	0.466	N/A	6/29	6/29	1/29	N/A
Radionuclides (pCi/L)						
Radon	248	N/A	3/17	N/A	N/A	N/A
Technetium-99	33.4	N/A	9/24	3/24	9/24	0/24
Volatiles (mg/L)						
1,1-Dichloroethene	0.003	N/A	5/22	N/A	5/22	0/22
cis-1,2-Dichloroethene	0.002	N/A	7/24	N/A	0/24	0/24
trans-1,2-Dichloroethene	0.0003	N/A	5/24	N/A	0/24	0/24
Trichloroethene	0.033	N/A	14/24	N/A	9/24	7/24

^a Frequency of detection is the number of detections of an analyte per number of analyses (includes regular and duplicate samples).

N/A = not applicable

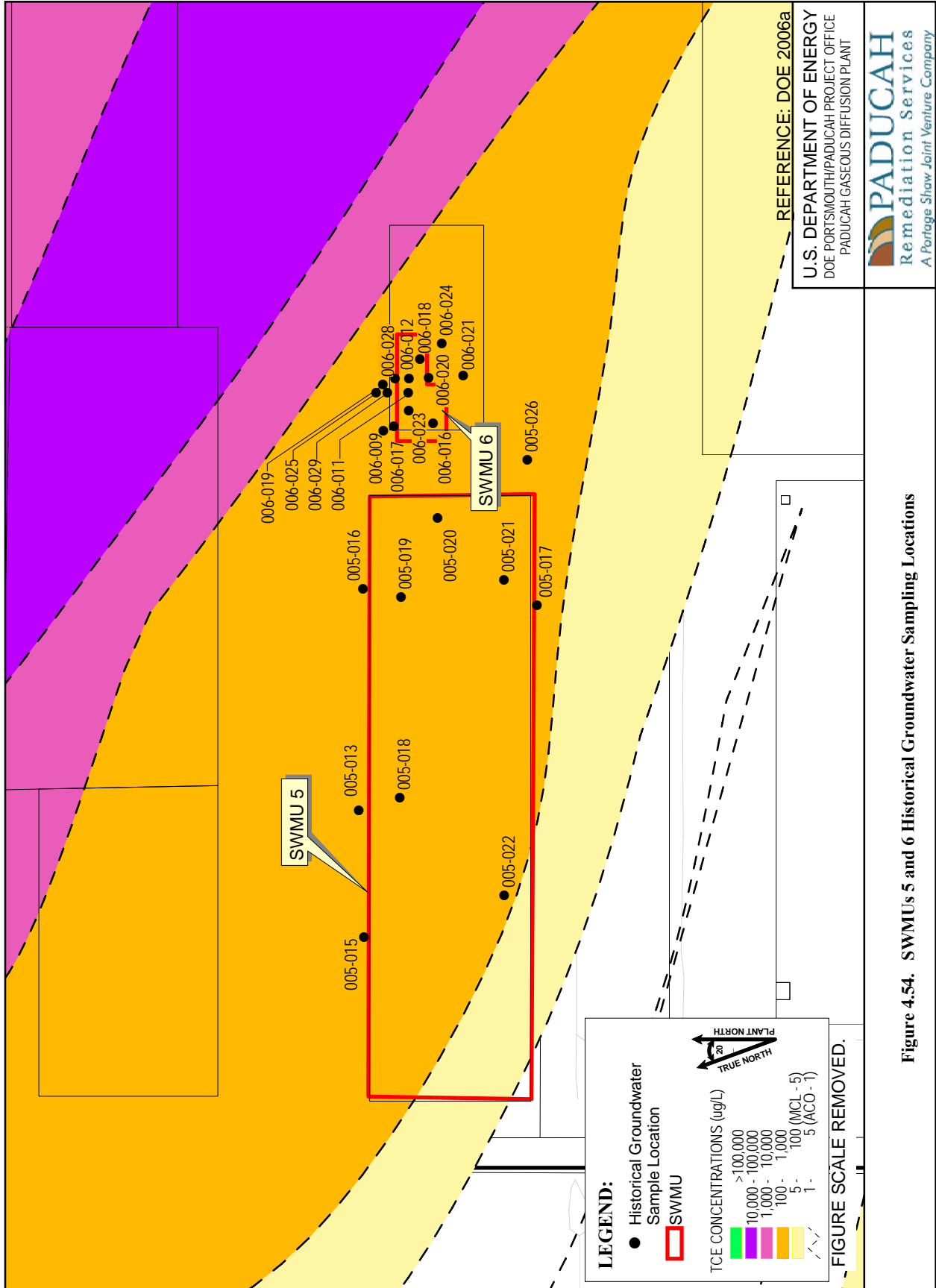


Figure 4.54. SWMUs 5 and 6 Historical Groundwater Sampling Locations

Table 4.29. SWMU 5 Locations of Groundwater Contaminants

Hydro Unit	Depth Interval	Analysis	Child Resident NAL	Bkgd	MCL	005-101	005-102	005-013	005-015	005-016	005-017	005-018	005-019	005-026	DG-002
	20-30	Radionuclides (pCi/L)													
		Radon	N/A	N/A	N/A					195U					
		Technetium-99	14	N/A	900					14	31				
		Volatiles (mg/L)													
		Acetone	0.0275	N/A	N/A					0.01U					
		<i>cis</i> -1,2-Dichloroethene	0.00273	N/A	0.07					0.005U	0.001U				
		Trichloroethene	0.0016	N/A	0.005					0.001U	0.001U				
		Metals (mg/L)													
	40-45	Aluminum	1.49	N/A	N/A	9.59	5.51								
		Arsenic	0.000035	N/A	0.01	0.01U	0.00182								
		Arsenic, Dissolved	0.000035	N/A	0.01	0.00192	0.00107								
		Barium	0.104	N/A	2	0.164	0.145								
		Barium, Dissolved	0.104	N/A	2	0.151	0.136								
		Beryllium	0.00264	N/A	0.004	0.001U	0.001U								
		Calcium	N/A	N/A	N/A	38.7	31.9								
		Calcium, Dissolved	N/A	N/A	N/A	39.1	31.3								
		Chromium	1.76	N/A	0.1	0.1U	0.01U								
		Cobalt	0.0906	N/A	N/A	0.01U	0.00548								
		Cobalt, Dissolved	0.0906	N/A	N/A	0.00837	0.00541								
		Copper	0.0557	N/A	1.3	0.2U	0.02U								
		Iron	0.449	N/A	N/A	34.7	37.9								
		Iron, Dissolved	0.449	N/A	N/A	24.8	29.4								
		Lead	0.015	N/A	0.015	0.00208	0.00164								
		Magnesium	N/A	N/A	N/A	15.6	12.9								
		Magnesium, Dissolved	N/A	N/A	N/A	15.3	12.8								
		Manganese	0.035	N/A	N/A	0.712	0.295								
		Manganese, Dissolved	0.035	N/A	N/A	0.626	0.322								
		Mercury	0.000444	N/A	0.002	0.00001U	0.00001U								
		Molybdenum	0.00753	N/A	N/A	0.0376	0.0386								
		Molybdenum, Dissolved	0.00753	N/A	N/A	0.029	0.0308								
		Nickel	0.0301	N/A	N/A	0.05U	0.0311								
		Nickel, Dissolved	0.0301	N/A	N/A	0.0372	0.0336								
		Selenium	0.00754	N/A	0.05	0.0102	0.00564								
		Selenium, Dissolved	0.00754	N/A	0.05	0.0099	0.00556								
		Sodium	N/A	N/A	N/A	43.5	38.9								

Table 4.29. SWMU 5 Locations of Groundwater Contaminants (Continued)

Hydro Unit	Depth Interval	Analysis	Child Resident NAL	Bkgd	MCL	005-101	005-102	005-013	005-015	005-016	005-017	005-018	005-019	005-026	DG-002	
UCRS	40-45	Sodium, Dissolved	N/A	N/A	N/A	44	38.7									
		Vanadium	0.00925	N/A	N/A	0.2U	0.02U									
		Zinc	0.45	N/A	N/A	0.2U	0.064									
		Zinc, Dissolved	0.45	N/A	N/A	0.0294	0.0556									
		Radionuclides (pCi/L)														
		Technetium-99	14	N/A	900	0.609U	78.4									
		Thorium-228	0.129	N/A	N/A	0.13U	0.142									
		Uranium-238	0.443	N/A	N/A	0.354	0.187									
		Metals (mg/L)														
		Aluminum	1.49	N/A	N/A	N/A			118				395	591		
		Arsenic	0.000035	N/A	0.01									0.014		
		Barium	0.104	N/A	2				0.866				3.4	10.5		
		Beryllium	0.00264	N/A	0.004				0.008				0.045	0.144		
		Calcium	N/A	N/A	N/A				44.5				66.9	202		
		Chromium	1.76	N/A	0.1				0.157				2.04	6.47		
		Cobalt	0.0906	N/A	N/A				0.06				0.266	1.82		
		Copper	0.0557	N/A	1.3				0.186				0.596	1.81		
		Iron	0.449	N/A	N/A				111				787	2,090		
		Lead	0.015	N/A	0.015								0.248	0.816		
		Magnesium	N/A	N/A	N/A				21.9				45.6	88.6		
	Manganese	0.035	N/A	N/A				1.64				11.4	54.8			
	Mercury	0.000444	N/A	0.002				0.0002U				0.0003	0.0025			
	Nickel	0.0301	N/A	N/A				0.1				0.482	1.37			
	Potassium	N/A	N/A	N/A				11.2				28	27.8			
	Selenium	0.00754	N/A	0.05				0.005U				0.005U	0.005			
	Sodium	N/A	N/A	N/A				24.5				23.7	56.9			
	Vanadium	0.00925	N/A	N/A				0.159				0.765	2.62			
	Zinc	0.45	N/A	N/A				0.556				2.44	6.77			
	Radionuclides (pCi/L)															
	Radon	N/A	N/A	N/A				348				267	361			
	Technetium-99	14	N/A	900				14U				15.8U	15.8U			
	Semivolatile (mg/L)															
	Bis(2-ethylhexyl)phthalate	0.00312	N/A	0.006				0.02U				0.005U	0.007			
	Diethyl phthalate	1.2	N/A	N/A				0.019				0.005U	0.005U			
	Pyrene	0.0182	N/A	N/A				0.023				0.005U	0.005U			

Table 4.29. SWMU 5 Locations of Groundwater Contaminants (Continued)

Hydro Unit	Depth Interval	Analysis	Child Resident NAL	Bkgd	MCL	005-101	005-102	005-1013	005-1015	005-1016	005-1017	005-1018	005-1019	005-1026	DG-002	
UCRS	60-61	Analysis														
		Volatiles (mg/L)														
		Acetone	0.0275	N/A	N/A	N/A							0.01U	0.01U		
		cis-1,2-Dichloroethene	0.00273	N/A	0.07				0.001U				0.005U	0.011		
	Trichloroethene	0.0016	N/A	0.005				0.001U				0.001U	0.001U			
	Metals (mg/L)															
	Aluminum	1.49	2.189	N/A				59.9						106		
	Barium	0.104	0.235	2				0.66						0.982		
	Beryllium	0.00264	0.004	0.004				0.005						0.006		
	Cadmium	0.000661	0.01	0.005				0.01U						0.01U		
	Calcium	N/A	41.238	N/A				17.4						17.9		
	Chromium	1.76	0.144	0.1				0.075						0.16		
	Cobalt	0.0906	0.045	N/A				0.044							0.071	
	Iron	0.449	5.03	N/A				91.5						109		
	Magnesium	N/A	16.262	N/A				10.8						13.1		
Manganese	0.035	0.119	N/A				1.73						7.16			
Mercury	0.000444	0.0002	0.002				0.0002U						0.0002U			
Nickel	0.0301	0.682	N/A				0.115						0.166			
Potassium	N/A	5.195	N/A				5.85						8.95			
Sodium	N/A	59.45	N/A				20						15.3			
Vanadium	0.00925	0.134	N/A				0.105						0.108			
Zinc	0.45	0.054	N/A				0.297						0.466			
Radionuclides (pCi/L)																
Radon	N/A	N/A	N/A				151U							153U		
Technetium-99	14	22.3	900				14U							13U		
RGA	63	Volatiles (mg/L)														
		1,1-Dichloroethene	0.000047	N/A	0.007				0.001U						0.001U	
		cis-1,2-Dichloroethene	0.00273	N/A	0.07				0.001U						0.001U	
		trans-1,2-Dichloroethene	0.00548	N/A	0.1				0.001U						0.001U	
	Trichloroethene	0.0016	N/A	0.005				0.001						0.001U		
	Metals (mg/L)															
	Aluminum	1.49	2.189	N/A				4.8						143		
	Barium	0.104	0.235	2				0.17						1.27		
	Beryllium	0.00264	0.004	0.004				0.005U						0.009		
	Cadmium	0.000661	0.01	0.005				0.01U						0.01U		
Calcium	N/A	41.238	N/A				18.1						18.7			
Chromium	1.76	0.144	0.1				0.05U						0.143			
Cobalt	0.0906	0.045	N/A				0.038						0.11			
68	Metals (mg/L)															
	Aluminum	1.49	2.189	N/A												
	Barium	0.104	0.235	2												
	Beryllium	0.00264	0.004	0.004												
	Cadmium	0.000661	0.01	0.005												
	Calcium	N/A	41.238	N/A												

Table 4.29. SWMU 5 Locations of Groundwater Contaminants (Continued)

Hydro Unit	Depth Interval	Analysis	Child Resident NAL	Bkgd	MCL	005-101	005-102	005-013	005-015	005-016	005-017	005-018	005-019	005-026	DG-002		
RGA	68	Iron	0.449	5.03	N/A			14.6						172			
		Magnesium	N/A	16.262	N/A			6.97						13.3			
		Manganese	0.035	0.119	N/A			2.43							14.5		
		Mercury	0.000444	0.0002	0.002			0.0002U							0.0002U		
		Nickel	0.0301	0.682	N/A			0.05U							0.21		
		Potassium	N/A	5.195	N/A			2U							9.71		
		Sodium	N/A	59.45	N/A			18.5							13.5		
		Vanadium	0.00925	0.134	N/A			0.1U							0.13		
		Zinc	0.45	0.054	N/A			0.2U							0.407		
		Radionuclides (pCi/L)															
		Radon	N/A	N/A	N/A	N/A			151U							153U	
		Technetium-99	14	22.3	900				16.5							14U	
		Volatiles (mg/L)															
		1,1-Dichloroethene	0.000047	N/A	0.007				0.001U							0.001U	
		cis-1,2-Dichloroethene	0.00273	N/A	0.07				0.001U							0.001U	
		trans-1,2-Dichloroethene	0.00548	N/A	0.1				0.001U							0.001U	
		Trichloroethene	0.0016	N/A	0.005				0.006							0.001U	
		Metals (mg/L)															
		Aluminum	1.49	2.189	N/A				13							8.48	
		Barium	0.104	0.235	2				0.776							0.207	
Beryllium	0.00264	0.004	0.004				0.005U							0.005U			
Cadmium	0.000661	0.01	0.005				0.01U							0.01U			
Calcium	N/A	41.238	N/A				18.9							11			
Chromium	1.76	0.144	0.1				0.05U							0.05U			
Cobalt	0.0906	0.045	N/A				0.047							0.026			
Iron	0.449	5.03	N/A				90.4							20.3			
Magnesium	N/A	16.262	N/A				7.32							4.79			
Manganese	0.035	0.119	N/A				6.52							3.53			
Mercury	0.000444	0.0002	0.002				0.0002U							0.0002U			
Nickel	0.0301	0.682	N/A				0.104							0.05U			
Potassium	N/A	5.195	N/A				2U							2U			
Sodium	N/A	59.45	N/A				17.3							12.3			
Vanadium	0.00925	0.134	N/A				0.1U							0.1U			
Zinc	0.45	0.054	N/A				0.2U							0.2U			
Radionuclides (pCi/L)																	
Radon	N/A	N/A	N/A				151U							153U			
Technetium-99	14	22.3	900				18.9U							14U	7U		

Table 4.29. SWMU 5 Locations of Groundwater Contaminants (Continued)

Hydro Unit	Depth Interval	Analysis	Child Resident NAL	Bkgd	MCL	005-101	005-102	005-013	005-015	005-016	005-017	005-018	005-019	005-026	DG-002		
RGA	70-73	Volatiles (mg/L)															
		1,1-Dichloroethene	0.000047	N/A	0.007			0.0006						0.001U	0.001U		
		<i>cis</i> -1,2-Dichloroethene	0.00273	N/A	0.07			0.0004							0.001U	0.005U	
		<i>trans</i> -1,2-Dichloroethene	0.00548	N/A	0.1			0.0002							0.001U	0.005U	
	Trichloroethene	0.0016	N/A	0.005			0.01							0.0001	0.001U		
	76-78	Metals (mg/L)															
		Aluminum	1.49	2.189	N/A				23						5.29		
		Barium	0.104	0.235	2				0.774						0.508		
		Beryllium	0.00264	0.004	0.004				0.005						0.005U		
		Cadmium	0.000661	0.01	0.005				0.01U						0.01U		
		Calcium	N/A	41.238	N/A				20.4						13.3		
		Chromium	1.76	0.144	0.1				0.05U						0.05U		
		Cobalt	0.0906	0.045	N/A				0.018						0.017		
		Iron	0.449	5.03	N/A				590						41.4		
		Magnesium	N/A	16.262	N/A				7.78						5.57		
		Manganese	0.035	0.119	N/A				5.97						3.98		
		Mercury	0.000444	0.002	0.002				0.0002U						0.0002U		
		Nickel	0.0301	0.682	N/A				0.24						0.05U		
		Potassium	N/A	5.195	N/A				2U						2U		
		Sodium	N/A	59.45	N/A				16.3						12.3		
		Vanadium	0.00925	0.134	N/A				0.1U						0.1U		
		Zinc	0.45	0.054	N/A				0.2U						0.2U		
		Radionuclides (pCi/L)	Radon	N/A	N/A	N/A			151U							153U	
			Technetium-99	14	22.3	900			31							14.9	33.4
			Volatiles (mg/L)														
	70-73	1,1-Dichloroethene	0.000047	N/A	0.007				0.002						0.001U	0.001U	
		<i>cis</i> -1,2-Dichloroethene	0.00273	N/A	0.07			0.002							0.0002	0.001U	
<i>trans</i> -1,2-Dichloroethene		0.00548	N/A	0.1			0.0002							0.001U	0.001U		
Trichloroethene		0.0016	N/A	0.005			0.033							0.008	0.001U		
Metals (mg/L)																	
81-83	Aluminum	1.49	2.189	N/A										5.68			
	Barium	0.104	0.235	2										1.93			
	Beryllium	0.00264	0.004	0.004										0.005U			
	Cadmium	0.000661	0.01	0.005										0.01U			
	Calcium	N/A	41.238	N/A										13.5			
Chromium	1.76	0.144	0.1										0.05U				

Table 4.29. SWMU 5 Locations of Groundwater Contaminants (Continued)

Hydro Unit	Depth Interval	Analysis	Child Resident NAL	Bkgd	MCL	005-101	005-102	005-013	005-015	005-016	005-017	005-018	005-019	005-026	DG-002		
RGA	81-83	Cobalt	0.0906	0.045	N/A									0.064			
		Iron	0.449	5.03	N/A									130			
		Magnesium	N/A	16.262	N/A										5.54		
		Manganese	0.035	0.119	N/A										9.93		
		Mercury	0.000444	0.0002	0.002										0.0002U		
		Nickel	0.0301	0.682	N/A										0.074		
		Potassium	N/A	5.195	N/A										2U		
		Sodium	N/A	59.45	N/A										12.2		
		Vanadium	0.00925	0.134	N/A										0.1U		
		Zinc	0.45	0.054	N/A										0.2U		
		Radionuclides (pCi/L)															
		Radon	N/A	N/A	N/A	N/A										217	
		Technetium-99	14	22.3	900											18.3	15.3
		Volatiles (mg/L)															
		1,1-Dichloroethene	0.000047	N/A	0.007									0.001U	0.001U		
		<i>cis</i> -1,2-Dichloroethene	0.00273	N/A	0.07									0.001U	0.001U		
		<i>trans</i> -1,2-Dichloroethene	0.00548	N/A	0.1									0.001U	0.001U		
		Trichloroethene	0.0016	N/A	0.005									0.003	0.001U		
Metals (mg/L)																	
	86-88	Aluminum	1.49	2.189	N/A			8.5						6.66			
		Barium	0.104	0.235	2			1.47						1.49			
		Beryllium	0.00264	0.004	0.004			0.005U						0.005U			
		Cadmium	0.000661	0.01	0.005			0.01U						0.01U			
		Calcium	N/A	41.238	N/A			19						13.8			
		Chromium	1.76	0.144	0.1			0.05U						0.05U			
		Cobalt	0.0906	0.045	N/A			0.012						0.034			
		Iron	0.449	5.03	N/A			120						215			
		Magnesium	N/A	16.262	N/A			7.54						5.53			
		Manganese	0.035	0.119	N/A			9.85						8.8			
		Mercury	0.000444	0.0002	0.002			0.0002U						0.0002U			
		Nickel	0.0301	0.682	N/A			0.06						0.15			
		Potassium	N/A	5.195	N/A			2U						2U			
		Sodium	N/A	59.45	N/A			16						11.8			
	Vanadium	0.00925	0.134	N/A			0.1U						0.1U				
	Zinc	0.45	0.054	N/A			0.2U						0.257				
Radionuclides (pCi/L)																	
Radon	N/A	N/A	N/A	N/A				155U						248			
Technetium-99	14	22.3	900					20						22.4	8.3U		

Table 4.29. SWMU 5 Locations of Groundwater Contaminants (Continued)

Hydro Unit	Depth Interval	Analysis	Child Resident NAL	Bkgd	MCL	005-101	005-102	005-1013	005-015	005-016	005-017	005-018	005-019	005-026	DG-002	
RGA	86-88	Volatiles (mg/L)														
		1,1-Dichloroethene	0.000047	N/A	0.007			0.003						0.001U	0.001U	
		<i>cis</i> -1,2-Dichloroethene	0.00273	N/A	0.07			0.002						0.0001	0.001U	
		<i>trans</i> -1,2-Dichloroethene	0.00548	N/A	0.1			0.0003						0.001U	0.001U	
		Trichloroethene	0.0016	N/A	0.005			0.02						0.005	0.001U	
	90-93	Metals (mg/L)														
		Aluminum	1.49	2.189	N/A				23						151	
		Barium	0.104	0.235	2				1.14						9.18	
		Beryllium	0.00264	0.004	0.004				0.005U						0.032	
		Cadmium	0.000661	0.01	0.005				0.01U						0.044	
		Calcium	N/A	41.238	N/A				17.7						34.9	
		Chromium	1.76	0.144	0.1				0.063						0.87	
		Cobalt	0.0906	0.045	N/A				0.019						0.248	
		Iron	0.449	5.03	N/A				348						2,160	
		Lead	0.015	0.129	0.015										0.655	
		Magnesium	N/A	16.262	N/A				8.78						23.3	
		Manganese	0.035	0.119	N/A				7.85						51.3	
		Mercury	0.000444	0.0002	0.002				0.0002U						0.0002	
		Nickel	0.0301	0.682	N/A				0.18						0.5U	
		Potassium	N/A	5.195	N/A				3.38						20.4	
		Sodium	N/A	59.45	N/A				15.3						10.8	
		Vanadium	0.00925	0.134	N/A				0.1U						1.21	
		Zinc	0.45	0.054	N/A				0.23						2U	
		Radionuclides (pCi/L)														
		Radon	N/A	N/A	N/A				155U						168U	
		Technetium-99	14	22.3	900				20						14U	17.9U
		Volatiles (mg/L)														
		1,1-Dichloroethene	0.000047	N/A	0.007				0.0008						0.001U	0.001U
		<i>cis</i> -1,2-Dichloroethene	0.00273	N/A	0.07				0.0009						0.001U	0.001U
		<i>trans</i> -1,2-Dichloroethene	0.00548	N/A	0.1				0.0002						0.001U	0.001U
		Trichloroethene	0.0016	N/A	0.005				0.015						0.0003	0.001U

Table 4.29. SWMU 5 Locations of Groundwater Contaminants (Continued)

Hydro Unit	Depth Interval	Analysis	Child Resident NAL	Bkgd	MCL	005-101	005-102	005-013	005-015	005-016	005-017	005-018	005-019	005-026	DG-002		
RGA	98	Metals (mg/L)															
		Aluminum	1.49	2.189	N/A			13.2							93.4		
		Barium	0.104	0.235	2			0.187							0.398		
		Beryllium	0.00264	0.004	0.004			0.01U							0.009		
		Cadmium	0.000661	0.01	0.005			0.01U							0.01U		
		Calcium	N/A	41.238	N/A			22.5							44.9		
		Chromium	1.76	0.144	0.1			0.05U							0.205		
		Cobalt	0.0906	0.045	N/A			0.018							0.038		
		Iron	0.449	5.03	N/A			54.7							312		
		Magnesium	N/A	16.262	N/A			10.4							28.2		
		Manganese	0.035	0.119	N/A			1.73							1.81		
		Mercury	0.000444	0.0002	0.002			0.0002U							0.0002U		
		Nickel	0.0301	0.682	N/A			0.05U							0.12		
		Potassium	N/A	5.195	N/A			4.5							22.8		
		Sodium	N/A	59.45	N/A			11.5							10.6		
		Vanadium	0.00925	0.134	N/A			0.101							0.473		
		Zinc	0.45	0.054	N/A			0.2U							0.403		
				Radionuclides (pCi/L)													
				Radon	N/A	N/A	N/A			169						168U	
				Technetium-99	14	22.3	900			16U						14U	
				Volatiles (mg/L)													
				1,1-Dichloroethene	0.000047	N/A	0.007			0.001U						0.001U	
				<i>cis</i> -1,2-Dichloroethene	0.00273	N/A	0.07			0.005U						0.001U	
				<i>trans</i> -1,2-Dichloroethene	0.00548	N/A	0.1			0.005U						0.001U	
				Trichloroethene	0.0016	N/A	0.005			0.0003						0.001U	

Blank cells indicate interval was not sampled for the specified analysis. Maximum value shown for each depth interval.

N/A = not applicable

Bold indicates result is greater than NAL value.

Italics indicates result is greater than MCL value.

Bold + Italics indicate result is greater than both NAL and MCL values.

Shading indicates result is greater than the background value.

4.7 SWMU 6

4.7.1 Subsurface Soils

Subsurface soil samples were collected from four angled borings at SWMU 6 as part of this RI (Figure 4.5). The screen of RI and historical data identified the contaminants listed in Table 4.30.

Each burial area within SWMU 6 received different types of waste. The contents buried within each area are summarized as follows:

- Area H—magnesium scrap
- Area I—exhaust hood blowers contaminated with perchloric acid
- Area J—contaminated aluminum scrap
- Area K—magnesium scrap
- Area L—contaminated modine cold trap

Metals analyses of subsurface soil samples from SWMU 6 rarely exceed screening criteria (both background and NALs, where applicable) for identifying contamination. Screening identified beryllium and vanadium as the most frequent metal contaminants above background (in 22 and 10 of 70 analyses, respectively). The distribution of beryllium in soil is shown in Figure 4.55. Most of the NAL exceedances of beryllium occur in a horizon at 40 to 55 ft bgs. There is uncertainty with the lower extent in the vicinity of historical soil borings 006-026 and 006-027 since no deeper samples were collected (the top of the RGA occurs at a depth of 58 to 60 ft). Figure 4.56 illustrates the vanadium distribution in soil at SWMU 6. While there appears to be some zones of higher concentration, exceedances are found at all depths of the UCRS.

Of the occurrences of aluminum detected above background levels, the majority represents samples collected beneath Area J (aluminum scrap). The SWMU 6 burial pits extended to a depth of approximately 6 to 8 ft. All five detections of aluminum above PGDP background were from a depth of 43 to 51 ft. The maximum aluminum result was 22,500 mg/kg from location 006-020 at 43 ft bgs. The screening process did not identify any radionuclides or organic compounds as potential contaminants for SWMU 6.

Table 4.31 lists the locations of the contaminant detections above screening levels.

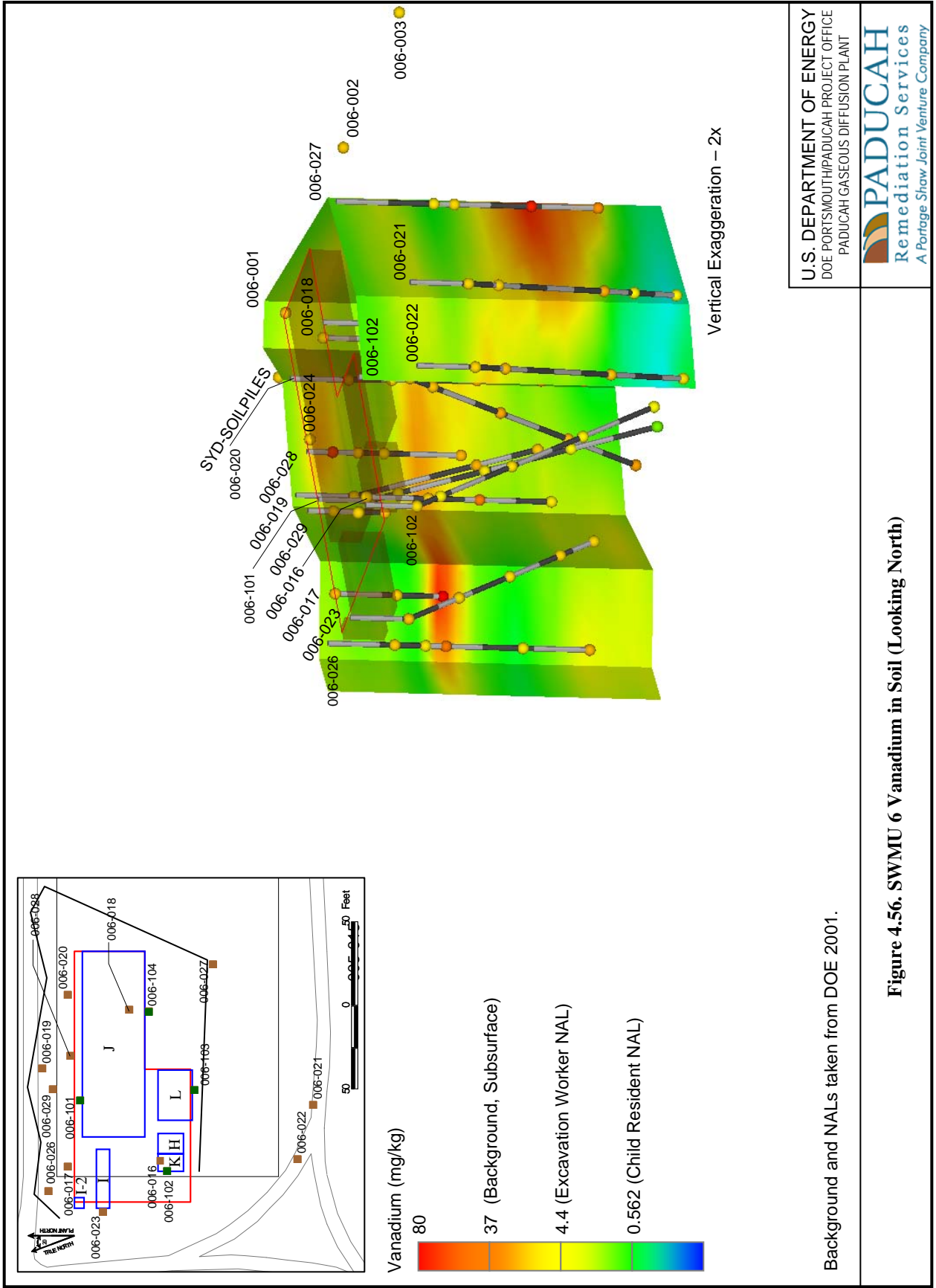
Table 4.30. SWMU 6 Subsurface Soil Contaminants

Analysis	Maximum Result		Frequency of Detection ^a	Frequency of Detection	
	Historical Data	RI Data		Above Background Value	Above Excavation Worker NAL
<i>Inorganics (mg/kg)</i>					
Aluminum	22,500	11,300	70/70	5/70	55/70
Arsenic	6.19	5.94	23/70	0/70	23/70
Barium	153	97.5	70/70	0/70	0/70
Beryllium	3.07	0.825	32/70	22/70	12/70
Calcium	14,900	1,410	70/70	2/70	N/A
Chromium	116	19	70/70	5/70	0/70
Cobalt	156	16.4	62/70	6/70	0/70
Copper	20.9	13	67/70	0/70	0/70
Iron	58,700	19,100	70/70	7/70	69/70
Lead	35.4	9.08	24/70	2/70	0/70
Magnesium	2,370	1,400	70/70	1/70	0/70
Manganese	1,550	315	70/70	1/70	64/70
Nickel	68.6	12.9	39/70	4/70	0/70
Potassium	1,330	N/A	46/48	5/48	N/A
Sodium	1,180	160	43/70	14/70	0/70
Uranium	N/A	1.23	5/22	0/22	0/22
Vanadium	79.1	38.5	69/70	10/70	69/70
Zinc	72.8	46.3	43/70	5/70	0/70
<i>Organics--Semivolatiles (mg/kg)</i>					
Bis(2-ethylhexyl)phthalate	0.6	N/A	2/44	N/A	0/44
Di-n-butyl phthalate	3.2	N/A	16/44	N/A	0/44
<i>Organics--Volatiles (mg/kg)</i>					
Acetone	0.085	N/A	6/38	N/A	0/38
Trichloroethene	0.0101	N/A	5/102	N/A	0/102
<i>Radionuclides(pCi/g)</i>					
Radium-226	0.907	N/A	6/26	0/26	6/26
Thorium-228	N/A	0.506	19/22	0/22	19/22
Thorium-230	N/A	0.341	11/22	0/22	0/22
Thorium-232	N/A	0.546	20/22	0/22	0/22
Thorium-234	1.41	ND	3/70	N/A	N/A
Uranium-234	ND	0.255	2/27	0/27	0/27
Uranium-238	ND	0.221	3/27	0/27	0/27

^a Frequency of detection is the number of detections of an analyte per number of analyses (includes regular and duplicate samples).

N/A = not applicable

ND = not detected



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Figure 4.56. SWMU 6 Vanadium in Soil (Looking North)

Background and NALs taken from DOE 2001.

Table 4.31. SWMU 6 Locations of Subsurface Soil Contaminants

Analysis	Depth (ft)	Excavation Worker NAL	Back ground	RIData							Historical Data									
				006-101	006-102	006-103	006-104	006-016	006-017	006-018	006-019	006-020	006-021	006-022	006-023	006-026	006-027	006-028	006-029	
Inorganics (mg/kg)	Aluminum	05	5,250	12,000	7,150	8,540	6,740	7,690	7,430	8,620	10,400	9,010	9,900	9,330	9,720	10,500	9,630	8,240	8,970	
		10-15	5,250	12,000													4,660	4,370	6,120	9,760
		15-20	5,250	12,000													3,510	10,300		
		20-25	5,250	12,000					3,630	5,080	5,420					8,730	4,060		4,260	
		30-35	5,250	12,000	5,400	8,630	4,970	6,270	3,600			10,700	11,300			4,060	7,460	3,910		
		35-40	5,250	12,000													13,100			
		40-45	5,250	12,000	7,740	1,890	8,750	5,940				17,300	22,500	13,100	4,960	11,100	10,300	12,100		
		50-55	5,250	12,000	2,900	6,630	11,300	5,830						11,300	2,230	8,750				
		60	5,250	12,000															5U	5.76
			05	0.324	7.9															5U
Arsenic	10-15	0.324	7.9	1.92	5.94	2.59	2.54	5U	5U	5U	5U	5U	5U	5U	5U	5U	5U	5U	5U	
	15-20	0.324	7.9																	
	20-25	0.324	7.9					5.5	5U	5U										
	30-35	0.324	7.9	2.38	2.76	3.15	3.24				5U	5U								
	35-40	0.324	7.9					5U												
	40-45	0.324	7.9	2.23	1.17	0.894U	0.967U				5U	5U	5U	5U	5U	5U	5U	5U	5U	
	50-55	0.324	7.9	0.935U	1.47	2.43	1.41													
	60	0.324	7.9																	
	05	272	170																	
	10-15	272	170	83.4	84.2	97.5	86.1	75.5	76.6	74.2	119	84.9	63	65.9	83.5	88.6	77.8	86.3		
15-20	272	170									28.2	35.9	25.8		17.6	67.6	98.6			
20-25	272	170													40.6	19.5				
30-35	272	170	14.6	18.4	13.9	19.1	17.3	20.7	37.8	25.1	29.4			17	14.1	16.3				
35-40	272	170									10.9				15.3	13.1				
40-45	272	170	52.5	7.61	63.8	39.3				153	114	69.4	65.4	104	108					
50-55	272	170	44.5	43.9	53.7	44.4					59.1	48.6	55.2	50.6						
60	272	170										32.8	51.1							
05	1.26	0.69																		
10-15	1.26	0.69	0.477U	0.479U	0.461U	0.479U	0.5U	0.5U	0.5U	0.5U	0.5U	0.5U	0.5U	0.5	0.56	0.59	2.62	0.5U		
15-20	1.26	0.69														0.5U	0.64	0.59		
20-25	1.26	0.69														0.5U				
30-35	1.26	0.69	0.478U	0.482U	0.448U	0.442U	0.74	1.51	1.17	0.81	0.88	0.88	0.52	0.67	0.59	1	0.5U			
35-40	1.26	0.69					0.5U									0.5U	0.6			
40-45	1.26	0.69	0.825	0.45U	0.447U	0.484U										0.5U				
50-55	1.26	0.69	0.467U	0.463U	0.451U	0.456U										0.5U	1.7			
60	1.26	0.69																		
05	n/a	6,100																		
10-15	n/a	6,100	950	1,370	961	945	1,180	1,180	1,410	1,070	980	4,940	5,800	14,900	1,530	1,930	463	1,800		
15-20	n/a	6,100										592	833	604		384	866	1,530		
20-25	n/a	6,100														264				
30-35	n/a	6,100	455	593	595	725	423	284	551	933	970		499		635	264	1,400			

Table 4.31. SWMU 6 Locations of Subsurface Soil Contaminants (Continued)

Analysis	Depth (ft)	Excavation Worker NAL	Back ground	RI Data							Historical Data								
				006-101	006-102	006-103	006-104	006-016	006-017	006-018	006-019	006-020	006-021	006-022	006-023	006-026	006-027	006-028	006-029
Calcium	35-40	n/a	6,100					362											
	40-45	n/a	6,100	1,370	143	1,410	989		1,820	2,150	1,600	1,830	1,770	455	270				
	50-55	n/a	6,100	192	424	888	800			1,400	512	1,130	1,380	1,380	1,270				
	60	n/a	6,100								273	1,040							
	05	476	43														17.5	9.06	
Chromium	10-15	476	43	13	19	10.2	10.4	12.2	12	20.5	11.3	13	13.3	15.5	15.3			8.17	12.8
	15-20	476	43												6.43	4.32			
	20-25	476	43					43.2	46.8	55.9				8.85	56.8	12.8			
	30-35	476	43	6.37	10.1	9.81	9.94				12.2	13.7		6.5		9.48			
	35-40	476	43					3.38							5.58	116			
Cobalt	40-45	476	43	16.6	13.7	7.43	8.71				19.2	21.3	19.8	14.4					
	50-55	476	43	5.32	12.4	9.2	16.8					18.1	13.5	18.1	23.1				
	60	476	43																
	05	1,110	13															6.81	4.57
	10-15	1,110	13	2.86	4.23	4.57	4.45	2.87	4.14	10.8	17.9	4.16	2.73	2.7	2.35			2.27	2.93
Copper	15-20	1,110	13									4.5	3.63	2U	1.98	2.02			
	20-25	1,110	13					2.86	7.53	8.47					4.48	1.96			
	30-35	1,110	13	3.06	2.41U	2.24U	2.21U				2.2	6.56		2.44		1.94			
	35-40	1,110	13					2.13							1.3	8.05			
	40-45	1,110	13	2.4U	2.25U	16.4	2.71				25	156	2.76	19.8	18.5				
Iron	50-55	1,110	13	3.08	2.71	2.26U	3.11					5.26	2.47	11.5	3.43				
	60	1,110	13										2.8	2.02					
	05	427	25															19.2	7.45
	10-15	427	25	6.28	13	7.28	6.74	5.49	4.97	7.81	8.05	7.13	6.53	20.9	5.77			5.63	7.61
	15-20	427	25									3.92	6.74	5.25	2U	2.02			
Iron	20-25	427	25					4.43	3.01	3.96					3.39	3.53			
	30-35	427	25	3.52	4.87	4.93	6.4				4.85	5.42		4.43					
	35-40	427	25					2U							2.51	2.46			
	40-45	427	25	10.2	2.46	4.73	2.55				9.94	10.8	13.8	9.78					
	50-55	427	25									7.57	3.42	8.12	9.7	9.76			
Iron	60	427	25	2,34U	5.17	7.74	5.27							6.15					
	05	2,170	28,000															54,200	10,700
	10-15	2,170	28,000	8,830	14,500	9,660	10,400	10,500	9,280	35,300	10,800	10,700	9,210	9,880	10,800			10,700	11,800
	15-20	2,170	28,000									7,520	10,000	11,500	6,270	4,160			
	20-25	2,170	28,000					24,800	58,700	33,700					11,500	20,200	9,200		
Iron	30-35	2,170	28,000	11,700	14,100	17,200	19,100				18,200	17,300			7,460			16,800	
	35-40	2,170	28,000					3,720							5,170	22,200			
	40-45	2,170	28,000	16,900	4,850	5,910	4,260				20,800	26,700	29,900	23,000	20,100				
	50-55	2,170	28,000									22,200	5,010	17,300	32,900	36,900			
	60	2,170	28,000	5,340	9,410	15,400	8,810						5,140	1,180					

Table 4.31. SWMU 6 Locations of Subsurface Soil Contaminants (Continued)

Analysis	Depth (ft)	Excavation Worker NAL	Back ground	RI Data							Historical Data								
				006-101	006-102	006-103	006-104	006-016	006-017	006-018	006-019	006-020	006-021	006-022	006-023	006-026	006-027	006-028	006-029
Lead	05	50	23																
	10-15	50	23	5.81	8.8	6.9	9.08	20U	20U	20U	20U	20U	20U	20U	20U	20U	20U	20U	
	15-20	50	23																
	20-25	50	23					20U	20U	20U	20U	20U	20U	20U	20U	20U	20U	20U	
	30-35	50	23	5.43	6.41	4.4	5.89												
	35-40	50	23					20U											
	40-45	50	23	5.88	2.34	8.92	3.7												
	50-55	50	23																
	60	50	23	2.27	4.31	4.73	4.45												
		05	n/a	2,100															
Magnesium	10-15	n/a	2,100	994	1,400	1,060	967	1,210	1,280	1,040	1,320	1,460	1,220	2,370	1,180	1,260	644	855	
	15-20	n/a	2,100									409	768	477	501	299	819	1,440	
	20-25	n/a	2,100					210	196	344					501	203			
	30-35	n/a	2,100	245	336	291	328				467	532			227	726		494	
	35-40	n/a	2,100					188								309		133	
	40-45	n/a	2,100	1,230	72	764	552				1,670	1,710	1,560	1,740	1,540				
	50-55	n/a	2,100									1,160	522	1,120	1,350	1,090			
	60	n/a	2,100	219	568	1,270	586						188	1,290					
	05	56.6	820																
	10-15	56.6	820	150	216	282	309	106	207	411	7,550	230	164	222	165	93.4	353	349	
15-20	56.6	820																	
20-25	56.6	820																	
30-35	56.6	820	103	57.1	19.7	27.1													
35-40	56.6	820					39.7												
40-45	56.6	820	59.2	35.6	150	60.5													
50-55	56.6	820																	
60	56.6	820	315	132	98.6	149													
05	216	22																	
10-15	216	22	9.65	12.9	10.7	10.5	7.35	9.61	5U	11.2	7.23	7.21	7.2	15.4	30	5U	5U	7.71	
15-20	216	22																	
20-25	216	22					7.38	5U	5U										
30-35	216	22	4.78U	4.82U	4.48U	4.42U													
35-40	216	22					5U												
40-45	216	22	11.8	4.5U	6.16	4.84U													
50-55	216	22																	
60	216	22	7.38	7.15	10.8	6.92													
05	n/a	950																	
10-15	n/a	950					305	374	308	424	514	265	387	387	407	366	403		
15-20	n/a	950																	
20-25	n/a	950					100U	100U	144		196	311	215	158	136				
30-35	n/a	950																	
35-40	n/a	950																	

Table 4.31. SWMU 6 Locations of Subsurface Soil Contaminants (Continued)

Analysis	Depth (ft)	Excavation Worker NAL	Back ground	RI Data							Historical Data							
				006-101	006-102	006-103	006-104	006-016	006-017	006-018	006-019	006-020	006-021	006-022	006-023	006-026	006-027	006-028
Organics--Semivolatiles (mg/kg)																		
Bis(2-ethylhexyl) phthalate	05	10.1	n/a															
	10-15	10.1	n/a					0.5U	0.5U	0.5U	0.5U	0.46U	0.46U	0.5U		0.5U	0.5U	
	15-20	10.1	n/a					0.5U	0.5U	0.5U		0.46U	0.47U	0.5U	0.5U			
	20-25	10.1	n/a					0.5U	0.5U					0.5U	0.5U			
	30-35	10.1	n/a					0.5U						0.5U	0.5U	0.5U		
	35-40	10.1	n/a											0.5U				
	40-45	10.1	n/a										0.47U	0.47U	0.6			
	50-55	10.1	n/a										0.46U	0.49	0.47U	0.5U	0.5U	
	05	1.520	n/a															
	10-15	1.520	n/a					1.7	0.5U	0.5U			0.46U	0.74	0.86	0.72	0.5U	0.67
15-20	1.520	n/a											1.5	0.92		0.5U	0.5U	
20-25	1.520	n/a					0.5U	0.5U							0.5U	0.5U		
30-35	1.520	n/a										0.86	0.48U			0.77	0.5U	
35-40	1.520	n/a																
40-45	1.520	n/a					0.5U					0.47U	1	1.6	2	0.5U	0.5U	
50-55	1.520	n/a										0.46U	3.2	0.82	0.5U	0.66		
Organics-- Volatiles (mg/kg)																		
Acetone	10-15	421	n/a					0.01U	0.01U	0.01U	0.01U	0.01U	0.01U	0.01U	0.01U	0.01U	0.01U	
	15-20	421	n/a									0.01U	0.01U	0.013				
	20-25	421	n/a					0.01U	0.01U	0.01U					0.085			
	30-35	421	n/a					0.01U	0.01U						0.01U			
	35-40	421	n/a					0.01U										
	40-45	421	n/a															
	50-55	421	n/a									0.039	0.01U	0.01U	0.01U	0.01U		
	60	421	n/a															
	05	3.25	n/a															0.392U
	10-15	3.25	n/a					0.337U	0.393U	0.418U	0.01U	0.0011	0.0038	0.01U	0.394U	0.351U	0.444U	0.351U
15-20	3.25	n/a										0.01U	0.0025	0.304U	0.28U	0.316U		
20-25	3.25	n/a					0.617U	0.274U	0.286U					0.369U	0.333U	0.294U		
30-35	3.25	n/a														0.444U		
35-40	3.25	n/a					0.357U							0.323U	0.405U			
40-45	3.25	n/a																
50-55	3.25	n/a												0.249U	0.327U			
60	3.25	n/a										0.0012U	0.01U	0.01U	0.01U			
Radionuclides (pCi/g)																		
Radium-226	10-15	0.033	1.5					0.58U	0.528U	0.575U	0.48U	0.555U						
	15-20	0.033	1.5					0.741	0.71	0.506U	0.506U							
	30-35	0.033	1.5					0.586U	0.555U			0.519U						
	40-45	0.033	1.5					0.907	0.605U	0.636U	0.515U	0.748U						

4.7.2 SWMU 6 Groundwater

UCRS groundwater samples were collected from all of the four angled borings installed at SWMU 6 as part of this RI. The WAG 3 RI sampled UCRS groundwater from 15 borings at SWMU 6 (Figure 4.13) (DOE 2000a). RI data were reviewed with historical data (primarily from the WAG 3 RI) to determine the contaminants listed in Table 4.32.

The SWMU 6 disposal pits are approximately 6 to 8 ft deep. Two temporary borings of the WAG 3 RI (006-011 and 006-012) provided groundwater samples from directly below the pits, from depths of 9–12 ft within the area of Pit J (used for contaminated aluminum scrap disposal). Metals, notably iron, and the radionuclides neptunium-237, technetium-99, uranium-234, and uranium-238 exceeded screening criteria. Samples from both borings contained PCB-1016, at levels of 0.05 to 0.26 mg/L. These were the only occurrences of organic contaminants at levels that exceed screening criteria in the UCRS groundwater samples from SWMU 6. UCRS groundwater samples from the locations 006-101, which angled beneath “Pit J,” and 006-029, which sampled directly north of “Pit J,” contained the highest levels of beryllium, cadmium, iron, lead, manganese, and mercury.

The HU2 interval of the UCRS occurs at approximate depths of 20 to 30 ft beneath SWMU 6. Groundwater samples from borings 006-017, 006-018, 006-025, 006-029, and 006-104 characterized contaminant levels in the HU2 interval. The samples from 006-018 and 006-104 were collected from directly below Area J. As discussed above, metals exceeded screening criteria with notably elevated levels of iron. The radionuclides technetium-99, uranium-234, and uranium-238 also exceeded screening criteria. These same contaminant trends persist through the HU3 interval. Ten temporary borings sampled groundwater in the HU3 interval: 006-019, 006-020, 006-021, 006-022, 006-023, and 006-028. (See Figure 4.13 for the location of these borings/sampling locations.)

RGA and McNairy groundwater samples were not collected at SWMU 6 as part of this RI; however, historical data were reviewed to identify the contaminants listed in Table 4.33. (The screen of analyses of McNairy groundwater samples from SWMU 6 determined that no groundwater contaminants are present.) RGA and McNairy groundwater samples were collected from the locations 006-025, north of the SWMU, and 006-024, located to the southeast. For those metals detected above screening levels in the RGA (all but manganese) and for TCE (the lone organic contaminant), the higher contaminant levels represent samples from 006-025. Iron and manganese continued to be the most common metals to exceed screening levels. TCE levels were greater than its MCL in nearly all RGA samples. The presence of TCE is due to the Northwest Plume; the west side of the plume passes beneath SWMU 6 (Figure 4.54). Boring 006-025 is located closer to the center of the Northwest Plume than boring 006-024.

Table 4.34 provides detail (depth, sample location, and analytical results) for SWMU 6 groundwater samples, including nondetects and detections above screening levels.

Table 4.32. SWMU 6 UCRS Groundwater Contaminants

Analysis	Historical Data	RI	Frequency of Detection ^a	Frequency of Detection		
				Above Background	Above NAL	Above MCL
Metals (mg/L)						
Aluminum	1,650	1,430	29/41	N/A	21/41	N/A
Arsenic	0.014	0.15	9/12	N/A	9/12	3/12
Arsenic, Dissolved	N/A	0.0021	2/2	N/A	2/2	0/2
Barium	9.59	42.9	36/41	N/A	35/41	4/41
Barium, Dissolved	N/A	0.545	2/2	N/A	2/2	0/2
Beryllium	0.09	0.0929	6/25	N/A	6/25	6/25
Cadmium	0.039	0.0288	5/21	N/A	5/21	3/21
Calcium	892	339	41/41	N/A	N/A	N/A
Calcium, Dissolved	N/A	109	2/2	N/A	N/A	N/A
Chromium	3	3.32	12/39	N/A	2/39	11/39
Cobalt	1.2	6.68	29/39	N/A	6/39	N/A
Cobalt, Dissolved	N/A	0.333	2/2	N/A	1/2	N/A
Copper	0.9	ND	10/37	N/A	9/37	0/37
Iron	2,640	2,110	35/41	N/A	33/41	N/A
Iron, Dissolved	N/A	61.1	2/2	N/A	2/2	N/A
Lead	2.03	2.02	7/7	N/A	5/7	5/7
Magnesium	190	192	41/41	N/A	0/41	N/A
Magnesium, Dissolved	N/A	45.3	2/2	N/A	0/2	N/A
Manganese	93	170	38/41	N/A	38/41	N/A
Manganese, Dissolved	N/A	5.44	2/2	N/A	2/2	N/A
Mercury	0.003	0.00279	7/25	N/A	4/25	2/25
Molybdenum	N/A	0.359	4/4	N/A	4/4	N/A
Molybdenum, Dissolved	N/A	0.154	2/2	N/A	2/2	N/A
Nickel	0.69	0.953	12/33	N/A	12/33	N/A
Nickel, Dissolved	N/A	0.0845	2/2	N/A	2/2	N/A
Potassium	40	N/A	18/37	N/A	N/A	N/A
Selenium, Dissolved	N/A	0.00574	1/2	N/A	0/2	0/2
Silver	ND	0.00367	2/17	N/A	0/17	N/A
Sodium	194	141	41/41	N/A	0/41	N/A
Sodium, Dissolved	N/A	139	2/2	N/A	0/2	N/A
Thallium	ND	0.0174	2/17	N/A	0/17	2/17
Uranium	N/A	0.315	3/4	N/A	3/4	2/4
Vanadium	3.34	ND	11/39	N/A	11/39	N/A
Zinc	4.16	10.8	17/37	N/A	11/37	N/A
Zinc, Dissolved	N/A	0.382	2/2	N/A	0/2	N/A
Radionuclides (pCi/L)						
Bismuth-214	N/A	529	4/4	N/A	N/A	N/A
Lead-214	N/A	509	4/4	N/A	N/A	N/A
Neptunium-237	219	ND	1/14	N/A	1/14	N/A
Technetium-99	2,920	104	14/16	N/A	14/16	6/16
Thorium-228	N/A	0.906	3/4	N/A	3/4	N/A
Thorium-230	N/A	0.299	1/4	N/A	0/4	N/A
Thorium-232	N/A	0.176	2/4	N/A	0/4	N/A
Uranium	2,320	46.8	3/5	N/A	N/A	N/A
Uranium-234	754	24	3/5	N/A	3/5	N/A
Uranium-235	N/A	1.07	5/4	N/A	N/A	N/A
Uranium-238	1,520	21.8	4/5	N/A	4/5	N/A

Table 4.32. SWMU 6 UCRS Groundwater Contaminants (Continued)

Analysis	Historical Data	RI	Frequency of Detection ^a	Frequency of Detection		
				Above Background	Above NAL	Above MCL
<i>PCBs (mg/L)</i>						
PCB-1016	0.255	ND	2/10	N/A	2/10	2/10
<i>Volatiles (mg/L)</i>						
Acetone	0.013	N/A	1/7	N/A	0/7	N/A
Trichloroethene	0.0002	N/A	1/16	N/A	0/16	0/16

^a Frequency of detection is the number of detections of an analyte per number of analyses (includes regular and duplicate samples).

N/A = not applicable

ND = not detected

Table 4.33. SWMU 6 RGA Groundwater Contaminants

Analysis	Historical Data	RI	Frequency of Detection ^a	Frequency of Detection		
				Above Background	Above NAL	Above MCL
Metals (mg/L)						
Aluminum	250	N/A	19/42	14/42	14/42	N/A
Arsenic	0.013	N/A	3/3	1/3	3/3	1/3
Barium	3.1	N/A	41/42	11/42	27/42	3/42
Beryllium	0.029	N/A	3/21	3/21	3/21	3/21
Cadmium	0.015	N/A	2/19	2/19	2/19	2/19
Calcium	127	N/A	42/42	1/42	N/A	N/A
Chromium	0.76	N/A	4/22	1/22	0/22	1/22
Cobalt	0.118	N/A	21/39	7/39	3/39	N/A
Iron	2,210	N/A	35/42	15/42	30/42	N/A
Lead	0.788	N/A	1/1	1/1	1/1	1/1
Magnesium	63.5	N/A	42/42	1/42	0/42	N/A
Manganese	27.1	N/A	42/42	42/42	42/42	N/A
Mercury	0.0012	N/A	1/18	1/18	1/18	0/18
Nickel	0.414	N/A	4/20	0/20	4/20	N/A
Potassium	38.1	N/A	14/32	2/32	N/A	N/A
Sodium	21.7	N/A	42/42	0/42	0/42	N/A
Vanadium	1.24	N/A	1/18	1/18	1/18	N/A
Zinc	0.406	N/A	6/25	6/25	0/25	N/A
Radionuclides (pCi/L)						
Technetium-99	119	N/A	9/18	7/18	9/18	0/18
Volatiles (mg/L)						
<i>cis</i> -1,2-Dichloroethene	0.006	N/A	10/18	N/A	1/18	0/18
<i>trans</i> -1,2-Dichloroethene	0.0003	N/A	5/18	N/A	0/18	0/18
Trichloroethene	0.74	N/A	18/18	N/A	17/18	15/18

^a Frequency of detection is the number of detections of an analyte per number of analyses (includes regular and duplicate samples).

N/A = not applicable

Table 4.34. SWMU 6 Locations of Groundwater Contaminants

Hydro Unit	Depth Interval	Analysis	Child Resident NAL	Bkgd	MCL	006-101	006-102	006-103	006-104	006-009	006-011	006-012	006-016	006-017	006-018	006-021	006-022	006-024	006-025	006-028	006-029
	9-12	Metals (mg/L)																			
		Aluminum	1.49	N/A	N/A						29.2	91.3									
		Arsenic	3.50E-05	N/A	0.01						0.005	0.005									
		Barium	0.104	N/A	2						0.134	0.683									
		Beryllium	0.00264	N/A	0.004						0.005U	0.005U									
		Cadmium	0.00066	N/A	0.005						0.01U	0.01U									
		Calcium	N/A	N/A	N/A						29.1	36.5									
		Chromium	1.76	N/A	0.1						0.05U	0.087									
		Cobalt	0.0906	N/A	N/A						0.01U	0.028									
		Copper	0.0557	N/A	1.3						0.05U	0.052									
		Iron	0.449	N/A	N/A						17.6	65.3									
		Magnesium	N/A	N/A	N/A						2.81	10.1									
		Manganese	0.035	N/A	N/A						0.43	1.98									
		Mercury	0.00044	N/A	0.002						0.0002U	0.0002U									
		Nickel	0.0301	N/A	N/A						0.05U	0.061									
		Potassium	N/A	N/A	N/A						29.5	37.9									
		Silver	0.0075	N/A	N/A						0.05U	0.05U									
		Sodium	N/A	N/A	N/A						70.4	84.4									
		Thallium	N/A	N/A	0.002						0.2U	0.2U									
		Vanadium	0.00925	N/A	N/A						0.1U	0.137									
		Zinc	0.45	N/A	N/A						0.2U	0.259									
		PCBs (mg/L)																			
		PCB-1016	4.70E-05	N/A	0.0005						0.255	0.053									
		Radionuclides (pCi/L)																			
		Neptunium-237	0.573	N/A	N/A						219	57.5U									
		Technetium-99	14	N/A	900						1.810	2.920									
		Uranium	N/A	N/A	N/A						2.320										
		Uranium-234	0.546	N/A	N/A						754										
		Uranium-238	0.443	N/A	N/A						1.520										
		Volatiles (mg/L)																			
		Trichloroethene	0.0016	N/A	0.005						0.001U	0.001U									
	21-22	Metals (mg/L)																			
		Aluminum	1.49	N/A	N/A									1.13							1.650
		Arsenic	3.50E-05	N/A	0.01									0.181							0.005
		Barium	0.104	N/A	2									0.005U							9.59
		Beryllium	0.00264	N/A	0.004									0.01U							0.09
		Cadmium	0.00066	N/A	0.005									0.01U							0.039
		Calcium	N/A	N/A	N/A									52.9							892
		Chromium	1.76	N/A	0.1									0.05U							3
		Cobalt	0.0906	N/A	N/A									0.017							1.2
		Copper	0.0557	N/A	1.3									0.05U							0.9
		Iron	0.449	N/A	N/A									0.869							2.640
		Lead	0.015	N/A	0.015																2.03
		Magnesium	N/A	N/A	N/A																190
		Manganese	0.035	N/A	N/A									1.54							93

Table 4.34. SWMU 6 Locations of Groundwater Contaminants (Continued)

Hydro Unit	Depth Interval	Analysis	Child Resident NAL	Bkgd	MCL	006-101	006-102	006-103	006-104	006-009	006-011	006-012	006-016	006-017	006-018	006-021	006-022	006-024	006-025	006-028	006-029
	27-30	Vanadium	0.00925	N/A	N/A			2U							1.02				0.563		
		Zinc	0.45	N/A	N/A			9.19							1.02				0.536		
		Zinc, Dissolved	0.45	N/A	N/A			0.382													
		Radionuclides (pCi/L)																			
		Bismuth-214	N/A	N/A	N/A				198												
		Lead-214	N/A	N/A	N/A				200												
		Neptunium-237	0.573	N/A	N/A			0.067U							30.2U						
		Technetium-99	14	N/A	900			17.3							55				14U		
		Thorium-228	0.129	N/A	N/A			0.866													
		Thorium-230	0.424	N/A	N/A			0.299													
		Thorium-232	0.382	N/A	N/A			0.176													
		Uranium	N/A	N/A	N/A			11.5													
		Uranium-234	0.546	N/A	N/A			5.53													
		Uranium-235	N/A	N/A	N/A			0.253													
		Uranium-238	0.443	N/A	N/A			5.76													
		Volatiles (mg/L)																			
		Trichloroethene	0.0016	N/A	0.005										0.001U						0.001U
	35-37	Metals (mg/L)																			
		Aluminum	1.49	N/A	N/A								62.4								
		Barium	0.104	N/A	2								0.496								
		Beryllium	0.00264	N/A	0.004								0.005U								
		Cadmium	0.00066	N/A	0.005								0.01U								
		Calcium	N/A	N/A	N/A								83.8								
		Chromium	1.76	N/A	0.1								0.125								
		Cobalt	0.0906	N/A	N/A								0.06								
		Copper	0.0557	N/A	1.3								0.05U								
		Iron	0.449	N/A	N/A								78.4								
		Magnesium	N/A	N/A	N/A								31.8								
		Manganese	0.035	N/A	N/A								2.02								
		Mercury	0.00044	N/A	0.002								0.0002U								
		Nickel	0.0301	N/A	N/A								0.052								
		Potassium	N/A	N/A	N/A								3.24								
		Silver	0.0075	N/A	N/A								0.05U								
		Sodium	N/A	N/A	N/A								82.7								
		Thallium	N/A	N/A	0.002								0.2U								
		Vanadium	0.00925	N/A	N/A								0.162								
		Zinc	0.45	N/A	N/A								0.2U								
		Radionuclides (pCi/L)																			
		Neptunium-237	0.573	N/A	N/A								50.2U								
		Technetium-99	14	N/A	900								690								930
		Volatiles (mg/L)																			
		Acetone	0.0275	N/A	N/A																0.013
		Trichloroethene	0.0016	N/A	0.005								0.0002								0.001U

Table 4.34. SWMU 6 Locations of Groundwater Contaminants (Continued)

Hydro Unit	Depth Interval	Analysis	Child Resident NAL	Bkgd	MCL	006-101	006-102	006-103	006-104	006-016	006-017	006-018	006-021	006-022	006-024	006-025	006-028	006-029
	45-46	Metals (mg/L)																
		Aluminum	1.49	N/A	N/A	1,430	7.14	26.2										
		Arsenic	3.50E-05	N/A	0.01	0.5U	0.01U	0.01U										
		Arsenic, Dissolved	3.50E-05	N/A	0.01		0.0021											
		Barium	0.104	N/A	2	42.9	0.555	0.398										
		Barium, Dissolved	0.104	N/A	2		0.433											
		Beryllium	0.00264	N/A	0.004	0.0229	0.001U	0.001U										
		Cadmium	0.00066	N/A	0.005	0.0288	0.0006U	0.000837										
		Calcium	N/A	N/A	N/A	339	113	45.1										
		Calcium, Dissolved	N/A	N/A	N/A		109											
		Chromium	1.76	N/A	0.1	5U	0.1U	0.1U										
		Cobalt	0.0906	N/A	N/A	6.68	0.0671	0.0372										
		Cobalt, Dissolved	0.0906	N/A	N/A		0.0395											
		Copper	0.0557	N/A	1.3	10U	0.2U	0.2U										
		Iron	0.449	N/A	N/A	2,110	124	62.2										
		Iron, Dissolved	0.449	N/A	N/A		61.1											
		Lead	0.015	N/A	0.015	2.02	0.00336	0.00808										
		Magnesium	N/A	N/A	N/A	192	46.1	22.2										
		Magnesium, Dissolved	N/A	N/A	N/A		45.3											
		Manganese	0.035	N/A	N/A	170	4.35	1.89										
		Manganese, Dissolved	0.035	N/A	N/A		4.31											
		Mercury	0.00044	N/A	0.002	0.00279	0.00001U	0.000047										
		Molybdenum	0.00753	N/A	N/A	0.0231	0.182	0.0703										
		Molybdenum, Dissolved	0.00753	N/A	N/A		0.154											
		Nickel	0.0301	N/A	N/A	2.5U	0.14	0.0523										
		Nickel, Dissolved	0.0301	N/A	N/A		0.0845											
		Selenium, Dissolved	0.00754	N/A	0.05													
		Silver	0.0075	N/A	N/A	0.00223	0.001U	0.001U										
		Sodium	N/A	N/A	N/A	99.6	141	112										
		Sodium, Dissolved	N/A	N/A	N/A		139											
		Thallium	N/A	N/A	0.002	0.0174	0.002U	0.002U										
		Uranium	0.00091	N/A	0.03	0.315	0.001U	0.00275										
		Vanadium	0.00925	N/A	N/A	10U	0.2U	0.2U										
		Zinc	0.45	N/A	N/A	10.8	0.812	0.497										
		Zinc, Dissolved	0.45	N/A	N/A		0.123											
		PCBs (mg/L)																
		PCB-1016	4.70E-05	N/A	0.0005	0.00025U		0.0001U										
		Radionuclides (pCi/L)																
		Bismuth-214	N/A	N/A	N/A	279	307	529										
		Lead-214	N/A	N/A	N/A	282	290	509										
		Neptunium-237	0.573	N/A	N/A	0.22U	0.065U	0.0634U										
		Technetium-99	14	N/A	900			104										
		Thorium-228	0.129	N/A	N/A	0.906	0.146U	0.18										
		Thorium-230	0.424	N/A	N/A	0.411U	0.107U	0.11U										
		Thorium-232	0.382	N/A	N/A	0.266U	0.0706U	0.0667										

Table 4.34. SWMU 6 Locations of Groundwater Contaminants (Continued)

Hydro Unit	Depth Interval	Analysis	Child Resident	Bkgd	MCL	006-101	006-102	006-103	006-104	006-009	006-011	006-012	006-016	006-017	006-018	006-021	006-022	006-024	006-025	006-028	006-029		
UCRS	45-46	Uranium	N/A	N/A	N/A	46.8	0.524U	0.53IU															
		Uranium-234	0.546	N/A	N/A	24	0.358U	0.359U															
		Uranium-235	N/A	N/A	N/A	1.07	0.046U	0.0498U															
		Uranium-238	0.443	N/A	N/A	21.8	0.12U	0.558															
	60-61	Metals (mg/L)																					
		Aluminum	1.49	N/A	N/A	N/A											25.2	130					
		Arsenic	3.50E-05	N/A	N/A	0.01											0.006	0.013					
		Barium	0.104	N/A	2												0.496	1.34					
		Beryllium	0.00264	N/A	0.004												0.01U	0.015					
		Cadmium	0.00066	N/A	0.005												0.01U	0.01U					
		Calcium	N/A	N/A	N/A	N/A											62.4	73.2					
		Chromium	1.76	N/A	0.1												1.07	1.67					
		Cobalt	0.0906	N/A	N/A	N/A											0.029	0.17					
		Copper	0.0557	N/A	1.3												0.223	0.386					
		Iron	0.449	N/A	N/A	N/A											158	389					
		Magnesium	N/A	N/A	N/A	N/A											25.7	34.9					
		Manganese	0.035	N/A	N/A	N/A											1.33	9.56					
		Mercury	0.00044	N/A	0.002												0.0002U	0.0002					
		Nickel	0.0301	N/A	N/A	N/A											0.38	0.69					
		Potassium	N/A	N/A	N/A	N/A											3.19	11.4					
		Silver	0.0075	N/A	N/A	N/A											0.05U	0.05U					
		Sodium	N/A	N/A	N/A	N/A											80.5	51.4					
		Thallium	N/A	N/A	0.002												0.2U	0.2U					
Vanadium	0.00925	N/A	N/A	N/A											0.076	0.348							
Zinc	0.45	N/A	N/A	N/A											0.611	1.14							
PCBs (mg/L)																							
		PCB-1016	4.70E-05	N/A	0.0005											0.00018U	0.00018U						
Radionuclides (pCi/L)																							
		Neptunium-237	0.573	N/A	N/A											65.8U	61.1U						
		Technetium-99	14	N/A	900											956	46.8						
Volatiles (mg/L)																							
		Acetone	0.0275	N/A	N/A											0.01U	0.01U						
		Trichloroethene	0.0016	N/A	0.005											0.001U	0.001U						
RGA	68	Metals (mg/L)																					
		Aluminum	1.49	2.189	N/A															41.8	56.3		
		Barium	0.104	0.235	2												0.488	0.632					
		Beryllium	0.00264	0.004	0.004												0.005U	0.006					
		Cadmium	0.00066	0.01	0.005												0.01U	0.01U					
		Calcium	N/A	41.238	N/A												15	13.7					
		Chromium	1.76	0.144	0.1												0.05U	0.053					
		Cobalt	0.0906	0.045	N/A												0.062	0.07					
		Iron	0.449	5.03	N/A												63.8	88.8					
		Magnesium	N/A	16.262	N/A												7.87	8.54					
		Manganese	0.035	0.119	N/A												5.58	4.25					
		Mercury	0.00044	0.0002	0.002												0.0002U	0.0002U					

Table 4.34. SWMU 6 Locations of Groundwater Contaminants (Continued)

Hydro Unit	Depth Interval	Analysis	Child Resident NAL	Bkgd	MCL	006-101	006-102	006-103	006-104	006-009	006-011	006-012	006-016	006-017	006-018	006-021	006-022	006-024	006-025	006-028	006-029			
RGA	68	Nickel	0.0301	0.682	N/A													0.05	0.05U					
		Potassium	N/A	5.195	N/A														3.96	5.63				
		Sodium	N/A	59.45	N/A														15.4	17.6				
		Vanadium	0.00925	0.134	N/A														0.1U	0.1U				
		Zinc	0.45	0.054	N/A														0.2U	0.26				
		Radionuclides (pCi/L)																						
			Technetium-99	14	22.3	900													14U	14U				
			Volatiles (mg/L)																					
			cis-1,2-Dichloroethene	0.00273	N/A	0.07													0.001U	0.001U				
			trans-1,2-Dichloroethene	0.00548	N/A	0.1													0.001U	0.001U				
			Trichloroethene	0.0016	N/A	0.005													0.006	0.0008				
			Metals (mg/L)																					
			Aluminum	1.49	2.189	N/A													23.6	3.35				
			Barium	0.104	0.235	2													0.565	0.17				
			Beryllium	0.00264	0.004	0.004													0.005U	0.005U				
			Cadmium	0.00066	0.01	0.005													0.01U	0.01U				
			Calcium	N/A	41.238	N/A													15.9	14.1				
			Chromium	1.76	0.144	0.1													0.05	0.05U				
			Cobalt	0.0906	0.045	N/A													0.09	0.013				
			Iron	0.449	5.03	N/A													78.1	14.4				
			Magnesium	N/A	16.262	N/A													7.19	6.05				
		Manganese	0.035	0.119	N/A													9.36	1.24					
		Mercury	0.00044	0.0002	0.002													0.0002U	0.0002U					
		Nickel	0.0301	0.682	N/A													0.111	0.05U					
		Potassium	N/A	5.195	N/A													3.05	2U					
		Sodium	N/A	59.45	N/A													13.8	14.7					
		Vanadium	0.00925	0.134	N/A													0.1U	0.1U					
		Zinc	0.45	0.054	N/A													0.23	0.2U					
		Radionuclides (pCi/L)																						
		Technetium-99	14	22.3	900													16.9	25.1					
		Volatiles (mg/L)																						
		cis-1,2-Dichloroethene	0.00273	N/A	0.07													0.005U	0.001U					
		trans-1,2-Dichloroethene	0.00548	N/A	0.1													0.005U	0.001U					
		Trichloroethene	0.0016	N/A	0.005													0.005	0.01					
		Metals (mg/L)																						
		Aluminum	1.49	2.189	N/A													1.37	14					
		Barium	0.104	0.235	2													0.478	0.983					
		Beryllium	0.00264	0.004	0.004													0.005U	0.005U					
		Cadmium	0.00066	0.01	0.005													0.01U	0.01U					
		Calcium	N/A	41.238	N/A													13.7	18.6					
		Chromium	1.76	0.144	0.1													0.05U	0.05U					
		Cobalt	0.0906	0.045	N/A													0.019	0.114					
		Iron	0.449	5.03	N/A													9.28	144					
		Magnesium	N/A	16.262	N/A													5.76	8.21					
		Manganese	0.035	0.119	N/A													4.49	11.6					
		77-78																						

Table 4.34. SWMU 6 Locations of Groundwater Contaminants (Continued)

Hydro Unit	Depth Interval	Analysis	Child Resident NAL	Bkgd	MCL	006-101	006-102	006-103	006-104	006-009	006-011	006-012	006-016	006-017	006-018	006-021	006-022	006-024	006-025	006-028	006-029
	77-78	Mercury	0.00044	0.0002	0.002													0.0002U	0.0002U		
		Nickel	0.0301	0.682	N/A													2U	2.54		
		Potassium	N/A	5.195	N/A													12.7	14.6		
		Sodium	N/A	59.45	N/A													0.1U	0.1U		
		Vanadium	0.00925	0.134	N/A													0.2U	0.255		
		Zinc	0.45	0.054	N/A													14U	14U		
		Radionuclides (pCi/L)																			
		Technetium-99	14	22.3	900													0.001U	0.001U		
		Volatiles (mg/L)																			
		cis-1,2-Dichloroethene	0.00273	N/A	0.07													0.006	0.015		
		trans-1,2-Dichloroethene	0.00548	N/A	0.1													0.006	0.015		
		Trichloroethene	0.0016	N/A	0.005													0.006	0.015		
	82-83	Metals (mg/L)																			
		Aluminum	1.49	2.189	N/A													19.4	6.04		
		Arsenic	3.50E-05	0.005	0.01													0.509	0.212		
		Barium	0.104	0.235	2													0.005U	0.005U		
		Beryllium	0.00264	0.004	0.004													0.01U	0.01U		
		Cadmium	0.00066	0.01	0.005													17.4	19.4		
		Calcium	N/A	41.238	N/A													0.05U	0.05U		
		Chromium	1.76	0.144	0.1													0.046	0.019		
		Cobalt	0.0906	0.045	N/A													52.7	46.6		
		Iron	0.449	5.03	N/A													7.82	8.02		
		Magnesium	N/A	16.262	N/A													5.58	2.72		
		Manganese	0.035	0.119	N/A													0.0002U	0.0002U		
		Mercury	0.00044	0.0002	0.002													0.055	0.05U		
		Nickel	0.0301	0.682	N/A													2.3	2U		
		Potassium	N/A	5.195	N/A													14.1	15.6		
		Sodium	N/A	59.45	N/A													0.1U	0.1U		
		Vanadium	0.00925	0.134	N/A													0.2U	0.2U		
		Zinc	0.45	0.054	N/A													14U	16.3U		
		Radionuclides (pCi/L)																			
		Technetium-99	14	22.3	900													0.0002	0.0003		
		Volatiles (mg/L)																			
		cis-1,2-Dichloroethene	0.00273	N/A	0.07													0.001U	0.0002		
		trans-1,2-Dichloroethene	0.00548	N/A	0.1													0.019	0.031		
		Trichloroethene	0.0016	N/A	0.005													0.019	0.031		
	88	Metals (mg/L)																			
		Aluminum	1.49	2.189	N/A													4.3	5.18		
		Arsenic	3.50E-05	0.005	0.01													2.02	0.4		
		Barium	0.104	0.235	2													0.005U	0.005U		
		Beryllium	0.00264	0.004	0.004													0.01U	0.01U		
		Cadmium	0.00066	0.01	0.005													17.4	22.7		
		Calcium	N/A	41.238	N/A													0.05U	0.05U		
		Chromium	1.76	0.144	0.1													0.017	0.017		
		Cobalt	0.0906	0.045	N/A													139	77.8		
		Iron	0.449	5.03	N/A													7.06	9.48		
		Magnesium	N/A	16.262	N/A																

RGA

Table 4.34. SWMU 6 Locations of Groundwater Contaminants (Continued)

Hydro Unit	Depth Interval	Analysis	Child Resident NAL	Bkgd	MCL	006-101	006-102	006-103	006-104	006-009	006-011	006-012	006-016	006-017	006-018	006-021	006-022	006-024	006-025	006-028	006-029			
RGA	88	Manganese	0.035	0.119	N/A													15.6	4.4					
		Mercury	0.00044	0.0002	0.002														0.0002U	0.0002U				
		Nickel	0.0301	0.682	N/A														0.05U	0.05U				
		Potassium	N/A	5.195	N/A														2U	2U				
		Sodium	N/A	59.45	N/A														14.3	17.1				
		Vanadium	0.00925	0.134	N/A														0.1U	0.1U				
		Zinc	0.45	0.054	N/A														0.256	0.2U				
		Radionuclides (pCi/L)																						
				Technetium-99	14	22.3	900													15.3	15U			
				Volatiles (mg/L)																				
			<i>cis</i> -1,2-Dichloroethene	0.00273	N/A	0.07													0.0003	0.0006				
			<i>trans</i> -1,2-Dichloroethene	0.00548	N/A	0.1													0.001U	0.0002				
			Trichloroethene	0.0016	N/A	0.005													0.029	0.06				
			Metals (mg/L)																					
			Aluminum	1.49	2.189	N/A													23.2	8.32				
			Barium	0.104	0.235	2													3.1	1.19				
			Beryllium	0.00264	0.004	0.004													0.006	0.005U				
			Cadmium	0.00066	0.01	0.005													0.011	0.01U				
			Calcium	N/A	41.238	N/A													22	24.6				
			Chromium	1.76	0.144	0.1													0.054	0.05U				
		Cobalt	0.0906	0.045	N/A													0.114	0.034					
		Iron	0.449	5.03	N/A													245	112					
		Magnesium	N/A	16.262	N/A													10.8	9.86					
		Manganese	0.035	0.119	N/A													27.1	11.1					
		Mercury	0.00044	0.0002	0.002													0.0002U	0.0002U					
		Nickel	0.0301	0.682	N/A													0.05U	0.05U					
		Potassium	N/A	5.195	N/A													4.41	2.24					
		Sodium	N/A	59.45	N/A													15.7	18.3					
		Vanadium	0.00925	0.134	N/A													0.1U	0.1U					
		Zinc	0.45	0.054	N/A													0.406	0.204					
		Radionuclides (pCi/L)																						
		Technetium-99	14	22.3	900													14U	56					
		Volatiles (mg/L)																						
		<i>cis</i> -1,2-Dichloroethene	0.00273	N/A	0.07													0.0008	0.001					
		<i>trans</i> -1,2-Dichloroethene	0.00548	N/A	0.1													0.001U	0.0003					
		Trichloroethene	0.0016	N/A	0.005													0.1	0.17					
		Metals (mg/L)																						
		Aluminum	1.49	2.189	N/A													4.72	10.6					
		Barium	0.104	0.235	2													0.079	0.184					
		Beryllium	0.00264	0.004	0.004													0.005U	0.005U					
		Cadmium	0.00066	0.01	0.005													0.01U	0.01U					
		Calcium	N/A	41.238	N/A													19.9	32.1					
		Chromium	1.76	0.144	0.1													0.05U	0.05U					

Table 4.34. SWMU 6 Locations of Groundwater Contaminants (Continued)

Hydro Unit	Depth Interval	Analysis	Child Resident NAL	Bkgd	MCL	006-101	006-102	006-103	006-104	006-009	006-011	006-012	006-016	006-017	006-018	006-021	006-022	006-024	006-025	006-028	006-029		
RGA	98	Cobalt	0.0906	0.045	N/A													0.01U	0.025				
		Iron	0.449	5.03	N/A														52.4	71.9			
		Magnesium	N/A	16.262	N/A														9.11	14.9			
		Manganese	0.035	0.119	N/A														0.433	1.93			
		Mercury	0.00044	0.0002	0.002														0.0002U	0.0002U			
		Nickel	0.0301	0.682	N/A														0.05U	0.414			
		Potassium	N/A	5.195	N/A														2.58	3.79			
		Sodium	N/A	59.45	N/A														0.1U	0.1U			
		Vanadium	0.00925	0.134	N/A														0.2U	0.2U			
		Zinc	0.45	0.054	N/A																		
		Radionuclides (pCi/L)																					
				Technetium-99	14	22.3	900													50	119		
	Volatiles (mg/L)																						
			cis-1,2-Dichloroethene	0.00273	N/A	0.07													0.002	0.006			
			trans-1,2-Dichloroethene	0.00548	N/A	0.1													0.001U	0.0003			
			Trichloroethene	0.0016	N/A	0.005													0.27	0.74			
	103	Metals (mg/L)																					
				Aluminum	1.49	2.189	N/A																
				Arsenic	3.50E-05	0.005	0.01																250
				Barium	0.104	0.235	2																0.013
				Beryllium	0.00264	0.004	0.004																2.41
			Cadmium	0.00066	0.01	0.005																0.029	
			Calcium	N/A	41.238	N/A																0.015	
			Chromium	1.76	0.144	0.1																127	
			Cobalt	0.0906	0.045	N/A																0.76	
			Iron	0.449	5.03	N/A																0.118	
			Lead	0.015	0.129	0.015																2.210	
			Magnesium	N/A	16.262	N/A																0.788	
			Manganese	0.035	0.119	N/A																63.5	
			Mercury	0.00044	0.0002	0.002																14.5	
			Nickel	0.0301	0.682	N/A																0.0012	
			Potassium	N/A	5.195	N/A																0.5U	
			Sodium	N/A	59.45	N/A																38.1	
			Vanadium	0.00925	0.134	N/A																13.7	
			Zinc	0.45	0.054	N/A																1.24	
																						2U	
Radionuclides (pCi/L)																							
		Technetium-99	14	22.3	900																41		
Volatiles (mg/L)																							
		cis-1,2-Dichloroethene	0.00273	N/A	0.07																0.0002		
		trans-1,2-Dichloroethene	0.00548	N/A	0.1																0.001U		
		Trichloroethene	0.0016	N/A	0.005																0.01		

Blank cells indicate interval was not sampled for the specified analysis. Maximum value shown for each depth interval.

N/A = not applicable

Bold indicates result is greater than NAL value.

Italics indicates result is greater than MCL value.

Bold + Italics indicate result is greater than both NAL and MCL values.

Shading indicates result is greater than the background value.

4.8 SWMU 7

4.8.1 Subsurface Soils

SWMU 7 consists of six discrete burial pit areas containing uranium-contaminated concrete, uranium-contaminated scrap metal and equipment, and empty uranium and magnesium powder drums. The SWMU 7 burial pits range from 6 to 15 ft deep. Sampling locations are shown in Figure 4.6.

Table 4.35 summarizes the contaminants detected in the subsurface at SWMU 7.

Metals concentrations in subsurface soil samples of SWMU 7 rarely exceed background levels. Uranium metal has been detected above background levels only at three locations (WB-9, 007-009, and 007-010). These locations characterize burial pits B and C, which contained uranium-contaminated noncombustible trash. The highest concentration of uranium at these locations is 45 mg/kg. This level is greater than five times the next highest level of 8.94 mg/kg. Figure 4.57 shows the uranium distribution at SWMU 7 and indicates that the extent of contamination is limited to shallow soil depths (5 to 10 ft bgs).

The screening process identified two VOCs as contaminants at SWMU 7: vinyl chloride and 1,1-DCE. Both were detected infrequently (1 and 2 detections in 69 analyses, respectively) at levels above the excavation worker NALs. Figure 4.58 presents the vinyl chloride distribution in subsurface soils. The elevated levels of vinyl chloride occur in a small area in the central portion of SWMU 7 near burial pit C. It was detected at approximately four times the NAL screening concentration in boring 007-007 at a depth of 30 ft. Uranium-238 is the most widely detected radionuclide contaminant above PGDP background levels in subsurface soils at SWMU 7; the maximum uranium-238 result is 150 pCi/g from WBP-12A.

Total Uranium was detected as high as 240 pCi/g from WBP-9A. Figure 4.59 provides the uranium-238 distribution in soils at SWMU 7. As expected, it is very similar to the uranium distribution presented in Figure 4.56, with most exceedances limited to depths less than 15 ft bgs. Subsurface soil samples for Pit E (located outside of the SWMU 7 boundary) at 10 ft depth contained arsenic concentrations in excess of screening levels. None of the other Pit E analyses documented metals or radionuclides above screening levels or the presence of any organic contaminants. Two test pits were excavated in SWMU 7 during the Phase II SI (CH2M HILL 1992). The excavations noted oily sheens on material being excavated and six drums were removed from one of the test pits. Samples from these pits identified many of the same inorganic, organic, and radionuclide contaminants reported in Table 4.34. In the test pits, uranium-234 ranged from 5 to 580 pCi/g; uranium-238 ranged from 35 to 2,100 pCi/g; PCB-1248 ranged from 0.54 to 1.20 mg/kg; and PCB-1254 ranged from 1.2 to 2 mg/kg.

Table 4.36 presents the locations of subsurface contaminants detected above screening levels.

Table 4.35. SWMU 7 Subsurface Soil Contaminants

Analysis	Maximum Result		Frequency of Detection ^a	Frequency of Detection	
	Historical Data	RI Data		Above Background Value	Above Excavation Worker NAL
<i>Inorganics (mg/kg)</i>					
Aluminum	16,000	11,000	69/69	1/69	42/69
Arsenic	ND	13.9	50/69	1/69	50/69
Barium	180	150	69/69	1/69	0/69
Beryllium	ND	1.55	7/69	3/69	1/69
Cadmium	0.18	ND	2/69	0/69	0/69
Calcium	3,300	1,590	69/69	0/69	N/A
Chromium	20	32.8	67/69	0/69	0/69
Cobalt	9.4	17.7	42/69	2/69	0/69
Copper	27	13	56/69	1/69	0/69
Iron	17,000	34,700	69/69	1/69	65/69
Lead	12	15.3	68/69	0/69	0/69
Magnesium	1,800	1,590	69/69	0/69	0/69
Manganese	1,200	628	69/69	1/69	44/69
Mercury	ND	0.057	12/69	0/69	0/69
Nickel	40	16.1	44/69	1/69	0/69
Selenium	0.72	ND	4/69	1/69	0/69
Sodium	330	342	28/69	1/69	0/69
Tin	5.1	N/A	8/8	N/A	0/8
Uranium	45	8.94	12/69	3/69	1/69
Vanadium	31	34	62/69	0/69	56/69
Zinc	47	53.9	33/69	0/69	0/69
<i>Organics--Semivolatiles (mg/kg)</i>					
1,2,4-Trichlorobenzene	0.077	ND	2/69	N/A	0/69
1,4-Dichlorobenzene	0.07	ND	2/77	N/A	0/77
2,4,5-Trichlorophenol	0.035	ND	1/69	N/A	0/69
2,4,6-Trichlorophenol	0.041	ND	1/69	N/A	0/69
2-Chlorophenol	0.039	ND	1/69	N/A	0/69
2-Methylphenol	0.02	N/A	1/8	N/A	0/8
4-Methylphenol	0.016	N/A	1/8	N/A	0/8
Acenaphthene	0.031	ND	1/69	N/A	0/69
Di-n-butyl phthalate	0.091	0.91	2/69	N/A	0/69
Di-n-octylphthalate	0.072	ND	1/69	N/A	0/69
Hexachlorobutadiene	0.058	ND	1/69	N/A	0/69
Hexachloroethane	0.034	ND	1/69	N/A	0/69
Pyrene	0.033	ND	1/69	N/A	0/69
<i>Organics--Volatiles (mg/kg)</i>					
1,1,1-Trichloroethane	ND	0.159	2/69	N/A	0/69
1,1,2-Trichloroethane	ND	0.149	2/69	N/A	0/69
1,1-Dichloroethane	ND	0.378	4/69	N/A	0/69
1,1-Dichloroethene	ND	1.66	4/69	N/A	2/69
1,2-Dichloroethane	ND	0.0163	2/69	N/A	0/69
Acetone	0.014	0.0154	8/69	N/A	0/69
<i>cis</i> -1,2-Dichloroethene	ND	0.684	15/69	N/A	0/69
Tetrachloroethene	ND	0.0062	1/69	N/A	0/69
Toluene	ND	0.0926	1/69	N/A	0/69
Trichloroethene	ND	0.26	11/69	N/A	0/69

Table 4.35. SWMU 7 Subsurface Soil Contaminants (Continued)

Analysis	Maximum Result		Frequency of Detection ^a	Frequency of Detection	
	Historical Data	RI Data		Above Background Value	Above Excavation Worker NAL
Vinyl chloride	ND	0.585	5/69	N/A	1/69
Organics--PCBs (mg/kg)					
PCB, Total	0.41	2.45	5/69	N/A	2/69
PCB-1248	0.41	ND	1/69	N/A	1/69
PCB-1254	0.054	ND	3/69	N/A	0/69
PCB-1260	ND	2.45	1/69	N/A	1/69
Radionuclides(pCi/g)					
Neptunium-237	0.06	0.0316	3/67	N/A	0/67
Plutonium-239	0.13	N/A	2/8	N/A	0/8
Plutonium-239/240	N/A	0.136	1/59	N/A	0/59
Technetium-99	5.28	8.23	19/67	5/67	0/67
Thorium-228	N/A	1.38	57/59	0/59	57/59
Thorium-230	3.7	1.34	39/67	3/67	2/67
Thorium-232	N/A	0.746	53/59	0/59	0/59
Thorium-234	N/A	7.52	12/59	N/A	N/A
Uranium	240	6.87	14/68	N/A	N/A
Uranium-234	115	1.34	41/76	10/76	8/76
Uranium-235	N/A	0.119	4/59	0/59	0/59
Uranium-235/236	1.03	N/A	8/8	6/8	3/8
Uranium-238	150	5.87	35/76	22/76	22/76

^a Frequency of detection is the number of detections of an analyte per number of analyses

(includes regular and duplicate samples).

N/A = not applicable

ND = not detected

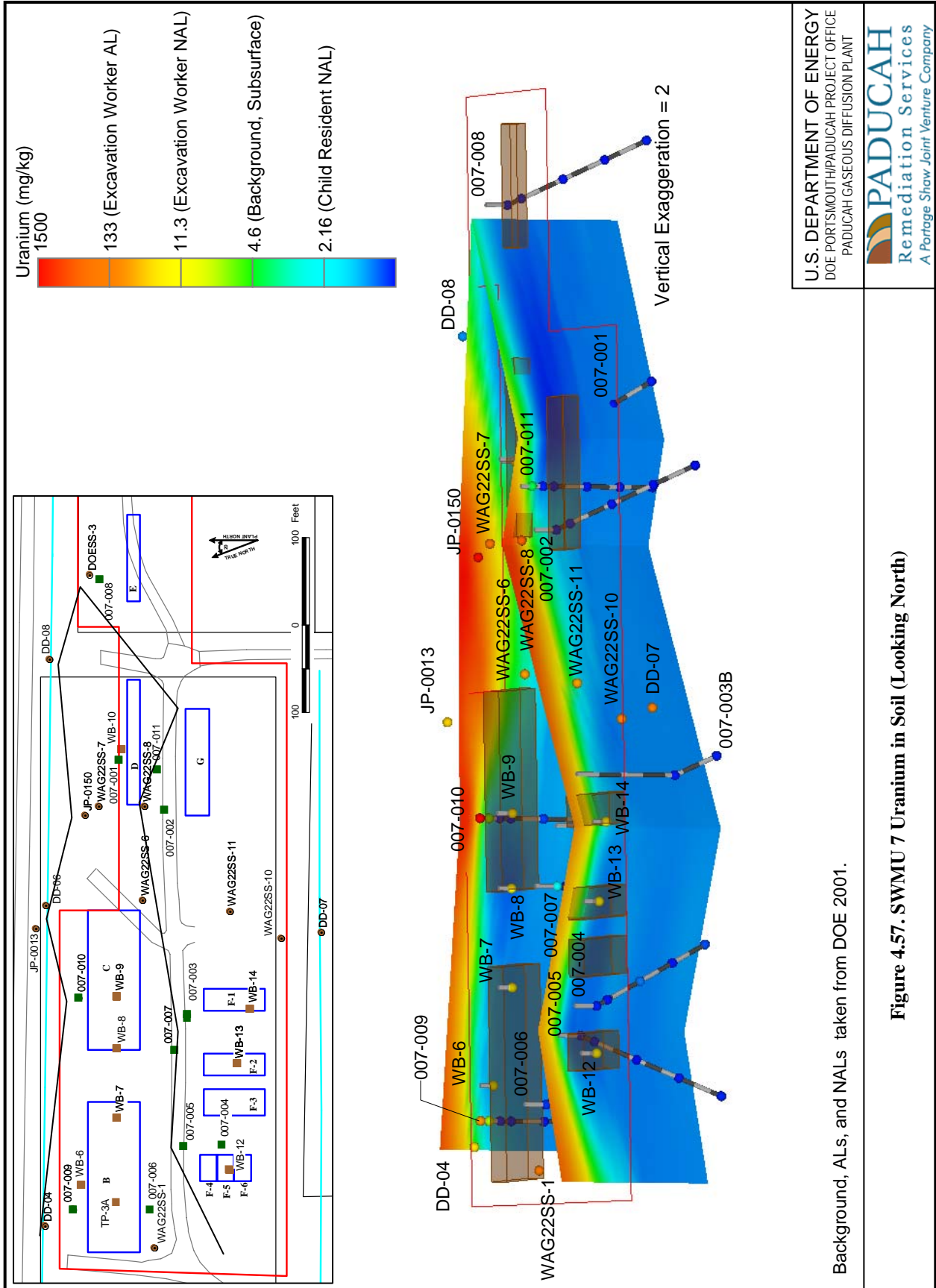
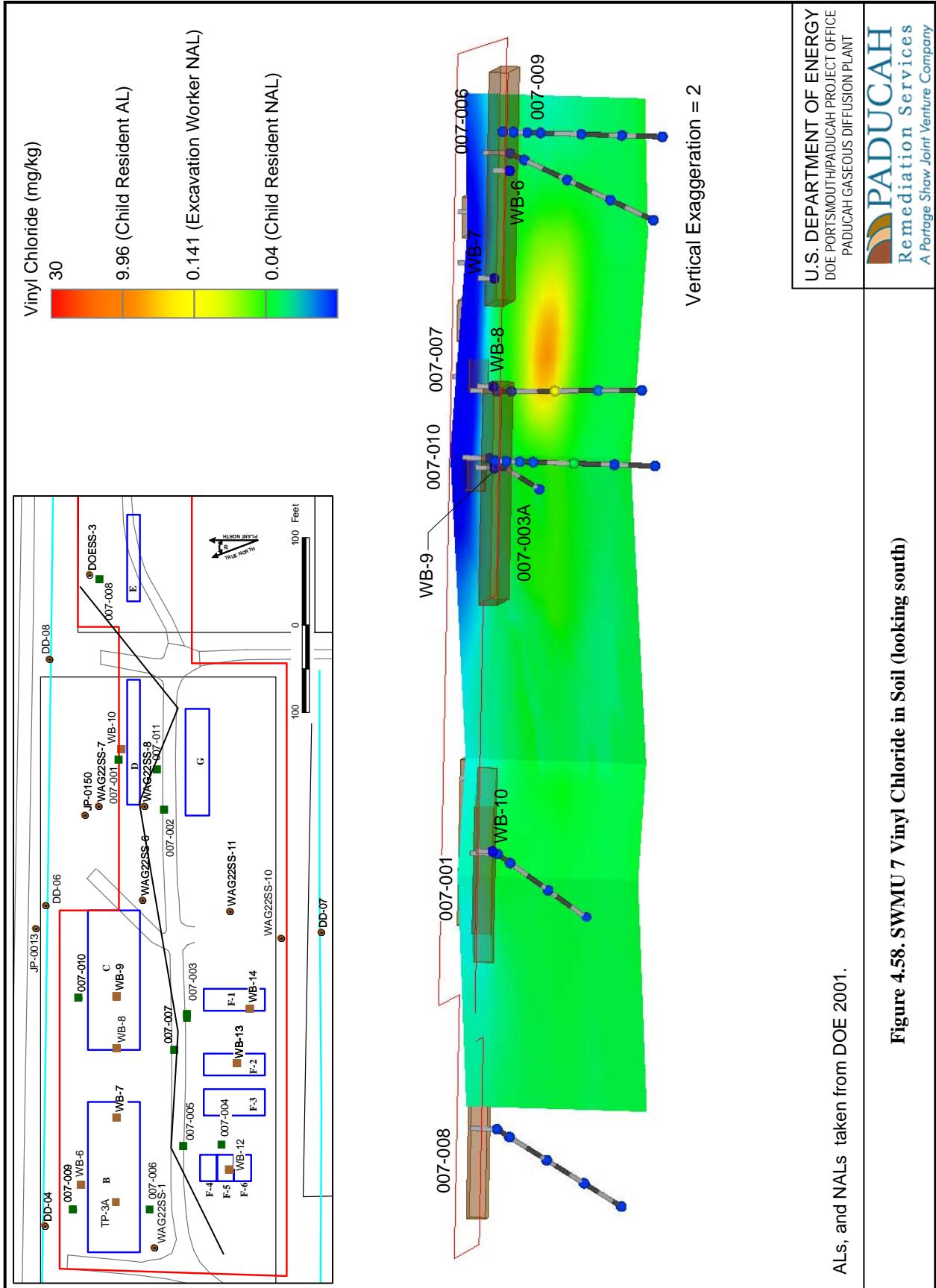


Figure 4.57. SWMU 7 Uranium in Soil (Looking North)



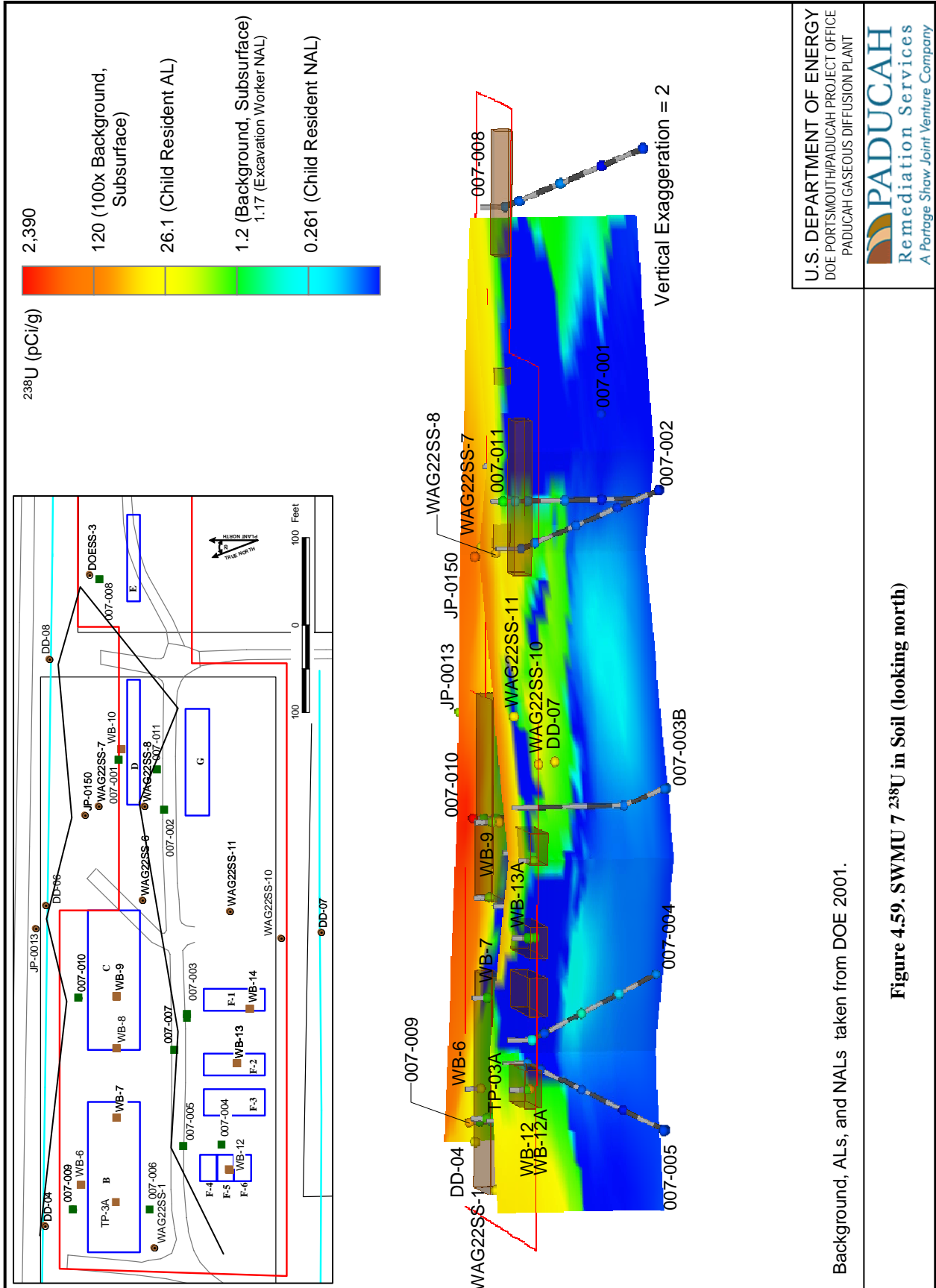


Figure 4.59. SWMU 7 ²³⁸U in Soil (looking north)

Table 4.36. SWMU 7 Locations of Subsurface Soil Contaminants

Analysis	Depth (ft)	Excavation Worker NAL	Background	RI Data												Historical Data													
				007-001	007-002	007-003A	007-003B	007-004	007-005	007-006	007-007	007-008	007-009	007-010	007-011	TP-3A	WB-10	WB-12	WB-13	WB-14	WB-6	WB-7	WB-8	WB-9	WB-12A	WB-13A	WBP-9A		
Aluminum	05-09	5,250	12,000	8,960	6,790	8,960		8,110	8,680	9,010	8,030	7,460	8,970	4,790	6,600	10,000	11,000	9,900	8,600	6,200	9,400	9,900	16,000						
	10-12	5,250	12,000	8,960	7,700	8,030		5,880	9,960	6,340	5,970	7,320	7,730	11,000	7,970														
	15	5,250	12,000	6,490	7,700	8,030		3,460	5,550	4,530	4,140	4,040	3,890	4,250	5,270														
	30	5,250	12,000	4,020	8,320	4,960		3,190	2,380	4,180	1,790	939	4,660	3,860	2,850														
	45	5,250	12,000	2,490	3,350		7,860	8,480	5,670	5,000	2,830	4,890	3,840	7,770															
	60	5,250	12,000	2,370	4,880																								
Arsenic	05-09	0.324	7.9	3.3	2.74	2.28		2.18	3.63	4.48	3.36	3.8	7.88	3.47	1.66	4.9U	3.7U	5.1U	3.9U	5.7U	2.8U	5.2U	3.7U						
	10-12	0.324	7.9	1.45	6.22	1.85		1.75	1.1	2.44	1.4	0.935U	1.22	1.35	2.68														
	15	0.324	7.9	0.917	2.5	1.18		4.57	0.895U	1.62	1.1	2.76	1.41	1.66	1.37														
	30	0.324	7.9	0.973	0.853U			1.24	1.1	0.97U	0.851U	1.02	0.863U	1.06	0.94U														
	45	0.324	7.9	0.968U	13.9		5.19	2.88	1.59	1.9	1.25	0.959U	1.99	1.49	2.44														
	60	0.324	7.9																										
Barium	05-09	272	170	98.2	84.5	102		65.6	86.6	86.4	92.1	150	129	111	50.1	79	180	93	130	50	120	170	97						
	10-12	272	170	72.1	95.5	67.1		82.4	56.5	77.2	64.6	77.7	73.7	87.1	71.1														
	15	272	170	13.9	51.7	13.8		22.2	15.9	19.2	13.3	12.7	21.8	18.2	23.1														
	30	272	170	8.86	11.4			11.2	15.3	9.31	23.1	6.14	31.8	14.5	9.24														
	45	272	170	18.1	38.3		53.8	51.3	40.1	146	52.1	17	41.8	19.2	71.2														
	60	272	170																										
Beryllium	05-09	1.26	0.69	0.468U	0.477U	0.495U		0.487U	0.485U	0.497U	0.469U	0.455U	0.464U	0.47U	0.48U	1.2U	1.3U	1.2U	1.1U	1.1U	1.2U	1.2U	1.3U						
	10-12	1.26	0.69	0.427U	0.5U	0.467U		0.46U	0.499U	0.435U	0.49U	0.467U	0.466U	0.469U	0.446U														
	15	1.26	0.69	0.44U	0.469U	0.483U		0.499U	0.448U	0.484U	0.439U	0.458U	0.473U	0.469U	0.471U														
	30	1.26	0.69	0.463U	0.426U		0.477U	0.461U	0.485U	0.426U	0.49U	0.432U	0.496U	0.488U	0.47U														
	45	1.26	0.69	0.484U	1.55		1	0.978	0.499U	0.588	0.512	0.479U	0.629	0.477U	0.542														
	60	1.26	0.69																										
Cadmium	05-09	15.2	0.21	1.87U	1.91U	1.98U		1.95U	1.94U	1.99U	1.88U	1.82U	1.86U	1.88U	1.92U	0.59U	0.25U	0.24U	0.25U	0.23U	0.11	0.24U	0.18						
	10-12	15.2	0.21	1.71U	2U	1.87U		1.84U	2U	1.74U	1.96U	1.87U	1.86U	1.88U	1.78U														
	15	15.2	0.21	1.76U	1.88U	1.93U		2U	1.79U	1.93U	1.76U	1.83U	1.89U	1.88U	1.89U														
	30	15.2	0.21	1.85U	1.71U		1.91U	1.84U	1.94U	1.7U	1.96U	1.73U	1.98U	1.95U	1.88U														
	45	15.2	0.21	1.94U	1.75U		1.94U	1.86U	2U	1.96U	1.86U	1.92U	2U	1.91U	1.84U														
	60	15.2	0.21	6.100																									
Calcium	05-09	n/a	6,100	839	709	626		362	672	923	719	1,590	1,340	780	637	740	1,600	1,600	1,200	750	1,900	2,000	3,300						
	10-12	n/a	6,100	876	813	992		923	1,160	1,030	936	1,080	1,000	1,260	1,280														
	15	n/a	6,100	704	810	610		376	740	777	752	474	694	850	826														
	30	n/a	6,100	393	555		372	557	195	629	233	113	1,220	568	298														
	45	n/a	6,100	196	573		1,120	1,220	544	1,210	545	313	699	259	778														
	60	n/a	6,100	43																									
Chromium	05-09	476	43	13	9.62	10.8		9.21	13	10.4	11.5	13.1	11.7	10.2	10.4	15	13	14	12	10	13	18	20						
	10-12	476	43	10.6	11.9	13.1		9.88	13	10.8	11.2	10.5	8.75	10.3	12.1														
	15	476	43	8.33	12.2	5.63		32.8	4.54	7.05	3.94	19.1	7.24	4.65	5.02														
	30	476	43	4.21	2.64			8.31	6.29	2.79	5.05	2.45U	3.12	14.1	3.51														
	45	476	43	8.07	10.5			11.3	13.9	6.68	5.9	6.67	3.88	5.84	8.24														
	60	476	43	3.46	3.75	5.3		2.97	4.7	3.6	7.7	7.85	2.32U	4.88	2.41														
Cobalt	05-09	1,110	13	2.47	6.41	4.56		2.3U	4	2.53	2.73	2.34U	2.33U	2.34U	3.33														
	10-12	1,110	13	2.2U	4.05	2.42U		5.26	2.24U	2.42U	2.2U	2.29U	2.36U	3.47	3.43														
	15	1,110	13	2.31U	2.13U		2.38U	2.31U	2.43U	3.99	2.45U	2.16U	2.48U	3.23	2.35U														
	30	1,110	13	3.77	15.5		2.43U	3.69	2.5U	17.7	3.52	2.4U	3.47	6.31															
	45	1,110	13	9.45	9.52	8.61		7.27	9.66	13	10.9	11.4	8.01	9.85	11.4														
	60	1,110	25	5.78	10.7	6		7.1	4.58	7.12	7.18	5.71	4.82	6.11	6.5														
Copper	05-09	427	25	2.2U	6.89	2.45		3.37	2.24U	2.45	2.79	2.97	2.36U	3.37	3.53														
	10-12	427	25	2.31U	2.25		2.38U	3.95	2.43U	2.83	2.45U	2.16U	3.84	4.03	2.35U														
	15	427	25	2.42U	9.22		11.4	11.3	4.95	9.46	5.13	2.4U	6.85	2.39U	8.09														
	30	427	25	10.9	9.52	8.61		7.27	9.66	13	10.9	11.4	8.01	9.85	11.4														
	45	427	25	2.42U	9.22		11.4	11.3	4.95	9.46	5.13	2.4U	6.85	2.39U	8.09														
	60	427	25	10.9	9.52	8.61		7.27	9.66	13	10.9	11																	

Table 4.36. SWMU 7 Locations of Subsurface Soil Contaminants (Continued)

Analysis	Depth (ft)	Excavation Worker NAL	Background	RI Data												Historical Data											
				007-001	007-002	007-003A	007-003B	007-004	007-005	007-006	007-007	007-008	007-009	007-010	007-011	TP-3A	WB-10	WB-12	WB-13	WB-14	WB-6	WB-7	WB-8	WB-9	WBP-12A	WBP-13A	WBP-9A
Technetium-99	15	57.9	2.8	1.74U	1.74U	1.81U	1.88U	1.88U	1.81U	1.68	2.69	1.77U	2.6														
	30	57.9	2.8	1.74U	1.74U	1.81U	1.88U	1.88U	1.81U	1.38U	2.35	1.77U	2.57														
	45	57.9	2.8	1.74U	1.74U	1.81U	1.88U	1.88U	1.81U	1.38U	2.97	1.77U	1.74														
	60	57.9	2.8	1.74U	1.74U	1.81U	1.88U	1.88U	1.81U	1.38U	2.21	1.77U	1.98														
	05-09	0.0357	1.6								0.397	1.38	0.337														
	10-12	0.0357	1.6	0.503	0.558	0.358			0.337	0.467	0.449	0.506	1.33	0.326													
Thorium-228	15	0.0357	1.6	0.397	0.371	0.354		0.32	0.46	0.366	0.357	1.33	0.327														
	30	0.0357	1.6	0.292	0.376	0.247		0.163	0.391	0.336	0.261	1.22	0.284														
	45	0.0357	1.6	0.207	0.176		0.112	0.506	0.118	0.409	0.077U	0.0845U	0.235	0.0923													
	60	0.0357	1.6	0.288			0.579	0.292	0.407	0.195	0.161	0.392	0.844	0.253													
	05-09	2.22	1.4								1.34	0.211	0.376							1.26	1.11	1.19	1.46	1.22	1.01	3.7	3.03
	10-12	2.22	1.4	0.442	0.521	0.237U		0.238U	0.294	0.471	0.351	0.446	0.462	0.359													
Thorium-230	15	2.22	1.4	0.284	0.375	0.24U		0.237U	0.239U	0.236U	0.369	0.232	0.299														
	30	2.22	1.4	0.24U	0.278	0.236U		0.239U	0.236U	0.24	0.252	0.11U	0.182														
	45	2.22	1.4	0.24U	0.24U		0.238U	0.299	0.237U	0.27	0.236U	0.24U	0.109U	0.109U													
	60	2.22	1.4	0.24U	0.24U		0.236U	0.238U	0.237U	0.555	0.237U	0.241U	0.142														
	05-09	1.95	1.5									0.423	0.205	0.42													
	10-12	1.95	1.5	0.42	0.454	0.28		0.32	0.365	0.535	0.417	0.535	0.479	0.54	0.342												
Thorium-232	15	1.95	1.5	0.306	0.396	0.217		0.271	0.36	0.285	0.473	0.301	0.409	0.37													
	30	1.95	1.5	0.278	0.387	0.213		0.174U	0.427	0.223	0.359	0.376	0.464	0.452													
	45	1.95	1.5	0.22	0.267		0.172U	0.507	0.171U	0.436	0.166U	0.264	0.205	0.163													
	60	1.95	1.5	0.205			0.522	0.317	0.35	0.746	0.191	0.166U	0.382	0.158	0.252												
	05-09	n/a	n/a									6.45	7.52	2.91													
	10-12	n/a	n/a	1.33U	1.12U	1.03U		1.1U	1.05U	0.986U	1.05U	1.01U	5.8	3.25													
Uranium	15	n/a	n/a	0.9U	0.992U	0.883U		0.952U	0.893U	0.967U	0.841U	1.57	1.23U														
	30	n/a	n/a	0.828U	0.857U	0.896U		0.797U	0.867U	0.89U	1.33	0.784U	1.15U														
	45	n/a	n/a	0.578	0.727U		0.687U	0.961U	0.632U	1.05U	0.704U	0.64U	0.9	0.952U													
	60	n/a	n/a	0.915U			0.957U	1.04U	0.933U	2.06U	0.876U	0.765U	1.2U	1.65	1.18U												
	05-09	n/a	n/a									3.76	6.87	3.98													
	10-12	n/a	n/a	0.301U	0.322U	0.301U		0.3U	0.299U	0.296U	0.297U	0.303U	0.301U	2.21	0.301U	4.9											
Uranium-234	15	n/a	n/a	0.301U	0.3U	0.301U		0.301U	0.299U	0.297U	0.295U	0.302U	0.3U	0.281U	0.3U												
	30	n/a	n/a	0.301U	0.304U	0.302U		0.301U	0.297U	0.301U	0.296U	0.301U	0.304U	0.281U	0.301U												
	45	n/a	n/a	0.302U	0.303U		0.304U	0.306U	0.299U	0.497	0.296U	0.302U	0.303U	0.306U	0.302U												
	60	n/a	n/a	0.303U			0.301U	0.302U	0.298U	0.296U	0.297U	0.302U	0.303U	0.281U	0.3U												
	05-09	2.84	2.4									1.34	0.903	1.2													
	10-12	2.84	2.4	0.148	0.141U	0.193		0.219	0.132U	0.212	0.182	0.142	0.304	0.319	0.139U	2.3											
Uranium-235	15	2.84	2.4	0.134U	0.134U	0.134U		0.134U	0.132U	0.131U	0.129U	0.134U	0.138U														
	30	2.84	2.4	0.134U	0.14	0.18		0.258	0.131U	0.132U	0.245	0.206	0.129U	0.138U													
	45	2.84	2.4	0.135U	0.135U		0.201	0.395	0.132U	0.239	0.13U	0.135U	0.14U	0.327	0.139U												
	60	2.84	2.4	0.135U	0.135U		0.326	0.219	0.131U	0.26	0.13U	0.173	0.129U	0.138U													
	05-09	0.455	0.14									0.0621	0.1	0.119													
	10-12	0.455	0.14	0.0372U	0.0452U	0.0375U		0.0371U	0.0381U	0.0372U	0.0381U	0.0392U	0.0376U	0.0509	0.0376U												
Uranium-235/236	15	0.455	0.14	0.0375U	0.0366U	0.0376U		0.0374U	0.0382U	0.0379U	0.037U	0.0382U	0.0379U	0.0382U	0.0379U												
	30	0.455	0.14	0.0372U	0.038U	0.0377U		0.0375U	0.0376U	0.0397U	0.0373U	0.0385U	0.0392U	0.0397U	0.0384U												
	45	0.455	0.14	0.0375U	0.038U		0.0386U	0.0396U	0.0383U	0.039U	0.0374U	0.0389U	0.0382U	0.0485U	0.0384U												
	60	0.455	0.14	0.0379U			0.0373U	0.0375U	0.0383U	0.0375U	0.0375U	0.0388U	0.0396U	0.0388U	0.0375U												
	05-09	0.455	0.14									0.07	0.12	0.34	0.35	0.86	0.17	0.88	1.03								
	10-12	1.17	1.2									2.36	5.87	2.66	8.4	2.02	4.07	12.3	10.8	23.2	4.01	15.2	15.0	1.8			
Uranium-238	15	1.17	1.2	0.147	0.136U	0.13U		0.335	0.129U	0.28	0.341	0.129U	0.262														
	30	1.17	1.2	0.13U	0.129U	0.13U		0.129U	0.129U	0.128U	0.128U	0.128U	0.181														
	45	1.17	1.2	0.13U	0.13U	0.13U		0.499	0.128U	0.129U	0.217	0.128U	0.125U	0.251	0.125U												
	60	1.17	1.2	0.13U	0.13U	0.13U		0.131U	0.225	0.129U	0.233	0.129U	0.125U	1.34	0.125U												
	05-09	1.17	1.2									0.13U	0.128U	0.197	0.129U	0.129U	0.129U	0.129U	0.129U	0.129U	0.129U	0.129U	0.129U	0.129U	0.129U	0.129U	
	10-12	1.17	1.2	0.147	0.136U	0.13U		0.335	0.129U	0.28	0.341	0.129U	0.262														

Blank cells indicate interval was not sampled for the specified analysis. Maximum value shown for each depth interval.

Bold indicates result is greater than NAL value.

Italics indicates result is greater than background value.

Bold + Italics indicate result is greater than both NAL and background values.

4.8.2 SWMU 7 Groundwater

The SWMU 7 waste pits containing various uranium-contaminated wastes are 8 to 15 ft deep. Seven temporary soil borings sampled groundwater from within and immediately below the waste pits: WB-7, WB-8, WB-9, WBP-9A, WB-12, WBP-12A, and WB-13 (Figure 4.14 shows groundwater sampling locations). Several metals, the uranium isotopes, and vinyl chloride were the primary contaminants that exceeded screening levels in these samples. These same contaminants were common throughout the thickness of the UCRS.

UCRS groundwater samples were collected from six of eight angled borings and three deep vertical borings installed at SWMU 7 as part of this RI. Several sources of historical UCRS groundwater data were available for SWMU 7, as follows:

- Wells MW186 and MW187 (with a period of record for 1995 through 2007, located immediately north of the burial pits);
- Temporary borings GW-01, GW-02, GW-03 (located on the perimeter of SWMU 7);, and WB-7, WB-8, WB-9, WB-12, and WB-13 (located within and immediately below the burial pits) of the SWMUs 7 and 30 RI (DOE 1998a);
- Temporary borings WBP-9A and WBP-12A (located within the burial pits) from a 1998 follow-up investigation of some SWMU 7 waste pits; and
- Temporary boring DG-005 (located to the north of SWMU 7) of the sitewide remedial evaluation for source areas (DOE 2000b).

RI data were reviewed with historical data to determine the UCRS contaminants listed in Table 4.37. Screening identified ten metals in UCRS groundwater samples from SWMU 7 at levels that exceed MCLs. Arsenic, iron, uranium, and manganese were the most frequently detected metals.

Organic contaminants in UCRS groundwater exceeding screening criteria at SWMU 7 consisted of seven VOCs. TCE and its reductive dechlorination products, *cis*-1,2-DCE and vinyl chloride, were the most frequently detected organic contaminants. The radionuclide contaminants present in the SWMU 7 UCRS groundwater samples were radon-222 and the uranium isotopes uranium-234 and uranium-238.

The HU2 interval is relatively thin beneath SWMU 7, at approximate depths of 20 to 25-to-30 ft bgs. Seven temporary soil borings, mostly located on the perimeter of the burial cells: 007-007, 007-010, 007-011, DG-005, GW-01, GW-02, and GW-03, and wells MW186 and MW187 provided groundwater samples from these depths. In addition to the radionuclides in the vicinity of the waste pits, significant levels of radon-222 were present. Organic contaminants from the HU2 interval included TCE, *cis*-1,2-DCE, and vinyl chloride. This corresponds to the soil contamination found in boring 007-007 near burial pit C. The relatively high level of the TCE degradation products, compared to TCE levels in this interval, is unusual at PGDP. It has been assumed, based on dissolved oxygen levels in a nearby shallow MW, that anaerobic degradation of TCE has occurred and still may occur within the UCRS. MW186, a nearby UCRS MW, has shown similar levels of TCE and *cis*-1,2-DCE. Analytical data from this well (dissolved oxygen, sulfides, total organic carbon, and TCE degradation products) indicate anaerobic degradation is likely. Because of the low connectivity of the thin HU2 sands and the high vertical hydraulic gradient, groundwater flow from this unit is primarily downward to the RGA.

Table 4.37. SWMU 7 UCRS Groundwater Contaminants

Analysis	Historical Data	RI	Frequency of Detection ^a	Frequency of Detection		
				Above Background	Above NAL	Above MCL
Metals (mg/L)						
Aluminum	1,300	1,980	16/17	N/A	14/17	N/A
Aluminum, Dissolved	N/A	0.517	1/9	N/A	0/9	N/A
Antimony	0.0027	ND	2/14	N/A	2/14	0/14
Arsenic	0.31	0.276	20/24	N/A	20/24	16/24
Arsenic, Dissolved	0.173	0.316	14/15	N/A	14/15	7/15
Barium	9.2	2.68	24/24	N/A	24/24	5/24
Barium, Dissolved	0.52	1.01	15/15	N/A	12/15	0/15
Beryllium	0.039	0.0379	11/24	N/A	8/24	8/24
Cadmium	0.03	0.00695	9/24	N/A	9/24	4/24
Calcium	4,600	143	21/24	N/A	N/A	N/A
Calcium, Dissolved	85.7	142	15/15	N/A	N/A	N/A
Chromium	1.5	2.43	14/22	N/A	1/22	8/22
Cobalt	0.52	0.303	11/24	N/A	5/24	N/A
Cobalt, Dissolved	ND	0.0571	9/15	N/A	0/15	N/A
Copper	1.8	ND	4/24	N/A	4/24	1/24
Iron	1,200	1,010	24/24	N/A	22/24	N/A
Iron, Dissolved	0.41	53.4	11/15	N/A	9/15	N/A
Lead	1.1	0.694	14/17	N/A	10/17	10/17
Magnesium	240	64.1	24/24	N/A	0/24	N/A
Magnesium, Dissolved	47.9	56.2	15/15	N/A	0/15	N/A
Manganese	28	8.73	24/24	N/A	24/24	N/A
Manganese, Dissolved	0.55	2.9	15/15	N/A	15/15	N/A
Mercury	0.0028	0.00117	9/21	N/A	5/21	1/21
Molybdenum	1.4	0.429	11/17	N/A	10/17	N/A
Molybdenum, Dissolved	N/A	0.0471	8/9	N/A	8/9	N/A
Nickel	7.6	0.703	16/22	N/A	15/22	N/A
Nickel, Dissolved	1.2	0.0753	11/14	N/A	9/14	N/A
Potassium	140	N/A	6/14	N/A	N/A	N/A
Selenium	ND	0.032	7/24	N/A	6/24	0/24
Selenium, Dissolved	N/A	0.0246	9/9	N/A	6/9	0/9
Silver	0.0058	ND	2/24	N/A	0/24	N/A
Sodium	300	234	23/24	N/A	0/24	N/A
Sodium, Dissolved	260	232	15/15	N/A	0/15	N/A
Thallium	0.051	0.0117	4/17	N/A	0/17	4/17
Tin	0.48	N/A	3/8	N/A	0/8	N/A
Uranium	83	0.239	28/117	N/A	28/117	21/117
Uranium, Dissolved	N/A	0.0366	2/9	N/A	2/9	1/9
Vanadium	1.8	1.72	8/17	N/A	7/17	N/A
Zinc	4	4.23	17/24	N/A	9/24	N/A
Zinc, Dissolved	0.04	1.12	10/15	N/A	3/15	N/A
Radionuclides (pCi/L)						
Neptunium-237	1.46	ND	5/24	N/A	1/24	N/A
Plutonium-239	1.72	N/A	5/9	N/A	3/9	N/A
Radon	N/A	262	1/1	N/A	N/A	N/A
Radon-222	801	N/A	6/7	N/A	6/7	N/A
Technetium-99	415	175	50/98	N/A	42/98	0/98
Thorium-228	ND	0.328	5/21	N/A	5/21	N/A

Table 4.37. SWMU 7 UCRS Groundwater Contaminants (Continued)

Analysis	Historical Data	RI	Frequency of Detection ^a	Frequency of Detection		
				Above Background	Above NAL	Above MCL
Thorium-230	6.61	0.151	11/30	N/A	9/30	N/A
Thorium-232	ND	0.137	2/21	N/A	0/21	N/A
Thorium-234	ND	149	2/15	N/A	N/A	N/A
Uranium	ND	146	5/13	N/A	N/A	N/A
Uranium-234	764	18.8	24/32	N/A	22/32	N/A
Uranium-235	0.59	2.28	10/26	N/A	2/26	N/A
Uranium-235/236	116	N/A	8/9	N/A	N/A	N/A
Uranium-238	4,910	125	27/32	N/A	22/32	N/A
Uranium-238, Dissolved	N/A	0.165	2/6	N/A	0/6	N/A
PCBs (mg/L)						
PCB, Total	N/A	0.00022	1/6	N/A	1/6	0/6
PCB-1016	0.00075	ND	1/14	N/A	1/14	1/14
PCB-1248	0.016	0.00022	4/14	N/A	4/14	3/14
PCB-1254	0.00089	ND	1/14	N/A	1/14	1/14
PCB-1260	0.00024	ND	1/14	N/A	1/14	0/14
Semivolatiles (mg/L)						
1,2,4-Trichlorobenzene	0.0017	ND	1/16	N/A	0/16	0/16
1,3-Dichlorobenzene	0.00055	ND	1/26	N/A	1/26	N/A
1,4-Dichlorobenzene	0.00074	ND	2/26	N/A	1/26	0/26
2,4-Dichlorophenol	0.0004	ND	1/16	N/A	0/16	N/A
2,4-Dimethylphenol	0.24	0.0066	6/16	N/A	3/16	N/A
2-Methylnaphthalene	0.0075	ND	3/16	N/A	0/16	N/A
2-Methylphenol	0.01	N/A	2/7	N/A	0/7	N/A
4-Chloro-3-methylphenol	0.0015	ND	1/16	N/A	N/A	N/A
4-Methylphenol	0.0017	N/A	2/7	N/A	0/7	N/A
Benzoic acid	0.0098	0.0062	3/16	N/A	0/16	N/A
Bis(2-ethylhexyl)phthalate	0.0015	0.0072	2/16	N/A	1/16	1/16
Dibenzofuran	0.00073	ND	1/16	N/A	0/16	N/A
Diethyl phthalate	0.0019	ND	5/16	N/A	0/16	N/A
Di-n-butyl phthalate	0.0011	N/A	2/7	N/A	0/7	N/A
Fluorene	0.00071	ND	2/16	N/A	0/16	N/A
Isophorone	0.0066	ND	1/16	N/A	0/16	N/A
Naphthalene	0.0042	ND	3/16	N/A	3/16	N/A
Phenanthrene	0.0022	ND	2/16	N/A	0/16	N/A
Pyrene	0.0012	ND	3/16	N/A	0/16	N/A
Volatiles (mg/L)						
1,1,1-Trichloroethane	0.0088	ND	1/99	N/A	0/99	0/99
1,1,2-Trichloro-1,2,2-	0.00095	N/A	1/10	N/A	0/10	N/A
1,1,2-Trichloroethane	ND	0.014	2/33	N/A	2/33	2/33
1,1-Dichloroethane	0.099	0.021	10/99	N/A	1/99	N/A
1,1-Dichloroethene	0.0029	0.0094	3/31	N/A	3/31	1/31
1,2-Dichloroethane	ND	0.04	2/33	N/A	2/33	2/33
4-Methyl-2-pentanone	0.0019	ND	2/20	N/A	0/20	N/A
Acetone	1.7	ND	7/20	N/A	4/20	N/A
Benzene	0.0078	0.012	10/43	N/A	10/43	6/43
Chloroethane	0.019	0.0098	3/21	N/A	3/21	N/A
cis-1,2-Dichloroethene	2.9	6.5	63/101	N/A	61/101	50/101
Dichlorodifluoromethane	N/A	0.015	1/10	N/A	0/10	N/A

Table 4.37. SWMU 7 UCRS Groundwater Contaminants (Continued)

Analysis	Historical Data	RI	Frequency of Detection ^a	Frequency of Detection		
				Above Background	Above NAL	Above MCL
Ethylbenzene	0.043	ND	5/93	N/A	3/93	0/93
Tetrachloroethene	0.00071	ND	1/74	N/A	1/74	0/74
Toluene	0.018	ND	8/99	N/A	0/99	0/99
Total Xylene	0.21	N/A	4/89	N/A	1/89	0/89
<i>trans</i> -1,2-Dichloroethene	0.0051	0.0054	3/100	N/A	0/100	0/100
Trichloroethene	2	12	89/100	N/A	87/100	86/100
Vinyl chloride	3.8	2.6	44/63	N/A	44/63	44/63

^a Frequency of detection is the number of detections of an analyte per number of analyses (includes regular and duplicate samples).

N/A = not applicable

ND = not detected

As discussed in Section 3.9.4, MW66 is an upper RGA well located between burial pits A and B of SWMUs 30 and 7, respectively. The analyses of groundwater samples from MW66 reveal abrupt rises or spikes of dissolved TCE (Figure 3.29) that correlate to periods of higher hydraulic head (TCE spikes often exceed 10,000 µg/L). This spiking behavior suggests a potential UCRS DNAPL source that releases contaminant mass in response to seasonal variations (more mass being released during times of higher hydraulic head). If this potential DNAPL source extended deeper into the RGA, the TCE trend should not fluctuate as much as observed.

The high-TCE spikes typically are limited to years where RGA water level exceeds 324 ft amsl. In MW66, the contact of the RGA and the overlying UCRS soils occurs at an approximate elevation of 318 ft amsl. The relationship between abrupt rises in TCE levels and high RGA water levels indicates the presence of a DNAPL source zone near the boundary of SWMUs 7 and 30 at an elevation of approximately 324 ft amsl in the silt/clay horizon that overlies the RGA. The SWMUs 7 and 30 RI report (DOE 1998a) also postulated a DNAPL source near burial pit B. The volume of soil potentially contaminated with TCE DNAPL at this SWMU is estimated to be approximately 9,375 yd³. This estimate assumes a source area that is 75 ft by 75 ft with a thickness of 45 ft (depth to top of RGA, which is 60 ft minus the estimated depth to base of waste at 15 ft). The volumetric extent of this potential DNAPL zone may be further refined for alternatives evaluation in the FS.

Five temporary soil borings of the RI: 007-001, 007-002, 007-003B, 007-008, and 007-009, sampled the HU3 interval at the perimeter of the burial pits. Metals, the uranium isotopes, and TCE and its degradation products were the primary contaminants exceeding screening levels.

RGA groundwater samples were collected from 10 ft intervals within the three deep vertical borings installed as part of this RI. Historical data for the RGA were available from MW185, MW339, MW340, and temporary soil borings DG-005, GW-01, GW-02, and GW-03. These data, together with historical data, were reviewed to identify the contaminants listed in Table 4.38.

The data review revealed the occurrence of 12 metal contaminants in the RGA groundwater samples from SWMU 7. As in the UCRS samples, arsenic, iron, and manganese were the most frequently detected groundwater contaminants. All of the SWMU 7 RGA organic groundwater contaminants were VOCs. TCE was the dominant organic contaminant. The RGA groundwater radionuclide contaminants of SWMU 7 consist of technetium-99, uranium-234, and uranium-238. Although a potential TCE DNAPL source is believed to exist near Pit B as discussed above, the primary occurrence of VOCs and technetium-99 in the RGA is largely due to the Northwest Plume, which passes beneath SWMU 7. Figure 4.60 shows the Northwest Plume TCE that passes beneath SWMUs 7 and 30.

Three locations, GWW-01, GWW-02, and GWW-03 were sampled for McNairy groundwater, close to the McNairy contact with the RGA, at SWMU 7. Table 4.39 summarizes the review of the McNairy groundwater analyses; TCE and chloroform were the only contaminants identified.

Table 4.40 provides detail (depth, sample location, and analytical results) for SWMU 7 groundwater samples, including nondetects and detections above screening levels.

Table 4.38. SWMU 7 RGA Groundwater Contaminants

Analysis	Historical Data	RI	Frequency of Detection ^a	Frequency of Detection		
				Above Background	Above NAL	Above MCL
Metals (mg/L)						
Aluminum	1,500	998	42/49	35/49	35/49	N/A
Antimony	0.0043	ND	3/22	0/22	3/22	0/22
Arsenic	0.42	0.481	48/55	44/55	48/55	32/55
Arsenic, Dissolved	0.12	0.0436	26/30	17/30	26/30	12/30
Barium	5.3	10.7	51/51	35/51	48/51	15/51
Barium, Dissolved	0.16	1.26	30/30	20/30	29/30	0/30
Beryllium	0.073	0.0732	33/49	22/49	25/49	22/49
Cadmium	0.016	0.02	24/47	10/47	24/47	11/47
Cadmium, Dissolved	ND	0.000809	4/30	0/30	4/30	0/30
Calcium	500	166	58/58	23/58	N/A	N/A
Calcium, Dissolved	35.2	160	30/30	6/30	N/A	N/A
Chromium	2	1.46	34/42	28/42	2/42	29/42
Cobalt	0.91	2.07	32/47	25/47	20/47	N/A
Cobalt, Dissolved	ND	0.108	24/30	10/30	2/30	N/A
Copper	1.3	ND	13/49	11/49	11/49	0/49
Iron	2,200	2,460	57/57	46/57	57/57	N/A
Iron (2+)	0.75	N/A	2/4	N/A	N/A	N/A
Iron, Dissolved	ND	57.6	22/30	20/30	20/30	N/A
Lead	1.6	0.489	35/39	13/39	27/39	27/39
Magnesium	120	79.1	56/56	23/56	0/56	N/A
Magnesium, Dissolved	14	72.9	30/30	4/30	0/30	N/A
Manganese	22	72.4	56/56	43/56	53/56	N/A
Manganese, Dissolved	0.1	12.8	29/30	25/30	26/30	N/A
Mercury	0.0008	0.000785	27/43	12/43	5/43	0/43
Molybdenum	0.33	0.0916	31/39	15/39	29/39	N/A
Molybdenum, Dissolved	ND	0.0269	19/26	0/26	12/26	N/A
Nickel	1.6	1.18	37/48	9/48	36/48	N/A
Nickel, Dissolved	0.25	0.0704	28/30	0/30	16/30	N/A
Potassium	89	N/A	15/28	10/28	N/A	N/A
Potassium, Dissolved	1.85	N/A	2/3	0/3	N/A	N/A
Selenium	0.00964	0.0377	15/47	15/47	9/47	0/47
Selenium, Dissolved	ND	0.0266	15/26	15/26	7/26	0/26
Silver	0.0057	0.00571	5/47	0/47	0/47	N/A
Sodium	46	320	55/61	4/61	0/61	N/A
Sodium, Dissolved	42.3	312	30/30	4/30	0/30	N/A
Strontium	0.09	N/A	12/12	N/A	0/12	N/A
Thallium	0.13	0.0106	15/41	5/41	0/41	15/41
Tin	1	N/A	10/13	N/A	3/13	N/A
Uranium	0.09	0.093	26/156	24/156	26/156	8/156
Uranium, Dissolved	ND	0.00132	2/26	0/26	2/26	0/26
Vanadium	2.7	ND	14/43	9/43	13/43	N/A
Zinc	9.8	4.28	33/47	32/47	25/47	N/A
Zinc, Dissolved	ND	0.555	23/30	17/30	2/30	N/A
Radionuclides (pCi/L)						
Americium-241	0.41	ND	1/35	N/A	1/35	N/A
Neptunium-237	1.55	ND	3/48	1/48	1/48	N/A
Plutonium-239	0.78	N/A	2/15	2/15	1/15	N/A

Table 4.38. SWMU 7 RGA Groundwater Contaminants (Continued)

Analysis	Historical Data	RI	Frequency of Detection ^a	Frequency of Detection		
				Above Background	Above NAL	Above MCL
Potassium-40	47.9	N/A	1/18	N/A	N/A	N/A
Radon-222	326	N/A	4/4	0/4	4/4	N/A
Technetium-99	5,116.9	812	137/141	130/141	133/141	43/141
Thorium-228	0.278	0.401	15/42	N/A	15/42	N/A
Thorium-230	8.42	ND	15/57	10/57	13/57	N/A
Thorium-232	ND	0.0936	4/42	N/A	0/42	N/A
Thorium-234	ND	141	2/33	N/A	N/A	N/A
Uranium	ND	17.8	24/30	N/A	N/A	N/A
Uranium-234	18.6	8.71	39/69	34/69	34/69	N/A
Uranium-235	ND	0.34	22/63	4/63	0/63	N/A
Uranium-235/236	2.07	N/A	11/15	7/15	N/A	N/A
Uranium-238	20.3	13.9	48/71	34/71	34/71	N/A
Semivolatiles (mg/L)						
2,4-Dimethylphenol	N/A	0.0085	4/24	N/A	0/24	N/A
Bis(2-ethylhexyl)phthalate	N/A	0.0052	2/24	N/A	2/24	0/24
Volatiles (mg/L)						
1,1,1-Trichloroethane	0.0013	ND	1/126	N/A	0/126	0/126
1,1,2-Trichloro-1,2,2-	0.0057	N/A	2/13	N/A	0/13	N/A
1,1,2-Trichloroethane	0.009	ND	4/56	N/A	4/56	1/56
1,1-Dichloroethane	0.0041	ND	2/126	N/A	0/126	N/A
1,1-Dichloroethene	0.0037	0.0052	12/63	N/A	12/63	0/63
1,2-Dichloroethane	0.0011	ND	2/56	N/A	2/56	0/56
2-Butanone	0.086	ND	1/40	N/A	0/40	N/A
2-Hexanone	0.0015	ND	1/40	N/A	0/40	N/A
4-Methyl-2-pentanone	0.0029	ND	1/40	N/A	0/40	N/A
Acetone	0.43	ND	7/40	N/A	4/40	N/A
Carbon tetrachloride	0.041	ND	7/56	N/A	7/56	5/56
Chloroform	0.012	ND	9/54	N/A	9/54	N/A
Chloromethane	0.014	ND	1/40	N/A	1/40	N/A
<i>cis</i> -1,2-Dichloroethene	2.1	0.58	31/107	N/A	31/107	13/107
Tetrachloroethene	0.0048	ND	6/65	N/A	6/65	0/65
Toluene	0.019	ND	3/126	N/A	0/126	0/126
<i>trans</i> -1,2-Dichloroethene	0.0038	ND	3/122	N/A	0/122	0/122
Trichloroethene	25	18	139/141	N/A	136/141	136/141
Vinyl chloride	ND	0.3	5/66	N/A	5/66	5/66

^a Frequency of detection is the number of detections of an analyte per number of analyses (includes regular and duplicate samples).

^b N/A = not applicable

ND = not detected

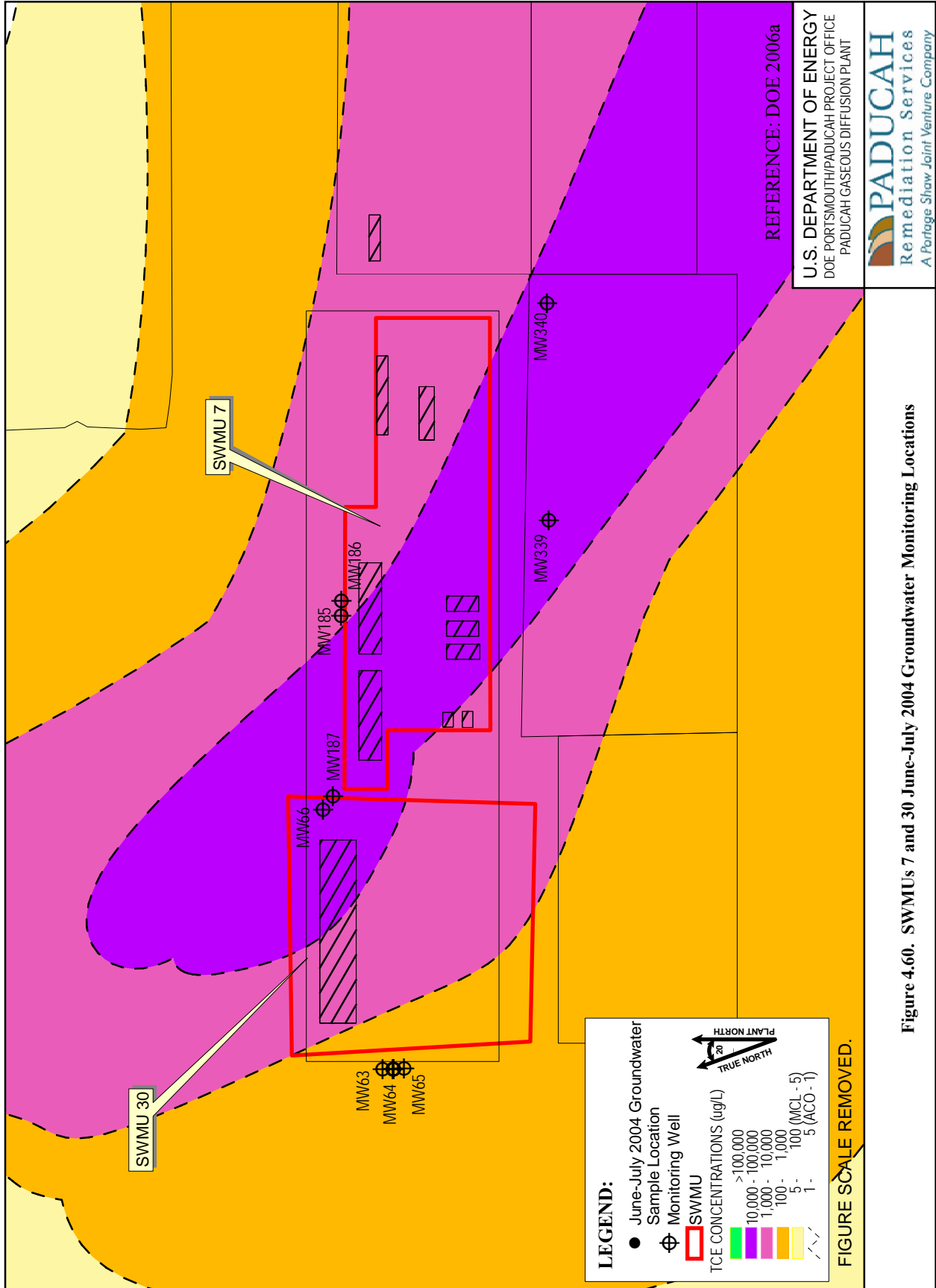


Figure 4.60. SWMUs 7 and 30 June-July 2004 Groundwater Monitoring Locations

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Table 4.39. SWMU 7 McNairy Groundwater Contaminants

Analysis	Historical Data	RI	Frequency of Detection ^a	Frequency of Detection		
				Above Background	Above NAL	Above MCL
Metals (mg/L)						
Aluminum	1,600	N/A	4/4	4/4	4/4	N/A
Arsenic	0.32	N/A	4/4	4/4	4/4	4/4
Barium	9.1	N/A	4/4	4/4	4/4	3/4
Beryllium	0.095	N/A	4/4	3/4	4/4	4/4
Cadmium	0.031	N/A	3/4	3/4	3/4	3/4
Calcium	920	N/A	4/4	4/4	N/A	N/A
Chromium	2.1	N/A	4/4	4/4	1/4	4/4
Cobalt	1.3	N/A	4/4	4/4	4/4	N/A
Copper	1.5	N/A	4/4	4/4	4/4	1/4
Iron	2,500	N/A	4/4	4/4	4/4	N/A
Lead	0.71	N/A	4/4	4/4	4/4	4/4
Magnesium	110	N/A	4/4	4/4	0/4	N/A
Manganese	54	N/A	4/4	4/4	4/4	N/A
Mercury	0.0012	N/A	4/4	4/4	1/4	0/4
Molybdenum	0.35	N/A	2/4	2/4	2/4	N/A
Nickel	1.4	N/A	4/4	4/4	4/4	N/A
Potassium	100	N/A	4/4	1/4	N/A	N/A
Selenium	0.065	N/A	2/4	2/4	2/4	1/4
Sodium	38	N/A	4/4	2/4	0/4	N/A
Thallium	0.088	N/A	3/4	0/4	0/4	3/4
Tin	1.1	N/A	4/4	N/A	2/4	N/A
Vanadium	3.8	N/A	4/4	4/4	4/4	N/A
Zinc	15	N/A	4/4	4/4	4/4	N/A
Radionuclides (pCi/L)						
Plutonium-239	0.81	N/A	1/4	1/4	1/4	N/A
Technetium-99	95.1	N/A	2/4	2/4	2/4	0/4
Thorium-230	2	N/A	4/4	1/4	4/4	N/A
Uranium-234	4.75	N/A	4/4	3/4	3/4	N/A
Uranium-235/236	0.52	N/A	3/4	2/4	N/A	N/A
Uranium-238	4.73	N/A	3/4	3/4	3/4	N/A
Volatiles (mg/L)						
Acetone	0.2	N/A	1/4	N/A	1/4	N/A
Chloroform	0.0038	N/A	3/4	N/A	3/4	N/A
cis-1,2-Dichloroethene	0.0056	N/A	3/4	N/A	1/4	0/4
Toluene	0.00078	N/A	2/4	N/A	0/4	0/4
Trichloroethene	0.32	N/A	4/4	N/A	4/4	4/4

^a Frequency of detection is the number of detections of an analyte per number of analyses (includes regular and duplicate samples).

N/A = not applicable

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Table 4.40. SWMU 7 Locations of Groundwater Contaminants

Hydro Unit	Depth Interval	Analysis	Child Resident NAL	Bkgd	MCL	007-001	007-002	007-003B	007-005	007-007	007-008	007-009	007-010	007-011	DG-005	GW-01	GW-02	GW-03	MW185	MW186	MW187	MW339	MW340	WB-12	WB-13	WB-7	WB-8	WB-9	WBP-12A	WBP-9A	
	7-9	Metals (mg/L)																													
		Aluminum	1.49	N/A	N/A																				9.6	410			1.300		
		Antimony	0.000564	N/A	0.006																				0.0026	0.003U			0.0027		
		Arsenic	0.000035	N/A	0.01																				0.0082	0.11			0.31		
		Barium	0.104	N/A	2																				0.14	2.8			9.2		
		Beryllium	0.00264	N/A	0.004																				0.0003U	0.015			0.039		
		Cadmium	0.000661	N/A	0.005																				0.004U	0.008			0.03		
		Calcium	N/A	N/A	N/A																				11U	400			4600		
		Chromium	1.76	N/A	0.1																				0.0083U	0.6			1.5		
		Cobalt	0.0906	N/A	N/A																				0.006U	0.19			0.46		
		Copper	0.0557	N/A	1.3																				0.01U	0.41			1.8		
		Iron	0.449	N/A	N/A																				10	420			1200		
		Lead	0.015	N/A	0.015																				0.0045U	0.32			0.73		
		Magnesium	N/A	N/A	N/A																				47	86			240		
		Manganese	0.035	N/A	N/A																				0.46	12			28		
		Mercury	0.000444	N/A	0.002																				0.002U	0.0006			0.0028		
		Molybdenum	0.00753	N/A	N/A																				1.4	0.063U			0.1U		
		Nickel	0.0301	N/A	N/A																				0.019	6.3			7.6		
		Potassium	N/A	N/A	N/A																				140	24			72		
		Selenium	0.00754	N/A	0.05																				0.005U	0.0093U			0.005U		
		Silver	0.0075	N/A	N/A																				0.007U	0.007U			0.007U		
		Sodium	N/A	N/A	N/A																				200	35U			240		
		Thallium	N/A	N/A	0.002																				0.01U	0.01U			0.01U		
		Tin	0.894	N/A	N/A																				0.05U	0.12			0.21		
		Uranium	0.000906	N/A	0.03																				0.88	83			14	0.15	
		Vanadium	0.00925	N/A	N/A																				0.022	0.7			1.8		
		Zinc	0.45	N/A	N/A																				0.075	1.6			4		
		PCBs (mg/L)																													
		PCB-1016	0.000468	N/A	0.0005																				0.00041U	0.00075			0.00041U		
		PCB-1248	0.000775	N/A	0.0005																				0.0083	0.00029U			0.00029U		
		PCB-1254	0.000194	N/A	0.0005																				0.000016U	0.000016U			0.00089		
		PCB-1260	0.000428	N/A	0.0005																				0.000025U	0.00024			0.00025U		
		Radionuclides (pCi/L)																													
		Neptunium-237	0.573	N/A	N/A																				0.36U	1.46			0.82U		
		Plutonium-239	0.286	N/A	N/A																				0.16U	1.72			0.44		
		Technetium-99	14	N/A	900																				187	415			92.1		
		Thorium-230	0.424	N/A	N/A																				2.83	2.65			5.98		
		Uranium-234	0.546	N/A	N/A																				38.8	764			555	4.68	
		Uranium-235	0.538	N/A	N/A																				4.93	116			56.4	0.59	
		Uranium-235/236	N/A	N/A	N/A																				192	4,910			1,760	43.6	
		Uranium-238	0.443	N/A	N/A																										
		Semivolatile (mg/L)																													
		1,2,4-Trichlorobenzene	0.00781	N/A	0.07																				0.0017				0.0011U		
		1,3-Dichlorobenzene	0.000241	N/A	N/A																				0.00055	0.00024U			0.00036U		
		1,4-Dichlorobenzene	0.000578	N/A	0.075																				0.00032U	0.00022U			0.00074		
		2,4-Dichlorophenol	0.0041	N/A	N/A																				0.00016U				0.00016U		
		2,4-Dimethylphenol	0.023	N/A	N/A																				0.026				0.0014		
		2-Methylnaphthalene	N/A	N/A	N/A																				0.0016				0.00029U		
		2-Methylphenol	0.0723	N/A	N/A																				0.0002U				0.0002U		
		4-Chloro-3-methylphenol	N/A	N/A	N/A																				0.0015				0.00028U		
		4-Methylphenol	0.00727	N/A	N/A																				0.0017				0.00018U		
		Benzoic acid	5.99	N/A	N/A																				0.0066U				0.0066U		
		Bis(2-ethylhexyl)phthalate	0.00312	N/A	0.006																				0.0015				0.0003U		
		Dibenzofuran	0.0011	N/A	N/A																				0.00011U				0.00011U		
		Diethyl phthalate	1.2	N/A	N/A																				0.00058				0.00094		
		Di-n-butyl phthalate	0.129	N/A	N/A																				0.01U				0.0011U		
		Fluorene	0.00972	N/A	N/A																				0.00022U				0.00022U		
		Isophorone	0.0547	N/A	N/A																				0.00022U				0.00022U		
		Naphthalene	0.000285	N/A	N/A																				0.0027				0.00029U		

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Table 4.40. SWMU 7 Locations of Groundwater Contaminants (Continued)

Hydro Unit	Depth Interval	Analysis	Child Resident NAL	Bkgd	MCL	007-001	007-002	007-003B	007-005	007-007	007-008	007-009	007-010	007-011	DG-005	GWW-01	GWW-02	GWW-03	MW185	MW186	MW187	MW339	MW340	WB-12	WB-13	WB-7	WB-8	WB-9	WBP-12A	WBP-9A	
	7-9	Phenanthrene	N/A	N/A	N/A																			0.00021U							
		Pyrene	0.0182	N/A	N/A																			0.00012							
		Volatiles (mg/L)																													
		1,1,1-Trichloroethane	0.0335	N/A	0.2																			0.00088	0.00036U						
		1,1,2-Trichloro-1,2,2-trifluoroethane	2.7	N/A	N/A																			0.00095	0.00036U						
		1,1,2-Trichloroethane	0.000238	N/A	0.005																			0.00053U	0.00053U						
		1,1-Dichloroethane	0.0363	N/A	N/A																			0.099	0.00078						
		1,1-Dichloroethene	0.000047	N/A	0.007																			0.00044U	0.00044U						
		1,2-Dichloroethane	0.000147	N/A	0.005																			0.00027U	0.00027U						
		4-Methyl-2-pentanone	0.00722	N/A	N/A																			0.0011U	0.0011U						
		Acetone	0.0275	N/A	N/A																			0.052	0.014						
		Benzene	0.000385	N/A	0.005																			0.0039	0.00038U						
		Chloroethane	0.00461	N/A	N/A																			0.016	0.0014U						
		cis-1,2-Dichloroethene	0.00273	N/A	0.07																			0.0022	0.00047U						
		Ethylbenzene	0.00468	N/A	0.7																			0.043	0.0016						
		Tetrachloroethene	0.000582	N/A	0.005																			0.00071	0.00035U						
		Toluene	0.0338	N/A	1																			0.0073	0.00047						
		Total Xylene	0.0653	N/A	10																			0.21	0.0073						
		trans-1,2-Dichloroethene	0.00548	N/A	0.1																			0.00048U	0.00048U						
		Trichloroethene	0.0016	N/A	0.005																			0.0013	0.00039U						
		Vinyl chloride	0.000035	N/A	0.002																			0.011	0.0013U						
	11-12	Metals (mg/L)																													
		Aluminum	1.49	N/A	N/A																					37	4.4U				
		Antimony	0.000564	N/A	0.006																				0.003U	0.003U					
		Arsenic	0.000035	N/A	0.01																				0.011	0.033					
		Barium	0.104	N/A	2																				0.4	0.19					
		Beryllium	0.00264	N/A	0.004																				0.0008U	0.0003U					
		Cadmium	0.000661	N/A	0.005																				0.004U	0.004U					
		Calcium	N/A	N/A	N/A																				28U	17U					
		Chromium	1.76	N/A	0.1																				0.039	0.015U					
		Cobalt	0.0906	N/A	N/A																				0.011U	0.0051U					
		Copper	0.0557	N/A	1.3																				0.093	0.022U					
		Iron	0.449	N/A	N/A																				32	3.9					
		Lead	0.015	N/A	0.015																				0.027	0.012U					
		Magnesium	N/A	N/A	N/A																				0.24	32					
		Manganese	0.035	N/A	N/A																				0.68	0.41					
		Mercury	0.000444	N/A	0.002																				0.0002U	0.0002U					
		Molybdenum	0.00753	N/A	N/A																				0.029U	0.17					
		Nickel	0.0301	N/A	N/A																				0.062	0.036					
		Potassium	N/A	N/A	N/A																				47	130					
		Selenium	0.00754	N/A	0.05																				0.005U	0.005U					
		Silver	0.0075	N/A	N/A																				0.007U	0.0058					
		Sodium	N/A	N/A	N/A																				130	300					
		Thallium	N/A	N/A	0.002																				0.01U	0.01U					
		Tin	0.894	N/A	N/A																				0.05U	0.05U					
		Uranium	0.000906	N/A	0.03																				0.25U	0.19U					
		Vanadium	0.00925	N/A	N/A																				0.057	0.0083U					
		Zinc	0.45	N/A	N/A																				0.14	0.13					
		PCBs (mg/L)																													
		PCB-1016	0.000468	N/A	0.0005																				0.00041U	0.00041U					
		PCB-1248	0.0000775	N/A	0.0005																				0.016	0.00089					
		PCB-1254	0.0000194	N/A	0.0005																				0.000016U	0.000016U					
		PCB-1260	0.0000428	N/A	0.0005																				0.000025U	0.000025U					
		Radionuclides (pCi/L)																													
		Neptunium-237	0.573	N/A	N/A																				0.46	0.1					
		Plutonium-239	0.286	N/A	N/A																				0.08	0.08					
		Technetium-99	14	N/A	900																				2.57	230					
		Thorium-230	0.424	N/A	N/A																				1.36	1.55					
		Uranium-234	0.546	N/A	N/A																				21.6	44.9					

Table 4.40. SWMU 7 Locations of Groundwater Contaminants (Continued)

Hydro Unit	Depth Interval	Analysis	Child Resident NAL	Bkgd	MCL	007-001	007-002	007-003B	007-005	007-007	007-008	007-009	007-010	007-011	DG-005	GW-01	GW-02	GW-03	MW185	MW186	MW187	MW339	MW340	WB-12	WB-13	WB-7	WB-8	WB-9	WBP-12A	WBP-9A	
	11-12	Uranium-235/236	N/A	N/A	N/A																					1.57	3.66				
		Uranium-238	0.443	N/A	N/A																					30.6	54.6			30.1	
		Semivolatiles (mg/L)																													
		1,2,4-Trichlorobenzene	0.00781	N/A	0.07																					0.0011U	0.0011U				
		1,3-Dichlorobenzene	0.000241	N/A	N/A																					0.00036U	0.00036U				
		1,4-Dichlorobenzene	0.000578	N/A	0.075																					0.00032U	0.00032U				
		2,4-Dichlorophenol	0.0041	N/A	N/A																					0.00016U	0.00016U				
		2,4-Dimethylphenol	0.023	N/A	N/A																					0.24	0.15				
		2-Methylnaphthalene	N/A	N/A	N/A																					0.0075	0.00094				
		2-Methylphenol	0.0723	N/A	N/A																					0.01	0.0035				
		4-Chloro-3-methylphenol	N/A	N/A	N/A																					0.00028U	0.00028U				
		4-Methylphenol	0.00727	N/A	N/A																					0.00018U	0.00018U				
		Benzoic acid	5.99	N/A	N/A																					0.0066U	0.0066U				
		Bis(2-ethylhexyl)phthalate	0.00312	N/A	0.006																					0.0003U	0.0003U				
		Dibenzofuran	0.0011	N/A	N/A																					0.00073	0.00011U				
		Diethyl phthalate	1.2	N/A	N/A																					0.00051	0.0019				
		Di-n-butyl phthalate	0.129	N/A	N/A																					0.0011	0.0011				
		Fluorene	0.00972	N/A	N/A																					0.00071	0.00041				
		Isophorone	0.0547	N/A	N/A																					0.00022U	0.00022U				
		Naphthalene	0.000285	N/A	N/A																					0.0042	0.0037				
		Phenanthrene	N/A	N/A	N/A																					0.0022	0.00041				
		Pyrene	0.0182	N/A	N/A																					0.00081	0.00025U				
		Volatiles (mg/L)																													
		1,1,1-Trichloroethane	0.0335	N/A	0.2																					0.00036U	0.00036U				
		1,1,2-Trichloro-1,2,2-trifluoroethane	2.7	N/A	N/A																					0.00036U	0.00036U				
		1,1,2-Trichloroethane	0.000238	N/A	0.005																					0.00053U	0.00053U				
		1,1-Dichloroethane	0.0363	N/A	N/A																					0.00063	0.02				
		1,1-Dichloroethene	0.000047	N/A	0.007																					0.00044U	0.00044U				
		1,2-Dichloroethane	0.000147	N/A	0.005																					0.00027U	0.00027U				
		4-Methyl-2-pentanone	0.00722	N/A	N/A																					0.0016	0.0011U				
		Acetone	0.0275	N/A	N/A																					0.0014U	0.0014U				
		Benzene	0.000385	N/A	0.005																					0.0021	0.003				
		Chloroethane	0.00461	N/A	N/A																					0.0014U	0.019				
		cis-1,2-Dichloroethene	0.00273	N/A	0.07																					0.0036	0.0018				
		Ethylbenzene	0.00468	N/A	0.7																					0.0053	0.012				
		Tetrachloroethene	0.000582	N/A	0.005																					0.00035U	0.00035U				
		Toluene	0.0338	N/A	1																					0.013	0.0038				
		Total Xylene	0.0653	N/A	10																					0.024	0.054				
		trans-1,2-Dichloroethene	0.00548	N/A	0.1																					0.00048U	0.00048U				
		Trichloroethene	0.0016	N/A	0.005																					0.00071	0.00039U				
		Vinyl chloride	0.000035	N/A	0.002																					0.018	0.0024				
	22-23	Metals (mg/L)																													
		Aluminum	1.49	N/A	N/A										1.200																
		Antimony	0.000564	N/A	0.006																						0.003U				
		Arsenic	0.000035	N/A	0.01										0.28												0.175				
		Arsenic, Dissolved	0.000035	N/A	0.01																						0.173				
		Barium	0.104	N/A	2										4.3												0.471				
		Barium, Dissolved	0.104	N/A	2																						0.52				
		Beryllium	0.00264	N/A	0.004										0.033												0.0002				
		Cadmium	0.000661	N/A	0.005										0.0006U												0.025U				
		Calcium	N/A	N/A	N/A										160												80				
		Calcium, Dissolved	N/A	N/A	N/A																						85.7				
		Chromium	1.76	N/A	0.1										0.91												0.063				
		Cobalt	0.0906	N/A	N/A										0.52												0.0075				
		Cobalt, Dissolved	0.0906	N/A	N/A																						0.045U				
		Copper	0.0557	N/A	1.3										0.46												0.025U				
		Iron	0.449	N/A	N/A										1.000												1				
		Iron, Dissolved	0.449	N/A	N/A																						0.41				

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Table 4.40. SWMU 7 Locations of Groundwater Contaminants (Continued)

Hydro Unit	Depth Interval	Analysis	Child Resident NAL	Bkgd	MCL	007-001	007-002	007-003B	007-005	007-007	007-008	007-009	007-010	007-011	DG-005	GW-01	GW-02	GW-03	MW185	MW186	MW187	MW339	MW340	WB-12	WB-13	WB-7	WB-8	WB-9	WBP-12A	WBP-9A
	22-23	Lead	0.015	N/A	0.015											1.1				0.0017U										
		Magnesium	N/A	N/A	N/A											88				43.5										
		Magnesium, Dissolved	N/A	N/A	N/A															47.9										
		Manganese	0.035	N/A	N/A											2.5				0.526										
		Manganese, Dissolved	0.035	N/A	N/A															0.55										
		Mercury	0.000444	N/A	0.002											0.0011				0.0002U										
		Molybdenum	0.00753	N/A	N/A											0.048U				0.0044U										
		Nickel	0.0301	N/A	N/A											0.53				0.046										
		Nickel, Dissolved	0.0301	N/A	N/A															0.1U										
		Potassium	N/A	N/A	N/A											33				10.5U										
		Selenium	0.00754	N/A	0.05											0.0038U				0.005U										
		Silver	0.0075	N/A	N/A											0.0045U				0.06U										
		Sodium	N/A	N/A	N/A											78				255										
		Sodium, Dissolved	N/A	N/A	N/A															260										
		Thallium	N/A	N/A	0.002											0.051				0.0037U										
		Tin	0.894	N/A	N/A											0.48				0.017U										
		Uranium	0.000906	N/A	0.03											0.0092U				0.01										
		Vanadium	0.00925	N/A	N/A											1.6				0.0017U										
		Zinc	0.45	N/A	N/A											1.4				0.034										
		Zinc, Dissolved	0.45	N/A	N/A															0.04										
		PCBs (mg/L)																												
		PCB-1016	0.000468	N/A	0.0005											0.00041U				0.00041U										
		PCB-1248	0.000775	N/A	0.0005											0.00029U				0.00029U										
		PCB-1254	0.000194	N/A	0.0005											0.000016U				0.000016U										
		PCB-1260	0.000428	N/A	0.0005											0.000023U				0.000023U										
		Radionuclides (pCi/L)																												
		Neptunium-237	0.573	N/A	N/A											0.07			0.29U											
		Plutonium-239	0.286	N/A	N/A											0.04U			0.42											
		Radon-222	0.866	N/A	N/A											19.5			13.4											
		Technetium-99	14	N/A	900															68.4										
		Thorium-228	0.129	N/A	N/A															0.66U										
		Thorium-230	0.424	N/A	N/A											6.61			1.56											
		Thorium-232	0.382	N/A	N/A															0.357U										
		Thorium-234	N/A	N/A	N/A															271U										
		Uranium	N/A	N/A	N/A															40U										
		Uranium-234	0.546	N/A	N/A											20.2			14.6											
		Uranium-235	0.538	N/A	N/A															16.6U										
		Uranium-235/236	N/A	N/A	N/A															0.08										
		Uranium-238	0.443	N/A	N/A											26.7			22.2											
		Semivolatile (mg/L)																												
		1,2,4-Trichlorobenzene	0.00781	N/A	0.07											0.0012U			0.0012U											
		1,3-Dichlorobenzene	0.000241	N/A	N/A											0.0004U			0.00024U											
		1,4-Dichlorobenzene	0.000578	N/A	0.075											0.00036U			0.00022U											
		2,4-Dichlorophenol	0.0041	N/A	N/A											0.00018U				0.0004										
		2,4-Dimethylphenol	0.023	N/A	N/A											0.00012U				0.0044										
		2-Methylnaphthalene	N/A	N/A	N/A											0.00032U				0.00029U										
		2-Methylphenol	0.0723	N/A	N/A											0.0022U				0.0002U										
		4-Chloro-3-methylphenol	N/A	N/A	N/A											0.00031U				0.00028U										
		4-Methylphenol	0.00727	N/A	N/A											0.0002U				0.00021										
		Benzoic acid	5.99	N/A	N/A											0.0098				0.0066U										
		Bis(2-ethylhexyl)phthalate	0.00312	N/A	0.006											0.0012U				0.0003U										
		Dibenzofuran	0.0011	N/A	N/A											0.00012U				0.00011U										
		Diethyl phthalate	1.2	N/A	N/A											0.0082U				0.00085										
		Di-n-butyl phthalate	0.129	N/A	N/A											0.0012U				0.0006U										
		Fluorene	0.00972	N/A	N/A											0.00024U				0.00022U										
		Isophorone	0.0547	N/A	N/A											0.00024U				0.0066										
		Naphthalene	0.000285	N/A	N/A											0.00032U				0.00029U										
		Phenanthrene	N/A	N/A	N/A											0.00023U				0.00021U										

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Table 4.40. SWMU 7 Locations of Groundwater Contaminants (Continued)

Hydro Unit	Depth Interval	Analysis	Child Resident NAL	Bkgd	MCL	007-001	007-002	007-003B	007-005	007-007	007-008	007-009	007-010	007-011	DG-005	GWW-01	GWW-02	GWW-03	MW185	MW186	MW187	MW339	MW340	WB-12	WB-13	WB-7	WB-8	WB-9	WBP-12A	WBP-9A
	22-23	Pyrene	0.0182	N/A	N/A										0.00049					0.00025U										
		Volatiles (mg/L)																												
		1,1,1-Trichloroethane	0.0335	N/A	0.2										0.00036U	0.00036U	0.00036U	0.00036U		0.5U										
		1,1,2-Trichloro-1,2,2-trifluoroethane	2.7	N/A	N/A										0.00036U	0.00036U	0.00036U	0.00036U		0.00036U										
		1,1,2-Trichloroethane	0.000238	N/A	0.005										0.00053U	0.00053U	0.00053U	0.00053U		0.05U										
		1,1-Dichloroethane	0.0363	N/A	N/A										0.00037U	0.00037U	0.00037U	0.00037U		0.012										
		1,1-Dichloroethene	0.000047	N/A	0.007										0.00044U	0.00044U	0.00044U	0.00044U		0.0029										
		1,2-Dichloroethane	0.000147	N/A	0.005										0.00027U	0.00027U	0.00027U	0.00027U		0.05U										
		4-Methyl-2-pentanone	0.00722	N/A	N/A										0.0011U	0.0011U	0.0011U	0.0011U		0.0019										
		Acetone	0.0275	N/A	N/A										0.15	0.038	1.7			0.0052										
		Benzene	0.000385	N/A	0.005										0.00038U	0.00038U	0.00038U	0.00038U		0.0078										
		Chloroethane	0.00461	N/A	N/A										0.0014U	0.0014U	0.0014U	0.0014U		0.0014U										
		cis-1,2-Dichloroethene	0.00273	N/A	0.07										0.00047U	0.00047U	0.00047U	0.00047U		2.9										
		Ethylbenzene	0.00468	N/A	0.7										0.00044U	0.00044U	0.00044U	0.00044U		0.00087										
		Tetrachloroethene	0.000582	N/A	0.005										0.00035U	0.00035U	0.00035U	0.00035U		0.05U										
		Toluene	0.0338	N/A	1										0.0038	0.00058	0.00044U	0.00044U		0.018										
		Total Xylene	0.0653	N/A	10										0.0013U	0.0013U	0.0013U	0.0013U		1U										
		trans-1,2-Dichloroethene	0.00548	N/A	0.1										0.00048U	0.00048U	0.00048U	0.00048U		0.0051										
		Trichloroethene	0.0016	N/A	0.005										0.00039U	0.00039U	0.00039U	0.00039U		0.69										
		Vinyl chloride	0.000035	N/A	0.002										0.0013U	0.0013U	0.0013U	0.0013U		3.8										
	26-32	Metals (mg/L)																												
		Aluminum	1.49	N/A	N/A				970	19.6			147	1.980							0.5									
		Aluminum, Dissolved	1.49	N/A	N/A				0.2U	0.2U			0.2U	0.517																
		Antimony	0.000564	N/A	0.006				0.005U	0.005U			0.005U								0.003U									
		Arsenic	0.000035	N/A	0.01				0.222	0.173			0.276	1U							0.011									
		Arsenic, Dissolved	0.000035	N/A	0.01				0.0031	0.133			0.316	0.01U							0.01									
		Barium	0.104	N/A	2				2.32	0.64			0.751	2.68							0.252									
		Barium, Dissolved	0.104	N/A	2				0.258	0.599			0.588	0.104							0.25									
		Beryllium	0.00264	N/A	0.004				0.034	0.001U			0.00595	0.0379							0.0003									
		Cadmium	0.000661	N/A	0.005				0.00525	0.0006U			0.00204	0.00695							0.025U									
		Calcium	N/A	N/A	N/A				66.7	113			141	103							40									
		Calcium, Dissolved	N/A	N/A	N/A				40.3	112			142	26.4							34.7									
		Chromium	1.76	N/A	0.1				2.43	0.1U			0.274								0.12									
		Cobalt	0.0906	N/A	N/A				0.303	0.0698			0.0475	1U							0.045U									
		Cobalt, Dissolved	0.0906	N/A	N/A				0.0238	0.0571			0.0316	0.0156							0.045U									
		Copper	0.0557	N/A	1.3				1U	0.2U			0.2U	20U							0.025U									
		Iron	0.449	N/A	N/A				922	75.7			142	1.010							2.9									
		Iron, Dissolved	0.449	N/A	N/A				34	53.4			26.4	1.06							0.36U									
		Lead	0.015	N/A	0.015				0.382	0.0053			0.0597	0.694							0.0018									
		Magnesium	N/A	N/A	N/A				42.8	46			59.5	59.4							17									
		Magnesium, Dissolved	N/A	N/A	N/A				17.2	45.1			56.2	9.71							15.2									
		Manganese	0.035	N/A	N/A				8.73	2.13			2.61	6.6							0.14									
		Manganese, Dissolved	0.035	N/A	N/A				1.71	1.7			2.9	0.862							0.22									
		Mercury	0.000444	N/A	0.002				0.00117	0.000017			0.00085	0.000769							0.0002U									
		Molybdenum	0.00753	N/A	N/A				0.429	0.0382			0.0348	0.00862							0.01U									
		Molybdenum, Dissolved	0.00753	N/A	N/A				0.0363	0.0404			0.0103	0.001U							0.01U									
		Nickel	0.0301	N/A	N/A				0.703	0.0908			0.112								0.7									
		Nickel, Dissolved	0.0301	N/A	N/A				0.0527	0.0753			0.053	0.05U							1.2									
		Potassium	N/A	N/A	N/A				0.05U	0.00822			0.032	0.05U							10.5U									
		Selenium	0.00754	N/A	0.05				0.00513	0.00936			0.0246	0.0182							0.005U									
		Selenium, Dissolved	0.00754	N/A	0.05				0.001U	0.001U			0.001U	0.001U							0.005									
		Silver	0.0075	N/A	N/A				33.8	116			148	42.7							130									
		Sodium	N/A	N/A	N/A				61.8	109			152	53.9							134									
		Sodium, Dissolved	N/A	N/A	N/A				0.00654	0.002U			0.002U	0.0117							0.0037U									
		Thallium	N/A	N/A	0.002																0.017U									
		Tin	0.894	N/A	N/A																0.017U									
		Uranium	0.000906	N/A	0.03				0.0475	0.00109			0.239	0.0726							0.00153									
		Uranium, Dissolved	0.000906	N/A	0.03				0.001U	0.001U			0.0366	0.001U							0.0018									
		Vanadium	0.00925	N/A	N/A																									

Table 4.40. SWMU 7 Locations of Groundwater Contaminants (Continued)

Hydro Unit	Depth Interval	Analysis	Child Resident NAL	Bkgd	MCL	007-001	007-002	007-003B	007-005	007-007	007-008	007-009	007-010	007-011	DG-005	GWW-01	GWW-02	GWW-03	MW185	MW186	MW187	MW339	MW340	WB-7	WB-8	WB-9	WBP-12A	WBP-9A	
	26-32	Zinc	0.45	N/A	N/A				2.93	0.769			1.17	20U							0.049								
		Zinc, Dissolved	0.45	N/A	N/A				0.0295	0.729			1	0.2U							0.035								
		PCBs (mg/L)																											
		PCB, Total	0.0000793	N/A	0.0005				0.0001U	0.0001U			0.00022																
		PCB-1016	0.0000468	N/A	0.0005				0.0001U	0.0001U			0.0001U																
		PCB-1248	0.0000775	N/A	0.0005				0.0001U	0.0001U			0.00022																
		PCB-1254	0.0000194	N/A	0.0005				0.0001U	0.0001U			0.0001U																
		PCB-1260	0.0000428	N/A	0.0005				0.0001U	0.0001U			0.0001U																
		Radionuclides (pCi/L)																											
		Neptunium-237	0.573	N/A	N/A				0.0682U	0.0725U			0.0689U	0.157U															
		Plutonium-239	0.286	N/A	N/A																								
		Radon-222	0.866	N/A	N/A																								
		Technetium-99	14	N/A	900				0.662U	3.27			19.6	20.5															
		Thorium-228	0.129	N/A	N/A				0.328	0.234			0.138U	0.353U															
		Thorium-230	0.424	N/A	N/A				0.151	0.137U			0.213U	0.452U															
		Thorium-232	0.382	N/A	N/A				0.0733U	0.117			0.0698U	0.223U															
		Thorium-234	N/A	N/A	N/A				26U	22U			64.4	149															
		Uranium	N/A	N/A	N/A				4.35	0.629U			146	20.9															
		Uranium-234	0.546	N/A	N/A				2.79	0.477U			18.8	8.39															
		Uranium-235	0.538	N/A	N/A				0.066	0.0497U			2.28	0.516															
		Uranium-235/236	N/A	N/A	N/A																								
		Uranium-238	0.443	N/A	N/A				2.1	0.265			125	12															
		Uranium-238, Dissolved	0.443	N/A	N/A				0.0945U	0.165																			
		Semivolatiles (mg/L)																											
		1,2,4-Trichlorobenzene	0.00781	N/A	0.07				0.0047U	0.0047U			0.0049U	0.0048U															
		1,3-Dichlorobenzene	0.000241	N/A	N/A				0.0047U	0.0047U			0.0049U	0.0048U															
		1,4-Dichlorobenzene	0.000578	N/A	0.075				0.0047U	0.0047U			0.0049U	0.0048U															
		2,4-Dichlorophenol	0.0041	N/A	N/A				0.0047U	0.0047U			0.0049U	0.0048U															
		2,4-Dimethylphenol	0.023	N/A	N/A				0.0047U	0.0047U			0.0049U	0.0048U															
		2-Methylnaphthalene	N/A	N/A	N/A				0.0047U	0.0047U			0.0049U	0.0048U															
		2-Methylphenol	0.0723	N/A	N/A				0.0047U	0.0047U			0.0049U	0.0048U															
		4-Chloro-3-methylphenol	N/A	N/A	N/A				0.0047U	0.0047U			0.0049U	0.0048U															
		4-Methylphenol	0.00727	N/A	N/A				0.0047U	0.0047U			0.0049U	0.0048U															
		Benzoic acid	5.99	N/A	N/A				0.0047U	0.0047U			0.0049U	0.0048U															
		Bis(2-ethylhexyl)phthalate	0.00312	N/A	0.006				0.0047U	0.0047U			0.0049U	0.0048U															
		Dibenzofuran	0.0011	N/A	N/A				0.0047U	0.0047U			0.0049U	0.0048U															
		Diethyl phthalate	1.2	N/A	N/A				0.0047U	0.0047U			0.0049U	0.0048U															
		Di-n-butyl phthalate	0.129	N/A	N/A				0.0047U	0.0047U			0.0049U	0.0048U															
		Fluorene	0.00972	N/A	N/A				0.0047U	0.0047U			0.0049U	0.0048U															
		Isophorone	0.0547	N/A	N/A				0.0047U	0.0047U			0.0049U	0.0048U															
		Naphthalene	0.000285	N/A	N/A				0.0047U	0.0047U			0.0049U	0.0048U															
		Phenanthrene	N/A	N/A	N/A				0.0047U	0.0047U			0.0049U	0.0048U															
		Pyrene	0.0182	N/A	N/A				0.0047U	0.0047U			0.0049U	0.0048U															
		Volatiles (mg/L)																											
		1,1,1-Trichloroethane	0.0335	N/A	0.2				0.005U	0.005U			1U	0.005U															
		1,1,2-Trichloro-1,2,2-trifluoroethane	2.7	N/A	N/A																								
		1,1,2-Trichloroethane	0.000238	N/A	0.005				0.005U	0.014			1U	0.005U															
		1,1-Dichloroethane	0.0363	N/A	N/A				0.005U	0.2U			1U	0.005U															
		1,1-Dichloroethene	0.000047	N/A	0.007				0.005U	0.2U			1U	0.005U															
		1,2-Dichloroethane	0.000147	N/A	0.005				0.005U	0.04			1U	0.005U															
		4-Methyl-2-pentanone	0.00722	N/A	N/A				0.01U	0.01U			2U	0.01U															
		Acetone	0.0275	N/A	N/A				0.01U	0.01U			2U	0.01U															
		Benzene	0.000385	N/A	0.005				0.005U	0.011			1U	0.005U															
		Chloroethane	0.00461	N/A	N/A				0.005U	0.005U			1U	0.005U															
		cis-1,2-Dichloroethene	0.00273	N/A	0.07				0.005U	3.4			6.5	0.005U															
		Dichlorodifluoromethane	0.018	N/A	N/A				0.005U	0.005U			1U	0.005U															

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Table 4.40. SWMU 7 Locations of Groundwater Contaminants (Continued)

Hydro Unit	Depth Interval	Analysis	Child Resident NAL	Bkgd	MCL	007-001	007-002	007-003B	007-005	007-007	007-008	007-009	007-010	007-011	DG-005	GW-01	GW-02	GW-03	MW185	MW186	MW187	MW339	MW340	WB-7	WB-8	WB-9	WBP-12A	WBP-9A	
	26-32	Ethylbenzene	0.00468	N/A	0.7				0.005U	0.005U			1U	0.005U							0.12U								
		Tetrachloroethene	0.000582	N/A	0.005				0.005U	0.005U			1U	0.005U							0.12U								
		Toluene	0.0338	N/A	1				0.005U	0.005U			1U	0.005U							0.12U								
		Total Xylene	0.0653	N/A	10																0.38U								
		trans-1,2-Dichloroethene	0.00548	N/A	0.1				0.005U	0.0054			1U	0.005U							0.12U								
		Trichloroethene	0.0016	N/A	0.005				0.001U	5.6			12	0.001U							2								
		Vinyl chloride	0.000035	N/A	0.002				0.002U	0.75			2.6	0.002U							0.05U								
	35	Radionuclides (pCi/L)																											
		Technetium-99	14	N/A	900										14U														
		Volatiles (mg/L)																											
		1,1-Dichloroethene	0.000047	N/A	0.007										0.001U														
		cis-1,2-Dichloroethene	0.00273	N/A	0.07										0.001U														
		trans-1,2-Dichloroethene	0.00548	N/A	0.1										0.001U														
		Trichloroethene	0.0016	N/A	0.005										0.001U														
		Vinyl chloride	0.000035	N/A	0.002										0.001U														
	43-45	Metals (mg/L)																											
		Aluminum	1.49	N/A	N/A	52.1		27.3			42.3																		
		Aluminum, Dissolved	1.49	N/A	N/A	0.2U		0.2U			0.2U																		
		Antimony	0.000564	N/A	0.006			0.005U			0.005U																		
		Arsenic	0.000035	N/A	0.01	0.0106		0.119			0.01U																		
		Arsenic, Dissolved	0.000035	N/A	0.01	0.00345		0.0986			0.00169																		
		Barium	0.104	N/A	2	0.16		0.188			0.182																		
		Barium, Dissolved	0.104	N/A	2	0.0937		0.122			0.0985																		
		Beryllium	0.00264	N/A	0.004	0.00166		0.001U			0.001U																		
		Cadmium	0.000661	N/A	0.005	0.0006U		0.000818			0.000654																		
		Calcium	N/A	N/A	N/A	19.7		33.1			9.49																		
		Calcium, Dissolved	N/A	N/A	N/A	18		34.2			8.44																		
		Chromium	1.76	N/A	0.1	0.102		0.1U			0.1U																		
		Cobalt	0.0906	N/A	N/A	0.0143		0.0143			0.0106																		
		Cobalt, Dissolved	0.0906	N/A	N/A	0.00715		0.0124			0.0062																		
		Copper	0.0557	N/A	1.3	0.2U		0.2U			0.2U																		
		Iron	0.449	N/A	N/A	59.7		40.2			25.2																		
		Iron, Dissolved	0.449	N/A	N/A	27.7		16.8			9.39																		
		Lead	0.015	N/A	0.015	0.0164		0.00638			0.0114																		
		Magnesium	N/A	N/A	N/A	8.52		12.3			4.44																		
		Magnesium, Dissolved	N/A	N/A	N/A	6.53		12.3			3.21																		
		Manganese	0.035	N/A	N/A	0.6		0.804			0.185																		
		Manganese, Dissolved	0.035	N/A	N/A	0.429		0.686			0.126																		
		Mercury	0.000444	N/A	0.002			0.000013			0.00001U																		
		Molybdenum	0.00753	N/A	N/A	0.054		0.0676			0.0199																		
		Molybdenum, Dissolved	0.00753	N/A	N/A	0.0323		0.0471			0.00896																		
		Nickel	0.0301	N/A	N/A	0.0757		0.0603			0.05U																		
		Nickel, Dissolved	0.0301	N/A	N/A	0.0402		0.0498			0.0158																		
		Selenium	0.00754	N/A	0.05	0.00708		0.0104			0.022																		
		Selenium, Dissolved	0.00754	N/A	0.05	0.00675		0.00619			0.0219																		
		Silver	0.0075	N/A	N/A	0.001U		0.001U			0.001U																		
		Sodium	N/A	N/A	N/A	37.4		48.3			45.9																		
		Sodium, Dissolved	N/A	N/A	N/A	36.3		46.9			44.5																		
		Thallium	N/A	N/A	0.002	0.002U		0.002U			0.002U																		
		Uranium	0.000906	N/A	0.03	0.0267		0.00137			0.00136																		
		Uranium, Dissolved	0.000906	N/A	0.03	0.001U		0.001U			0.001U																		
		Vanadium	0.00925	N/A	N/A	0.2U		0.2U			0.2U																		
		Zinc	0.45	N/A	N/A	1.54		0.529			0.29																		
		Zinc, Dissolved	0.45	N/A	N/A	1.12		0.414			0.2																		
		PCBs (mg/L)																											
		PCB, Total	0.000793	N/A	0.0005			0.0001U			0.0001U																		
		PCB-1016	0.0000468	N/A	0.0005			0.0001U			0.0001U																		
		PCB-1248	0.0000775	N/A	0.0005			0.0001U			0.0001U																		
		PCB-1254	0.0000194	N/A	0.0005			0.0001U			0.0001U																		
		PCB-1260	0.0000428	N/A	0.0005			0.0001U			0.0001U																		

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Table 4.40. SWMU 7 Locations of Groundwater Contaminants (Continued)

Hydro Unit	Depth Interval	Analysis	Child Resident NAL	Bkgd	MCL	007-001	007-002	007-003B	007-005	007-007	007-008	007-009	007-010	007-011	DG-005	GWW-01	GWW-02	GWW-03	MW185	MW186	MW187	MW339	MW340	WB-12	WB-13	WB-7	WB-8	WB-9	WBP-12A	WBP-9A
	43-45	Radionuclides (pCi/L)																												
		Neptunium-237	0.573	N/A	N/A	0.0613U		0.0666U			0.0583U																			
		Radon	N/A	N/A	N/A	262		0.689U			4.17																			
		Technetium-99	14	N/A	900	22.7		0.145U			0.205																			
		Thorium-228	0.129	N/A	N/A	0.145U		0.131U			0.156U																			
		Thorium-230	0.424	N/A	N/A	0.135U		0.0766U			0.0731U																			
		Thorium-232	0.382	N/A	N/A	0.0765U		26.5U			21U																			
		Thorium-234	N/A	N/A	N/A	20.5U		0.62U			0.548U																			
		Uranium	N/A	N/A	N/A	0.613U		0.474U			0.422U																			
		Uranium-234	0.546	N/A	N/A	0.947		0.0516U			0.0493U																			
		Uranium-235	N/A	N/A	N/A	0.0516U		0.119			0.385																			
		Uranium-238	0.443	N/A	N/A	1.42		0.0987U			0.0903U																			
		Uranium-238, Dissolved	0.443	N/A	N/A	0.0881U																								
		Semivolatiles (mg/L)																												
		1,2,4-Trichlorobenzene	0.00781	N/A	0.07	0.0048U		0.0047U			0.0048U																			
		1,3-Dichlorobenzene	0.000241	N/A	N/A	0.0048U		0.0047U			0.0048U																			
		1,4-Dichlorobenzene	0.000578	N/A	0.075	0.0048U		0.0047U			0.0048U																			
		2,4-Dichlorophenol	0.0041	N/A	N/A	0.0048U		0.0047U			0.0048U																			
		2,4-Dimethylphenol	0.023	N/A	N/A	0.0048U		0.0047U			0.0048U																			
		2-Methylnaphthalene	N/A	N/A	N/A	0.0048U		0.0047U			0.0048U																			
		4-Chloro-3-methylphenol	N/A	N/A	N/A	0.0048U		0.0047U			0.0048U																			
		Benzoic acid	5.99	N/A	N/A	0.0056		0.0047U			0.0048U																			
		Bis(2-ethylhexyl)phthalate	0.00312	N/A	0.006	0.0048U		0.0047U			0.0048U																			
		Dibenzofuran	0.0011	N/A	N/A	0.0048U		0.0047U			0.0048U																			
		Diethyl phthalate	1.2	N/A	N/A	0.0048U		0.0047U			0.0048U																			
		Fluorene	0.00972	N/A	N/A	0.0048U		0.0047U			0.0048U																			
		Isophorone	0.0547	N/A	N/A	0.0048U		0.0047U			0.0048U																			
		Naphthalene	0.000285	N/A	N/A	0.0048U		0.0047U			0.0048U																			
		Phenanthrene	N/A	N/A	N/A	0.0048U		0.0047U			0.0048U																			
		Pyrene	0.0182	N/A	N/A	0.0048U		0.0047U			0.0048U																			
		Volatiles (mg/L)																												
		1,1,1-Trichloroethane	0.0335	N/A	0.2	0.005U		0.005U			0.005U																			
		1,1,2-Trichloroethane	0.000238	N/A	0.005	0.005U		0.005U			0.005U																			
		1,1-Dichloroethane	0.0363	N/A	N/A	0.005U		0.005U			0.005U																			
		1,1-Dichloroethene	0.000047	N/A	0.007	0.005U		0.005U			0.005U																			
		1,2-Dichloroethane	0.000147	N/A	0.005	0.005U		0.005U			0.005U																			
		4-Methyl-2-pentanone	0.00722	N/A	N/A	0.01U		0.01U			0.01U																			
		Acetone	0.0275	N/A	N/A	0.01U		0.01U			0.01U																			
		Benzene	0.000385	N/A	0.005	0.005U		0.005U			0.005U																			
		Chloroethane	0.00461	N/A	N/A	0.005U		0.005U			0.005U																			
		cis-1,2-Dichloroethene	0.00273	N/A	0.07	0.005U		0.005U			0.005U																			
		Dichlorodifluoromethane	0.018	N/A	N/A	0.005U		0.005U			0.005U																			
		Ethylbenzene	0.00468	N/A	0.7	0.005U		0.005U			0.005U																			
		Tetrachloroethene	0.000582	N/A	0.005	0.005U		0.005U			0.005U																			
		Toluene	0.0338	N/A	1	0.005U		0.005U			0.005U																			
		trans-1,2-Dichloroethene	0.00548	N/A	0.1	0.005U		0.005U			0.005U																			
		Trichloroethene	0.0016	N/A	0.005	0.01U		0.001U			0.003U																			
		Vinyl chloride	0.000035	N/A	0.002	0.002U		0.002U			0.002U																			
	50-53	Metals (mg/L)																												
		Aluminum	1.49	N/A	N/A		5.46					1.85																		
		Aluminum, Dissolved	1.49	N/A	N/A		0.2U				0.2U																			
		Antimony	0.000564	N/A	0.006	0.005U		0.005U			0.005U																			
		Arsenic	0.000035	N/A	0.01	0.01U		0.01U			0.01U																			
		Arsenic, Dissolved	0.000035	N/A	0.01	0.00727		0.00727			0.00727																			
		Barium	0.104	N/A	2	1.42		1.42			1.69																			
		Barium, Dissolved	0.104	N/A	2	0.127		0.127			1.01																			
		Beryllium	0.00264	N/A	0.004	0.0214		0.0214			0.00524																			
		Cadmium	0.000661	N/A	0.005	0.00275		0.00275			0.00136																			

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Table 4.40. SWMU 7 Locations of Groundwater Contaminants (Continued)

Hydro Unit	Depth Interval	Analysis	Child Resident NAL	Bkgd	MCL	007-001	007-002	007-003B	007-005	007-007	007-008	007-009	007-010	007-011	DG-005	GWW-01	GWW-02	GWW-03	MW185	MW186	MW187	MW339	MW340	WB-12	WB-13	WB-7	WB-8	WB-9	WBP-12A	WBP-9A
	50-53	Calcium	N/A	N/A	N/A		37.8					143																		
		Calcium, Dissolved	N/A	N/A	N/A		24.5					132																		
		Chromium	1.76	N/A	0.1		1.7																							
		Chromium	0.0906	N/A	N/A		0.153					0.1U																		
		Cobalt	0.0906	N/A	N/A		0.0238					0.0163																		
		Cobalt, Dissolved	0.0557	N/A	1.3		2U					2U																		
		Copper	0.449	N/A	N/A		490					115																		
		Iron	0.449	N/A	N/A		23.4					10.1																		
		Iron, Dissolved	0.015	N/A	0.015		0.218					0.074																		
		Lead	N/A	N/A	N/A		27.3					64.1																		
		Magnesium	N/A	N/A	N/A		8.48					53.6																		
		Magnesium, Dissolved	0.035	N/A	N/A		3.29					1.08																		
		Manganese	0.035	N/A	N/A		0.757					0.00158																		
		Manganese, Dissolved	0.000444	N/A	0.002		0.00001U					0.00542																		
		Mercury	0.00753	N/A	N/A		0.227					0.00875																		
		Molybdenum	0.00753	N/A	N/A		0.0155					0.0211																		
		Molybdenum, Dissolved	0.0301	N/A	N/A		0.5U					0.0225																		
		Nickel	0.0301	N/A	N/A		0.0329					0.0183																		
		Nickel, Dissolved	0.00754	N/A	0.05		0.0128					0.001U																		
		Selenium	0.00754	N/A	0.05		0.0116					234																		
		Selenium, Dissolved	0.0075	N/A	N/A		0.001U					232																		
		Silver	N/A	N/A	N/A		28.5					0.002U																		
		Sodium	N/A	N/A	N/A		35.1					0.0112																		
		Sodium, Dissolved	N/A	N/A	N/A		0.00569					0.00407																		
		Thallium	N/A	N/A	0.002		2U					2U																		
		Thallium	0.000906	N/A	0.03		0.039					2U																		
		Uranium	0.000906	N/A	0.03		0.001U					2U																		
		Uranium, Dissolved	0.00925	N/A	N/A		4.23					0.14																		
		Vanadium	0.45	N/A	N/A		0.386					0.0001U																		
		Zinc	0.45	N/A	N/A							0.0001U																		
		Zinc, Dissolved	0.45	N/A	N/A							0.0001U																		
		PCBs (mg/L)																												
		PCB, Total	0.000793	N/A	0.0005							0.0001U																		
		PCB-1016	0.000468	N/A	0.0005							0.0001U																		
		PCB-1248	0.000775	N/A	0.0005							0.0001U																		
		PCB-1254	0.000194	N/A	0.0005							0.0001U																		
		PCB-1260	0.0000428	N/A	0.0005							0.0001U																		
		Radionuclides (pCi/L)																												
		Neptunium-237	0.573	N/A	N/A		0.0729U					0.09U																		
		Technetium-99	14	N/A	900		175					79.1																		
		Thorium-228	0.129	N/A	N/A		0.318					0.307																		
		Thorium-230	0.424	N/A	N/A		0.141U					0.27U																		
		Thorium-232	0.382	N/A	N/A		0.137					0.13U																		
		Thorium-234	N/A	N/A	N/A		29.4U					20.8U																		
		Uranium	N/A	N/A	N/A		13.8					8.83																		
		Uranium-234	0.546	N/A	N/A		4.98					4.39																		
		Uranium-235	N/A	N/A	N/A		0.328					0.161																		
		Uranium-238	0.443	N/A	N/A		8.51					4.28																		
		Uranium-238, Dissolved	0.443	N/A	N/A		0.112																							
		Semivolatile (mg/L)																												
		1,2,4-Trichlorobenzene	0.00781	N/A	0.07		0.0048U					0.0048U																		
		1,3-Dichlorobenzene	0.000241	N/A	N/A		0.0048U					0.0048U																		
		1,4-Dichlorobenzene	0.000578	N/A	0.075		0.0048U					0.0048U																		
		2,4-Dichlorophenol	0.0041	N/A	N/A		0.0048U					0.0048U																		
		2,4-Dimethylphenol	0.023	N/A	N/A		0.0048U					0.0066																		
		2-Methylnaphthalene	N/A	N/A	N/A		0.0048U					0.0048U																		
		4-Chloro-3-methylphenol	N/A	N/A	N/A		0.0048U					0.0048U																		
		Benzoic acid	5.99	N/A	N/A		0.0062					0.0048U																		
		Bis(2-ethylhexyl)phthalate	0.00312	N/A	0.006		0.0072					0.0048U																		
		Dibenzofuran	0.0011	N/A	N/A		0.0048U					0.0048U																		
		Diethyl phthalate	1.2	N/A	N/A		0.0048U					0.0048U																		

UCRS

Table 4.40. SWMU 7 Locations of Groundwater Contaminants (Continued)

Hydro Unit	Depth Interval	Analysis	Child Resident NAL	Bkgd	MCL	007-001	007-002	007-003B	007-005	007-007	007-008	007-009	007-010	007-011	DG-005	GW-01	GW-02	GW-03	MW185	MW186	MW187	MW339	MW340	WB-12	WB-13	WB-7	WB-8	WB-9	WBP-12A	WBP-9A		
	60	Zinc	0.45	0.054	N/A								0.348	2U																		
		Zinc, Dissolved	0.45	0.049	N/A								0.0524	0.2U																		
		Radionuclides (pCi/L)																														
		Americium-241	0.371	N/A	N/A								0.155U	0.0581U																		
		Neptunium-237	0.573	0.8	N/A								0.101U	0.0668U																		
		Technetium-99	14	22.3	900								1.28	31.1	14U																	
		Thorium-228	0.129	N/A	N/A								0.192U	0.141U																		
		Thorium-230	0.424	1.1	N/A								0.319U	0.176U																		
		Thorium-232	0.382	N/A	N/A								0.109U	0.0928U																		
		Thorium-234	N/A	N/A	N/A								19.9U	21.4U																		
		Uranium	N/A	N/A	N/A								16.4	3.83																		
		Uranium-234	0.546	0.7	N/A								2.25	1.82																		
		Uranium-235	N/A	0.3	N/A								0.279	0.0925																		
		Uranium-238	0.443	0.7	N/A								13.9	1.91																		
		Semivolatiles (mg/L)																														
		2,4-Dimethylphenol	0.023	N/A	N/A								0.0048U	0.0048U																		
		Bis(2-ethylhexyl)phthalate	0.00312	N/A	0.006								0.0048U	0.0048U																		
		Volatiles (mg/L)																														
		1,1,1-Trichloroethane	0.0335	N/A	0.2								0.005U	0.005U																		
		1,1,2-Trichloroethane	0.000238	N/A	0.005								0.005U	0.005U																		
		1,1-Dichloroethane	0.0363	N/A	N/A								0.005U	0.005U																		
		1,1-Dichloroethene	0.000047	N/A	0.007								0.005U	0.005U	0.001U																	
		1,2-Dichloroethane	0.000147	N/A	0.005								0.005U	0.005U																		
		2-Butanone	0.0868	N/A	N/A								0.01U	0.01U																		
		2-Hexanone	N/A	N/A	N/A								0.01U	0.01U																		
		4-Methyl-2-pentanone	0.00722	N/A	N/A								0.01U	0.01U																		
		Acetone	0.0275	N/A	N/A								0.01U	0.01U																		
		Carbon tetrachloride	0.000181	N/A	0.005								0.005U	0.005U																		
		Chloroform	0.000287	N/A	N/A								0.005U	0.005U																		
		Chloromethane	0.00167	N/A	N/A								0.005U	0.005U																		
		cis-1,2-Dichloroethene	0.00273	N/A	0.07								0.073	0.005U	0.001U																	
		Tetrachloroethene	0.000582	N/A	0.005								0.005U	0.005U																		
		Toluene	0.0338	N/A	1								0.005U	0.005U																		
		trans-1,2-Dichloroethene	0.00548	N/A	0.1								0.005U	0.005U	0.001U																	
		Trichloroethene	0.0016	N/A	0.005								0.1	0.0068	0.0014																	
		Vinyl chloride	0.000035	N/A	0.002								0.016	0.002U	0.001U																	
		Metals (mg/L)																														
	65-66	Aluminum	1.49	2.189	N/A								100	360																		
		Antimony	0.000564	0.06	0.006								0.004																			
		Arsenic	0.000035	0.005	0.01								0.041	0.085																		
		Barium	0.104	0.235	2								0.6	3.5																		
		Beryllium	0.00264	0.004	0.004								0.0051	0.017																		
		Cadmium	0.000661	0.01	0.005								0.0003U	0.0044																		
		Calcium	N/A	41.238	N/A								190	65																		
		Chromium	1.76	0.144	0.1								0.41	0.37																		
		Cobalt	0.0906	0.045	N/A								0.078	0.24																		
		Copper	0.0557	0.036	1.3								0.68	0.23																		
		Iron	0.449	5.03	N/A								360	400																		
		Lead	0.015	0.129	0.015								0.11	0.21																		
		Magnesium	N/A	16.262	N/A								14	36																		
		Manganese	0.035	0.119	N/A								4.9	22																		
		Mercury	0.000444	0.0002	0.002								0.0001	0.0001																		
		Molybdenum	0.00753	0.05	N/A								0.33	0.029U																		
		Nickel	0.0301	0.682	N/A								0.35	0.44																		
		Potassium	N/A	5.195	N/A								15	38																		
		Selenium	0.00754	0.005	0.05								0.0038U	0.0038U																		
		Silver	0.0075	0.011	N/A								0.0045U	0.0045U																		
		Sodium	N/A	59.45	N/A								37	24																		
		Thallium	N/A	0.056	0.002								0.011U	0.0037U																		

RGA

Table 4.40. SWMU 7 Locations of Groundwater Contaminants (Continued)

Hydro Unit	Depth Interval	Analysis	Child Resident NAL	Bkgd	MCL	007-001	007-002	007-003B	007-005	007-007	007-008	007-009	007-010	007-011	DG-005	GWW-01	GWW-02	GWW-03	MW185	MW186	MW187	MW339	MW340	WB-12	WB-13	WB-7	WB-8	WB-9	WBP-12A	WBP-9A					
	65-66	Tin	0.894	N/A	N/A											0.23	0.23																		
		Uranium	0.000906	0.002	0.03											0.0092U	0.0092U																		
		Vanadium	0.00925	0.134	N/A											0.14	0.4																		
		Zinc	0.45	0.054	N/A											4.2	2.1																		
		Radionuclides (pCi/L)																																	
		Neptunium-237	0.573	0.8	N/A											0.02U	-0.026U																		
		Plutonium-239	0.286	0.1	N/A											0.06U	-0.696U																		
		Technetium-99	14	22.3	900										14U	6.38	147																		
		Thorium-230	0.424	1.1	N/A											2.16	8.26																		
		Uranium-234	0.546	0.7	N/A											1.66	2.75																		
		Uranium-235/236	N/A	0.3	N/A											0.29	0.22																		
		Uranium-238	0.443	0.7	N/A											1.51	2.9																		
		Volatiles (mg/L)																																	
		1,1,1-Trichloroethane	0.0335	N/A	0.2											0.00036U	0.00036U																		
		1,1,2-Trichloro-1,2,2-trifluoroethane	2.7	N/A	N/A											0.00036U	0.00036U																		
		1,1,2-Trichloroethane	0.000238	N/A	0.005											0.00053U	0.00053U																		
		1,1-Dichloroethane	0.0363	N/A	N/A											0.00037U	0.00037U																		
		1,1-Dichloroethene	0.000047	N/A	0.007										0.001U	0.00044U	0.00044U																		
		1,2-Dichloroethane	0.000147	N/A	0.005											0.00027U	0.00027U																		
		2-Butanone	0.0868	N/A	N/A											0.0034U	0.0034U																		
		2-Hexanone	N/A	N/A	N/A											0.0011U	0.0011U																		
		4-Methyl-2-pentanone	0.00722	N/A	N/A											0.0011U	0.0011U																		
		Acetone	0.0275	N/A	N/A											0.024	0.16																		
		Carbon tetrachloride	0.000181	N/A	0.005											0.00035U	0.00035U																		
		Chloroform	0.000287	N/A	N/A											0.00041U	0.00095																		
		Chloromethane	0.00167	N/A	N/A											0.00037U	0.00037U																		
		cis-1,2-Dichloroethene	0.00273	N/A	0.07										0.001U	0.00047U	0.0029																		
		Tetrachloroethene	0.000582	N/A	0.005											0.00035U	0.0012																		
		Toluene	0.0338	N/A	1											0.00044U	0.00044U																		
		trans-1,2-Dichloroethene	0.00548	N/A	0.1										0.001U	0.00048U	0.00048U																		
		Trichloroethene	0.00016	N/A	0.005										0.021	0.00039U	0.00048U																		
		Vinyl chloride	0.000035	N/A	0.002										0.001U	0.00013U	0.0013U																		
	68-70	Metals (mg/L)																																	
		Aluminum	1.49	2.189	N/A							173	82.7	422	0.2U																				
		Antimony	0.000564	0.06	0.006								0.005U	0.0348	0.1U	0.006																			
		Arsenic	0.000035	0.005	0.01							0.1U	0.0348	0.1U	0.006																				
		Arsenic, Dissolved	0.000035	0.005	0.01							0.00455	0.00173	0.00476																					
		Barium	0.104	0.235	2							2.24	1.25	2.8	0.1																				
		Barium, Dissolved	0.104	0.2	2							0.451	0.395	0.301																					
		Beryllium	0.00264	0.004	0.004							0.00994	0.0077	0.041	0.005U																				
		Cadmium	0.000661	0.01	0.005							0.00329	0.0018	0.0102	0.005U																				
		Cadmium, Dissolved	0.000661	0.01	0.005							0.00066U	0.0006U	0.000899																					
		Calcium	N/A	41.238	N/A							44.6	17.3	61.9	29.7																				
		Calcium, Dissolved	N/A	38.166	N/A							34.2	14.7	38.4																					
		Chromium	1.76	0.144	0.1							0.163	0.124	0.269	0.05U																				
		Cobalt	0.0906	0.045	N/A							0.0405	0.0537	0.0349	0.01U																				
		Cobalt, Dissolved	0.0906	0.045	N/A							0.0405	0.0537	0.0349																					
		Copper	0.0557	0.036	1.3							2U	0.2U	2U	0.05U																				
		Iron	0.449	5.03	N/A							216	189	938	4.15																				
		Iron, Dissolved	0.449	0.267	N/A							16	5.11	9.14																					
		Lead	0.015	0.129	0.015							0.0935	0.0355	0.309																					
		Magnesium	N/A	16.262	N/A							27.1	10.4	37.2	11																				
		Magnesium, Dissolved	N/A	16.215	N/A							14.6	5.58	12.6																					
		Manganese	0.035	0.119	N/A							9.31	6.75	20.5	0.1																				
		Manganese, Dissolved	0.035	0.068	N/A							4.42	4.01	8.32																					
		Mercury	0.000444	0.0002	0.002							0.00098	0.00041	0.000251	0.0002U																				
		Molybdenum	0.00753	0.05	N/A							0.0916	0.079	0.0459																					
		Molybdenum, Dissolved	0.																																

Table 4.40. SWMU 7 Locations of Groundwater Contaminants (Continued)

Hydro Unit	Depth Interval	Analysis	Child Resident NAL	Bkgd	MCL	007-001	007-002	007-003B	007-005	007-007	007-008	007-009	007-010	007-011	DG-005	GW-01	GW-02	GW-03	MW185	MW186	MW187	MW339	MW340	WB-12	WB-13	WB-7	WB-8	WB-9	WBP-12A	WBP-9A						
	68-70	Nickel	0.0301	0.682	N/A							0.0391	0.104	0.584	0.05U			0.12																		
		Nickel, Dissolved	0.0301	N/A	N/A							0.0391	0.104	0.584	0.05U			0.12																		
		Potassium	N/A	5.195	N/A										2U			12																		
		Selenium	0.00754	0.005	0.05							0.00768	0.005U	0.005U	0.005U			0.0038U																		
		Selenium, Dissolved	0.00754	0.005	0.05							0.005U	0.005U	0.00673	0.005U			0.0045U																		
		Silver	0.0075	0.011	N/A							0.001U	0.001U	0.001U	0.05U			0.0045U																		
		Sodium	N/A	59.45	N/A							40.6	21.3	0.5U	34.4			23																		
		Sodium, Dissolved	N/A	60.433	N/A							41.5	21.9	32.5	0.07			23																		
		Strontium	0.901	N/A	N/A										0.07																					
		Thallium	N/A	0.056	0.002							0.00224	0.002U	0.00586	0.2U			0.0074U																		
		Tin	0.894	N/A	N/A										0.2U			0.093U																		
		Uranium	0.000906	0.002	0.03							0.0118	0.03	0.031				0.0092U																		
		Uranium, Dissolved	0.000906	0.002	0.03							0.001U	0.001U	0.001U				0.0092U																		
		Vanadium	0.00925	0.134	N/A							2U	0.2U	2U	0.1U			0.15																		
		Zinc	0.45	0.054	N/A							2U	0.473	2.46	0.2U			0.99																		
		Zinc, Dissolved	0.45	0.049	N/A							0.31	0.0799	0.047				0.99																		
		Radionuclides (pCi/L)																																		
		Americium-241	0.371	N/A	N/A							0.0685U	0.09U	0.0601U																						
		Neptunium-237	0.573	0.8	N/A							0.0749U	0.0743U	0.0645U				0.03U																		
		Plutonium-239	0.286	0.1	N/A													0.01U																		
		Technetium-99	14	22.3	900							812	7.89	16.7	127			-2.7U																		
		Thorium-228	0.129	N/A	N/A							0.184U	0.116U	0.155																						
		Thorium-230	0.424	1.1	N/A							0.242U	0.194U	0.172U																						
		Thorium-232	0.382	N/A	N/A							0.146U	0.0652U	0.0836U																						
		Thorium-234	N/A	N/A	N/A							22.8U	19.5U	24.8U																						
		Uranium	N/A	N/A	N/A							11.5	9.09	9.56																						
		Uranium-234	0.546	0.7	N/A							4.51	1.26	4.39				18.6																		
		Uranium-235	N/A	0.3	N/A							0.233	0.153	0.194																						
		Uranium-235/236	N/A	0.3	N/A																															
		Uranium-238	0.443	0.7	N/A							6.71	7.68	4.97				2.07																		
		Semivolatiles (mg/L)																																		
		2,4-Dimethylphenol	0.023	N/A	N/A							0.0048U	0.0049U	0.0047U																						
		Bis(2-ethylhexyl)phthalate	0.00312	N/A	0.006							0.0048U	0.0049U	0.0052																						
		Volatiles (mg/L)																																		
		1,1,1-Trichloroethane	0.0335	N/A	0.2							0.5U	0.005U	0.005U				0.00036U																		
		1,1,2-Trichloro-1,2,2-trifluoroethane	2.7	N/A	N/A													0.00036U																		
		1,1,2-Trichloroethane	0.000238	N/A	0.005							0.5U	0.005U	0.005U				0.00053U																		
		1,1-Dichloroethane	0.0363	N/A	N/A							0.5U	0.005U	0.005U				0.00037U																		
		1,1-Dichloroethene	0.000047	N/A	0.007							0.5U	0.005U	0.005U	0.001U			0.00044U																		
		1,2-Dichloroethane	0.000147	N/A	0.005							0.5U	0.005U	0.005U				0.00027U																		
		2-Butanone	0.0868	N/A	N/A							1U	0.01U	0.01U				0.0034U																		
		2-Hexanone	N/A	N/A	N/A							1U	0.01U	0.01U				0.0011U																		
		4-Methyl-2-pentanone	0.00722	N/A	N/A							1U	0.01U	0.01U				0.0011U																		
		Acetone	0.0275	N/A	N/A							1U	0.01U	0.01U				0.0049U																		
		Carbon tetrachloride	0.000181	N/A	0.005							0.5U	0.005U	0.005U				0.00035U																		
		Chloroform	0.000287	N/A	N/A							0.5U	0.005U	0.005U				0.00041U																		
		Chloromethane	0.00167	N/A	N/A							0.5U	0.005U	0.005U				0.00037U																		
		cis-1,2-Dichloroethene	0.00273	N/A	0.07							0.5U	0.0084	0.005U	0.001U			0.00047U																		
		Tetrachloroethene	0.000582	N/A	0.005							0.5U	0.005U	0.005U				0.00035U																		
		Toluene	0.0338	N/A	1							0.5U	0.005U	0.005U				0.00044U																		
		trans-1,2-Dichloroethene	0.00548	N/A	0.1							0.5U	0.005U	0.005U	0.001U			0.00048U																		
		Trichloroethene	0.0016	N/A	0.005							14	0.18	0.091	0.12			0.00039U																		
		Vinyl chloride	0.000035																																	

Table 4.40. SWMU 7 Locations of Groundwater Contaminants (Continued)

Hydro Unit	Depth Interval	Analysis	Child Resident NAL	Bkgd	MCL	007-001	007-002	007-003B	007-005	007-007	007-008	007-009	007-010	007-011	DG-005	GW-01	GW-02	GW-03	MW185	MW186	MW187	MW339	MW340	WB-12	WB-13	WB-7	WB-8	WB-9	WBP-12A	WBP-9A
	73-78	Barium	0.104	0.235	2											5.3	0.61	4.7	0.162											
		Barium, Dissolved	0.104	0.2	2											0.073	0.003	0.07	0.015U											
		Beryllium	0.00264	0.004	0.004											0.011	0.0003U	0.016	0.025U											
		Cadmium	0.000661	0.01	0.005															0.025U										
		Cadmium, Dissolved	0.000661	0.01	0.005															0.025U										
		Calcium	N/A	41.238	N/A											500	71	400	39											
		Calcium, Dissolved	N/A	38.166	N/A														30.4											
		Chromium	1.76	0.144	0.1											1.3	0.28	2	0.51											
		Cobalt	0.0906	0.045	N/A											0.54	0.059	0.91	0.045U											
		Cobalt, Dissolved	0.0906	0.045	N/A														0.045U											
		Copper	0.0557	0.036	1.3											0.88	0.56	1.1	0.025											
		Iron	0.449	5.03	N/A											1.700	270	2.200	9.1											
		Iron (2+)	N/A	N/A	N/A														0.02U											
		Iron, Dissolved	0.449	0.267	N/A														0.36U											
		Lead	0.015	0.129	0.015											1.6	0.036	0.74	0.0017U											
		Magnesium	N/A	16.262	N/A											120	16	75	11.9											
		Magnesium, Dissolved	N/A	16.215	N/A														11.6											
		Manganese	0.035	0.119	N/A											18	5.2	16	0.11											
		Manganese, Dissolved	0.035	0.068	N/A														0.1											
		Mercury	0.000444	0.0002	0.002											0.0008		0.0008	0.0002U											
		Molybdenum	0.00753	0.05	N/A											0.085U	0.13U	0.18	0.015U											
		Nickel	0.0301	0.682	N/A											1.1	0.33	1.6	0.34											
		Nickel, Dissolved	0.0301	0.305	N/A														0.25											
		Potassium	N/A	5.195	N/A											72	8.7	89	2U											
		Potassium, Dissolved	N/A	4.096	N/A														10.5U											
		Selenium	0.00754	0.005	0.05											0.0038U	0.0038U	0.0038U	0.005U											
		Silver	0.0075	0.011	N/A											0.0045U	0.0045U	0.0045U	0.0057											
		Sodium	N/A	59.45	N/A											45	26	30	31.2											
		Sodium, Dissolved	N/A	60.433	N/A														28.7											
		Thallium	N/A	0.056	0.002											0.11	0.0037U	0.13	0.0037U											
		Tin	0.894	N/A	N/A											0.76	0.17	1	0.033U											
		Uranium	0.000906	0.002	0.03											0.0092U	0.0092U	0.0092U	0.05U											
		Vanadium	0.00925	0.134	N/A											1.6	0.097	2.7	0.0031											
		Zinc	0.45	0.054	N/A											4	4.9	9.8	0.031											
		Zinc, Dissolved	0.45	0.049	N/A														0.03U											
		Radionuclides (pCi/L)																												
		Americium-241	0.371	N/A	N/A														44.1U											
		Neptunium-237	0.573	0.8	N/A											-0.054U	-0.063U	0.03U	0.989U											
		Plutonium-239	0.286	0.1	N/A											0U	0.06U	0.08U	-0.12U											
		Potassium-40	N/A	N/A	N/A														443U											
		Radon-222	0.866	626	N/A														326											
		Technetium-99	14	22.3	900											143	1.310	106	15	1260										
		Thorium-228	0.129	N/A	N/A														0.249U											
		Thorium-230	0.424	1.1	N/A														0.66											
		Thorium-232	0.382	N/A	N/A											1.53	6.31	0.9	0.351U											
		Thorium-234	N/A	N/A	N/A														150U											
		Uranium	N/A	N/A	N/A														40U											
		Uranium-234	0.546	0.7	N/A														0.52											
		Uranium-235	0.538	0.3	N/A											3.36	0.08	7.95	2.2U											
		Uranium-235/236	N/A	0.3	N/A											0.37	0.02	1.01	0.05U											
		Uranium-238	0.443	0.7	N/A											5.3	-0.159U	5.46	10.3U											
		Volatiles (mg/L)																												
		1,1,1-Trichloroethane	0.0335	N/A	0.2											0.00036U	0.00036U	0.00036U	0.5U											
		1,1,2-Trichloro-1,2,2-trifluoroethane	2.7	N/A	N/A											0.00036U	0.00036U	0.00036U	0.0057											
		1,1,2-Trichloroethane	0.00238	N/A	0.005											0.004	0.00053U	0.00053U	0.002											
		1,1-Dichloroethane	0.0363	N/A	N/A											0.00037U	0.00037U	0.00037U	0.0041											
		1,1-Dichloroethene	0.000047	N/A	0.007											0.001U	0.00044U	0.00044U	0.0013											
		1,2-Dichloroethane	0.000147	N/A	0.005											0.00027U	0.00027U	0.00027U	0.0011											

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Table 4.40. SWMU 7 Locations of Groundwater Contaminants (Continued)

Hydro Unit	Depth Interval	Analysis	Child Resident NAL	Bkgd	MCL	007-001	007-002	007-003B	007-005	007-007	007-008	007-009	007-010	007-011	DG-005	GW-01	GW-02	GW-03	MW185	MW186	MW187	MW339	MW340	WB-12	WB-13	WB-7	WB-8	WB-9	WBP-12A	WBP-9A	
	73-78	2-Butanone	0.0868	N/A	N/A											0.0034U	0.086	0.0034U	0.25U												
		2-Hexanone	N/A	N/A	N/A											0.0011U	0.0015	0.0011U	0.25U												
		4-Methyl-2-pentanone	0.00722	N/A	N/A											0.0011U	0.0029	0.0011U	0.25U												
		Acetone	0.0275	N/A	N/A											0.013	0.26	0.028U	0.25U												
		Carbon tetrachloride	0.000181	N/A	0.0005											0.006	0.00035U	0.00035U	0.041												
		Chloroform	0.000287	N/A	N/A											0.0034	0.0015	0.00041U	0.0031												
		Chloromethane	0.00167	N/A	N/A											0.00037U	0.014	0.00037U	0.12U												
		cis-1,2-Dichloroethene	0.00273	N/A	0.07											0.001U	0.011	0.0039	0.00047U	0.49											
		Tetrachloroethene	0.000582	N/A	0.0005											0.0024	0.00035U	0.00035U	0.0019												
		Toluene	0.0338	N/A	1											0.00044U	0.019	0.00044U	0.5U												
		trans-1,2-Dichloroethene	0.00548	N/A	0.1											0.001U	0.00048U	0.00048U	0.0038												
		Trichloroethene	0.0016	N/A	0.005											0.18	4.8	0.28	0.00058	4.6											
		Vinyl chloride	0.000035	N/A	0.002											0.001U	0.0013U	0.0013U	0.1U												
		Metals (mg/L)																													
		Aluminum	1.49	2.189	N/A							35.8	11.4	488	0.53																
		Antimony	0.000564	0.06	0.006							0.005U	0.005U																		
		Arsenic	0.000035	0.005	0.01							0.0187	0.0226	0.481	0.01																
		Arsenic, Dissolved	0.000035	0.005	0.01							0.00384	0.0174	0.00561																	
		Barium	0.104	2.35	2							0.588	0.285	10.7	0.13																
		Barium, Dissolved	0.104	0.2	2							0.296	0.239	0.616																	
		Beryllium	0.00264	0.004	0.004							0.0024	0.001U	0.063	0.005U																
		Cadmium	0.000661	0.01	0.005							0.00108	0.0006U	0.02	0.005U																
		Cadmium, Dissolved	0.000661	0.01	0.005							0.0006U	0.0006U	0.000724																	
		Calcium	N/A	41.238	N/A							37.5	31.8	53.2	28.5																
		Calcium, Dissolved	N/A	38.166	N/A							35.2	31.8	36.6																	
		Chromium	1.76	0.144	0.1							0.239	0.1U	1.08	0.05U																
		Cobalt	0.0906	0.045	N/A							0.036	0.01U	2.07	0.01U																
		Cobalt, Dissolved	0.0906	0.045	N/A							0.0141	0.00543	0.108																	
		Copper	0.0557	0.036	1.3							0.2U	0.2U	2U	0.05U																
		Iron	0.449	5.03	N/A							108	47.7	2460	29.5																
		Iron, Dissolved	0.449	0.267	N/A							18.8	16.8	0.05U																	
		Lead	0.015	0.129	0.015							0.0232	0.00364	0.393																	
		Magnesium	N/A	16.262	N/A							16.8	13.7	34	10.5																
		Magnesium, Dissolved	N/A	16.215	N/A							14.4	13.1	12.7																	
		Manganese	0.035	0.119	N/A							2.85	0.506	72.4	0.42																
		Manganese, Dissolved	0.035	0.068	N/A							1.81	0.367	12.8																	
		Mercury	0.000444	0.0002	0.002							0.00037	0.0001U	0.000354	0.0002U																
		Molybdenum	0.00753	0.05	N/A							0.0842	0.0347	0.0415																	
		Molybdenum, Dissolved	0.00753	0.05	N/A							0.0194	0.00936	0.001U																	
		Nickel	0.0301	0.682	N/A							0.109	0.05U	1.15	0.06																
		Nickel, Dissolved	0.0301	0.305	N/A							0.0499	0.0289	0.0233																	
		Potassium	N/A	5.195	N/A							0.005U	0.00572	0.00505	0.005U																
		Selenium	0.00754	0.005	0.05							0.005U	0.00569	0.00692																	
		Selenium, Dissolved	0.00754	0.005	0.05							0.001U	0.001U	0.00571	0.05U																
		Silver	0.0075	0.011	N/A							42.8	37.8	0.5U	34.6																
		Sodium	N/A	59.45	N/A							43.9	38.1	35.6																	
		Sodium, Dissolved	N/A	60.433	N/A							0.002U	0.002U	0.0084	0.2U																
		Strontium	0.901	N/A	N/A										0.08																
		Thallium	N/A	0.056	0.002							0.00392	0.00866	0.0858																	
		Uranium	0.000906	0.002	0.03							0.001U	0.00132	0.001U																	
		Uranium, Dissolved	0.000906	0.002	0.03							0.2U	0.2U	2U	0.1U																
		Vanadium	0.00925	0.134	N/A							1.33	0.273	4.12	0.2U																
		Zinc	0.45	0.054	N/A							0.555	0.16	0.0381																	
		Zinc, Dissolved	0.45	0.049	N/A																										
		Radionuclides (pCi/L)																													
		Americium-241	0.371	N/A	N/A							0.0907U	0.0891U	0.059U																	
		Neptunium-237	0.573	0.8	N/A							0.0587U	0.0594U	0.0639U																	
		Technetium-99	14	22.3	900							530	129	45.7	138																
		Thorium-228	0.129	N/A	N/A							0.294	0.116U	0.196																	

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Table 4.40. SWMU 7 Locations of Groundwater Contaminants (Continued)

Hydro Unit	Depth Interval	Analysis	Child Resident NAL	Bkgd	MCL	007-001	007-002	007-003B	007-005	007-007	007-008	007-009	007-010	007-011	DG-005	GW-01	GW-02	GW-03	MW185	MW186	MW187	MW339	MW340	WB-12	WB-13	WB-7	WB-8	WB-9	WBP-12A	WBP-9A	
	80	Thorium-230	0.424	1.1	N/A							0.192U	0.193U	0.165U																	
		Thorium-232	0.382	N/A	N/A							0.0975U	0.0936	0.082U																	
		Thorium-234	N/A	N/A	N/A							19.4U	20.2U	26U																	
		Uranium	N/A	N/A	N/A							6.32	2.17	15.3																	
		Uranium-234	0.546	0.7	N/A							2.34	0.361U	6.79																	
		Uranium-235	N/A	0.3	N/A							0.14	0.0562U	0.322																	
		Uranium-238	0.443	0.7	N/A							3.83	1.81	8.14																	
		Semivolatiles (mg/L)																													
		2,4-Dimethylphenol	0.023	N/A	N/A							0.0048U	0.0049U	0.0047U																	
		Bis(2-ethylhexyl)phthalate	0.00312	N/A	0.006							0.0048U	0.0049U	0.0047U																	
		Volatiles (mg/L)																													
		1,1,1-Trichloroethane	0.0335	N/A	0.2							1U	0.1U	0.005U																	
		1,1,2-Trichloroethane	0.000238	N/A	0.005							1U	0.1U	0.005U																	
		1,1-Dichloroethane	0.0363	N/A	N/A							1U	0.1U	0.005U																	
		1,1-Dichloroethene	0.000047	N/A	0.007							1U	0.1U	0.005U	0.0005																
		1,2-Dichloroethane	0.000147	N/A	0.005							1U	0.1U	0.005U																	
		2-Butanone	0.0868	N/A	N/A							2U	0.2U	0.01U																	
		2-Hexanone	N/A	N/A	N/A							2U	0.2U	0.01U																	
		4-Methyl-2-pentanone	0.00722	N/A	N/A							2U	0.2U	0.01U																	
		Acetone	0.0275	N/A	N/A							2U	0.2U	0.01U																	
		Carbon tetrachloride	0.000181	N/A	0.005							1U	0.1U	0.005U																	
		Chloroform	0.000287	N/A	N/A							1U	0.1U	0.005U																	
		Chloromethane	0.00167	N/A	N/A							1U	0.1U	0.005U																	
		cis-1,2-Dichloroethene	0.00273	N/A	0.07							1U	0.1U	0.005U	0.003																
		Tetrachloroethene	0.000582	N/A	0.005							1U	0.1U	0.005U																	
		Toluene	0.0338	N/A	1							1U	0.1U	0.005U																	
		trans-1,2-Dichloroethene	0.00548	N/A	0.1							1U	0.1U	0.005U	0.001U																
		Trichloroethene	0.0016	N/A	0.005							18	1.7	0.12	0.24																
		Vinyl chloride	0.000035	N/A	0.002							0.4U	0.04U	0.002U	0.001U																
	85-87	Metals (mg/L)																													
		Aluminum	1.49	2.189	N/A											930	18	570													
		Antimony	0.000564	0.06	0.006											0.0043															
		Arsenic	0.000035	0.005	0.01											0.25	0.091	0.18													
		Barium	0.104	0.235	2											5	0.45	3.1													
		Beryllium	0.00264	0.004	0.004											0.068	0.006	0.031													
		Cadmium	0.000661	0.01	0.005											0.012	0.0003U	0.0003U													
		Calcium	N/A	41.238	N/A											470	44	280													
		Chromium	1.76	0.144	0.1											1.3	0.13	0.86													
		Cobalt	0.0906	0.045	N/A											0.49	0.037	0.55													
		Copper	0.0557	0.036	1.3											1.3	0.1	0.54													
		Iron	0.449	5.03	N/A											1,600	340	890													
		Lead	0.015	0.129	0.015											1.6	0.019	0.35													
		Magnesium	N/A	16.262	N/A											110	15	43													
		Manganese	0.035	0.119	N/A											17	4.2	14													
		Mercury	0.000444	0.0002	0.002											0.0005	0.087U	0.16													
		Molybdenum	0.00753	0.05	N/A											0.1U	0.087U	0.16													
		Nickel	0.0301	0.682	N/A											1.2	0.23	0.58													
		Potassium	N/A	5.195	N/A											51	3.3	49													
		Selenium	0.00754	0.005	0.05											0.0038U	0.0038U	0.0038U													
		Silver	0.0075	0.011	N/A											0.0045U	0.0045U	0.0045U													
		Sodium	N/A	59.45	N/A											46	37	27													
		Thallium	N/A	0.056	0.002											0.063	0.0055U	0.04													
		Tin	0.894	N/A	N/A											0.75	0.23	0.46													
		Uranium	0.000906	0.002	0.03											0.0092U	0.063	0.0092U													
		Vanadium	0.00925	0.134	N/A											1.3	0.082	1.1													
		Zinc	0.45	0.054	N/A											6.5	2.1	3.6													
		Radionuclides (pCi/L)																													
		Neptunium-237	0.573	0.8	N/A											0.04	0.01U	0.05U													
		Plutonium-239	0.286	0.1	N/A											0U	0.13	0.78													

RGA

Table 4.40. SWMU 7 Locations of Groundwater Contaminants (Continued)

Hydro Unit	Depth Interval	Analysis	Child Resident NAL	Bkgd	MCL	007-001	007-002	007-003B	007-005	007-007	007-008	007-009	007-010	007-011	DG-005	GW-01	GW-02	GW-03	MW185	MW186	MW187	MW339	MW340	WB-12	WB-13	WB-7	WB-8	WB-9	WBP-12A	WBP-9A		
	85-87	Technetium-99	14	22.3	900										177	633	56.9	3.4U														
		Thorium-230	0.424	1.1	N/A											1.56	8.42	1.78														
		Uranium-234	0.546	0.7	N/A											14.5	1.25	6.98														
		Uranium-235/236	N/A	0.3	N/A											1.36	0.08	0.85														
		Uranium-238	0.443	0.7	N/A											20.3	1.25	8														
		Volatiles (mg/L)																														
		1,1,1-Trichloroethane	0.0335	N/A	0.2										0.1U	0.00036U	0.00036U	0.00036U														
		1,1,2-Trichloro-1,2,2,2-trifluoroethane	2.7	N/A	N/A											0.00036U	0.00036U	0.00036U														
		1,1,2-Trichloroethane	0.000238	N/A	0.005										0.1U	0.00037U	0.00037U	0.00037U														
		1,1-Dichloroethane	0.0363	N/A	N/A										0.0006	0.00075	0.00044U	0.00044U														
		1,1-Dichloroethene	0.000047	N/A	0.007										0.00027U	0.00027U	0.00027U	0.00027U														
		1,2-Dichloroethane	0.000147	N/A	0.005										0.2U	0.0034U	0.0034U	0.0034U														
		2-Butanone	0.0868	N/A	N/A										0.2U	0.0011U	0.0011U	0.0011U														
		2-Hexanone	N/A	N/A	N/A										0.2U	0.0011U	0.0011U	0.0011U														
		4-Methyl-2-pentanone	0.00722	N/A	N/A										0.2U	0.013	0.14	0.023U														
		Acetone	0.0275	N/A	N/A										0.2U	0.016	0.00035U	0.00035U														
		Carbon tetrachloride	0.000181	N/A	0.005											0.0041	0.0032	0.00041U														
		Chloroform	0.000287	N/A	N/A										0.1U	0.00037U	0.00037U	0.00037U														
		Chloromethane	0.00167	N/A	N/A										0.003	0.01	0.0062	0.00047U														
		cis-1,2-Dichloroethene	0.00273	N/A	0.07										0.1U	0.00091	0.00035U	0.00035U														
		Tetrahydroethene	0.000582	N/A	0.005										0.1U	0.00044U	0.00025	0.00044U														
		Toluene	0.0338	N/A	1										0.1U	0.00048U	0.00048U	0.00048U														
		trans-1,2-Dichloroethene	0.00548	N/A	0.1										0.32	2.9	0.35	0.0012														
		Trichloroethene	0.0016	N/A	0.005										0.001U	0.0013U	0.0013U	0.0013U														
		Vinyl chloride	0.000035	N/A	0.002																											
	90-91	Metals (mg/L)																														
		Aluminum	1.49	2.189	N/A							29	5.78	998	0.9		500															
		Antimony	0.000564	0.06	0.006							0.005U	0.005U																			
		Arsenic	0.000035	0.005	0.01							0.07	0.0292	0.22	0.01		0.2															
		Arsenic, Dissolved	0.000035	0.005	0.01							0.0436	0.0157	0.00608																		
		Barium	0.104	0.235	2							1.58	0.275	8.54	0.26																	
		Barium, Dissolved	0.104	0.2	2							1.26	0.239	0.354																		
		Beryllium	0.00264	0.004	0.004							0.00276	0.00144	0.0732	0.005U																	
		Cadmium	0.000661	0.01	0.005							0.00131	0.0006U	0.0197	0.005U																	
		Cadmium, Dissolved	0.000661	0.01	0.005							0.00066U	0.0006U	0.0006U																		
		Calcium	N/A	41.238	N/A							166	31.1	61.3	32.2		110															
		Calcium, Dissolved	N/A	38.166	N/A							160	31.4	34.8																		
		Chromium	1.76	0.144	0.1							0.206	0.0275	1.46	0.05U																	
		Cobalt	0.0906	0.045	N/A							0.0914	0.0072	1.32	0.01U																	
		Cobalt, Dissolved	0.0906	0.045	N/A							0.0587	0.00448	0.0668																		
		Copper	0.0557	0.036	1.3							0.2U	0.02U	2U	0.05U																	
		Iron	0.449	5.03	N/A							162	86.7	2.410	8.88																	
		Iron, Dissolved	0.449	0.267	N/A							57.6	18.2	0.057																		
		Lead	0.015	0.129	0.015							0.0177	0.00355	0.489																		
		Magnesium	N/A	16.262	N/A							79.1	13.2	38.8	12.9																	
		Magnesium, Dissolved	N/A	16.215	N/A							72.9	13.1	12.3																		
		Manganese	0.035	0.119	N/A							9.4	0.445	58.9	0.06																	
		Manganese, Dissolved	0.035	0.068	N/A							8.95	0.338	7.24																		
		Mercury	0.000444	0.0002	0.002							0.00046	0.0001U	0.000785	0.0002U																	
		Molybdenum	0.00753	0.05	N/A							0.0889	0.0282	0.0239																		
		Molybdenum, Dissolved	0.00753	0.05	N/A							0.0269	0.00434	0.001U																		
		Nickel	0.0301	0.682	N/A							0.15	0.0441	1.18	0.05U																	
		Nickel, Dissolved	0.0301	0.305	N/A							0.0704	0.0343	0.0171																		
		Potassium	N/A	5.195	N/A							0.0377	0.0055	0.005U	2U																	
		Selenium	0.00754	0.005	0.05							0.0266	0.00508	0.00775																		
		Selenium, Dissolved	0.00754	0.005	0.05							0.001U	0.001U	0.00289	0.05U																	
		Silver	0.0075	0.011	N/A							320	41.4	0.5U	37.4																	
		Sodium	N/A																													

Table 4.40. SWMU 7 Locations of Groundwater Contaminants (Continued)

Hydro Unit	Depth Interval	Analysis	Child Resident NAL	Bkgd	MCL	007-001	007-002	007-003B	007-005	007-007	007-008	007-009	007-010	007-011	DG-005	GW-01	GW-02	GW-03	MW185	MW186	MW187	MW339	MW340	WB-12	WB-13	WB-7	WB-8	WB-9	WBP-12A	WBP-9A	
	90-91	Strontium	0.901	N/A	N/A							0.002U	0.002U	0.0106	0.09		0.087														
		Thallium	N/A	0.056	0.002										0.2U		0.92														
		Tin	0.894	N/A	N/A												0.0092U														
		Uranium	0.000906	0.002	0.03							0.00413	0.00197	0.093																	
		Uranium, Dissolved	0.000906	0.002	0.03							0.001U	0.001U	0.001U																	
		Vanadium	0.00925	0.134	N/A							0.2U	0.02U	2U	0.1U		0.83														
		Zinc	0.45	0.054	N/A							1.05	0.333	4.28	0.2U		5.4														
		Zinc, Dissolved	0.45	0.049	N/A							0.257	0.208	0.0312																	
		Radionuclides (pCi/L)																													
		Americium-241	0.371	N/A	N/A							0.0991U	0.0901U	0.0628U																	
		Neptunium-237	0.573	0.8	N/A							0.0992U	0.0666U	0.0605U			0.01U														
		Plutonium-239	0.286	0.1	N/A												0.01U														
		Technetium-99	14	22.3	900							221	193	64	187		122														
		Thorium-228	0.129	N/A	N/A							0.401	0.103U	0.187																	
		Thorium-230	0.424	1.1	N/A							0.231U	0.171U	0.168U			1.38														
		Thorium-232	0.382	N/A	N/A							0.157U	0.057U	0.0849U																	
		Thorium-234	N/A	N/A	N/A							21.7U	19.9U	141																	
		Uranium	N/A	N/A	N/A							2.72	7.03	17.8																	
		Uranium-234	0.546	0.7	N/A							1.08	1.44	8.71			10.2														
		Uranium-235	N/A	0.3	N/A							0.0639	0.121	0.34																	
		Uranium-235/236	N/A	0.3	N/A												1.55														
		Uranium-238	0.443	0.7	N/A							1.7	5.47	8.74			5.72														
		Semivolatiles (mg/L)																													
		2,4-Dimethylphenol	0.023	N/A	N/A							0.0085	0.0048U	0.0047U																	
		Bis(2-ethylhexyl)phthalate	0.00312	N/A	0.006							0.0048U	0.0048U	0.0047U																	
		Volatiles (mg/L)																													
		1,1,1-Trichloroethane	0.0335	N/A	0.2							0.12U	0.25U	0.005U			0.00036U														
		1,1,2-Trichloro-1,2,2-trifluoroethane	2.7	N/A	N/A												0.00036U														
		1,1,2-Trichloroethane	0.000238	N/A	0.005							0.12U	0.25U	0.005U			0.00033U														
		1,1-Dichloroethane	0.0363	N/A	N/A							0.12U	0.25U	0.005U			0.00037U														
		1,1-Dichloroethene	0.00047	N/A	0.007							0.12U	0.25U	0.0052	0.0008		0.00044U														
		1,2-Dichloroethane	0.000147	N/A	0.005							0.12U	0.25U	0.005U			0.00027U														
		2-Butanone	0.0868	N/A	N/A							0.25U	0.5U	0.01U			0.0034U														
		2-Hexanone	N/A	N/A	N/A							0.25U	0.5U	0.01U			0.0011U														
		4-Methyl-2-pentanone	0.00722	N/A	N/A							0.25U	0.5U	0.01U			0.0011U														
		Acetone	0.0275	N/A	N/A							0.25U	0.5U	0.01U			0.43														
		Carbon tetrachloride	0.000181	N/A	0.005							0.12U	0.25U	0.005U			0.00035U														
		Chloroform	0.000287	N/A	N/A							0.12U	0.25U	0.005U			0.0045														
		Chloromethane	0.00167	N/A	N/A							0.12U	0.25U	0.005U			0.00037U														
		cis-1,2-Dichloroethene	0.00273	N/A	0.07							0.58	0.25U	0.005U	0.003		0.0064														
		Tetrachloroethene	0.000582	N/A	0.005							0.12U	0.25U	0.005U			0.00035U														
		Toluene	0.0338	N/A	1							0.12U	0.25U	0.005U			0.0025														
		trans-1,2-Dichloroethene	0.00548	N/A	0.1							0.12U	0.25U	0.005U	0.001U		0.00048U														
		Trichloroethene	0.0016	N/A	0.005							1.7	3.6	0.2	0.26		0.41														
		Vinyl chloride	0.000035	N/A	0.002							0.3	0.1U	0.002U	0.001U		0.0013U														
	95-96	Metals (mg/L)																													
		Aluminum	1.49	2.189	N/A											670							1.36								
		Antimony	0.000564	0.06	0.006											0.003							0.005U								
		Arsenic	0.000035	0.005	0.01											0.42							0.00337								
		Arsenic, Dissolved	0.000035	0.005	0.01																		0.00149								
		Barium	0.104	0.235	2											3.5							0.148								
		Barium, Dissolved	0.104	0.2	2																		0.144								
		Beryllium	0.00264	0.004	0.004											0.062							0.0006	0.0003							
		Cadmium	0.000661	0.01	0.005											0.016							0.001U								
		Cadmium, Dissolved	0.000661	0.01	0.005																		0.001U								
		Calcium	N/A	41.238	N/A											160							43	32							
		Calcium, Dissolved	N/A	38.166	N/A																		35.2								
		Chromium	1.76	0.144	0.1											1.3							0.0723								

Table 4.40. SWMU 7 Locations of Groundwater Contaminants (Continued)

Hydro Unit	Depth Interval	Analysis	Child Resident NAL	Bkgd	MCL	007-001	007-002	007-003B	007-005	007-007	007-008	007-009	007-010	007-011	DG-005	GW-01	GW-02	GW-03	MW185	MW186	MW187	MW339	MW340	WB-12	WB-13	WB-7	WB-8	WB-9	WBP-12A	WBP-9A
	95-96	Cobalt	0.0906	0.045	N/A											0.31					0.001U									
		Cobalt, Dissolved	0.0906	0.045	N/A																	0.001U								
		Copper	0.0557	0.036	1.3											0.91						0.02U	0.025U							
		Iron	0.449	5.03	N/A											2.100						24	11							
		Iron (2+)	N/A	N/A	N/A																	0.75	0.02U							
		Iron, Dissolved	0.449	0.267	N/A																	0.1U								
		Lead	0.015	0.129	0.015											0.65						0.005U								
		Magnesium	N/A	16.262	N/A											58						17	14							
		Magnesium, Dissolved	N/A	16.215	N/A																	14								
		Manganese	0.035	0.119	N/A											20						1.9	0.22							
		Manganese, Dissolved	0.035	0.068	N/A																	0.00917								
		Mercury	0.000444	0.0002	0.002											0.0003						0.0002U								
		Molybdenum	0.00753	0.05	N/A											0.17						0.00634								
		Molybdenum, Dissolved	0.00753	0.05	N/A																	0.001U								
		Nickel	0.0301	0.682	N/A											0.98						0.0224								
		Nickel, Dissolved	0.0301	0.305	N/A																	0.0257								
		Potassium	N/A	5.195	N/A											47						3.2	1.9							
		Potassium, Dissolved	N/A	4.096	N/A																	1.85								
		Selenium	0.00754	0.005	0.05											0.0038U						0.00964								
		Selenium, Dissolved	0.00754	0.005	0.05																	0.005U								
		Silver	0.0075	0.011	N/A											0.0045U						0.001U								
		Sodium	N/A	59.45	N/A											38						42.9	44.1							
		Sodium, Dissolved	N/A	60.433	N/A																	42.3								
		Thallium	N/A	0.056	0.002											0.077														
		Tin	0.894	N/A	N/A											1														
		Uranium	0.000906	0.002	0.03											0.0092U						0.09	0.05U							
		Uranium, Dissolved	0.000906	0.002	0.03																	0.001U								
		Vanadium	0.00925	0.134	N/A											1.1						0.018	0.014							
		Zinc	0.45	0.054	N/A											5.4						0.02U								
		Zinc, Dissolved	0.45	0.049	N/A																	0.02U								
		Radionuclides (pCi/L)																												
		Americium-241	0.371	N/A	N/A																	46.5U	0.41							
		Neptunium-237	0.573	0.8	N/A											0.03						7.55	0.989U							
		Plutonium-239	0.286	0.1	N/A											-0.28U						0.13U	0.08U							
		Potassium-40	N/A	N/A	N/A																	362U	47.9							
		Technetium-99	14	22.3	900										118	1.550						5.116.9	747							
		Thorium-228	0.129	N/A	N/A																	0.255U	0.278							
		Thorium-230	0.424	1.1	N/A											1.37						0.76	0.34							
		Thorium-232	0.382	N/A	N/A																	0.269U	0.419U							
		Thorium-234	N/A	N/A	N/A																	269U	282U							
		Uranium	N/A	N/A	N/A																	40U	40U							
		Uranium-234	0.546	0.7	N/A											6.34						0.48	0.45							
		Uranium-235	0.538	0.3	N/A																	2.2U	2.2U							
		Uranium-235/236	N/A	0.3	N/A											0.83						0.01U	0.03U							
		Uranium-238	0.443	0.7	N/A											6.27						10.4U	0.18							
		Volatiles (mg/L)																												
		1,1,1-Trichloroethane	0.0335	N/A	0.2											0.0013						2.5U	0.5U							
		1,1,2-Trichloro-1,2,2-trifluoroethane	2.7	N/A	N/A											0.00036U						1U	0.5U							
		1,1,2-Trichloroethane	0.000238	N/A	0.005											0.009						2.5U	0.5U							
		1,1-Dichloroethane	0.0363	N/A	N/A											0.00037U						1U	0.5U							
		1,1-Dichloroethene	0.000047	N/A	0.007										0.0009	0.0037						1U	0.5U							
		1,2-Dichloroethane	0.000147	N/A	0.005											0.00068						1U	0.5U							
		2-Butanone	0.0868	N/A	N/A											0.0034U							0.5U							
		2-Hexanone	N/A	N/A	N/A											0.0011U							0.5U							
		4-Methyl-2-pentanone	0.00722	N/A	N/A											0.0011U							0.5U							
		Acetone	0.0275	N/A	N/A											0.016U							0.5U							
		Carbon tetrachloride	0.000181	N/A	0.005											0.0047							1U	0.5U						
		Chloroform	0.000287	N/A	N/A											0.012						1U	0.5U							

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Table 4.40. SWMU 7 Locations of Groundwater Contaminants (Continued)

Hydro Unit	Depth Interval	Analysis	Child Resident NAL	Bkgd	MCL	007-001	007-002	007-003B	007-005	007-007	007-008	007-009	007-010	007-011	DG-005	GWW-01	GWW-02	GWW-03	MW185	MW186	MW187	MW339	MW340	WB-12	WB-13	WB-7	WB-8	WB-9	WBP-12A	WBP-9A				
	95-96	Chloromethane	0.00167	N/A	N/A										0.0037U	0.00037U						0.25U												
		cis-1,2-Dichloroethene	0.00273	N/A	0.07										0.0039	0.023						2.1	0.013											
		Tetrachloroethene	0.000582	N/A	0.005											0.0048						1U	0.5U											
		Toluene	0.0338	N/A	1											0.00044U						2.5U	0.5U											
		trans-1,2-Dichloroethene	0.00548	N/A	0.1										0.001U	0.00048U						1U	0.00049											
		Trichloroethene	0.0016	N/A	0.005										0.42	14						25	6.5											
		Vinyl chloride	0.000035	N/A	0.002										0.001U	0.0013U						0.4U	0.2U											
	100	Metals (mg/L)																																
		Aluminum	1.49	2.189	N/A								7.62																					
		Antimony	0.000564	0.06	0.006								0.005U																					
		Arsenic	0.000035	0.005	0.01								0.01U																					
		Arsenic, Dissolved	0.000035	0.005	0.01								0.001U																					
		Barium	0.104	0.235	2								0.174																					
		Barium, Dissolved	0.104	0.2	2								0.13																					
		Beryllium	0.00264	0.004	0.004								0.001U																					
		Cadmium	0.000661	0.01	0.005								0.0006U																					
		Cadmium, Dissolved	0.000661	0.01	0.005								0.0006U																					
		Calcium	N/A	41.238	N/A								17.8																					
		Calcium, Dissolved	N/A	38.166	N/A								17.2																					
		Chromium	1.76	0.144	0.1								0.1U																					
		Cobalt	0.0906	0.045	N/A								0.015																					
		Cobalt, Dissolved	0.0906	0.045	N/A								0.0121																					
		Copper	0.0557	0.036	1.3								0.2U																					
		Iron	0.449	5.03	N/A								42.7																					
		Iron, Dissolved	0.449	0.267	N/A								22.1																					
		Lead	0.015	0.129	0.015								0.00515																					
		Magnesium	N/A	16.262	N/A								8.21																					
		Magnesium, Dissolved	N/A	16.215	N/A								7.44																					
		Manganese	0.035	0.119	N/A								0.516																					
		Manganese, Dissolved	0.035	0.068	N/A								0.437																					
		Mercury	0.000444	0.0002	0.002								0.00001U																					
		Molybdenum	0.00753	0.05	N/A								0.0237																					
		Molybdenum, Dissolved	0.00753	0.05	N/A								0.0142																					
		Nickel	0.0301	0.682	N/A								0.05U																					
		Nickel, Dissolved	0.0301	0.305	N/A								0.0311																					
		Selenium	0.00754	0.005	0.05								0.005U																					
		Selenium, Dissolved	0.00754	0.005	0.05								0.005U																					
		Silver	0.0075	0.011	N/A								0.001U																					
		Sodium	N/A	59.45	N/A								15.9																					
		Sodium, Dissolved	N/A	60.433	N/A								16.1																					
		Thallium	N/A	0.056	0.002								0.002U																					
		Uranium	0.000906	0.002	0.03								0.00374																					
		Uranium, Dissolved	0.000906	0.002	0.03								0.001U																					
		Vanadium	0.00925	0.134	N/A								0.2U																					
		Zinc	0.45	0.054	N/A								0.37																					
		Zinc, Dissolved	0.45	0.049	N/A								0.274																					
		Radionuclides (pCi/L)																																
		Americium-241	0.371	N/A	N/A								0.0894U																					
		Neptunium-237	0.573	0.8	N/A								0.0584U																					
		Technetium-99	14	22.3	900								36.6																					
		Thorium-228	0.129	N/A	N/A								0.155																					
		Thorium-230	0.424	1.1	N/A								0.193U																					
		Thorium-232	0.382	N/A	N/A								0.0673																					
		Thorium-234	N/A	N/A	N/A								20U																					
		Uranium	N/A	N/A	N/A								7.17																					
		Uranium-234	0.546	0.7	N/A								1.11																					
		Uranium-235	N/A	0.3	N/A								0.125																					
		Uranium-238	0.443	0.7	N/A								5.94																					

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Table 4.40. SWMU 7 Locations of Groundwater Contaminants (Continued)

Hydro Unit	Depth Interval	Analysis	Child Resident NAL	Bkgd	MCL	007-001	007-002	007-003B	007-005	007-007	007-008	007-009	007-010	007-011	DG-005	GWW-01	GWW-02	GWW-03	MW185	MW186	MW187	MW339	MW340	WB-12	WB-13	WB-7	WB-8	WB-9	WBP-12A	WBP-9A		
McNairy	111	Metals (mg/L)																														
		Aluminum	1.49	0.687	N/A												470	560	300													
		Arsenic	0.00035	0.005	0.01												0.21	0.13	0.12													
		Barium	0.104	0.296	2												1.9	3.5	2.3													
		Beryllium	0.00264	0.017	0.004												0.026	0.06	0.015													
		Cadmium	0.000661	0.01	0.005												0.028	0.011	0.0003U													
		Calcium	N/A	38.858	N/A												66	100	110													
		Chromium	1.76	0.06	0.1												1.7	1.4	0.45													
		Cobalt	0.0906	0.096	N/A												0.2	0.35	0.22													
		Copper	0.0557	0.057	1.3												1.5	0.78	0.26													
		Iron	0.449	18.36	N/A												1.900	1.900	550													
		Lead	0.015	0.05	0.015												0.39	0.29	0.16													
		Magnesium	N/A	13.418	N/A												37	45	28													
		Manganese	0.035	0.941	N/A												12	79	15													
		Mercury	0.000444	0.0002	0.002												0.0004	0.0003	0.0003													
		Molybdenum	0.00753	0.05	N/A												0.35	0.065U	0.085U													
		Nickel	0.0301	0.109	N/A												1.2	0.83	0.26													
		Potassium	N/A	55.752	N/A												37	32	29													
		Selenium	0.00754	0.005	0.05												0.065	0.0038U	0.039													
		Sodium	N/A	29.2	N/A												13	38	27													
		Thallium	N/A	0.644	0.002												0.037U	0.071	0.023													
		Tin	0.894	N/A	N/A												0.92	0.89	0.31													
		Vanadium	0.00925	0.126	N/A												0.95	0.81	0.76													
		Zinc	0.45	0.142	N/A												15	4.6	1.7													
				Radionuclides (pCi/L)																												
				Plutonium-239	0.286	0.2	N/A											-0.53U	-0.142U	0.19U												
				Technetium-99	14	20.6	900											7.9U	95.1	1.3U												
				Thorium-230	0.424	1.5	N/A											1.38	2	0.8												
				Uranium-234	0.546	0.3	N/A											0.26	4.75	4.26												
				Uranium-235/236	N/A	0.2	N/A											0.02U	0.52	0.36												
				Uranium-238	0.443	0.3	N/A											0.17U	4.73	3.94												
				Volatiles (mg/L)																												
				Acetone	0.0275	N/A	N/A											0.0073U	0.2	0.014U												
				Chloroform	0.000287	N/A	N/A											0.00084	0.0038	0.00041U												
				cis-1,2-Dichloroethene	0.00273	N/A	0.07											0.0011	0.0056	0.00047U												
				Toluene	0.0338	N/A	1											0.00052	0.00078	0.00044U												
				Trichloroethene	0.0016	N/A	0.005											0.11	0.32	0.021												

Blank cells indicate interval was not sampled for the specified analysis. Maximum value shown for each depth interval.

N/A = not applicable

Bold indicates result is greater than NAL value.

Italics indicates result is greater than MCL value.

Bold + Italics indicate result is greater than both NAL and MCL values.

Shading indicates result is greater than the background value.

4.9 SWMU 30

4.9.1 Subsurface Soils

SWMU 30 was used to burn combustible trash, which may have contained uranium contamination. Ash and debris then were buried in a pit contained within the SWMU (excavated to a depth of approximately 12 ft). Table 4.41 summarizes the subsurface soil contaminants for SWMU 30 determined by the data review. Figure 4.7 depicts the sample locations.

As in neighboring SWMU 7, metals concentrations in subsurface soil samples of SWMU 30 rarely exceed background levels. Iron, manganese, and vanadium are the most frequent metals to be detected above the excavation worker NALs. Figure 4.61 illustrates the distribution of vanadium in subsurface soils at SWMU 30. There was only one detection above background, which occurred at boring WB-1 in the 5 to 10 ft sample. Concentrations above the excavation worker NALs extended throughout the depth of the UCRS.

Few organic compounds are present in subsurface soils at SWMU 30. The screening steps identified benzo(a)pyrene and Total PAHs as organic contaminants, at frequencies of one in 26 analyses for both. Analyses of the RI samples did not detect PCBs, but a review of historical data identified four PCB-1260 detections (locations WB-1, WB-4, WB-5, and WB-11) and one PCB-1254 detection (location WB-3), all at depths of 6 to 7 ft and distributed across the SWMU. The highest level, 0.18 mg/kg of PCB-1260, was detected in a sample from 6 ft bgs at location WB-1 from within the area of Burial Pit A. Two test pits were excavated in SWMU 30 during the Phase II SI (CH2M HILL 1992). The excavations found ash, bricks, and railroad rails in the test pits. Samples from these pits identified many of the same inorganic, organic, and radionuclide contaminants reported in Table 4.40. In the test pit samples, uranium-234 ranged from 0.26 to 44 pCi/L and uranium-238 ranged from 0.3 to 62 pCi/L.

The uranium isotopes uranium-234 (maximum 20.6 pCi/g), uranium-235/236 (maximum 0.55 pCi/g), and uranium-238 (maximum 37.4 pCi/g) are the only radionuclide contaminants at depths of 10 ft or less. Figure 4.62 provides the uranium-238 distribution in soils, which indicates the limited extent of uranium contamination with all exceedances above background being less than 10 ft bgs.

Locations containing subsurface soil contaminants detected above screening levels are presented in Table 4.42.

Table 4.41. SWMU 30 Subsurface Soil Contaminants

Analysis	Maximum Result		Frequency of Detection ^a	Frequency of Detection	
	Historical Data	RI Data		Above Background Value	Above Excavation Worker NAL
<i>Inorganics (mg/kg)</i>					
Aluminum	19,000	9,270	25/25	1/25	22/25
Arsenic	ND	4.03	18/25	0/25	18/25
Barium	150	126	25/25	0/25	0/25
Beryllium	ND	1.48	7/25	5/25	2/25
Calcium	11,000	1,770	25/25	2/25	N/A
Chromium	49	17.2	25/25	1/25	0/25
Cobalt	14	12.2	17/25	1/25	0/25
Copper	35	18.9	24/25	2/25	0/25
Iron	29,000	22,200	25/25	1/25	25/25
Lead	18	14.3	25/25	0/25	0/25
Magnesium	2,000	1,670	25/25	0/25	0/25
Manganese	1,200	486	25/25	1/25	18/25
Mercury	0.12	0.026	5/25	0/25	0/25
Nickel	32	16.9	21/25	2/25	0/25
Selenium	1	ND	3/25	1/25	0/25
Sodium	ND	187	8/25	0/25	0/25
Tin	8.6	N/A	4/4	N/A	0/4
Uranium	ND	2.03	11/25	0/25	0/25
Vanadium	40	19.3	24/25	1/25	17/25
Zinc	67	59.5	16/25	1/25	0/25
<i>Organics--Semivolatiles (mg/kg)</i>					
1,2,4-Trichlorobenzene	0.033	ND	1/26	N/A	0/26
1,4-Dichlorobenzene	0.025	ND	1/31	N/A	0/31
2-Chlorophenol	0.023	ND	1/26	N/A	0/26
Acenaphthene	0.017	ND	1/26	N/A	0/26
Benz(a)anthracene	0.038	ND	3/26	N/A	0/26
Benzo(a)pyrene	0.052	ND	1/26	N/A	1/26
Benzo(b)fluoranthene	0.067	ND	2/26	N/A	0/26
Benzo(k)fluoranthene	0.026	ND	1/26	N/A	0/26
Benzoic acid	ND	0.53	6/26	N/A	0/26
Chrysene	0.035	ND	3/26	N/A	0/26
Fluoranthene	0.054	ND	5/26	N/A	0/26
Phenanthrene	0.015	ND	2/26	N/A	0/26
Pyrene	0.059	ND	4/26	N/A	0/26
<i>Organics--Volatiles (mg/kg)</i>					
1,1-Dichloroethene	ND	0.005	1/26	N/A	0/26
Acetone	0.0065	0.00973	3/26	N/A	0/26
Methylene chloride	0.0018	ND	1/26	N/A	0/26
Toluene	0.0011	ND	2/26	N/A	0/26
Trichloroethene	ND	0.0374	1/26	N/A	0/26
<i>Organics--PCBs (mg/kg)</i>					
PCB, Total	0.18	ND	5/26	N/A	1/26
PCB-1254	0.028	ND	1/26	N/A	0/26
PCB-1260	0.18	ND	4/26	N/A	1/26

Table 4.41. SWMU 30 Subsurface Soil Contaminants (Continued)

Analysis	Maximum Result		Frequency of Detection ^a	Frequency of Detection	
	Historical Data	RI Data		Above Background Value	Above Excavation Worker NAL
Radionuclides(pCi/g)					
Neptunium-237	0.06	ND	2/26	N/A	0/26
Plutonium-239	0.19	N/A	4/5	N/A	0/5
Technetium-99	6.79	ND	5/26	1/26	0/26
Thorium-228	N/A	0.617	21/21	0/21	21/21
Thorium-230	1.44	0.571	17/26	3/26	0/26
Thorium-232	N/A	0.626	21/21	0/21	0/21
Thorium-234	N/A	2.88	7/21	N/A	N/A
Uranium	59	2.97	8/23	N/A	N/A
Uranium-234	20.6	2.46	19/28	5/28	4/28
Uranium-235	N/A	0.0852	1/21	0/21	0/21
Uranium-235/236	0.55	N/A	5/5	3/5	1/5
Uranium-238	37.4	0.77	16/28	6/28	6/28

^a Frequency of detection is the number of detections of an analyte per number of analyses

(includes regular and duplicate samples).

N/A = not applicable

ND = not detected

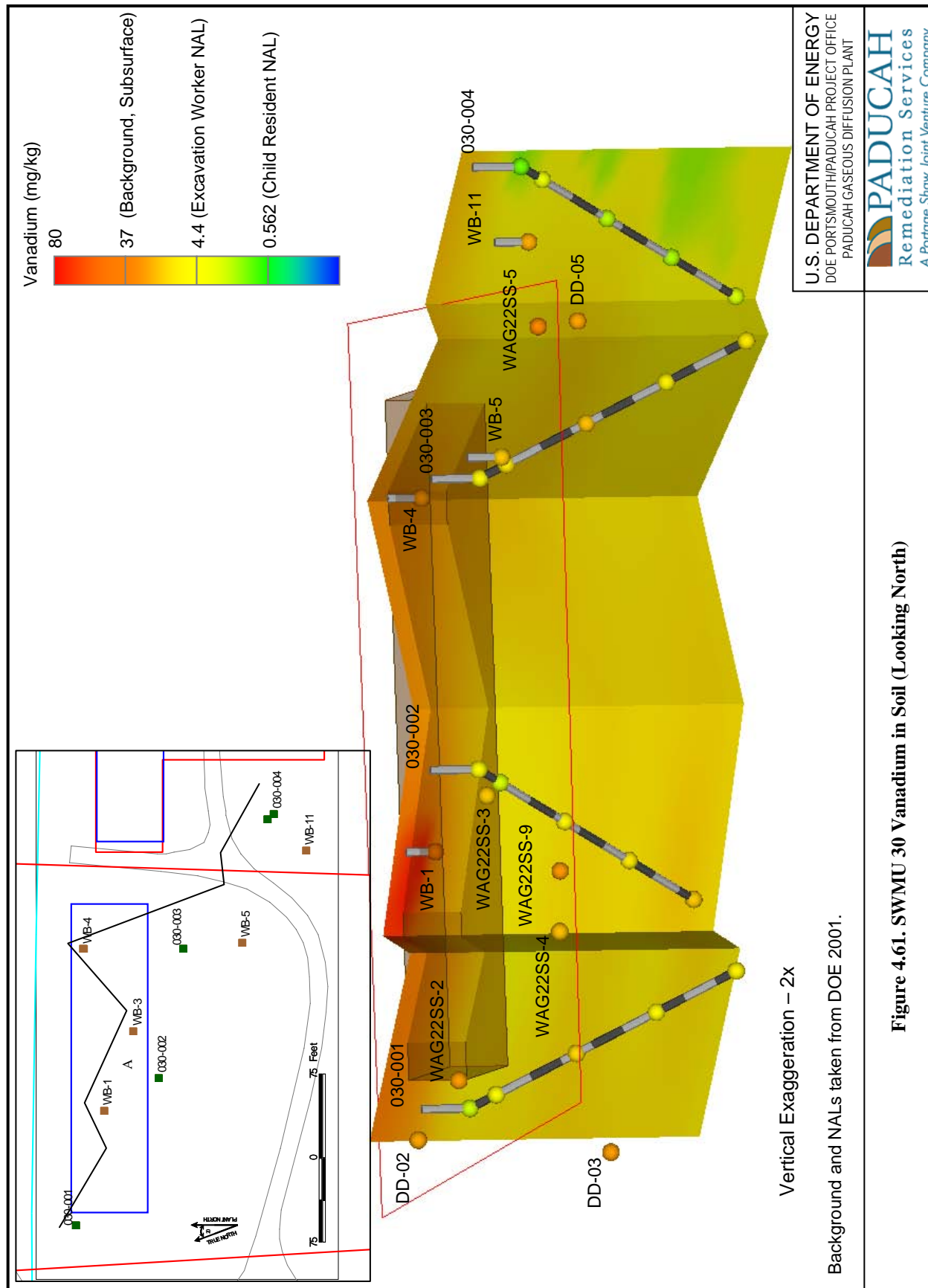


Figure 4.61. SWMU 30 Vanadium in Soil (Looking North)

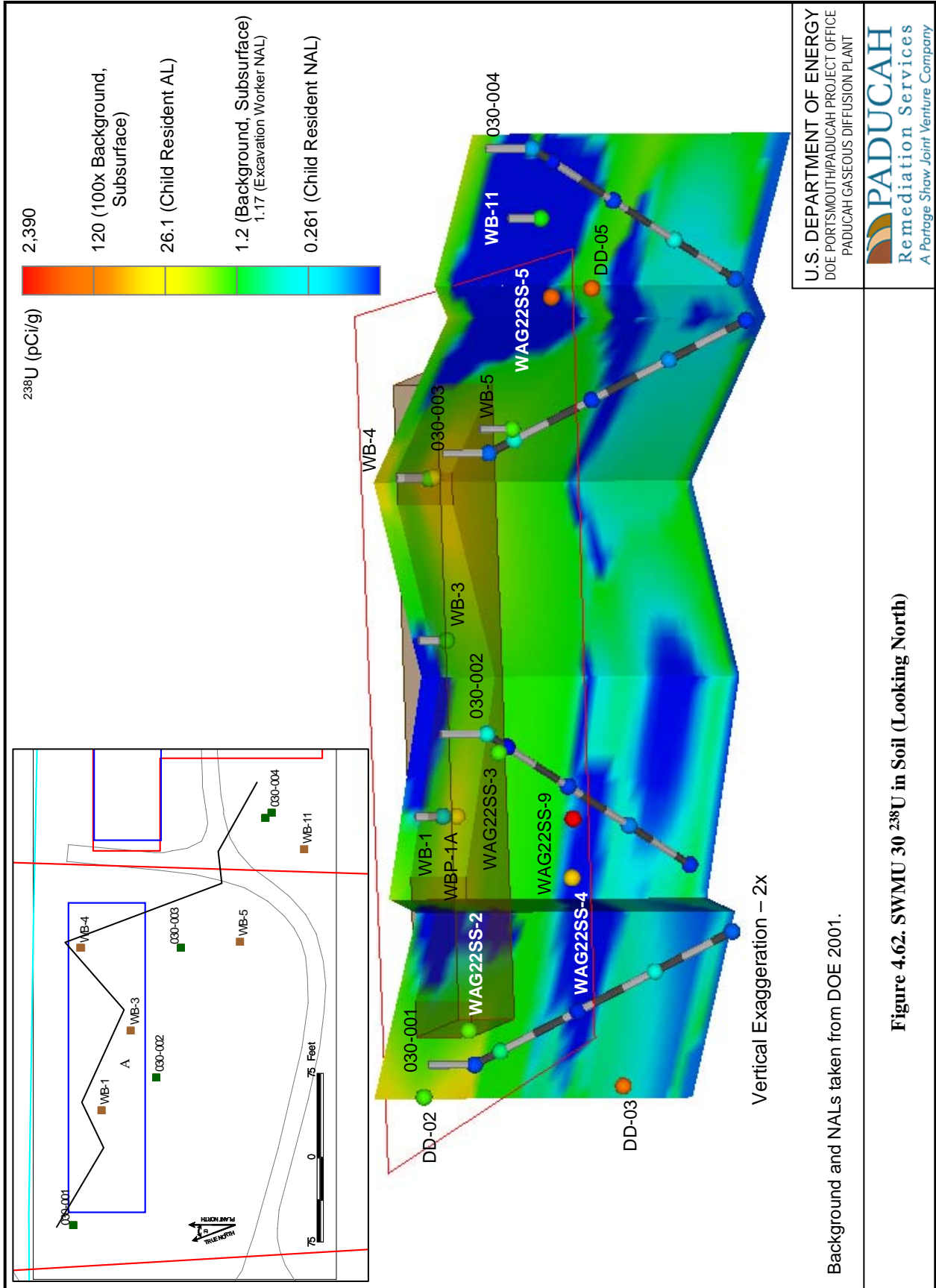


Figure 4.62. SWMU 30 ²³⁸U in Soil (Looking North)

Table 4.42. SWMU 30 Locations of Subsurface Soil Contaminants

Analysis	Depth (ft)	Excavation Worker NAL	Background	RI Data				Historical Data								
				030-001	030-002	030-003	030-004	WB-1	WB-11	WB-3	WB-4	WB-5	WBP-1A	WBP-4A		
Inorganics (mg/kg) Aluminum	05-09	5,250	12,000					19,000	9,100				12,000	11,000		
	10-12	5,250	12,000	7,620	7,430	8,920	8,970									
	15	5,250	12,000	7,700	8,190	4,420	8,110									
	30	5,250	12,000	4,330	6,070	7,150	8,330									
	45	5,250	12,000	8,080	8,750	7,160	3,740									
	60	5,250	12,000	7,060	6,610	8,940	9,270									
Arsenic	05-09	0.324	7.9					6.3U	9.7U				12U	5.1U		
	10-12	0.324	7.9	0.905U	2.09	3.59	1.69									
	15	0.324	7.9	2.62	1.43	4.03	0.946U									
	30	0.324	7.9	1.52	0.954	3.85	0.942U									
	45	0.324	7.9	2.64	3.51	1.67	0.898									
	60	0.324	7.9	2.36	3.76	3.89	2.17									
Barium	05-09	272	170					130	110				150	97		
	10-12	272	170	81.5	55.5	105	94.5									
	15	272	170	35.7	54.5	116	78.7									
	30	272	170	17.1	31.9	27.6	30.7									
	45	272	170	49.4	67.3	58.5	22.2									
	60	272	170	57.2	39.2	126	97.6									
Beryllium	05-09	1.26	0.69					1.4U	1.2U				1.2U	1.2U		
	10-12	1.26	0.69	0.452U	0.442U	0.483U	0.453U									
	15	1.26	0.69	0.49U	0.426U	0.441U	0.473U									
	30	1.26	0.69	0.49U	0.465U	0.478U	0.471U									
	45	1.26	0.69	0.484	0.909	0.438U	0.439U									
	60	1.26	0.69	0.806	1.48	1.41	0.61									
Calcium	05-09	n/a	6,100					10,000	1,500				6,000	11,000		
	10-12	n/a	6,100	1,250	1,530	674	748									
	15	n/a	6,100	987	1,040	1,160	1,210									
	30	n/a	6,100	566	1,030	911	1,270									
	45	n/a	6,100	1,370	1,770	1,140	723									
	60	n/a	6,100	811	1,120	1,320	1,610									
Chromium	05-09	476	43					49	14				32	35		
	10-12	476	43	12.8	13.4	11.9	10.7									
	15	476	43	7.31	12.3	8.65	12.7									
	30	476	43	11.6	7.05	13.2	6.45									
	45	476	43	14.9	17.2	11.3	3.84									
	60	476	43	8.23	9.86	13.8	15.7									

Table 4.42. SWMU 30 Locations of Subsurface Soil Contaminants (Continued)

Analysis	Depth (ft)	Excavation Worker NAL	Background	RI Data				Historical Data							
				030-001	030-002	030-003	030-004	WB-1	WB-11	WB-3	WB-4	WB-5	WBP-1A	WBP-4A	
Cobalt	05-09	1,110	13					5.6	14			12	6.4		
	10-12	1,110	13	2,26U	2,27	5,93	2,56								
	15	1,110	13	5,15	2,66	2,9	2,36U								
	30	1,110	13	2,45U	2,33U	2,39U	2,36U								
	45	1,110	13	2,57	3,57	2,19U	2,19U								
	60	1,110	13	11,5	4,73	12,2	5,9	20	14			35	33		
Copper	05-09	427	25												
	10-12	427	25	6,27	9,66	11,4	8,15								
	15	427	25	4,2	4,79	18,9	5,35								
	30	427	25	3,83	2,57	3,87	2,36U								
	45	427	25	10,7	10,2	4,44	2,62								
	60	427	25	7,05	8,27	12,4	9,99	21,000	17,000			29,000	14,000		
Iron	05-09	2,170	28,000												
	10-12	2,170	28,000	5,940	8,960	13,100	9,400								
	15	2,170	28,000	16,200	7,690	11,300	5,890								
	30	2,170	28,000	10,800	7,900	18,600	6,620								
	45	2,170	28,000	17,700	22,200	11,100	5,020								
	60	2,170	28,000	16,200	21,600	20,000	15,400								
Lead	05-09	50	23					14	18			18	16		
	10-12	50	23	5,22	6	8,13	3,15								
	15	50	23	7,39	6,67	6,61	5,47								
	30	50	23	5,58	4,27	5,51	4,17								
	45	50	23	9,62	8,08	6,17	4,77								
	60	50	23	5,09	4,89	14,3	5,55								
Magnesium	05-09	n/a	2,100					2,000	990			1,300	1,300		
	10-12	n/a	2,100	1,050	1,430	1,090	1,280								
	15	n/a	2,100	689	917	1,140	1,120								
	30	n/a	2,100	271	526	405	627								
	45	n/a	2,100	1,410	1,620	557	348								
	60	n/a	2,100	907	1,020	1,400	1,670	340	1,200			740	230		
Manganese	05-09	56,6	820												
	10-12	56,6	820	66	69,4	188	161								
	15	56,6	820	92,1	62,2	92,9	36,6								
	30	56,6	820	24,8	15,6	26,1	15,7								
	45	56,6	820	70,6	73,1	30,8	22,8								
	60	56,6	820	163	72,3	486	171								
Mercury	05-09	1,17	0,13					0,12	0,12U			0,12U	0,12U		
	10-12	1,17	0,13	0,02	0,019U	0,026	0,019U								
	15	1,17	0,13	0,019U	0,02U	0,018U	0,017U								
	30	1,17	0,13	0,02U	0,019U	0,02U	0,018U								
	45	1,17	0,13	0,02	0,018U	0,019U	0,019U								
	60	1,17	0,13	0,017U	0,019	0,017U	0,019U								

Table 4.42. SWMU 30 Locations of Subsurface Soil Contaminants (Continued)

Analysis	Depth (ft)	Excavation Worker NAL	Background	RI Data				Historical Data								
				030-001	030-002	030-003	030-004	WB-1	WB-11	WB-3	WB-4	WB-5	WB-1A	WBP-4A		
Nickel	05-09	216	22					25	16							
	10-12	216	22	9.35	12	12.9	11.3									
	15	216	22	9.38	6.14	14.9	8.92									
	30	216	22	4.9U	4.65U	4.92	4.71U									
	45	216	22	10.1	12.4	5.07	4.39U									
	60	216	22	10.6	14.7	15.5	16.9									
Selenium	05-09	71.3	0.7					0.69	0.6							
	10-12	71.3	0.7	0.905U	0.885U	0.965U	0.907U									
	15	71.3	0.7	0.98U	0.852U	0.882U	0.946U									
	30	71.3	0.7	0.981U	0.931U	0.955U	0.942U									
	45	71.3	0.7	0.88U	0.999U	0.876U	0.878U									
	60	71.3	0.7	0.948U	0.978U	0.887U	0.953U									
Sodium	05-09	n/a	340					150U	180U							
	10-12	n/a	340	170	154	178	176									
	15	n/a	340	98U	154	162	187									
	30	n/a	340	98.1U	93.1U	95.5U	101									
	45	n/a	340	88U	99.9U	87.6U	87.8U									
	60	n/a	340	94.8U	97.8U	88.7U	95.3U									
Tin	05-09	3150	n/a					4.7	4.6							
	10-12	11.3	4.6					14U	12U							
Uranium	05-09	11.3	4.6													
	10-12	11.3	4.6	0.905U	1.07	1.17	0.907U									
	15	11.3	4.6	1.72	0.852U	1.05	0.946U									
	30	11.3	4.6	0.981U	0.931U	0.99	0.942U									
	45	11.3	4.6	2.03	1.79	0.958	0.878U									
	60	11.3	4.6	0.948U	0.978U	1.1	1.05									
Vanadium	05-09	4.4	37					40	25							
	10-12	4.4	37	3.69	5.41	5.66	2.27U									
	15	4.4	37	4.35	3.82	10.4	4.08									
	30	4.4	37	13.8	6.38	19.3	3.33									
	45	4.4	37	5.78	9.32	7.3	3.21									
	60	4.4	37	6.82	17.6	8.87	3.79									
Zinc	05-09	2,660	60					52	26							
	10-12	2,660	60	18.1U	25.1	33.1	32.6									
	15	2,660	60	19.6U	17U	29.7	20.9									
	30	2,660	60	19.6U	18.6U	19.1U	18.8U									
	45	2,660	60	44.1	47	17.5U	17.6U									
	60	2,660	60	35.7	41.4	59.5	59.1									

Table 4.42. SWMU 30 Locations of Subsurface Soil Contaminants (Continued)

Analysis	Depth (ft)	Excavation Worker NAL	Background	RI Data				Historical Data								
				030-001	030-002	030-003	030-004	WB-1	WB-11	WB-3	WB-4	WB-5	WBP-1A	WBP-4A		
Organics--Semi-volatiles (mg/kg) 1,2,4-Trichlorobenzene	05-09	83	n/a					0.01U	0.033	0.01U	0.01U	0.01U	0.01U			
	10-12	83	n/a	0.49U	0.47U	0.49U	0.48U									
	15	83	n/a	0.47U	0.48U	0.47U	0.48U									
	30	83	n/a	0.46U	0.49U	0.47U	0.47U									
	45	83	n/a	0.48U	0.49U	0.47U	0.48U									
	60	83	n/a	0.46U	0.48U	0.47U	0.48U									
1,4-Dichlorobenzene	05-09	5.35	n/a					0.009U	0.025	0.009U	0.009U	0.009U	0.009U			
	10-12	5.35	n/a	0.49U	0.47U	0.49U	0.48U									
	15	5.35	n/a	0.47U	0.48U	0.47U	0.48U									
	30	5.35	n/a	0.46U	0.49U	0.47U	0.47U									
	45	5.35	n/a	0.48U	0.49U	0.47U	0.48U									
	60	5.35	n/a	0.46U	0.48U	0.47U	0.48U									
2-Chlorophenol	05-09	21.2	n/a					0.011U	0.023	0.011U	0.011U	0.011U	0.011U			
	10-12	21.2	n/a	0.49U	0.47U	0.49U	0.48U									
	15	21.2	n/a	0.47U	0.48U	0.47U	0.48U									
	30	21.2	n/a	0.46U	0.49U	0.47U	0.47U									
	45	21.2	n/a	0.48U	0.49U	0.47U	0.48U									
	60	21.2	n/a	0.46U	0.48U	0.47U	0.48U									
Acenaphthene	05-09	350	n/a					0.008U	0.017	0.008U	0.008U	0.008U	0.008U			
	10-12	350	n/a	0.49U	0.47U	0.49U	0.48U									
	15	350	n/a	0.47U	0.48U	0.47U	0.48U									
	30	350	n/a	0.46U	0.49U	0.47U	0.47U									
	45	350	n/a	0.48U	0.49U	0.47U	0.48U									
	60	350	n/a	0.46U	0.48U	0.47U	0.48U									
Benz(a)anthracene	05-09	0.232	n/a					0.008U	0.038	0.008U	0.008U	0.026	0.017			
	10-12	0.232	n/a	0.49U	0.47U	0.49U	0.48U									
	15	0.232	n/a	0.47U	0.48U	0.47U	0.48U									
	30	0.232	n/a	0.46U	0.49U	0.47U	0.47U									
	45	0.232	n/a	0.48U	0.49U	0.47U	0.48U									
	60	0.232	n/a	0.46U	0.48U	0.47U	0.48U									
Benzo(a)pyrene	05-09	0.0232	n/a					0.009U	0.052	0.009U	0.009U	0.009U	0.009U			
	10-12	0.0232	n/a	0.49U	0.47U	0.49U	0.48U									
	15	0.0232	n/a	0.47U	0.48U	0.47U	0.48U									
	30	0.0232	n/a	0.46U	0.49U	0.47U	0.47U									
	45	0.0232	n/a	0.48U	0.49U	0.47U	0.48U									
	60	0.0232	n/a	0.46U	0.48U	0.47U	0.48U									

Table 4.42. SWMU 30 Locations of Subsurface Soil Contaminants (Continued)

Analysis	Depth (ft)	Excavation Worker NAL	Background	RI Data				Historical Data							
				030-001	030-002	030-003	030-004	WB-1	WB-11	WB-3	WB-4	WB-5	WBP-1A	WBP-4A	
Benzo(b)fluoranthene	05-09	232	n/a					0.01U	0.067	0.01U	0.041	0.01U			
	10-12	232	n/a	0.49U	0.47U	0.49U	0.48U								
	15	232	n/a	0.47U	0.48U	0.47U	0.48U								
	30	232	n/a	0.46U	0.49U	0.47U	0.47U								
	45	232	n/a	0.48U	0.49U	0.47U	0.48U								
	60	232	n/a	0.46U	0.48U	0.47U	0.48U	0.01U	0.026	0.01U	0.01U	0.01U	0.01U		
Benzo(k)fluoranthene	05-09	232	n/a												
	10-12	232	n/a	0.49U	0.47U	0.49U	0.48U								
	15	232	n/a	0.47U	0.48U	0.47U	0.48U								
	30	232	n/a	0.46U	0.49U	0.47U	0.47U								
	45	232	n/a	0.48U	0.49U	0.47U	0.48U								
	60	232	n/a	0.46U	0.48U	0.47U	0.48U	0.025U	0.025U	0.025U	0.025U	0.025U	0.025U		
Benzoic acid	05-09	60,700	n/a												
	10-12	60,700	n/a	0.49U	0.47U	0.49U	0.48U								
	15	60,700	n/a	0.51	0.49	0.47U	0.48U								
	30	60,700	n/a	0.53	0.49U	0.47U	0.47U								
	45	60,700	n/a	0.5	0.5	0.47U	0.48U								
	60	60,700	n/a	0.52	0.48U	0.47U	0.48U	0.01U	0.035	0.01U	0.025	0.012			
Chrysene	05-09	232	n/a												
	10-12	232	n/a	0.49U	0.47U	0.49U	0.48U								
	15	232	n/a	0.47U	0.48U	0.47U	0.48U								
	30	232	n/a	0.46U	0.49U	0.47U	0.47U								
	45	232	n/a	0.48U	0.49U	0.47U	0.48U								
	60	232	n/a	0.46U	0.48U	0.47U	0.48U	0.02	0.054	0.012	0.039	0.028			
Fluoranthene	05-09	242	n/a												
	10-12	242	n/a	0.49U	0.47U	0.49U	0.48U								
	15	242	n/a	0.47U	0.48U	0.47U	0.48U								
	30	242	n/a	0.46U	0.49U	0.47U	0.47U								
	45	242	n/a	0.48U	0.49U	0.47U	0.48U								
	60	242	n/a	0.46U	0.48U	0.47U	0.48U	0.008U	0.012	0.008U	0.008U	0.015			
Phenanthrene	05-09	n/a	n/a												
	10-12	n/a	n/a	0.49U	0.47U	0.49U	0.48U								
	15	n/a	n/a	0.47U	0.48U	0.47U	0.48U								
	30	n/a	n/a	0.46U	0.49U	0.47U	0.47U								
	45	n/a	n/a	0.48U	0.49U	0.47U	0.48U								
	60	n/a	n/a	0.46U	0.48U	0.47U	0.48U	0.016U	0.041	0.018	0.036	0.059			
Pyrene	05-09	181	n/a												
	10-12	181	n/a	0.49U	0.47U	0.49U	0.48U								
	15	181	n/a	0.47U	0.48U	0.47U	0.48U								
	30	181	n/a	0.46U	0.49U	0.47U	0.47U								
	45	181	n/a	0.48U	0.49U	0.47U	0.48U								
	60	181	n/a	0.46U	0.48U	0.47U	0.48U								

Table 4.42. SWMU 30 Locations of Subsurface Soil Contaminants (Continued)

Analysis	Depth (ft)	Excavation Worker NAL	Background	RI Data				Historical Data								
				030-001	030-002	030-003	030-004	WB-1	WB-11	WB-3	WB-4	WB-5	WBP-1A	WBP-4A		
Organics--Volatiles (mg/kg) 1,1-Dichloroethene	05-09	0.119	n/a					0.00078U	0.00078U	0.00078U	0.00078U	0.00078U	0.00078U			
	10-12	0.119	n/a	0.00498U	0.00505U	0.00499U	0.00504U									
	15	0.119	n/a	0.00503U	0.00497U	0.00501U	0.00504U									
	30	0.119	n/a	0.00504U	0.005U	0.00503U	0.00498U									
	45	0.119	n/a	0.00499U	0.00502U	0.00498U	0.00496U									
	60	0.119	n/a	0.00499U	0.005U	0.00497U	0.00503U	0.0041U	0.018U	0.0041U	0.0041U	0.0041U	0.0065			
Acetone	05-09	421	n/a													
	10-12	421	n/a	0.00498U	0.00641	0.00973	0.00504U									
	15	421	n/a	0.00503U	0.00497U	0.00501U	0.00504U									
	30	421	n/a	0.00504U	0.005U	0.00503U	0.00498U									
	45	421	n/a	0.00499U	0.00502U	0.00498U	0.00496U									
	60	421	n/a	0.00499U	0.00499U	0.00497U	0.00503U	0.007U	0.006U	0.0018	0.0019U	0.0019U				
Methylene chloride	05-09	15.7	n/a													
	10-12	15.7	n/a	0.00498U	0.00505U	0.00499U	0.00504U									
	15	15.7	n/a	0.00503U	0.00497U	0.00501U	0.00504U									
	30	15.7	n/a	0.00504U	0.005U	0.00503U	0.00498U									
	45	15.7	n/a	0.00499U	0.00502U	0.00498U	0.00496U									
	60	15.7	n/a	0.00499U	0.00499U	0.00497U	0.00503U	0.0011	0.00075U	0.00075U	0.0011	0.00075U				
Toluene	05-09	272	n/a													
	10-12	272	n/a	0.00498U	0.00505U	0.00499U	0.00504U									
	15	272	n/a	0.00503U	0.00497U	0.00501U	0.00504U									
	30	272	n/a	0.00504U	0.005U	0.00503U	0.00498U									
	45	272	n/a	0.00499U	0.00502U	0.00498U	0.00496U									
	60	272	n/a	0.00499U	0.00499U	0.00497U	0.00503U	0.00056U	0.00056U	0.00056U	0.00056U	0.00056U				
Trichloroethene	05-09	3.25	n/a													
	10-12	3.25	n/a	0.00498U	0.00505U	0.00499U	0.00504U									
	15	3.25	n/a	0.00503U	0.00497U	0.00501U	0.00504U									
	30	3.25	n/a	0.0374	0.005U	0.00503U	0.00498U									
	45	3.25	n/a	0.00499U	0.00502U	0.00498U	0.00496U									
	60	3.25	n/a	0.00499U	0.00499U	0.00497U	0.00503U									
Organics--PCBs (mg/kg) PCB, Total	05-09	0.168	n/a					0.18	0.02	0.028	0.049	0.065				
	10-12	0.168	n/a	0.1U	0.09U	0.1U	0.09U									
	15	0.168	n/a	0.09U	0.1U	0.09U	0.09U									
	30	0.168	n/a	0.09U	0.1U	0.09U	0.09U									
	45	0.168	n/a	0.1U	0.1U	0.1U	0.09U									
	60	0.168	n/a	0.09U	0.09U	0.1U	0.1U									

Table 4.42. SWMU 30 Locations of Subsurface Soil Contaminants (Continued)

Analysis	Depth (ft)	Excavation Worker NAL	Background	RI Data				Historical Data								
				030-001	030-002	030-003	030-004	WB-1	WB-11	WB-3	WB-4	WB-5	WBP-1A	WBP-4A		
PCB-1254	05-09	0.168	n/a					0.00071U	0.00071U	0.028	0.00071U	0.00071U	0.00071U			
	10-12	0.168	n/a	0.1U	0.09U	0.1U	0.09U									
	15	0.168	n/a	0.09U	0.1U	0.09U	0.09U									
	30	0.168	n/a	0.09U	0.1U	0.09U	0.09U									
	45	0.168	n/a	0.1U	0.1U	0.1U	0.09U									
	60	0.168	n/a	0.09U	0.09U	0.1U	0.1U									
PCB-1260	05-09	0.168	n/a					0.18	0.02	0.00084U	0.049	0.065				
	10-12	0.168	n/a	0.1U	0.09U	0.1U	0.09U									
	15	0.168	n/a	0.09U	0.1U	0.09U	0.09U									
	30	0.168	n/a	0.09U	0.1U	0.09U	0.09U									
	45	0.168	n/a	0.1U	0.1U	0.1U	0.09U									
	60	0.168	n/a	0.09U	0.09U	0.1U	0.1U									
Radionuclides (pCi/g)																
Neptunium-237	05-09	0.328	n/a					0.04U	0.02U	0.05	0.03U	0.06				
	10-12	0.328	n/a	0.043U	0.0412U	0.0407U	0.0412U									
	15	0.328	n/a	0.0405U	0.0398U	0.0413U	0.0414U									
	30	0.328	n/a	0.0407U	0.0438U	0.0406U	0.0402U									
	45	0.328	n/a	0.0434U	0.0415U	0.0427U	0.0399U									
	60	0.328	n/a	0.0401U	0.0396U	0.0404U	0.0415U									
Plutonium-239 Technetium-99	05-09	1.63	n/a					0.08	0U	0.05	0.08	0.19				
	10-12	57.9	2.8					6.79	0.24	0.82	0.12	1.75				
	15	57.9	2.8	1.92U	1.92U	1.71U	1.71U									
	30	57.9	2.8	1.92U	1.92U	1.71U	1.71U									
	45	57.9	2.8	1.92U	1.92U	1.71U	1.71U									
	60	57.9	2.8	1.92U	1.92U	1.71U	1.71U									
Thorium-228	10-12	0.0357	1.6	0.453	0.393	0.334	0.465									
	15	0.0357	1.6	0.301	0.412	0.378	0.302									
	30	0.0357	1.6	0.223	0.181	0.32	0.226									
	45	0.0357	1.6	0.617	0.43	0.401	0.187									
	60	0.0357	1.6	0.339	0.546	0.463	0.432									
	05-09	2.22	1.4					1.43	1.27	1.21	1.44	1.41				
Thorium-230	10-12	2.22	1.4	0.38	0.445	0.311	0.365									
	15	2.22	1.4	0.263	0.119U	0.301	0.24U									
	30	2.22	1.4	0.198	0.138U	0.236U	0.236U									
	45	2.22	1.4	0.571	0.374	0.239U	0.235U									
	60	2.22	1.4	0.323	0.314	0.236U	0.236U									

Table 4.42. SWMU 30 Locations of Subsurface Soil Contaminants (Continued)

Analysis	Depth (ft)	Excavation Worker NAL	Background	RI Data				Historical Data									
				030-001	030-002	030-003	030-004	WB-1	WB-11	WB-3	WB-4	WB-5	WBP-1A	WBP-4A			
Thorium-232	10-12	1.95	1.5	0.401	0.445	0.299	0.4										
	15	1.95	1.5	0.374	0.321	0.3	0.299										
	30	1.95	1.5	0.308	0.203	0.263	0.189										
	45	1.95	1.5	0.626	0.432	0.41	0.191										
	60	1.95	1.5	0.461	0.509	0.436	0.39										
Thorium-234	10-12	n/a	n/a	2.22	2.88	0.912U	0.922U										
	15	n/a	n/a	2.56	0.813U	0.957U	0.867U										
	30	n/a	n/a	0.799U	1.58	0.893U	0.778U										
	45	n/a	n/a	1.08U	2.27	0.886U	1.36										
	60	n/a	n/a	1.26	0.961U	1.05U	0.999U										
Uranium	05-09	n/a	n/a													34	59
	10-12	n/a	n/a	0.294U	1.02	0.297U	0.35										
	15	n/a	n/a	1.3	0.293U	2.97	0.297U										
	30	n/a	n/a	0.292U	0.294U	0.299U	0.299U										
	45	n/a	n/a	0.815	0.294U	0.299U	0.298U										
Uranium-234	05-09	2.84	2.4	0.651	0.292U	0.3U	0.298U		0.2	2.26	2.24	6.56	5.87	8.1	20.6		
	10-12	2.84	2.4	0.131U	0.592	0.131U	0.15										
	15	2.84	2.4	0.5	0.131U	2.46	0.131U										
	30	2.84	2.4	0.13U	0.131U	0.133U	0.132U										
	45	2.84	2.4	0.429	0.203	0.206	0.163										
Uranium-235	05-09	2.84	2.4	0.517	0.131U	0.173	0.16										
	10-12	0.455	0.14	0.0386U	0.0373U	0.0379U	0.0371U										
	15	0.455	0.14	0.0383U	0.0376U	0.0852	0.0382U										
	30	0.455	0.14	0.0376U	0.0382U	0.038U	0.0385U										
	45	0.455	0.14	0.0415U	0.0382U	0.0381U	0.0383U										
Uranium-235/236	05-09	0.455	0.14	0.0379U	0.0372U	0.0388U	0.038U		0.02	0.14	0.16	0.55	0.4				
	10-12	1.17	1.2						0.38	2.74	2.92	10.3	8.2	25.3	37.4		
	15	1.17	1.2	0.125U	0.397	0.128U	0.181										
	30	1.17	1.2	0.77	0.125U	0.424	0.128U										
	45	1.17	1.2	0.124U	0.125U	0.129U	0.129U										
Uranium-238	05-09	1.17	1.2	0.357	0.149	0.163	0.333										
	10-12	1.17	1.2	0.124U	0.124U	0.129U	0.128U										

Blank cells indicate interval was not sampled for the specified analysis. Maximum value shown for each depth interval.

Bold indicates result is greater than NAL value.

Italics indicates result is greater than background value.

Bold + Italics indicate result is greater than both NAL and background values.

4.9.2 SWMU 30 Groundwater

UCRS groundwater samples were collected from one of the four angled borings installed at SWMU 30 as part of this RI (030-003, located south of Burial Pit A). The UCRS groundwater samples in the historic data set represent MW64, on the west side of the SWMU, (for the period of record 1995 to 2007); the SWMUs 7 and 30 RI, temporary borings WB-1, WB-4, and WB-5, both within and south of Burial Pit A (DOE 1998a); and a 1998 follow-up investigation of the main SWMU 30 waste pit, temporary boring WBP-4A. RI data were reviewed with historical data to identify the UCRS contaminants listed in Table 4.43. Groundwater sampling locations are shown in Figure 4.15.

Only a limited number of samples were available to characterize the UCRS groundwater at SWMU 30. Temporary borings WB-1, WB-4, WB-5, and WBP-4A sampled the depth interval of the buried waste at SWMU 30. (Pit A of SWMU 30 is approximately 12 ft deep.) Borings WB-1, WB-4, and WBP-4A are located within the burial pit. MW64 and 030-003 characterize the HU2 interval of the UCRS, found at approximately 20 to 30 ft bgs at SWMU 30. The suite of contaminants was similar at the depth of the waste pits and within the HU2 interval. Screening of the sample analyses revealed nine metal contaminants: arsenic, cadmium, iron, lead, manganese, molybdenum, nickel, uranium, and vanadium. All but cadmium were detected at levels exceeding screening criteria in 50% or more of the samples. TCE was detected at three locations (MW64, WB-1, and WB-4); one location exceeded the screening level (MW64 up to 0.45 mg/L). Benzene (at location WB-4) and vinyl chloride (at locations WB-1 and WB-4) also were detected above screening levels. The uranium isotopes uranium-234 and uranium-238 frequently exceeded screening levels in the SWMU 30 UCRS groundwater samples.

RGA and McNairy groundwater samples were not collected at SWMU 30 as part of this RI. Historical data were reviewed for RGA groundwater to determine the contaminants listed in Table 4.44. All of the SWMU 30 RGA groundwater samples were from MWs shown in Figure 4.15. MW63, MW66, and MW245 sample the upper RGA. MW65 is a lower RGA well.

The RGA groundwater samples from SWMU 30 contained five metal contaminants: arsenic, iron, lead, manganese, and uranium. Of the organic analytes, only TCE was detected frequently above screening levels, in all four RGA groundwater MWs (Figure 4.60). Tetrachloroethene was detected at only one location, MW66, at 0.32 mg/L, which is above the screening level. Radon-222 and technetium-99 were the most frequently detected radionuclide contaminants. All technetium-99 analyses above the MCL represented samples from MW66.

No McNairy groundwater data were available.

Table 4.45 lists all SWMU 30 locations with groundwater contamination.

Table 4.43. SWMU 30 UCRS Groundwater Contaminants

Analysis	Historical Data	RI	Frequency of Detection ^a	Frequency of Detection		
				Above Background	Above NAL	Above MCL
Metals (mg/L)						
Aluminum	53	7.46	4/4	N/A	3/4	N/A
Antimony	0.0053	ND	2/4	N/A	2/4	0/4
Arsenic	0.067	ND	2/4	N/A	2/4	2/4
Barium	0.52	0.262	4/4	N/A	4/4	0/4
Barium, Dissolved	N/A	0.195	1/1	N/A	1/1	0/1
Cadmium	0.011	ND	1/4	N/A	1/4	1/4
Calcium	57	66.6	3/4	N/A	N/A	N/A
Calcium, Dissolved	N/A	63.7	1/1	N/A	N/A	N/A
Chromium	0.087	ND	3/4	N/A	0/4	0/4
Cobalt	ND	0.0248	1/4	N/A	0/4	N/A
Cobalt, Dissolved	N/A	0.0181	1/1	N/A	0/1	N/A
Copper	0.39	ND	3/4	N/A	2/4	0/4
Iron	51	38.3	4/4	N/A	4/4	N/A
Iron, Dissolved	N/A	18.2	1/1	N/A	1/1	N/A
Lead	0.23	0.00357	3/4	N/A	2/4	2/4
Magnesium	37	22.6	4/4	N/A	0/4	N/A
Magnesium, Dissolved	N/A	21.7	1/1	N/A	0/1	N/A
Manganese	0.97	2.87	4/4	N/A	3/4	N/A
Manganese, Dissolved	N/A	2.76	1/1	N/A	1/1	N/A
Mercury	0.0007	0.00001	3/3	N/A	2/3	0/3
Molybdenum	0.14	0.111	2/4	N/A	2/4	N/A
Molybdenum, Dissolved	N/A	0.108	1/1	N/A	1/1	N/A
Nickel	0.14	ND	3/4	N/A	3/4	N/A
Nickel, Dissolved	N/A	0.0362	1/1	N/A	1/1	N/A
Potassium	56	N/A	3/3	N/A	N/A	N/A
Selenium	0.0087	ND	1/4	N/A	1/4	0/4
Sodium	55	43.3	2/4	N/A	0/4	N/A
Sodium, Dissolved	N/A	45.4	1/1	N/A	0/1	N/A
Tin	0.026	N/A	2/3	N/A	0/3	N/A
Uranium	0.17	0.15	3/6	N/A	3/6	3/6
Uranium, Dissolved	N/A	0.122	1/1	N/A	1/1	1/1
Vanadium	0.095	ND	2/4	N/A	2/4	N/A
Zinc	0.28	ND	3/4	N/A	0/4	N/A
Zinc, Dissolved	N/A	0.0418	1/1	N/A	0/1	N/A
Radionuclides (pCi/L)						
Bismuth-214	N/A	323	1/1	N/A	N/A	N/A
Lead-214	N/A	316	1/1	N/A	N/A	N/A
Neptunium-237	5.57	ND	2/4	N/A	2/4	N/A
Plutonium-239	4.21	N/A	3/3	N/A	2/3	N/A
Technetium-99	402	3.63	3/4	N/A	2/4	0/4
Thorium-230	11.7	ND	3/4	N/A	3/4	N/A
Uranium	N/A	28.2	1/1	N/A	N/A	N/A
Uranium-234	2,220	9.84	5/5	N/A	5/5	N/A
Uranium-235	N/A	0.491	1/1	N/A	0/1	N/A
Uranium-235/236	169	N/A	2/3	N/A	N/A	N/A
Uranium-238	2710	17.8	5/5	N/A	4/5	N/A
Uranium-234, Dissolved	N/A	18.5	1/1	N/A	1/1	N/A

Table 4.43. SWMU 30 UCRS Groundwater Contaminants (Continued)

Analysis	Historical Data	RI	Frequency of Detection ^a	Frequency of Detection		
				Above Background	Above NAL	Above MCL
Uranium-235, Dissolved	N/A	1.25	1/1	N/A	1/1	N/A
Uranium-238, Dissolved	N/A	33.9	1/1	N/A	1/1	N/A
PCBs (mg/L)						
PCB-1260	0.0029	ND	1/3	N/A	1/3	1/3
Semivolatiles (mg/L)						
1,2,4-Trichlorobenzene	0.0016	ND	1/4	N/A	0/4	0/4
1,2-Dichlorobenzene	0.012	ND	4/8	N/A	0/8	0/8
1,3-Dichlorobenzene	0.021	ND	4/8	N/A	4/8	N/A
1,4-Dichlorobenzene	0.067	ND	4/8	N/A	4/8	0/8
2,4-Dichlorophenol	0.00074	ND	1/4	N/A	0/4	N/A
2,4-Dimethylphenol	0.064	ND	2/4	N/A	1/4	N/A
2-Methylnaphthalene	0.00043	ND	2/4	N/A	0/4	N/A
2-Methylphenol	0.0012	N/A	1/3	N/A	0/3	N/A
4-Methylphenol	0.00082	N/A	1/3	N/A	0/3	N/A
Acenaphthene	0.00035	ND	1/4	N/A	0/4	N/A
Diethyl phthalate	0.00045	ND	1/4	N/A	0/4	N/A
Di-n-octylphthalate	0.0018	ND	1/4	N/A	1/4	N/A
Fluoranthene	0.00027	ND	1/4	N/A	0/4	N/A
Naphthalene	0.00072	ND	1/4	N/A	1/4	N/A
Phenanthrene	0.0033	ND	1/4	N/A	0/4	N/A
Pyrene	0.00077	ND	2/4	N/A	0/4	N/A
Volatiles (mg/L)						
1,1-Dichloroethane	0.072	ND	2/5	N/A	1/5	N/A
1,2-Dichloroethane	0.003	ND	2/5	N/A	2/5	0/5
Acetone	0.012	0.027	4/5	N/A	0/5	N/A
Benzene	0.0054	ND	2/5	N/A	2/5	1/5
Chlorobenzene	0.093	ND	2/5	N/A	2/5	0/5
Chloroethane	0.12	ND	2/5	N/A	2/5	N/A
Chloromethane	0.0094	ND	1/5	N/A	1/5	N/A
<i>cis</i> -1,2-Dichloroethene	0.067	ND	3/5	N/A	2/5	0/5
Ethylbenzene	0.011	ND	1/5	N/A	1/5	0/5
Toluene	0.0082	ND	2/5	N/A	0/5	0/5
Total Xylene	0.03	N/A	1/4	N/A	0/4	0/4
<i>trans</i> -1,2-Dichloroethene	0.0017	ND	1/5	N/A	0/5	0/5
Trichloroethene	0.45	ND	4/6	N/A	4/6	2/6
Vinyl chloride	0.0086	ND	2/5	N/A	2/5	2/5

^a Frequency of detection is the number of detections of an analyte per number of analyses (includes regular and duplicate samples).

N/A = not applicable

ND = not detected

Table 4.44. SWMU 30 RGA Groundwater Contaminants

Analysis	Historical Data	RI	Frequency of Detection ^a	Frequency of Detection		
				Above Background	Above NAL	Above MCL
Metals (mg/L)						
Aluminum	143	N/A	58/71	38/71	45/71	N/A
Aluminum, Dissolved	3.94	N/A	18/66	17/66	3/66	N/A
Arsenic	0.0123	N/A	1/4	1/4	1/4	1/4
Arsenic, Dissolved	0.00815	N/A	1/1	1/1	1/1	0/1
Barium	1.47	N/A	19/23	2/23	13/23	0/23
Barium, Dissolved	1.06	N/A	16/20	2/20	12/20	0/20
Beryllium	0.0004	N/A	3/23	0/23	0/23	0/23
Cadmium	0.00224	N/A	1/20	0/20	1/20	0/20
Calcium	85	N/A	80/80	2/80	N/A	N/A
Calcium, Dissolved	78.2	N/A	76/76	2/76	N/A	N/A
Chromium	0.5	N/A	7/23	3/23	0/23	5/23
Cobalt	0.334	N/A	3/23	2/23	1/23	N/A
Cobalt, Dissolved	0.311	N/A	2/11	2/11	1/11	N/A
Copper	0.031	N/A	2/24	0/24	0/24	0/24
Iron	226	N/A	66/79	28/79	61/79	N/A
Iron (2+)	0.0353	N/A	1/1	N/A	N/A	N/A
Iron, Dissolved	54.6	N/A	25/76	24/76	16/76	N/A
Lead	0.432	N/A	3/7	1/7	1/7	1/7
Magnesium	29.6	N/A	80/80	3/80	0/80	N/A
Magnesium, Dissolved	25.8	N/A	76/76	1/76	0/76	N/A
Manganese	39.9	N/A	64/79	56/79	58/79	N/A
Manganese, Dissolved	38.2	N/A	57/76	56/76	57/76	N/A
Molybdenum	0.00281	N/A	1/15	0/15	0/15	N/A
Molybdenum, Dissolved	0.00119	N/A	1/4	0/4	0/4	N/A
Nickel	0.47	N/A	10/23	0/23	8/23	N/A
Nickel, Dissolved	0.203	N/A	6/10	0/10	6/10	N/A
Potassium	7.72	N/A	11/76	1/76	N/A	N/A
Potassium, Dissolved	3.93	N/A	12/24	0/24	N/A	N/A
Sodium	162	N/A	80/80	8/80	0/80	N/A
Sodium, Dissolved	150	N/A	76/76	7/76	0/76	N/A
Uranium	0.19	N/A	4/128	2/128	4/128	2/128
Vanadium	0.064	N/A	3/5	0/5	1/5	N/A
Vanadium, Dissolved	0.071	N/A	2/2	0/2	2/2	N/A
Zinc	0.041	N/A	3/23	0/23	0/23	N/A
Radionuclides (pCi/L)						
Neptunium-237	0.2	N/A	2/24	0/24	0/24	N/A
Potassium-40	275	N/A	2/21	N/A	N/A	N/A
Radium-226	0.52	N/A	2/8	0/8	2/8	0/8
Radon	319.99	N/A	8/9	N/A	N/A	N/A
Radon-222	632	N/A	43/44	1/44	43/44	N/A
Technetium-99	2,911	N/A	210/279	175/279	194/279	39/279
Thorium-230	0.79	N/A	4/33	0/33	4/33	N/A
Uranium	911	N/A	1/8	N/A	N/A	N/A
Uranium-234	448	N/A	4/11	1/11	2/11	N/A
Uranium-238	441	N/A	1/14	1/14	1/14	N/A

Table 4.44. SWMU 30 RGA Groundwater Contaminants (Continued)

Analysis	Historical Data	RI	Frequency of Detection ^a	Frequency of Detection		
				Above Background	Above NAL	Above MCL
<i>Volatiles (mg/L)</i>						
1,1,2-Trichloro-1,2,2-	0.00088	N/A	1/3	N/A	0/3	N/A
4-Methyl-2-pentanone	0.0025	N/A	1/4	N/A	0/4	N/A
Acetone	0.0081	N/A	3/4	N/A	0/4	N/A
Chloroform	0.001	N/A	1/41	N/A	1/41	N/A
<i>cis</i> -1,2-Dichloroethene	0.0226	N/A	5/237	N/A	4/237	0/237
Tetrachloroethene	0.32	N/A	1/193	N/A	1/193	1/193
Toluene	0.011	N/A	2/278	N/A	0/278	0/278
Trichloroethene	15	N/A	252/277	N/A	250/277	232/277

^a Frequency of detection is the number of detections of an analyte per number of analyses (includes regular and duplicate samples).

N/A = not applicable

Table 4.45. SWMU 30 Locations of Groundwater Contaminants

Hydro Unit	Depth Interval	Analysis	Child Resident NAL	Bkgd	MCL	030-003	MW245	MW63	MW64	MW65	MW66	WB-1	WB-4	WB-5	WBP-4A
	8-10	Metals (mg/L)													
		Aluminum	1.49	N/A	N/A							9.8	53		
		Antimony	0.000564	N/A	0.006							0.0053	0.0047		
		Arsenic	0.000035	N/A	0.01							0.067	0.03		
		Barium	0.104	N/A	2							0.26	0.52		
		Cadmium	0.000661	N/A	0.005							0.011	0.004U		
		Calcium	N/A	N/A	N/A							24U	57		
		Chromium	1.76	N/A	0.1							0.023	0.087		
		Cobalt	0.0906	N/A	N/A							0.006U	0.016U		
		Copper	0.0557	N/A	1.3							0.39	0.25		
		Iron	0.449	N/A	N/A							10	51		
		Lead	0.015	N/A	0.015							0.23	0.084		
		Magnesium	N/A	N/A	N/A							32	37		
		Manganese	0.035	N/A	N/A							0.25	0.97		
		Mercury	0.000444	N/A	0.002							0.0005	0.0007		
		Molybdenum	0.00753	N/A	N/A							0.088U	0.14		
		Nickel	0.0301	N/A	N/A							0.076	0.14		
		Potassium	N/A	N/A	N/A							56	18		
		Selenium	0.00754	N/A	0.05							0.005U	0.005U		
		Sodium	N/A	N/A	N/A							55U	28U		
		Tin	0.894	N/A	N/A							0.05U	0.024		
		Uranium	0.000906	N/A	0.03							0.36U	0.37U		0.17
		Vanadium	0.00925	N/A	N/A							0.02	0.095		
		Zinc	0.45	N/A	N/A							0.21	0.28		
		PCBs (mg/L)													
		PCB-1260	0.0000428	N/A	0.0005								0.0029		
		Radionuclides (pCi/L)													
		Neptunium-237	0.573	N/A	N/A							5.57	3.5		
		Plutonium-239	0.286	N/A	N/A							4.21	0.18		
		Technetium-99	14	N/A	900							402	88.7		
		Thorium-230	0.424	N/A	N/A							11.7	2.07		
		Uranium-234	0.546	N/A	N/A							2.220	106		20.3
		Uranium-235/236	N/A	N/A	N/A							169	9.77		
		Uranium-238	0.443	N/A	N/A							2.710	247		53.6
		Semivolatiles (mg/L)													
		1,2,4-Trichlorobenzene	0.00781	N/A	0.07							0.0016	0.0012U		
		1,2-Dichlorobenzene	0.0166	N/A	0.6							0.0028	0.012	0.0002U	
		1,3-Dichlorobenzene	0.000241	N/A	N/A							0.0037	0.021	0.00024U	
		1,4-Dichlorobenzene	0.000578	N/A	0.075							0.016	0.067	0.00022U	
		2,4-Dichlorophenol	0.0041	N/A	N/A							0.00074	0.00018U		
		2,4-Dimethylphenol	0.023	N/A	N/A							0.064	0.0008		
		2-Methylnaphthalene	N/A	N/A	N/A							0.00043	0.00034		
		2-Methylphenol	0.0723	N/A	N/A							0.0012	0.00022U		

Table 4.45. SWMU 30 Locations of Groundwater Contaminants (Continued)

Hydro Unit	Depth Interval	Analysis	Child Resident NAL	Bkgd	MCL	030-003	MW245	MW63	MW64	MW65	MW66	WB-1	WB-4	WB-5	WBP-4A		
UCRS	8-10	4-Methylphenol	0.00727	N/A	N/A							0.00082	0.0002U				
		Acenaphthene	0.0136	N/A	N/A							0.00021U	0.00035				
		Diethyl phthalate	1.2	N/A	N/A							0.00023U	0.00045				
		Di-n-octylphthalate	0.000684	N/A	N/A							0.0018	0.00017U				
		Fluoranthene	0.0226	N/A	N/A							0.00023U	0.00027				
		Naphthalene	0.000285	N/A	N/A							0.00072	0.00032U				
		Phenanthrene	N/A	N/A	N/A							0.0033	0.00023U				
		Pyrene	0.0182	N/A	N/A							0.00069	0.00077				
		Volatiles (mg/L)															
				1,1-Dichloroethane	0.0363	N/A	N/A							0.072	0.019	0.00037U	
				1,2-Dichloroethane	0.000147	N/A	0.005							0.003	0.0015	0.00027U	
				Acetone	0.0275	N/A	N/A							0.0067	0.01	0.012	
				Benzene	0.000385	N/A	0.005							0.0025	0.0054	0.00038U	
				Chlorobenzene	0.00466	N/A	0.1							0.016	0.093	0.00049U	
				Chloroethane	0.00461	N/A	N/A							0.12	0.033	0.0014U	
				Chloromethane	0.00167	N/A	N/A							0.00037U	0.0094	0.00037U	
				cis-1,2-Dichloroethene	0.00273	N/A	0.07							0.067	0.015	0.00047U	
				Ethylbenzene	0.00468	N/A	0.7							0.011	0.00044U	0.00044U	
				Toluene	0.0338	N/A	1							0.0082	0.0012	0.00044U	
		Total Xylene	0.0653	N/A	10							0.03	0.0013U	0.0013U			
		trans-1,2-Dichloroethene	0.00548	N/A	0.1							0.00048U	0.0017	0.00048U			
		Trichloroethene	0.0016	N/A	0.005							0.0022	0.0037	0.00039U			
		Vinyl chloride	0.000035	N/A	0.002							0.0086	0.0048	0.0013U			
	23	Metals (mg/L)															
		Aluminum	1.49	N/A	N/A	7.46											
		Antimony	0.000564	N/A	0.006	0.005U											
		Arsenic	0.000035	N/A	0.01	0.01U											
		Barium	0.104	N/A	2	0.262											
		Barium, Dissolved	0.104	N/A	2	0.195											
		Cadmium	0.000661	N/A	0.005	0.0006U											
		Calcium	N/A	N/A	N/A	66.6											
		Calcium, Dissolved	N/A	N/A	N/A	63.7											
		Chromium	1.76	N/A	0.1	0.1U											
		Cobalt	0.0906	N/A	N/A	0.0248											
		Cobalt, Dissolved	0.0906	N/A	N/A	0.0181											
		Copper	0.0557	N/A	1.3	0.2U											
		Iron	0.449	N/A	N/A	38.3											
		Iron, Dissolved	0.449	N/A	N/A	18.2											
		Lead	0.015	N/A	0.015	0.00357											
		Magnesium	N/A	N/A	N/A	22.6											
		Magnesium, Dissolved	N/A	N/A	N/A	21.7											
		Manganese	0.035	N/A	N/A	2.87											
		Manganese, Dissolved	0.035	N/A	N/A	2.76											

Table 4.45. SWMU 30 Locations of Groundwater Contaminants (Continued)

Hydro Unit	Depth Interval	Analysis	Child Resident NAL	Bkgd	MCL	030-003	MW245	MW63	MW64	MW65	MW66	WB-1	WB-4	WB-5	WBP-4A		
UCRS	23	Mercury	0.000444	N/A	0.002	0.00001											
		Molybdenum	0.00753	N/A	N/A	0.111											
		Molybdenum, Dissolved	0.00753	N/A	N/A	0.108											
		Nickel	0.0301	N/A	N/A	N/A	0.05U										
		Nickel, Dissolved	0.0301	N/A	N/A	N/A	0.0362										
		Selenium	0.00754	N/A	N/A	0.05	0.005U										
		Sodium	N/A	N/A	N/A	N/A	43.3										
		Sodium, Dissolved	N/A	N/A	N/A	N/A	45.4										
		Uranium	0.000906	N/A	N/A	0.03	0.15										
		Uranium, Dissolved	0.000906	N/A	N/A	0.03	0.122										
		Vanadium	0.00925	N/A	N/A	N/A	0.2U										
		Zinc	0.45	N/A	N/A	N/A	0.2U										
		Zinc, Dissolved	0.45	N/A	N/A	N/A	0.0418										
		PCBs (mg/L)															
				PCB-1260	0.0000428	N/A	0.0005	0.0001U									
		Radionuclides (pCi/L)															
				Bismuth-214	N/A	N/A	N/A	323									
				Lead-214	N/A	N/A	N/A	316									
				Neptunium-237	0.573	N/A	N/A	0.0639U									
				Technetium-99	14	N/A	N/A	900	3.63								
				Thorium-230	0.424	N/A	N/A	0.13U									
				Uranium	N/A	N/A	N/A	28.2									
				Uranium-234	0.546	N/A	N/A	9.84									
		Uranium-235	N/A	N/A	N/A	0.491											
		Uranium-238	0.443	N/A	N/A	17.8											
		Uranium-234, Dissolved	0.546	N/A	N/A	18.5											
		Uranium-235, Dissolved	N/A	N/A	N/A	1.25											
		Uranium-238, Dissolved	0.443	N/A	N/A	33.9											
Semivolatiles (mg/L)																	
		1,2,4-Trichlorobenzene	0.00781	N/A	0.07	0.0049U											
		1,2-Dichlorobenzene	0.0166	N/A	0.6	0.0049U											
		1,3-Dichlorobenzene	0.000241	N/A	N/A	0.0049U											
		1,4-Dichlorobenzene	0.000578	N/A	0.075	0.0049U											
		2,4-Dichlorophenol	0.0041	N/A	N/A	0.0049U											
		2,4-Dimethylphenol	0.023	N/A	N/A	0.0049U											
		2-Methylnaphthalene	N/A	N/A	N/A	0.0049U											
		Acenaphthene	0.0136	N/A	N/A	0.0049U											
		Diethyl phthalate	1.2	N/A	N/A	0.0049U											
		Di-n-octylphthalate	0.000684	N/A	N/A	0.0049U											
		Fluoranthene	0.0226	N/A	N/A	0.0049U											
		Naphthalene	0.000285	N/A	N/A	0.0049U											
		Phenanthrene	N/A	N/A	N/A	0.0049U											
		Pyrene	0.0182	N/A	N/A	0.0049U											

Table 4.45. SWMU 30 Locations of Groundwater Contaminants (Continued)

Hydro Unit	Depth Interval	Analysis	Child Resident NAL	Bkgd	MCL	030-003	MW245	MW63	MW64	MW65	MW66	WB-1	WB-4	WB-5	WBP-4A		
UCRS	23	Volatiles (mg/L)															
		1,1-Dichloroethane	0.0363	N/A	N/A	0.005U											
		1,2-Dichloroethane	0.000147	N/A	0.005	0.005U											
		Acetone	0.0275	N/A	N/A	0.027											
		Benzene	0.000385	N/A	0.005	0.005U											
		Chlorobenzene	0.00466	N/A	0.1	0.005U											
		Chloroethane	0.00461	N/A	N/A	0.005U											
		Chloromethane	0.00167	N/A	N/A	0.005U											
		cis-1,2-Dichloroethene	0.00273	N/A	0.07	0.005U											
		Ethylbenzene	0.00468	N/A	0.7	0.005U											
		Toluene	0.0338	N/A	1	0.005U											
		trans-1,2-Dichloroethene	0.00548	N/A	0.1	0.005U											
		Trichloroethene	0.0016	N/A	0.005	0.001U											
		Vinyl chloride	0.000035	N/A	0.002	0.002U											
		Metals (mg/L)															
			33	Aluminum	1.49	N/A	N/A				0.26						
				Antimony	0.000564	N/A	0.006				0.0024U						
				Arsenic	0.000035	N/A	0.01				0.0026U						
				Barium	0.104	N/A	2				0.27						
				Cadmium	0.000661	N/A	0.005				0.0003U						
				Calcium	N/A	N/A	N/A				40						
				Chromium	1.76	N/A	0.1				0.068						
				Cobalt	0.0906	N/A	N/A				0.005U						
		Copper	0.0557	N/A	1.3				0.019								
		Iron	0.449	N/A	N/A				0.74								
		Lead	0.015	N/A	0.015				0.0017U								
		Magnesium	N/A	N/A	N/A				13								
		Manganese	0.035	N/A	N/A				0.017								
		Molybdenum	0.00753	N/A	N/A				0.0044U								
		Nickel	0.0301	N/A	N/A				0.083								
		Potassium	N/A	N/A	N/A				1.6								
		Selenium	0.00754	N/A	0.05				0.0087								
		Sodium	N/A	N/A	N/A				55								
		Tin	0.894	N/A	N/A				0.026								
		Uranium	0.000906	N/A	0.03				0.11U								
		Vanadium	0.00925	N/A	N/A				0.0023U								
		Zinc	0.45	N/A	N/A				0.014								
PCBs (mg/L)																	
		PCB-1260	0.0000428	N/A	0.0005				0.000025U								

Table 4.45. SWMU 30 Locations of Groundwater Contaminants (Continued)

Hydro Unit	Depth Interval	Analysis	Child Resident NAL	Bkgd	MCL	030-003	MW245	MW63	MW64	MW65	MW66	WB-1	WB-4	WB-5	WBP-4A		
UCRS	33	Radionuclides (pCi/L)															
		Neptunium-237	0.573	N/A	N/A					-0.397U							
		Plutonium-239	0.286	N/A	N/A					0.76							
		Technetium-99	14	N/A	900					-0.59U							
		Thorium-230	0.424	N/A	N/A					0.44							
		Uranium-234	0.546	N/A	N/A					0.58							
		Uranium-235/236	N/A	N/A	N/A					0.02U							
		Uranium-238	0.443	N/A	N/A					0.44							
		Semivolatiles (mg/L)															
		1,2,4-Trichlorobenzene	0.00781	N/A	N/A	0.07					0.0011U						
		1,2-Dichlorobenzene	0.0166	N/A	N/A	0.6					0.00032U						
		1,3-Dichlorobenzene	0.000241	N/A	N/A	N/A					0.00036U						
		1,4-Dichlorobenzene	0.000578	N/A	N/A	0.075					0.00032U						
		2,4-Dichlorophenol	0.0041	N/A	N/A	N/A					0.00016U						
		2,4-Dimethylphenol	0.023	N/A	N/A	N/A					0.00011U						
		2-Methylnaphthalene	N/A	N/A	N/A	N/A					0.00029U						
		2-Methylphenol	0.0723	N/A	N/A	N/A					0.0002U						
		4-Methylphenol	0.00727	N/A	N/A	N/A					0.00018U						
		Acenaphthene	0.0136	N/A	N/A	N/A					0.00021U						
		Diethyl phthalate	1.2	N/A	N/A	N/A					0.00023U						
		Di-n-octylphthalate	0.000684	N/A	N/A	N/A					0.00015U						
		Fluoranthene	0.0226	N/A	N/A	N/A					0.00023U						
		Naphthalene	0.000285	N/A	N/A	N/A					0.00029U						
		Phenanthrene	N/A	N/A	N/A	N/A					0.00021U						
		Pyrene	0.0182	N/A	N/A	N/A					0.00025U						
		Volatiles (mg/L)															
		1,1-Dichloroethane	0.0363	N/A	N/A	N/A					0.00037U						
		1,2-Dichloroethane	0.000147	N/A	N/A	0.005					0.00027U						
		Acetone	0.0275	N/A	N/A	N/A					0.0014U						
		Benzene	0.000385	N/A	N/A	0.005					0.00038U						
		Chlorobenzene	0.00466	N/A	N/A	0.1					0.00049U						
		Chloroethane	0.00461	N/A	N/A	N/A					0.0014U						
		Chloromethane	0.00167	N/A	N/A	N/A					0.00037U						
cis-1,2-Dichloroethene	0.00273	N/A	N/A	0.07					0.00067								
Ethylbenzene	0.00468	N/A	N/A	0.7					0.00044U								
Toluene	0.0338	N/A	N/A	1					0.00044U								
Total Xylene	0.0653	N/A	N/A	10					0.0013U								
trans-1,2-Dichloroethene	0.00548	N/A	N/A	0.1					0.00048U								
Trichloroethene	0.0016	N/A	N/A	0.005					0.45								
Vinyl chloride	0.000035	N/A	N/A	0.002					0.0013U								

Table 4.45. SWMU 30 Locations of Groundwater Contaminants (Continued)

Hydro Unit	Depth Interval	Analysis	Child Resident NAL	Bkgd	MCL	030-003	MW245	MW63	MW64	MW65	MW66	WB-1	WB-4	WB-5	WBP-4A
	60-64	Metals (mg/L)													
		Aluminum	1.49	2.189	N/A			0.32			2.56				
		Aluminum, Dissolved	1.49	0.311	N/A			0.0026U			0.63U				
		Arsenic	0.000035	0.005	0.01			0.0026U			0.0026U				
		Barium	0.104	0.235	2			0.095			0.156				
		Barium, Dissolved	0.104	0.2	2			0.09			0.171				
		Beryllium	0.00264	0.004	0.004			0.0003			0.0004				
		Cadmium	0.000661	0.01	0.005			0.025U			0.025U				
		Calcium	N/A	41.238	N/A			23.2			30.7				
		Calcium, Dissolved	N/A	38.166	N/A			22			31.5				
		Chromium	1.76	0.144	0.1			0.5			0.0045				
		Cobalt	0.0906	0.045	N/A			0.0061			0.045U				
		Cobalt, Dissolved	0.0906	0.045	N/A			0.045U			0.45U				
		Copper	0.0557	0.036	1.3			0.013			0.025U				
		Iron	0.449	5.03	N/A			3			3.13				
		Iron (2+)	N/A	N/A	N/A						0.0353				
		Iron, Dissolved	0.449	0.267	N/A			0.36U			0.36U				
		Lead	0.015	0.129	0.015			0.0065			0.0017U				
		Magnesium	N/A	16.262	N/A			6.99			10.2				
		Magnesium, Dissolved	N/A	16.215	N/A			6.4			11.9				
		Manganese	0.035	0.119	N/A			0.041			0.037				
		Manganese, Dissolved	0.035	0.068	N/A			0.02U			0.04				
		Molybdenum	0.00753	0.05	N/A			0.025U			0.055U				
		Molybdenum, Dissolved	0.00753	0.05	N/A						0.055U				
		Nickel	0.0301	0.682	N/A			0.47			0.1				
		Nickel, Dissolved	0.0301	0.305	N/A			0.203			0.101				
		Potassium	N/A	5.195	N/A			10.5U			2U				
		Potassium, Dissolved	N/A	4.096	N/A			10.5U			10.5U				
		Sodium	N/A	59.45	N/A			18.9			21				
		Sodium, Dissolved	N/A	60.433	N/A			17.2			26.9				
		Uranium	0.000906	0.002	0.03			0.19			0.19				
		Vanadium	0.00925	0.134	N/A			0.005			0.064				
		Vanadium, Dissolved	0.00925	0.134	N/A						0.071				
		Zinc	0.45	0.054	N/A			0.03U			0.03U				
		Radionuclides (pCi/L)													
		Neptunium-237	0.573	0.8	N/A			0.07			1.01U				
		Potassium-40	N/A	N/A	N/A			275			469U				
		Radon-222	0.866	626	N/A			356			632				
		Technetium-99	14	22.3	900			22.5			2911				
		Thorium-230	0.424	1.1	N/A			0.79			0.5				
		Uranium	N/A	N/A	N/A			911			1.11U				
		Uranium-234	0.546	0.7	N/A			448			0.57				
		Uranium-238	0.443	0.7	N/A			441			8.01U				

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Table 4.45. SWMU 30 Locations of Groundwater Contaminants (Continued)

Hydro Unit	Depth Interval	Analysis	Child Resident NAL	Bkgd	MCL	030-003	MW245	MW63	MW64	MW65	MW66	WB-1	WB-4	WB-5	WBP-4A
	60-64	Semivolatiles (mg/L)													
		1,1,2-Trichloro-1,2,2-trifluoroethane	2.7	N/A	N/A		143	0.00036U			0.00088				
		4-Methyl-2-pentanone	0.00722	N/A	N/A		3.94	0.0011U			0.2U				
		Acetone	0.0275	N/A	N/A		0.0123	0.0054			0.0057				
		Chloroform	0.000287	N/A	N/A		0.00815	0.005U			0.001				
		cis-1,2-Dichloroethene	0.00273	N/A	0.07		1.47	0.005U			0.0226				
		Tetrachloroethene	0.000582	N/A	0.005		1.06	0.005U			0.32				
		Toluene	0.0338	N/A	1		0.025U	0.005U			0.0021				
		Trichloroethene	0.0016	N/A	0.005		0.00224	0.028			15				
	75	Metals (mg/L)													
		Aluminum	1.49	2.189	N/A		85								
		Aluminum, Dissolved	1.49	0.311	N/A		78.2								
		Arsenic	0.000035	0.005	0.01		0.2U								
		Arsenic, Dissolved	0.000035	0.005	0.01		0.334								
		Barium	0.104	0.235	2		0.311								
		Barium, Dissolved	0.104	0.2	2		0.031								
		Beryllium	0.00264	0.004	0.004		226								
		Cadmium	0.000661	0.01	0.005		54.6								
		Calcium	N/A	41.238	N/A		0.432								
		Calcium, Dissolved	N/A	38.166	N/A		29.6								
		Chromium	1.76	0.144	0.1		25.8								
		Cobalt	0.0906	0.045	N/A		39.9								
		Cobalt, Dissolved	0.0906	0.045	N/A		38.2								
		Copper	0.0557	0.036	1.3		0.00281								
		Iron	0.449	5.03	N/A		0.00119								
		Iron, Dissolved	0.449	0.267	N/A		0.137								
		Lead	0.015	0.129	0.015		0.0661								
		Magnesium	N/A	16.262	N/A		7.72								
		Magnesium, Dissolved	N/A	16.215	N/A		3.93								
		Manganese	0.035	0.119	N/A		162								
		Manganese, Dissolved	0.035	0.068	N/A		150								
		Molybdenum	0.00753	0.05	N/A		0.001U								
		Molybdenum, Dissolved	0.00753	0.05	N/A		0.035								
		Nickel	0.0301	0.682	N/A		0.00906								
		Nickel, Dissolved	0.0301	0.305	N/A		0.002								
		Potassium	N/A	5.195	N/A		0.054								
		Potassium, Dissolved	N/A	4.096	N/A		0.002								
		Sodium	N/A	59.45	N/A		0.035								
		Sodium, Dissolved	N/A	60.433	N/A		0.001U								
		Uranium	0.000906	0.002	0.03		0.035								
		Zinc	0.45	0.054	N/A										

Table 4.45. SWMU 30 Locations of Groundwater Contaminants (Continued)

Hydro Unit	Depth Interval	Analysis	Child Resident NAL	Bkgd	MCL	030-003	MW245	MW63	MW64	MW65	MW66	WB-1	WB-4	WB-5	WBP-4A
	75	Radionuclides (pCi/L)													
		Neptunium-237	0.573	0.8	N/A		0.2								
		Radium-226	0.1	0.6	5		0.52								
		Radon	N/A	N/A	N/A		319.99								
		Radon-222	0.866	626	N/A		394								
		Technetium-99	14	22.3	900		63.6								
		Thorium-230	0.424	1.1	N/A		0.73U								
		Volatiles (mg/L)													
		Chloroform	0.0000287	N/A	N/A		0.025U								
		cis-1,2-Dichloroethene	0.00273	N/A	0.07		0.025U								
		Tetrachloroethene	0.000582	N/A	0.005		0.025U								
		Toluene	0.0338	N/A	1		0.025U								
		Trichloroethene	0.0016	N/A	0.005		0.21								
	91	Metals (mg/L)													
		Aluminum	1.49	2.189	N/A					0.015U					
		Arsenic	0.000035	0.005	0.01					0.0026U					
		Barium	0.104	0.235	2					0.051					
		Barium, Dissolved	0.104	0.2	2					0.07U					
		Beryllium	0.00264	0.004	0.004					0.0003					
		Cadmium	0.000661	0.01	0.005					0.025U					
		Calcium	N/A	41.238	N/A					15.2					
		Calcium, Dissolved	N/A	38.166	N/A					15.2					
		Chromium	1.76	0.144	0.1					0.0025					
		Cobalt	0.0906	0.045	N/A					0.045U					
		Cobalt, Dissolved	0.0906	0.045	N/A					0.045U					
		Copper	0.0557	0.036	1.3					0.025U					
		Iron	0.449	5.03	N/A					0.22					
		Iron, Dissolved	0.449	0.267	N/A					0.36U					
		Lead	0.015	0.129	0.015					0.0017U					
		Magnesium	N/A	16.262	N/A					6.55					
		Magnesium, Dissolved	N/A	16.215	N/A					6.5					
		Manganese	0.035	0.119	N/A					0.0042					
		Manganese, Dissolved	0.035	0.068	N/A					0.02U					
		Molybdenum	0.00753	0.05	N/A					0.015U					
		Nickel	0.0301	0.682	N/A					0.014					
		Nickel, Dissolved	0.0301	0.305	N/A					0.1U					
		Potassium	N/A	5.195	N/A					10.5U					
		Potassium, Dissolved	N/A	4.096	N/A					10.5U					
		Sodium	N/A	59.45	N/A					13.7					
		Sodium, Dissolved	N/A	60.433	N/A					13.2					
		Uranium	0.000906	0.002	0.03					0.00171					
		Vanadium	0.00925	0.134	N/A					0.0017U					
		Zinc	0.45	0.054	N/A					0.041					

RGA

Table 4.45. SWMU 30 Locations of Groundwater Contaminants (Continued)

Hydro Unit	Depth Interval	Analysis	Child Resident NAL	Bkgd	MCL	030-003	MW245	MW63	MW64	MW65	MW66	WB-1	WB-4	WB-5	WBP-4A		
RGA	91	Radionuclides (pCi/L)															
		Neptunium-237	0.573	0.8	N/A						0.989U						
		Potassium-40	N/A	N/A	N/A						458U						
		Radon-222	0.866	626	N/A						475						
		Technetium-99	14	22.3	900						37.11						
		Thorium-230	0.424	1.1	N/A						0.47						
		Uranium	N/A	N/A	N/A						40U						
		Uranium-234	0.546	0.7	N/A						0.17						
		Uranium-238	0.443	0.7	N/A						10.4U						
		Volatiles (mg/L)															
		1,1,2-Trichloro-1,2,2-trifluoroethane	2.7	N/A	N/A							0.00036U					
		4-Methyl-2-pentanone	0.00722	N/A	N/A							0.0025					
		Acetone	0.0275	N/A	N/A							0.0081					
		Chloroform	0.0000287	N/A	N/A							0.005U					
		cis-1,2-Dichloroethene	0.00273	N/A	N/A	0.07						0.0056					
		Tetrachloroethene	0.000582	N/A	N/A	0.005						0.005U					
		Toluene	0.0338	N/A	N/A	1						0.011					
		Trichloroethene	0.0016	N/A	N/A	0.005						0.096					

Blank cells indicate interval was not sampled for the specified analysis. Maximum value shown for each depth interval.

N/A = not applicable

Bold indicates result is greater than NAL value.

Italics indicates result is greater than MCL value.

Bold + Italics indicate result is greater than both NAL and MCL values.

Shading indicates result is greater than the background value.

4.10 SWMU 145

4.10.1 Subsurface Soils

The RI collected subsurface soil samples from seven angled borings at SWMU 145 (Figure 4.8). These seven angled borings and historic boring DG029 characterize the landfilled waste area of SWMU 145. Historic samples that characterize the fill (waste and soils) of the abandoned NSDD channel derive from 11 historic soil borings. Table 4.46 lists the contaminants identified by a review of the RI and historical data for the landfill area; Table 4.47 lists the contaminants identified for the area of the abandoned NSDD channel.

Locations of SWMU 145 subsurface soil contaminants are listed in Table 4.48. The “145-#” series borings were sampled for the BGOU RI. All samples of these borings and historic boring DG-029 characterized the landfill area. The other historic borings—A2, A10, NSD030, and the “NST#” series borings—characterize the abandoned channel of the NSDD.

The metal detected predominantly above screening levels in subsurface soils in the area of the SWMU 145 landfill is antimony. One third of the samples had an antimony level that exceeded background and the excavation worker NAL criteria. Figure 4.63 presents the antimony distribution in soil. The antimony “hot spot” shown in this figure in the southwest corner, near historical borings DG-029 and NSD030, is the result of modeling the contaminant distribution using a high laboratory reporting limit (antimony was not detected at the reporting limit in the sample). As explained in Appendix D, the laboratory reporting limit was used for modeling, and in this instance, the reporting limit was 20 mg/kg. Antimony concentrations, for the most part, exceeded the background value in the landfill area throughout the depth of the shallow soils to the top of the RGA. The only other metal that was frequently present at concentrations above the NAL (but rarely exceeded background) in soils of the landfill area was aluminum.

Arsenic was the most frequent metal to exceed the excavation worker NAL in samples from the abandoned channel of the NSDD. Figure 4.64 shows the arsenic distribution in soil at SWMU 145. The vertical extent of arsenic contamination is an uncertainty. All of the historic samples that characterized the former ditch channel were collected from depths of 15 ft or less. Uranium and beryllium were other commonly detected metals at levels above background and the excavation worker NAL. The extent of the depth of contamination by these metals also is uncertain.

Of the organics in subsurface soils, only PCBs were detected in one sample from the landfill area (145-103, from the depth interval 8 to 10 ft), at 0.33 mg/kg. PCBs were detected at levels above NAL criteria at four historical sampling locations within the abandoned channel of the former NSDD (NST1S01, NST2S02, NSD030, and A10, all at depths of 2 to 4 ft). The maximum detected PCB result was 12.5 mg/kg from A10.

Table 4.46. SWMU 145 (Landfill Area) Subsurface Soil Contaminants

Analysis	Maximum Result		Frequency of Detection ^a	Frequency of Detection	
	Historical Data	RI Data		Above Background Value	Above Excavation Worker NAL
<i>Inorganics (mg/kg)</i>					
Aluminum	16,200	12,700	42/42	3/42	35/42
Antimony	ND	20.2	15/42	15/42	15/42
Arsenic	ND	7.88	30/42	0/42	30/42
Barium	226	183	42/42	2/42	0/42
Beryllium	1.88	1.24	10/42	10/42	3/42
Cadmium	ND	2.47	1/42	1/42	0/42
Calcium	2330	20,900	42/42	1/42	N/A
Chromium	25.1	27.9	42/42	0/42	0/42
Cobalt	15	20.5	24/42	2/42	0/42
Copper	11.1	81.8	40/42	1/42	0/42
Iron	25,200	23,000	42/42	0/42	42/42
Lead	ND	27.9	36/42	1/42	0/42
Lithium	9.95	N/A	6/6	N/A	0/6
Magnesium	2,160	2,070	42/42	1/42	0/42
Manganese	626	733	42/42	0/42	30/42
Mercury	ND	0.4	6/42	1/42	0/42
Molybdenum	N/A	3.39	2/36	N/A	0/36
Nickel	23.1	98.9	33/42	3/42	0/42
Potassium	1,300	N/A	6/6	2/6	N/A
Sodium	281	228	30/42	0/42	0/42
Strontium	22	N/A	6/6	N/A	0/6
Uranium	N/A	1.55	9/36	0/36	0/36
Vanadium	29.1	24.3	22/32	0/32	21/32
Zinc	87.8	81.6	15/42	2/42	0/42
<i>Organics--Volatiles (mg/kg)</i>					
2-Butanone	ND	0.0283	3/45	N/A	0/45
Acetone	ND	0.116	10/45	N/A	0/45
<i>Organics--Semivolatiles (mg/kg)</i>					
Di-n-butyl phthalate	0.81	N/A	1/7	N/A	0/7
<i>Organics--PCBs (mg/kg)</i>					
PCB, Total	ND	0.33	1/38	N/A	1/38
PCB-1254	ND	0.33	1/38	N/A	1/38
<i>Radionuclides(pCi/g)</i>					
Technetium-99	ND	1.83	1/45	0/45	0/45
Thorium-228	N/A	0.775	36/37	0/37	36/37
Thorium-230	N/A	0.534	34/37	0/37	0/37
Thorium-232	N/A	0.727	36/37	0/37	0/37
Thorium-234	ND	4.7	8/45	N/A	N/A
Uranium	N/A	0.795	1/37	N/A	N/A
Uranium-234	N/A	0.405	17/37	0/37	0/37
Uranium-238	N/A	0.378	17/37	0/37	0/37

a Frequency of detection is the number of detections of an analyte per number of analyses

(includes regular and duplicate samples).

N/A = not applicable

ND = not detected

**Table 4.47. SWMU 145 (Old North-South Diversion Ditch Area)
Subsurface Soil Contaminants**

Analysis	Maximum Result		Frequency of Detection ^a	Frequency of Detection	
	Historical Data	RI Data		Above Background Value	Above Excavation Worker NAL
<i>Inorganics (mg/kg)</i>					
Aluminum	11,500	N/A	3/3	0/3	3/3
Arsenic	21.9	N/A	9/17	4/17	9/17
Barium	300	N/A	17/17	1/17	1/17
Beryllium	2.08	N/A	13/17	5/17	2/17
Cadmium	2.36	N/A	1/17	1/17	0/17
Calcium	5330	N/A	3/3	0/3	N/A
Chromium	120	N/A	17/17	2/17	0/17
Cobalt	9.72	N/A	2/3	0/3	0/3
Copper	135	N/A	3/3	1/3	0/3
Iron	11,000	N/A	3/3	0/3	3/3
Lead	46.7	N/A	5/17	5/17	0/17
Lithium	8.37	N/A	3/3	N/A	0/3
Magnesium	1260	N/A	3/3	0/3	0/3
Manganese	572	N/A	3/3	0/3	3/3
Mercury	0.47	N/A	1/17	1/17	0/17
Nickel	101	N/A	3/3	1/3	0/3
Silver	19.6	N/A	1/17	1/17	0/17
Uranium	311	N/A	15/17	7/17	7/17
Vanadium	27.7	N/A	3/3	0/3	3/3
Zinc	261	N/A	8/8	3/8	0/8
<i>Organics--Volatiles (mg/kg)</i>					
1,2-Dimethylbenzene	0.036	N/A	2/16	N/A	0/16
4-Methyl-2-pentanone	0.015	N/A	1/16	N/A	0/16
Acetone	0.055	N/A	9/16	N/A	0/16
Ethylbenzene	0.018	N/A	3/16	N/A	0/16
m,p-Xylene	0.04	N/A	2/16	N/A	0/16
Toluene	0.037	N/A	5/16	N/A	0/16
<i>Organics--PCBs (mg/kg)</i>					
PCB, Total	12.5	N/A	4/17	N/A	5/17
PCB-1254	1.9	N/A	2/17	N/A	2/17
PCB-1260	12.5	N/A	2/17	N/A	2/17
<i>Radionuclides (pCi/g)</i>					
Actinium-228	5.05	N/A	12/14	N/A	N/A
Americium-241	1.956	N/A	2/19	N/A	1/19
Bismuth-212	4.705	N/A	13/15	N/A	N/A
Bismuth-214	5.561	N/A	14/14	N/A	N/A
Cesium-137	1.057	N/A	11/19	4/19	7/19
Lead-210	2.646	N/A	1/15	N/A	N/A
Lead-212	4.629	N/A	13/13	N/A	N/A
Lead-214	5.456	N/A	14/14	N/A	N/A
Neptunium-237	1.169	N/A	7/33	N/A	3/33
Plutonium-239/240	10.1	N/A	6/18	N/A	1/18
Potassium-40	52.21	N/A	15/16	1/16	N/A
Protactinium-231	2.645	N/A	8/13	N/A	N/A
Protactinium-233	0.372	N/A	2/15	N/A	N/A
Protactinium-234m	260	N/A	11/13	N/A	N/A

**Table 4.47. SWMU 145 (Old North-South Diversion Ditch Area)
Subsurface Soil Contaminants (Continued)**

Analysis	Maximum Result		Frequency of Detection ^a	Frequency of Detection	
	Historical Data	RI Data		Above Background Value	Above Excavation Worker NAL
Radium-223	0.5283	N/A	1/12	N/A	N/A
Radium-226	0.16	N/A	1/11	0/11	1/11
Radium-228	0.3315	N/A	2/11	N/A	N/A
Strontium-90	5	N/A	6/14	0/14	3/14
Technetium-99	281	N/A	11/18	11/18	5/18
Thallium-208	0.346	N/A	12/15	N/A	N/A
Thorium-228	1.92	N/A	22/31	1/31	22/31
Thorium-230	193	N/A	18/18	6/18	5/18
Thorium-232	2.282	N/A	23/29	2/29	2/29
Thorium-234	260	N/A	12/14	N/A	N/A
Uranium	593	N/A	10/13	N/A	N/A
Uranium-233/234	4.7	N/A	5/5	1/5	N/A
Uranium-234	254	N/A	19/26	13/26	12/26
Uranium-235	2.2	N/A	16/17	7/17	3/17
Uranium-236	0.12	N/A	1/5	N/A	N/A
Uranium-238	326	N/A	26/31	18/31	18/31

^a Frequency of detection is the number of detections of an analyte per number of analyses (includes regular and duplicate samples).

N/A = not applicable

ND = not detected

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Table 4.48. SWMU 145 Locations of Subsurface Soil Contaminants

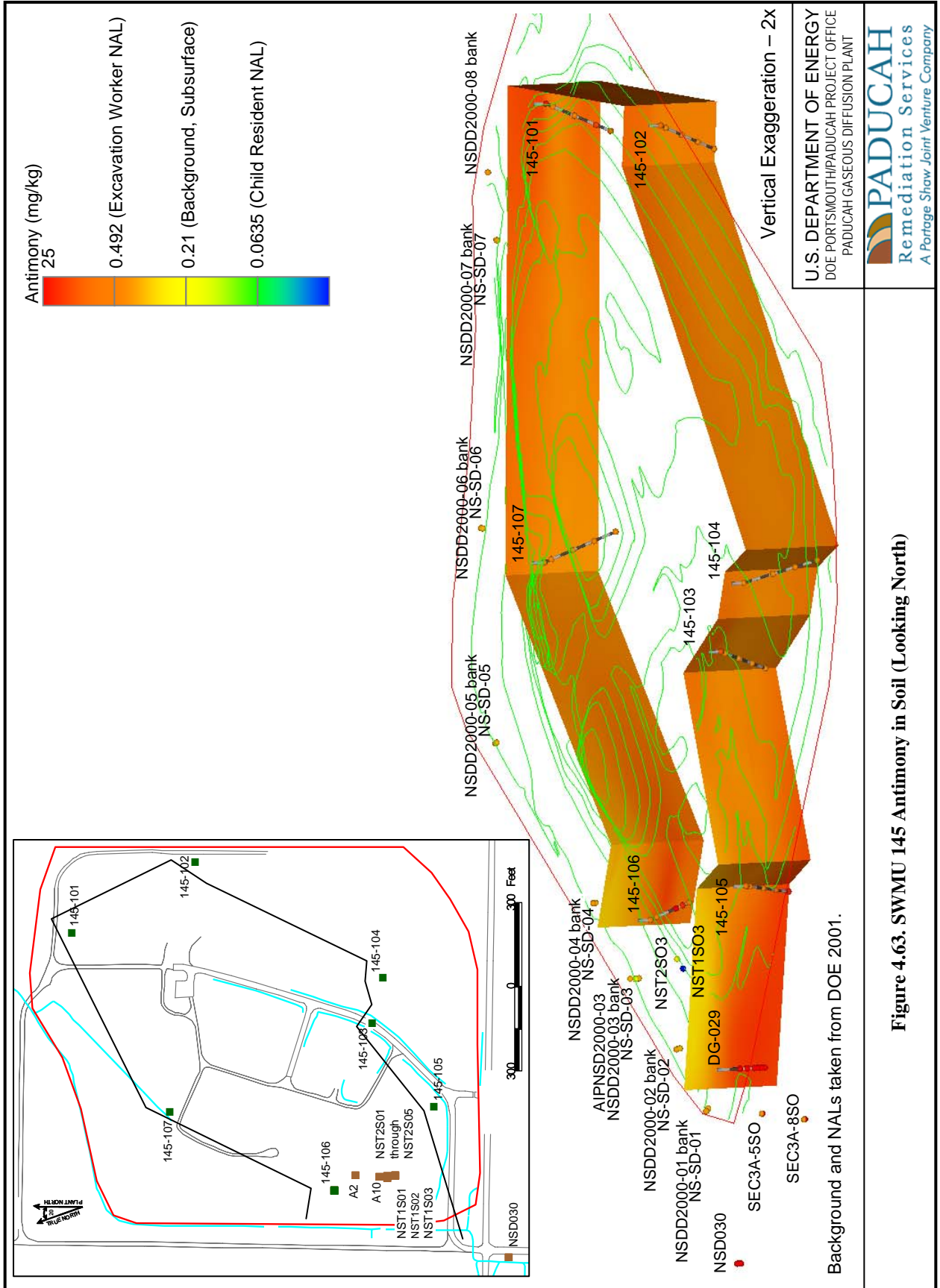
Analysis	Depth (ft)	Excavation Worker NAL	Background	RI Data							Historical Data													
				145-101	145-102	145-103	145-104	145-105	145-106	145-107	DG-029	A10	A2	NSD030	NST1S01	NST1S02	NST1S03	NST2S01	NST2S02	NST2S03	NST2S04	NST2S05		
Inorganics (mg/kg)	01-04	5.250	12,000	7,400	10,900	10,600	9,660	12,700	6,100	8,350	11,500													
	08-10	5.250	12,000	7,380	6,470	6,770			8,090	4,190	10,200													
	11-15	5.250	12,000																					
	19-25	5.250	12,000																					
	30-34	5.250	12,000	7,500	5,030	6,150	7,200	7,200	5,310	9,720	6,840	8,180												
	35-37	5.250	12,000																					
Antimony	01-04	5.250	12,000	7,170	5,580	4,170	6,340	7,070	7,730	5,770	7,200													
	08-10	0.492	0.21	7,480	5,830	5,620	8,030	2,370	3,670															
	11-15	0.492	0.21	18.1	11.5	13.6	9.68U	10.9	9.82U	9.81														
	19-25	0.492	0.21	9.57U	9.42U	9.59	11	9.83U	9.52U															
	30-34	0.492	0.21	9.83U	9.53U	8.76U	9.67U	9.62U	20.2	9.54U	20U													
	35-37	0.492	0.21						16.6		20U													
Arsenic	01-04	0.492	0.21	14.7	9.83U	9.2U	9.9U	12.5	11.1	9.79U	20U													
	08-10	0.324	7.9	13.2	9.57U	9.42U	10.7	17.7		9.45U														
	11-15	0.324	7.9	3.71	7.18	7.16	3.33	6.45	3.32	7.88	5.03	5U												
	19-25	0.324	7.9	1.19	2.87	6.67		3.57	2.04	3.25	21.9													
	30-34	0.324	7.9	2.01	2.36	2.36	3.39	2.16	3.68	0.894U	5U													
	35-37	0.324	7.9	6.98	1.57	1.72	1.92	2.81	3.1	0.895U	5U													
Barium	01-04	272	170	0.983U	1.49	3.16	2.63	3.92		0.926U														
	08-10	272	170	94.4	103	91.7	97.3	150	79	89.7	82.3	122	133											
	11-15	272	170	47.8	97.8	73.7		162	84.4	54.8														
	19-25	272	170																					
	30-34	272	170	23.3	32.8	82	95.3	30.6	64	40.3	41.7													
	35-37	272	170	38.2	38.9	36.5	56.5	83.2	98.5	28.1	226													
Beryllium	01-04	1.26	0.69	0.494U	0.492U	0.487U	0.499U	0.477U	0.812	0.447U	0.5U													
	08-10	1.26	0.69	0.449U	0.487U	0.484U	0.465U	0.482U	0.474U	0.466U	0.6	0.78												
	11-15	1.26	0.69	0.488U	0.486U	0.488U		0.497U	0.456U	0.487U	2.08													
	19-25	1.26	0.69	0.494U	0.492U	0.487U	0.499U	0.477U	0.812	0.447U														
	30-34	1.26	0.69	0.495U	0.478U	0.484U	0.454U	0.827	0.745	0.448U	1.48													
	35-37	1.26	0.69	0.492U	0.474U	1.24	0.706	0.798		0.463U														
Cadmium	01-04	15.2	0.21	1.79U	1.95U	1.94U	1.86U	1.93U	1.9U	1.86U	2U	2U	2U											
	08-10	15.2	0.21	1.95U	1.95U	1.95U	1.99U	1.82U	1.82U	1.95U	2U	2U	2U											
	11-15	15.2	0.21	1.97U	1.89U	1.95U	1.99U	1.77U	1.85U															
	19-25	15.2	0.21	1.97U	1.97U	1.95U	1.99U	1.98U	1.79U	1.79U	2U													
	30-34	15.2	0.21	2.47	1.91U	1.94U	1.82U	1.82U	1.79U	1.94U	2U													
	35-37	15.2	0.21	1.97U	1.89U	1.95U	1.99U	1.77U	1.85U															
Calcium	01-04	n/a	6,100	1,040	1,090	1,020	1,180	1,080	1,700	1,260	1,080													
	08-10	n/a	6,100	1,210	420	20,900	4,970	547	1,040	4,540														
	11-15	n/a	6,100	1,240	1,050	1,020	1,630	901	544															
	19-25	n/a	6,100																					
	30-34	n/a	6,100	1,040	1,090	1,020	1,180	1,080	1,700	1,260	1,080													
	35-37	n/a	6,100	968	674	649	962	1,400	1,360	1,430	2,330													

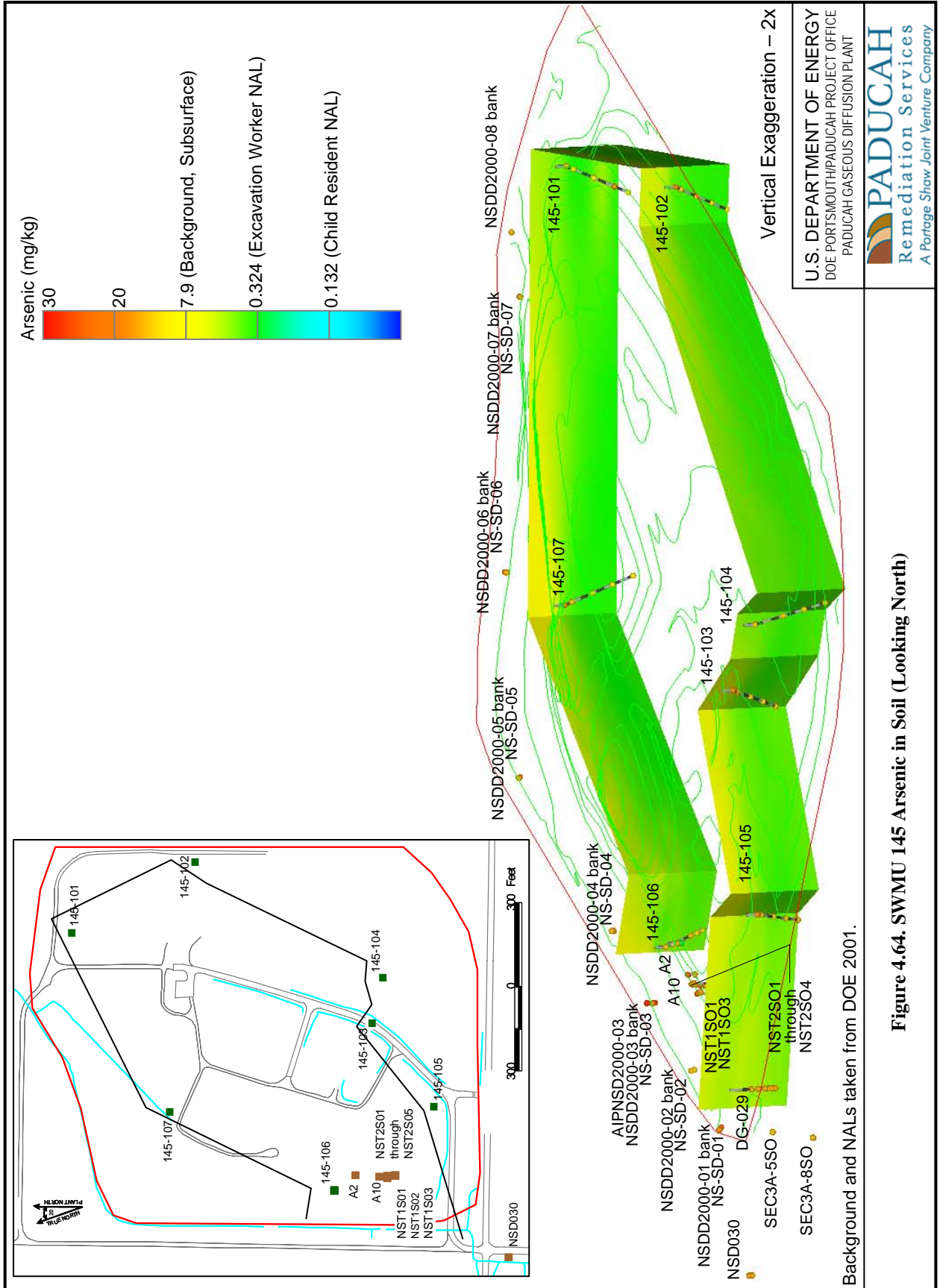
Table 4-48. SWMU 145 Locations of Subsurface Soil Contaminants (Continued)

Analysis	Depth (ft)	Excavation Worker NAL	Background	RI Data							Historical Data											
				145-101	145-102	145-103	145-104	145-105	145-106	145-107	DG-029	A10	A2	NSD030	NST1S01	NST1S02	NST1S03	NST2S01	NST2S02	NST2S03	NST2S04	NST2S05
Manganese	30-34	56.6	820	32.9	41.9	178	222	18.9	113	55.8	52.9											
	35-37	56.6	820						116		104											
	40-45	56.6	820	60.1	27.7	16.5	19.6	144	242	13.4	626											
Mercury	55-60	56.6	820	59	219	733	80.1	178		54.4												
	01-04	1.17	0.13																			
	08-10	1.17	0.13	0.017U	0.02U	0.019	0.016U	0.02	0.024	0.017												
Molybdenum	11-15	1.17	0.13	0.019U	0.016U	0.02U		0.019U	0.017U	0.02	0.2U											
	19-25	1.17	0.13	0.015U	0.4	0.019U	0.019U	0.019U	0.02U	0.019U	0.2U											
	30-34	1.17	0.13	0.019U	0.014U	0.02U	0.017U	0.02U	0.015U	0.017U	0.2U											
Nickel	35-37	1.17	0.13	0.019U	0.018U	0.018U	0.02U	0.018U	0.02U	0.019U	0.2U											
	40-45	1.17	0.13	0.019U	0.018U	0.018U	0.02U	0.018U	0.02U	0.019U	0.2U											
	55-60	1.17	0.13	0.019U	0.018U	0.018U	0.02U	0.018U	0.02U	0.019U	0.2U											
Potassium	08-10	66	n/a	2.44U	2.44U	3.39	2.32U	2.41U	2.37U	2.33U												
	11-15	66	n/a	2.44U	2.43U	2.44U	2.49U	2.48U	2.48U	2.23U												
	30-34	66	n/a	2.47U	2.42	2.44U	2.49U	2.39U	2.48U	2.23U												
Silver	35-37	66	n/a	2.47U	2.39U	2.42U	2.27U	2.27U	2.27U	2.24U												
	40-45	66	n/a	2.46U	2.37U	2.44U	2.48U	2.22U	2.24U	2.32U												
	55-60	66	n/a	2.46U	2.37U	2.44U	2.48U	2.22U	2.24U	2.32U												
Sodium	01-04	216	22																			
	08-10	216	22	11	10.3	98.9	9.53	13.3	6.57	9.76												
	11-15	216	22	7.48	11.2	6.54	13.3	4.56U	6.79													
Strontium	19-25	216	22	4.94U	4.92	6.34	7.68	6.15	13.5	4.47U	9.31											
	35-37	216	22					12.5	22.8													
	40-45	216	22	8.69	5.28	4.84U	4.81	8.77	13.4	4.48U	23.7											
Uranium	55-60	216	22	8.24	5.35	16.9	11.5	11.1		4.63U												
	19-25	n/a	950								128											
	30-34	n/a	950								444											
Vanadium	35-37	n/a	950								1300											
	40-45	n/a	950								941											
	01-04	41.2	2.7																			
Zinc	08-10	41.2	2.7	2.24U		2.42U	2.32U		2.37U	2.33U												
	11-15	41.2	2.7	2.44U		2.44U	2.48U	2.22U	2.27U	2.24U												
	19-25	41.2	2.7	2.47U		2.44U	2.49U		2.48U	2.44U												
Cadmium	30-34	41.2	2.7	2.47U		2.44U	2.49U		2.48U	2.23U												
	35-37	41.2	2.7	2.47U		2.44U	2.49U		2.48U	2.23U												
	40-45	41.2	2.7	2.47U		2.44U	2.49U		2.48U	2.23U												
Chromium	55-60	41.2	2.7	2.46U		2.44U	2.48U	2.22U	2.27U	2.24U												
	08-10	n/a	340	132	122	202	154	228	94.9U	179												
	11-15	n/a	340	148	121	111	159	115	205													
Copper	19-25	n/a	340								200U											
	30-34	n/a	340	145	153	97.4U	163	99.4	159	168												
	35-37	n/a	340								212											
Lead	40-45	n/a	340	123	95.7U	96.8U	122	90.9U	108	133												
	55-60	n/a	340	99.9	114	97.7U	99.4U	88.7U		92.6U												
	19-25	5.330	n/a								6.61											
Cobalt	30-34	5.330	n/a								8.63											
	35-37	5.330	n/a								19.5											
	40-45	5.330	n/a								22											
Iron	01-04	11.3	4.6																			
	08-10	11.3	4.6	0.897U	1.15	0.993	0.929U	0.963U	0.976	0.932U												
	11-15	11.3	4.6	0.975U	0.973U	0.975U	0.997U	0.994U	0.912U	0.974U												
Manganese	30-34	11.3	4.6	0.987U	0.984U	0.974U	0.997U	0.954U	1.55	0.894U												
	35-37	11.3	4.6	0.987U	0.984U	0.974U	0.997U	0.954U	1.55	0.894U												
	40-45	11.3	4.6	0.99U	0.957U	0.968U	0.908U	0.909U	0.908U	1.16												
Zinc	55-60	11.3	4.6	1.16	0.947U	1.41	0.999	0.887U		0.926U												
	01-04	4.4	37																			
	08-10	4.4	37	13.8		2.42U	2.32U		24.3	22.5												
Cadmium	11-15	4.4	37	10.4		2.44U			3.92	14.7												
	19-25	4.4	37								12.6											

Table 4-48. SWMU 145 Locations of Subsurface Soil Contaminants (Continued)

Analysis	Depth (ft)	Excavation Worker NAL	Background	RI Data										Historical Data														
				145-101	145-102	145-103	145-104	145-105	145-106	145-107	DG-029	A10	A2	NSD030	NST1S01	NST1S02	NST1S03	NST2S01	NST2S02	NST2S03	NST2S04	NST2S05						
m,p-Xylene	30-34	963	n/a	0.00994U	0.01U	0.00994U	0.00991U	0.00995U	0.00996U	0.0101U	0.01U	0.01U	0.01U	0.01U	0.01U													
	35-37	963	n/a						0.01U																			
	40-45	963	n/a	0.01U	0.0101U	0.00991U	0.01U	0.01U	0.00997U	0.01U	0.01U	0.01U	0.01U	0.01U	0.01U													
	55-60	963	n/a	0.01U	0.0101U	0.00992U	0.00993U	0.00996U																				
	01-04	272	n/a															0.037	0.018	0.025	0.018							
	08-10	272	n/a	0.00501U	0.005U	0.00504U	0.00501U	0.00499U	0.00499U	0.00499U	0.00505U	0.00505U	0.00505U	0.00505U	0.01U	0.01U	0.01U	0.01U	0.01U	0.01U	0.01U	0.01U	0.01U	0.01U	0.01U			
Toluene	11-15	272	n/a	0.00504U	0.00499U	0.00504U	0.00497U	0.005U	0.00502U	0.00497U	0.00497U	0.00497U	0.00497U	0.01U	0.01U	0.01U	0.01U	0.01U	0.01U	0.01U	0.01U	0.01U	0.01U	0.01U	0.01U			
	19-25	272	n/a	0.00497U	0.00501U	0.00497U	0.00496U	0.00498U	0.00498U	0.00504U	0.00504U	0.00504U	0.00504U	0.01U	0.01U	0.01U	0.01U	0.01U	0.01U	0.01U	0.01U	0.01U	0.01U	0.01U	0.01U			
	30-34	272	n/a	0.00497U	0.00501U	0.00497U	0.00496U	0.00498U	0.00498U	0.00504U	0.00504U	0.00504U	0.00504U	0.01U	0.01U	0.01U	0.01U	0.01U	0.01U	0.01U	0.01U	0.01U	0.01U	0.01U	0.01U			
	35-37	272	n/a						0.00502U																			
	40-45	272	n/a	0.005U	0.00505U	0.00495U	0.005U	0.005U	0.00499U	0.005U	0.005U	0.005U	0.005U	0.01U	0.01U	0.01U	0.01U	0.01U	0.01U	0.01U	0.01U	0.01U	0.01U	0.01U	0.01U			
	55-60	272	n/a	0.00502U	0.00503U	0.00496U	0.00496U	0.00498U																				
Organics--Semivolatile (mg/kg)																												
Di-n-butyl phthalate																												
19-25	1,520	n/a															0.5U											
30-34	1,520	n/a															0.5U											
35-37	1,520	n/a															0.81											
40-45	1,520	n/a															0.5U											
Organics--PCBs (mg/kg)																												
PCB, Total																												
01-04	0.168	n/a	n/a	0.09U	0.1U	0.33	0.08U	0.07U	0.09U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	12.5	0.1U	1.4	1.4	1.9	1.9	1U	1U	0.8U	1.1	0.1U	0.1U
08-10	0.168	n/a	n/a	0.09U	0.09U	0.09U	0.09U	0.08U	0.09U	0.09U	0.09U	0.09U	0.09U	0.09U	0.09U	0.09U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U
11-15	0.168	n/a	n/a	0.09U	0.09U	0.09U	0.09U	0.08U	0.09U	0.09U	0.09U	0.09U	0.09U	0.09U	0.09U	0.09U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U
19-25	0.168	n/a	n/a	0.08U	0.09U	0.09U	0.1U	0.08U	0.09U	0.09U	0.09U	0.09U	0.09U	0.09U	0.09U	0.09U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U
30-34	0.168	n/a	n/a	0.08U	0.09U	0.09U	0.09U	0.08U	0.09U	0.09U	0.09U	0.09U	0.09U	0.09U	0.09U	0.09U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U
35-37	0.168	n/a	n/a																									
40-45	0.168	n/a	n/a	0.09U	0.08U	0.08U	0.09U	0.09U	0.09U	0.09U	0.09U	0.09U	0.09U	0.09U	0.09U	0.09U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U
55-60	0.168	n/a	n/a	0.08U	0.09U	0.09U	0.09U	0.08U	0.09U	0.09U	0.09U	0.09U	0.09U	0.09U	0.09U	0.09U	0.1U	0.1U	0.9	0.9	0.9	1.9	1U	1U	0.8U	1U		
PCB-1254																												
01-04	0.168	n/a	n/a	0.09U	0.1U	0.33	0.08U	0.07U	0.09U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	12.5	0.1U	0.5	0.5	0.1U	0.1U	0.1U	0.1U	0.8U	1U		
08-10	0.168	n/a	n/a	0.09U	0.09U	0.09U	0.09U	0.08U	0.09U	0.09U	0.09U	0.09U	0.09U	0.09U	0.09U	0.09U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U
11-15	0.168	n/a	n/a	0.09U	0.09U	0.09U	0.09U	0.08U	0.09U	0.09U	0.09U	0.09U	0.09U	0.09U	0.09U	0.09U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U
19-25	0.168	n/a	n/a	0.08U	0.09U	0.09U	0.1U	0.08U	0.09U	0.09U	0.09U	0.09U	0.09U	0.09U	0.09U	0.09U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U
30-34	0.168	n/a	n/a	0.08U	0.09U	0.09U	0.09U	0.08U	0.09U	0.09U	0.09U	0.09U	0.09U	0.09U	0.09U	0.09U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U
35-37	0.168	n/a	n/a																									
40-45	0.168	n/a	n/a	0.09U	0.08U	0.08U	0.09U	0.09U	0.09U	0.09U	0.09U	0.09U	0.09U	0.09U	0.09U	0.09U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U
55-60	0.168	n/a	n/a	0.08U	0.09U	0.09U	0.09U	0.08U	0.09U	0.09U	0.09U	0.09U	0.09U	0.09U	0.09U	0.09U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U
PCB-1260																												
01-04	0.168	n/a	n/a	0.09U	0.1U	0.33	0.08U	0.07U	0.09U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	12.5	0.1U	0.5	0.5	0.1U	0.1U	0.1U	0.1U	0.8U	1U		
08-10	0.168	n/a	n/a	0.09U	0.09U	0.09U	0.09U	0.08U	0.09U	0.09U	0.09U	0.09U	0.09U	0.09U	0.09U	0.09U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U
11-15	0.168	n/a	n/a	0.09U	0.09U	0.09U	0.09U	0.08U	0.09U	0.09U	0.09U	0.09U	0.09U	0.09U	0.09U	0.09U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U
19-25	0.168	n/a	n/a	0.08U	0.09U	0.09U	0.1U	0.08U	0.09U	0.09U	0.09U	0.09U	0.09U	0.09U	0.09U	0.09U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U
30-34	0.168	n/a	n/a	0.08U	0.09U	0.09U	0.09U	0.08U	0.09U	0.09U	0.09U	0.09U	0.09U	0.09U	0.09U	0.09U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U
35-37	0.168	n/a	n/a																									
40-45	0.168	n/a	n/a	0.09U	0.08U	0.08U	0.09U	0.09U	0.09U	0.09U	0.09U	0.09U	0.09U	0.09U	0.09U	0.09U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U
55-60	0.168	n/a	n/a	0.08U	0.09U	0.09U	0.09U	0.08U	0.09U	0.09U	0.09U	0.09U	0.09U	0.09U	0.09U	0.09U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U	0.1U
Radionuclides (pCi/g)																												
Actinium-228																												
01-04	n/a	n/a	n/a														0.66	0.64	0.4131	0.6114	0.4929	0.2238U	0.4131	0.6114	0.4131	0.6114	1.934	
08-10	n/a	n/a	n/a														0.59	0.59	0.9062	0.9286	0.9062	0.243U	0.1137U	0.2051U	0.9286	0.9286	5.05	
11-15	n/a	n/a	n/a														0.1U	0.1U	0.1137U	0.2051U	0.1802U	0.243U	0.1137U	0.2051U	0.9286	0.9286	5.05	
01-04	1.74	n/a	n/a	0.0417U	0.0418U	0.0499U	0.0504U	0.0498U	0.0422U	0.0417U	0.0417U	0.0422U	0.0422U	0.0422U	0.0417U	0.0417U	0.1U	0.11U	0.1137U	0.2051U	0.1802U	0.243U	0.1137U	0.2051U	0.9286	0.9286	5.05	
08-10	1.74	n/a	n/a	0.0417U	0.0418U	0.0499U	0.0504U	0.0498U	0.0422U	0.0417U	0.0417U	0.0422U	0.0422U	0.0422U	0.0417U	0.0417U	0.1U	0.11U	0.1137U	0.2051U	0.1802U	0.243U	0.1137U	0.2051U	0.9286	0.9286	5.05	
11-15	1.74	n/a	n/a	0.0423U	0.0417U	0.0499U	0.0511U	0.05																				





Screening did not identify any radionuclides above both background and excavation worker NALs in subsurface soils of the landfill area. Radionuclides in subsurface soils of the abandoned NSDD channel at SWMU 145 include americium-241, cesium-137, technetium-99, thorium isotopes, and uranium isotopes. Figure 4.65 shows the uranium distribution at SWMU 145. This figure shows the elevated concentrations of uranium in shallow samples collected near the previous location of the NSDD. Figure 4.66 presents the thorium-228 distribution in soils at SWMU 145. Thorium is widely distributed (above NALs) throughout the entire thickness of the UCRS, but localized hot spots occur near the buried reach of the NSDD. The highest detected activity from the database for SWMU 145 was 193 pCi/g from historic boring NSD030, at a depth of 1 to 4 ft bgs.

4.10.2 SWMU 145 Groundwater

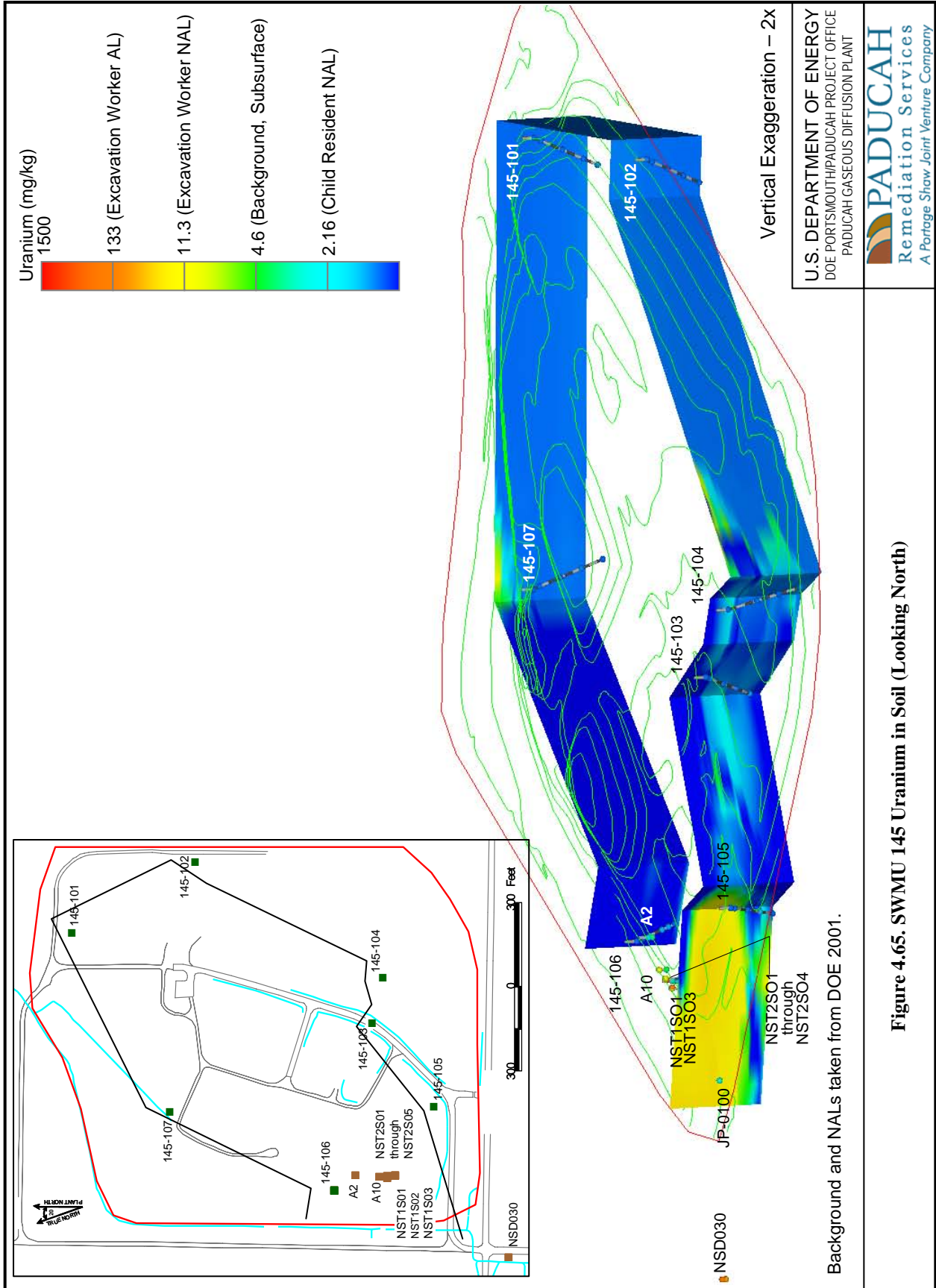
The collection of UCRS groundwater samples was attempted at each of the seven angled borings installed at SWMU 145; however, sufficient groundwater was not available for sampling. Historical data from seven MWs in the area were reviewed to identify the UCRS contaminants listed in Table 4.49. The UCRS wells with groundwater data for SWMU 145 (for the period 1995 through 2006) were MW180, MW182, MW371, MW386, MW390, MW393, and MW396 (all are located peripheral to the burial area) (Figure 4.16).

Screening of the SWMU 145 analyses determined six metals that exceed contaminant criteria in UCRS groundwater. Iron and manganese were common groundwater contaminants. Arsenic and uranium accounted for most of the other metal exceedances.

The VOC chloroform (in a sample from MW386) was the only UCRS groundwater organic contaminant to exceed screening criteria. Analyses detected TCE in samples from six locations; however, the maximum detected result (0.002 mg/L or 2 µg/L) was less than the screening level. PCB-1260 was detected at MW371 at 0.00007 mg/L (0.07 µg/L).

Uranium contamination in the UCRS groundwater was found primarily at location MW182. Samples from MW182 accounted for most of the detections of uranium-234 above screening levels, as well as one of the detections of uranium-238 as a groundwater contaminant. The isotope uranium-238 also was detected above screening levels at location MW180.

RGA and McNairy groundwater samples were not collected at SWMU 145 as part of this RI. Historical data for the period 1995 through 2006 were reviewed for RGA and McNairy groundwater from 25 RGA MWs in the area and one temporary boring that sampled the McNairy Formation to identify the contaminants listed in Table 4.50.



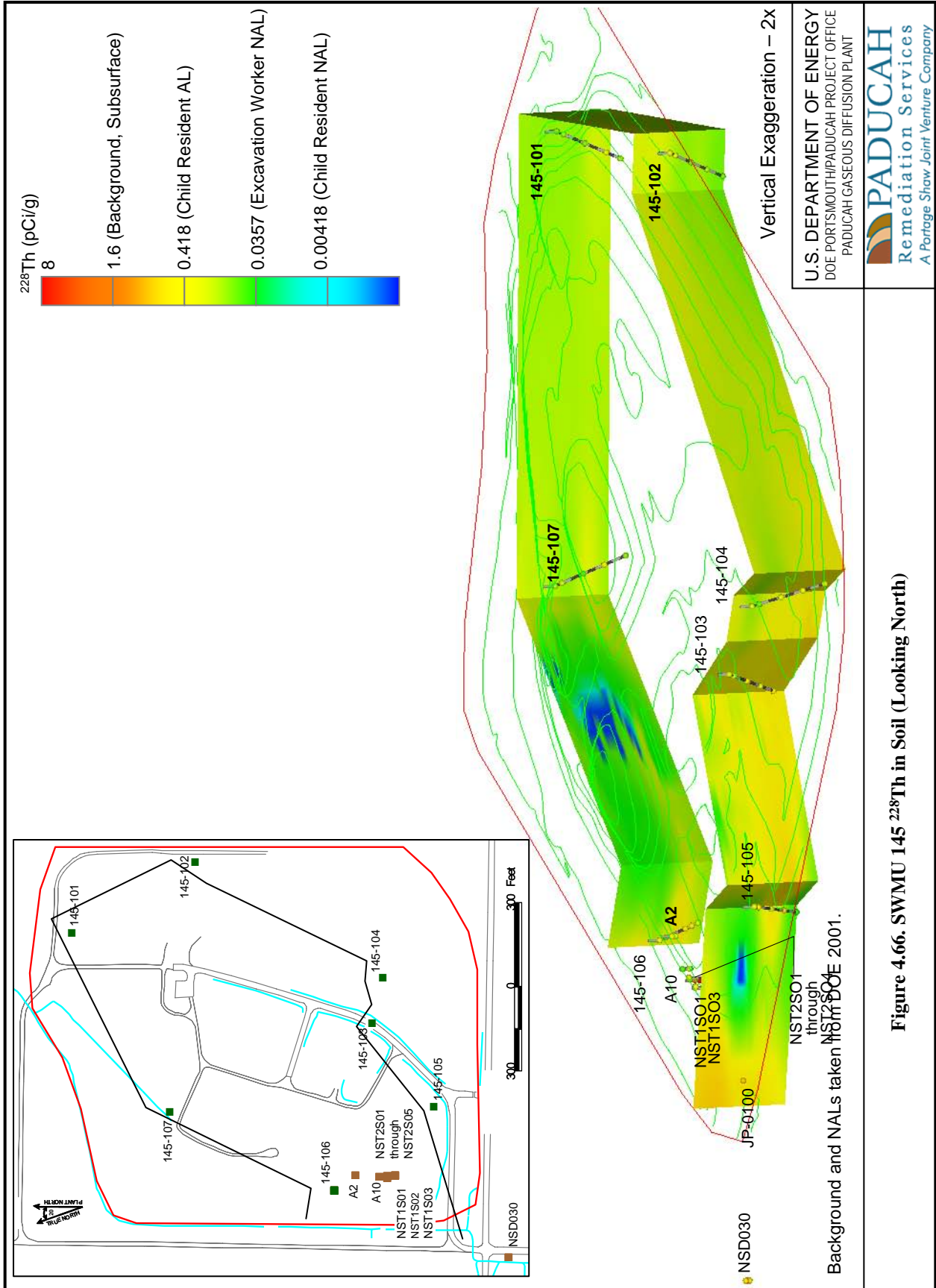


Figure 4.66. SWMU 145 ²²⁸Th in Soil (Looking North)

Table 4.49. SWMU 145 UCRS Groundwater Contaminants

Analysis	Historical Data	RI	Frequency of Detection ^a	Frequency of Detection		
				Above Background	Above NAL	Above MCL
Metals (mg/L)						
Aluminum	15	N/A	37/116	N/A	14/116	N/A
Aluminum, Dissolved	0.346	N/A	1/2	N/A	0/2	N/A
Arsenic	0.0189	N/A	42/68	N/A	42/68	2/68
Arsenic, Dissolved	0.0158	N/A	4/4	N/A	4/4	2/4
Barium	0.897	N/A	113/113	N/A	84/113	0/113
Barium, Dissolved	0.816	N/A	111/112	N/A	80/112	0/112
Beryllium	0.0011	N/A	1/113	N/A	0/113	0/113
Boron	0.07	N/A	1/111	N/A	0/111	N/A
Calcium	165	N/A	113/113	N/A	N/A	N/A
Calcium, Dissolved	121	N/A	2/2	N/A	N/A	N/A
Chromium	0.197	N/A	5/118	N/A	0/118	2/118
Cobalt	0.067	N/A	43/113	N/A	0/113	N/A
Copper	0.031	N/A	5/118	N/A	0/118	0/118
Iron	33.6	N/A	115/118	N/A	98/118	N/A
Iron, Dissolved	28.1	N/A	3/4	N/A	3/4	N/A
Lead	0.0133	N/A	6/100	N/A	0/100	0/100
Magnesium	36.1	N/A	113/113	N/A	0/113	N/A
Magnesium, Dissolved	33.8	N/A	2/2	N/A	0/2	N/A
Manganese	4.53	N/A	96/112	N/A	79/112	N/A
Manganese, Dissolved	3.75	N/A	2/2	N/A	2/2	N/A
Molybdenum	0.00508	N/A	27/111	N/A	0/111	N/A
Nickel	0.595	N/A	23/118	N/A	10/118	N/A
Nickel, Dissolved	0.442	N/A	2/4	N/A	2/4	N/A
Phosphorous	0.18	N/A	2/2	N/A	2/2	N/A
Phosphorous, Dissolved	0.12	N/A	2/2	N/A	2/2	N/A
Potassium	12.3	N/A	90/113	N/A	N/A	N/A
Potassium, Dissolved	13	N/A	1/2	N/A	N/A	N/A
Selenium	0.0276	N/A	38/113	N/A	37/113	0/113
Selenium, Dissolved	0.0113	N/A	2/2	N/A	1/2	0/2
Silicon	14.4	N/A	1/1	N/A	N/A	N/A
Sodium	176	N/A	113/113	N/A	0/113	N/A
Sodium, Dissolved	87.3	N/A	2/2	N/A	0/2	N/A
Uranium	0.6	N/A	54/161	N/A	54/161	27/161
Uranium, Dissolved	0.51	N/A	12/72	N/A	12/72	4/72
Vanadium	0.038	N/A	3/111	N/A	3/111	N/A
Zinc	0.078	N/A	4/118	N/A	0/118	N/A
Radionuclides (pCi/L)						
Potassium-40	361	N/A	3/14	N/A	N/A	N/A
Radium-226	0.898	N/A	6/88	N/A	6/88	0/88
Radium-228	6.25	N/A	2/28	N/A	N/A	N/A
Radon-222	519	N/A	2/2	N/A	2/2	N/A
Technetium-99	269	N/A	51/146	N/A	51/146	0/146
Thorium-228	0.185	N/A	1/13	N/A	1/13	N/A
Thorium-230	0.86	N/A	1/70	N/A	1/70	N/A
Thorium-232	1.73	N/A	2/15	N/A	2/15	N/A
Thorium-234	1,010	N/A	3/48	N/A	N/A	N/A
Uranium	2,170	N/A	2/5	N/A	N/A	N/A

Table 4.49. SWMU 145 UCRS Groundwater Contaminants (Continued)

Analysis	Historical Data	RI	Frequency of Detection ^a	Frequency of Detection		
				Above Background	Above NAL	Above MCL
Uranium-234	840	N/A	3/5	N/A	3/5	N/A
Uranium-235	53.8	N/A	2/11	N/A	2/11	N/A
Uranium-238	1,270	N/A	5/7	N/A	5/7	N/A
PCBs (mg/L)						
PCB-1260	0.00007	N/A	1/40	N/A	1/40	0/40
Semivolatiles (mg/L)						
2-Methylnaphthalene	0.0006	N/A	1/10	N/A	0/10	N/A
3-Methylphenol	0.28	N/A	5/10	N/A	2/10	N/A
Bis(2-ethylhexyl)phthalate	0.015	N/A	9/10	N/A	3/10	2/10
Butyl benzyl phthalate	0.006	N/A	1/10	N/A	0/10	N/A
Di-n-butyl phthalate	0.012	N/A	8/10	N/A	0/10	N/A
Phenol	0.003	N/A	2/10	N/A	0/10	N/A
Volatiles (mg/L)						
1,3-Dimethylbenzene	0.008	N/A	4/10	N/A	0/10	0/10
2-Butanone	0.012	N/A	4/111	N/A	0/111	N/A
Acetone	0.22	N/A	7/107	N/A	5/107	N/A
Benzene	0.005	N/A	1/144	N/A	1/144	0/144
Chloroform	0.003	N/A	2/77	N/A	2/77	N/A
Ethylbenzene	0.002	N/A	1/144	N/A	0/144	0/144
Methylene chloride	0.013	N/A	5/111	N/A	1/111	1/111
Total Xylene	0.008	N/A	4/144	N/A	0/144	0/144
Trichloroethene	0.002	N/A	5/144	N/A	3/144	0/144

^a Frequency of detection is the number of detections of an analyte per number of analyses (includes regular and duplicate samples).

N/A = not applicable

Table 4.50. SWMU 145 RGA Groundwater Contaminants

Analysis	Historical Data	RI	Frequency of Detection ^a	Frequency of Detection		
				Above Background	Above NAL	Above MCL
<i>Metals (mg/L)</i>						
Aluminum	57.2	N/A	153/466	26/466	34/466	N/A
Aluminum, Dissolved	0.2	N/A	17/36	0/36	0/36	N/A
Antimony	0.01	N/A	3/407	0/407	3/407	3/407
Antimony, Dissolved	0.019	N/A	7/26	0/26	7/26	6/26
Arsenic	0.0246	N/A	173/232	34/232	173/232	16/232
Arsenic, Dissolved	0.0162	N/A	6/6	6/6	6/6	3/6
Barium	2.25	N/A	770/804	267/804	698/804	1/804
Barium, Dissolved	2.15	N/A	723/735	405/735	661/735	1/735
Beryllium	0.002	N/A	29/804	0/804	0/804	0/804
Beryllium, Dissolved	0.002	N/A	13/28	0/28	0/28	0/28
Boron	1.54	N/A	38/450	N/A	16/450	N/A
Boron, Dissolved	1.61	N/A	20/28	N/A	N/A	N/A
Cadmium	0.00111	N/A	12/791	0/791	4/791	0/791
Calcium	80	N/A	469/469	21/469	N/A	N/A
Calcium, Dissolved	64.4	N/A	39/39	6/39	N/A	N/A
Cerium	0.08	N/A	26/39	N/A	N/A	N/A
Cerium, Dissolved	0.08	N/A	13/26	N/A	N/A	N/A
Chromium	5.4	N/A	293/821	156/821	19/821	180/821
Chromium, Dissolved	0.198	N/A	29/504	1/504	0/504	1/504
Cobalt	0.574	N/A	189/804	37/804	13/804	N/A
Cobalt, Dissolved	0.01	N/A	15/41	0/41	0/41	N/A
Copper	0.178	N/A	48/804	14/804	7/804	0/804
Copper, Dissolved	0.256	N/A	16/41	14/41	1/41	0/41
Gallium	0.09	N/A	26/39	N/A	N/A	N/A
Gallium, Dissolved	0.09	N/A	13/26	N/A	N/A	N/A
Iron	117	N/A	600/803	139/803	447/803	N/A
Iron, Dissolved	0.3	N/A	14/58	13/58	0/58	N/A
Lead	0.0167	N/A	26/381	0/381	2/381	2/381
Lithium	0.08	N/A	26/39	N/A	26/39	N/A
Lithium, Dissolved	0.08	N/A	13/26	N/A	N/A	N/A
Magnesium	25.7	N/A	469/469	18/469	0/469	N/A
Magnesium, Dissolved	25.7	N/A	39/39	4/39	0/39	N/A
Manganese	36.5	N/A	369/469	177/469	260/469	N/A
Manganese, Dissolved	0.246	N/A	34/36	7/36	17/36	N/A
Mercury	0.0004	N/A	5/791	5/791	0/791	0/791
Mercury, Dissolved	0.0003	N/A	1/32	1/32	0/32	0/32
Molybdenum	0.117	N/A	115/450	8/450	57/450	N/A
Molybdenum, Dissolved	0.03	N/A	15/28	0/28	15/28	N/A
Nickel	1.89	N/A	351/804	30/804	282/804	N/A
Nickel, Dissolved	0.5	N/A	23/52	2/52	23/52	N/A
Potassium	87.5	N/A	417/469	28/469	N/A	N/A
Potassium, Dissolved	10	N/A	18/28	3/28	N/A	N/A
Selenium	0.0134	N/A	182/762	180/762	48/762	0/762
Silicon	7.13	N/A	2/2	N/A	N/A	N/A
Silicon, Dissolved	6.81	N/A	2/2	N/A	N/A	N/A
Silver	0.03	N/A	16/804	13/804	13/804	N/A
Silver, Dissolved	0.03	N/A	13/41	0/41	13/41	N/A

Table 4.50. SWMU 145 RGA Groundwater Contaminants (Continued)

Analysis	Historical Data	RI	Frequency of Detection ^a	Frequency of Detection		
				Above Background	Above NAL	Above MCL
Sodium	178	N/A	802/803	71/803	0/803	N/A
Sodium, Dissolved	71.8	N/A	65/65	2/65	0/65	N/A
Strontium	0.168	N/A	45/45	N/A	0/45	N/A
Strontium, Dissolved	0.167	N/A	32/32	N/A	N/A	N/A
Thorium	0.05	N/A	26/39	N/A	0/39	N/A
Thorium, Dissolved	0.05	N/A	13/26	N/A	N/A	N/A
Titanium	0.12	N/A	27/39	N/A	0/39	N/A
Titanium, Dissolved	0.06	N/A	13/26	N/A	N/A	N/A
Uranium	0.00417	N/A	18/791	6/791	18/791	0/791
Uranium, Dissolved	0.00146	N/A	3/211	0/211	3/211	0/211
Vanadium	0.219	N/A	81/806	3/806	74/806	N/A
Vanadium, Dissolved	0.142	N/A	43/63	2/63	43/63	N/A
Zinc	0.46	N/A	72/804	13/804	1/804	N/A
Zinc, Dissolved	0.186	N/A	17/43	15/43	0/43	N/A
Zirconium	0.02	N/A	26/39	N/A	0/39	N/A
Zirconium, Dissolved	0.02	N/A	13/26	N/A	N/A	N/A
Radionuclides (pCi/L)						
Iodine-131	164	N/A	1/407	N/A	N/A	N/A
Potassium-40	47.2	N/A	1/1	N/A	N/A	N/A
Radium	0.541	N/A	3/160	N/A	N/A	N/A
Radium-224	0.313	N/A	1/102	N/A	N/A	N/A
Radium-226	2.68	N/A	36/338	7/338	36/338	0/338
Radium-228	13.2	N/A	8/91	N/A	N/A	N/A
Strontium-90	11.2	N/A	2/407	N/A	2/407	N/A
Technetium-99	621	N/A	328/784	204/784	266/784	0/784
Thorium-230	0.665	N/A	3/186	0/186	3/186	N/A
Thorium-232	0.468	N/A	1/1	N/A	1/1	N/A
Thorium-234	4,490	N/A	8/143	N/A	N/A	N/A
Pesticides/PCBs (mg/L)						
Heptachlor	0.0000059	N/A	1/32	N/A	0/32	0/32
PCB, Total	0.00787	N/A	25/149	N/A	33/149	8/149
PCB-1016	0.001184	N/A	16/163	N/A	16/163	3/163
PCB-1242	0.00787	N/A	10/164	N/A	9/164	5/164
PCB-1260	0.00017	N/A	1/162	N/A	1/162	0/162
Semivolatiles (mg/L)						
1,2-Dichlorobenzene	0.000057	N/A	1/796	N/A	0/796	0/796
1,4-Dichlorobenzene	0.000062	N/A	1/796	N/A	0/796	0/796
3-Methylphenol	1.4	N/A	5/31	N/A	5/31	N/A
Bis(2-	0.002	N/A	1/40	N/A	0/40	N/A
Bis(2-ethylhexyl)phthalate	0.007	N/A	21/40	N/A	4/40	1/40
Butyl benzyl phthalate	0.006	N/A	1/33	N/A	0/33	N/A
Chrysene	0.0006	N/A	1/39	N/A	0/39	N/A
Diethyl phthalate	0.0006	N/A	3/40	N/A	0/40	N/A
Dimethyl phthalate	0.001	N/A	1/40	N/A	0/40	N/A
Di-n-butyl phthalate	0.038	N/A	19/40	N/A	0/40	N/A

Table 4.50. SWMU 145 RGA Groundwater Contaminants (Continued)

Analysis	Historical Data	RI	Frequency of Detection ^a	Frequency of Detection		
				Above Background	Above NAL	Above MCL
<i>Volatiles (mg/L)</i>						
1,1,1-Trichloroethane	0.004	N/A	1/793	N/A	0/793	0/793
1,2-Dichloroethene	0.001	N/A	1/4	N/A	0/4	N/A
1,3,5-Trimethylbenzene	0.0002	N/A	1/2	N/A	0/2	N/A
1,3-Dimethylbenzene	0.005	N/A	5/24	N/A	0/24	0/24
1,4-Dioxane	0.0006	N/A	3/34	N/A	0/34	N/A
2-Butanone	0.01	N/A	18/773	N/A	0/773	N/A
Acetone	5.8	N/A	40/744	N/A	6/744	N/A
Benzene	0.00014	N/A	1/792	N/A	0/792	0/792
Bromomethane	0.000062	N/A	2/793	N/A	0/793	N/A
Carbon disulfide	0.002	N/A	1/793	N/A	0/793	N/A
Chlorobenzene	0.001	N/A	3/793	N/A	0/793	0/793
Chloroform	0.004	N/A	11/222	N/A	11/222	N/A
Chloromethane	0.00018	N/A	12/793	N/A	0/793	N/A
<i>cis</i> -1,2-Dichloroethene	0.00053	N/A	11/800	N/A	0/800	0/800
Dichlorodifluoromethane	0.000074	N/A	1/708	N/A	0/708	N/A
Ethanol	0.35	N/A	3/493	N/A	N/A	N/A
Ethylbenzene	0.001	N/A	3/793	N/A	0/793	0/793
<i>m,p</i> -Xylene	0.00015	N/A	3/13	N/A	N/A	0/13
Methylene chloride	0.014	N/A	18/793	N/A	2/793	2/793
Tetrachloroethene	0.002	N/A	4/793	N/A	2/793	0/793
Toluene	0.002	N/A	12/793	N/A	0/793	0/793
Total Xylene	0.005	N/A	7/793	N/A	0/793	0/793
<i>trans</i> -1,2-Dichloroethene	0.000054	N/A	1/800	N/A	0/800	0/800
<i>trans</i> -1,3-Dichloropropene	0.00017	N/A	1/793	N/A	N/A	N/A
Trichloroethene	0.033	N/A	466/814	N/A	382/814	215/814
Trichlorofluoromethane	0.00064	N/A	5/791	N/A	0/791	N/A

^a Frequency of detection is the number of detections of an analyte per number of analyses (includes regular and duplicate samples).

N/A = not applicable

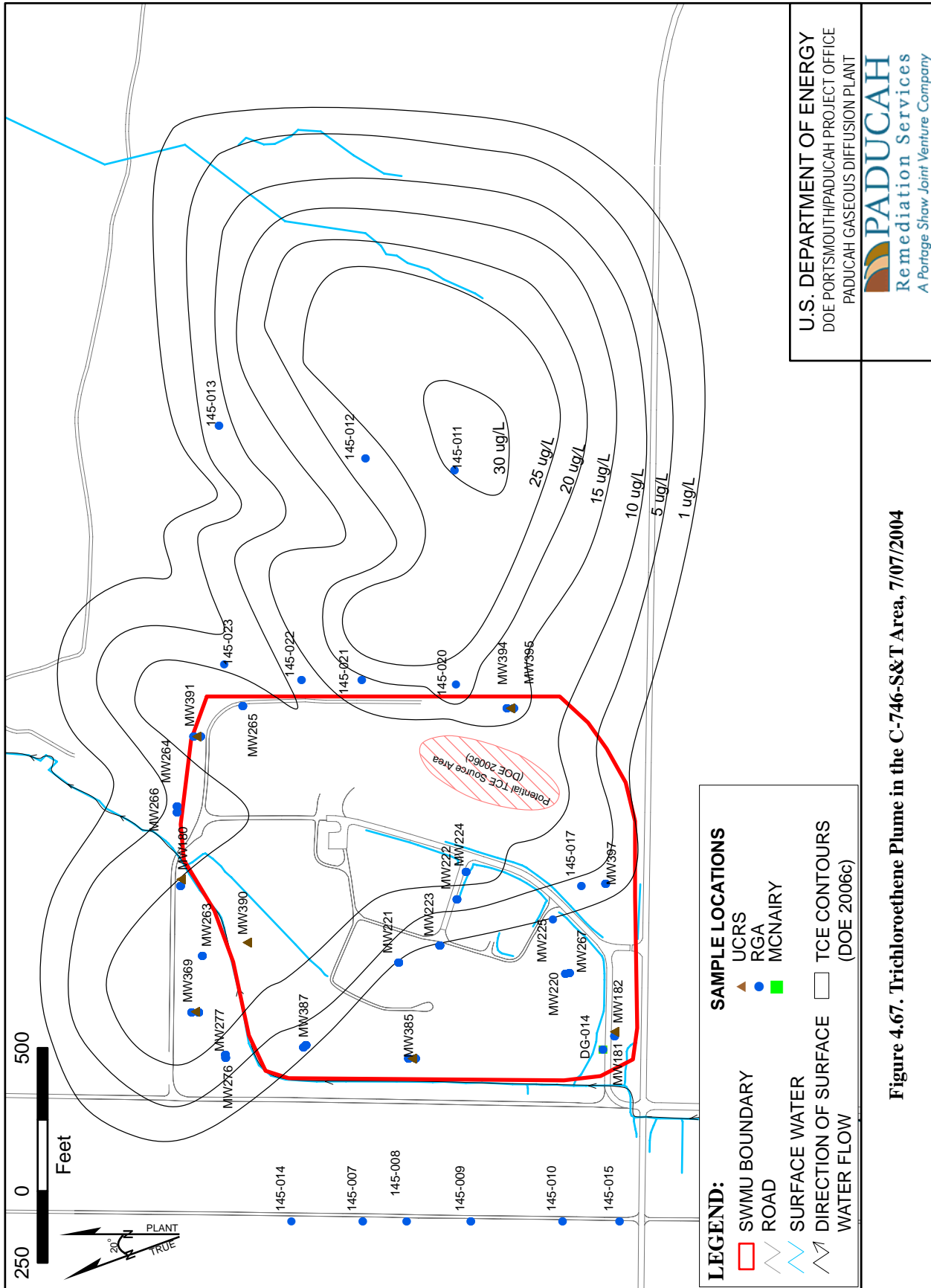
Arsenic, chromium, iron, and manganese were detected above background levels in RGA groundwater at a frequency of over 10%. A summer 2004 SI of the SWMU 145 area (DOE 2006c) assessed whether TCE in RGA groundwater in the area was leaching from the site landfills or originated from upgradient sources. The field investigation measured water levels in 13 RGA wells and sampled 18 temporary borings to characterize the lithology and dissolved VOC levels of the RGA in the SWMU 145 area. This SI postulated the presence of a small UCRS TCE source in SWMU 145 (Figure 4.67); an upgradient source was not responsible for the RGA TCE contamination in the SWMU 145 area wells. The assumed location of the source was based on TCE concentrations in the upper, middle, and lower RGA from established MWs and temporary borings (Figure 4.67). RGA groundwater flow directions (Figure 4.68) were used to approximate the general location of a UCRS source that would result in the observed RGA TCE distribution in the borings and wells. Because of the low concentrations, this source is not considered to be a TCE DNAPL source. Subsequent and ongoing groundwater monitoring conducted for the downgradient C-746-U Landfill demonstrates that TCE concentrations in RGA groundwater in the area remain low and relatively stable. TCE concentrations in MW394 (URGA) and MW395 (LRGA), located downgradient of the postulated source area (Figure 4.68), have shown a seasonally cyclic but consistent downward trend, from concentrations that ranged from 10 to 20 µg/L in 1994 to concentrations that ranged from 5 to 10 µg/L in 2008 and 2009 monitoring data. TCE concentrations in MW391 (URGA) and MW392 (LRGA), located at the northern boundary of SWMU 145 (Figure 4.68) have shown a seasonally cyclic but consistent upward trend, from concentrations that ranged from 5 to 10 µg/L in 1994 to concentrations that range from 15 to 20 µg/L in recent data. However, the most recent TCE data from these wells (2008-2009) have shown stable or slightly decreasing TCE concentrations. The reported TCE concentrations at MW391/MW392 are the highest recorded in the area. Other RGA monitoring wells in the area, located primarily to the north or along the western edge of SWMU 145, have shown mostly stable or decreasing TCE concentrations, some just slightly above, but most below, the MCL of 5 µg/L.

Seven RGA MWS of the C-746-S&T Landfills have produced samples with PCB contamination. The highest detected levels have been 0.001 mg/L PCB-1016 and 0.008 mg/L PCB-1242.

Strontium-90 was the only radionuclide to exceed screening criteria (in 2 of 408 analyses) in the RGA samples for SWMU 145, which may represent false positives. Documentation of strontium and cesium at PGDP in *Study of Plutonium and Fission Products*, KYL-20, July 1995, identified cesium-137 and strontium-89; and *Historical Impact of Reactor Tails on the Paducah Cascade*, KY/L-1239, March 1984, identified cesium-137 and strontium-90 in feed plant ash.

While the quantity of strontium-90 that came to Paducah from other sites (Hanford, Savannah River, and others) is a trace quantity, it has been seen in samples collected from C-410 ash. It is logical to assume that wherever cesium-137 was found, strontium-90 could be found as well. Cesium-137 and strontium-90 have half-lives of 30.2 years and 28.8 years, respectively, and still are being found at the site, while other fission products with shorter half-lives are detected less frequently [if at all at their minimum detectable concentration (MDC)] and at lower concentrations.

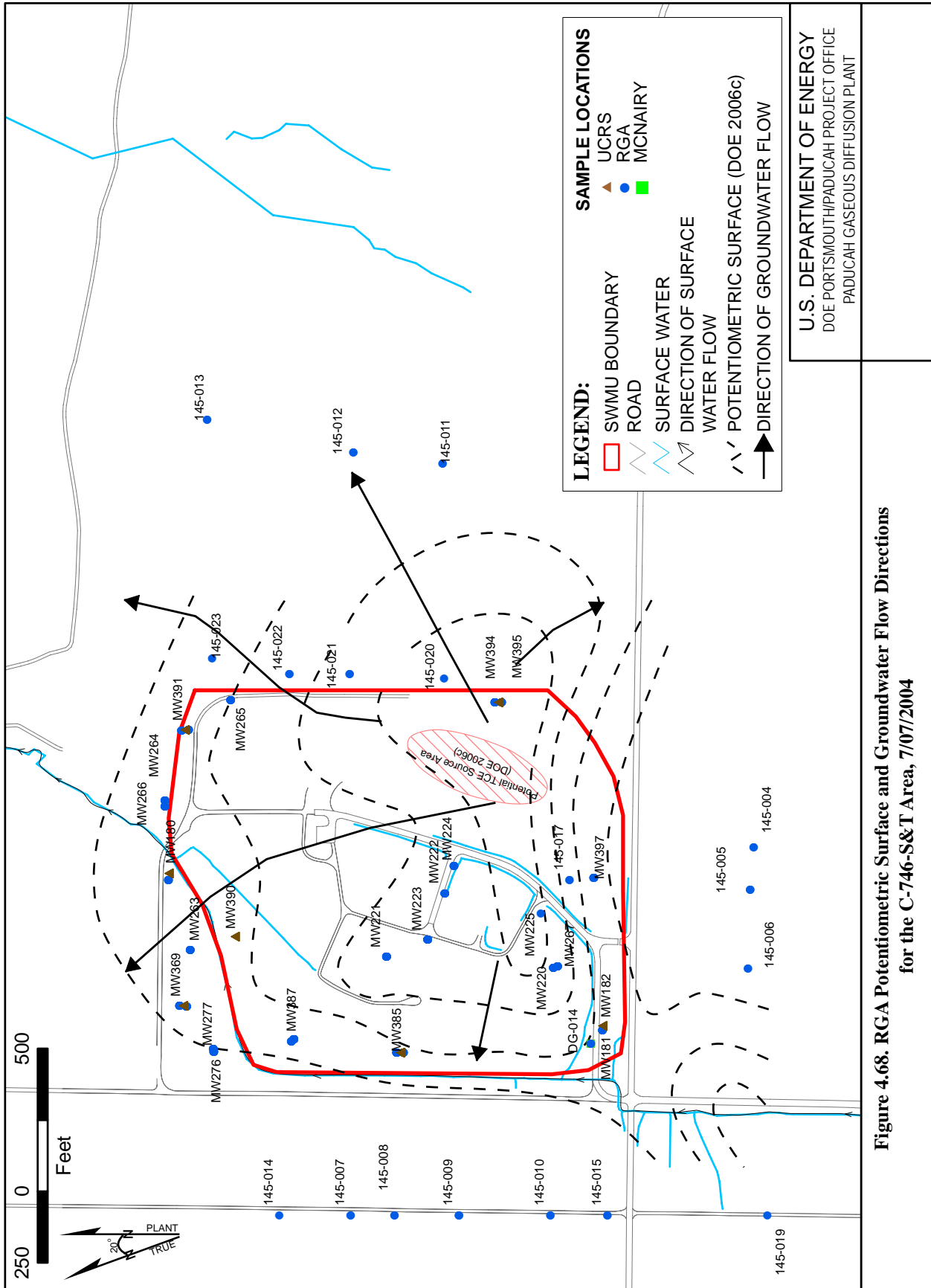
Table 4.51 shows the locations of all SWMU 145 groundwater contaminants.



U.S. DEPARTMENT OF ENERGY
DOE PORTSMOUTH/PADUCAH PROJECT OFFICE
PADUCAH GASEOUS DIFFUSION PLANT



Figure No. BGOUISWU145source.apr
DATE 08-05-09



U.S. DEPARTMENT OF ENERGY
DOE PORTSMOUTH/PADUCAH PROJECT OFFICE
PADUCAH GASEOUS DIFFUSION PLANT

Figure No. BGOUISWMMU145_wtrflw.apr
DATE 08-05-09

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Table 4.51. SWMU 145 Locations of Groundwater Contaminants

Hydro Unit	Depth Interval	Analysis	Child Resident NAL	Bkgd	MCL	MW180	MW182	MW371	MW386	MW390	MW393	MW396
	20	Metals (mg/L)										
		Aluminum	1.49	N/A	N/A		3.47					
		Aluminum, Dissolved	1.49	N/A	N/A		0.346					
		Arsenic	0.000035	N/A	0.01		0.01					
		Barium	0.104	N/A	2		0.897					
		Barium, Dissolved	0.104	N/A	2		0.816					
		Beryllium	0.00264	N/A	0.004		0.0011					
		Boron	0.136	N/A	N/A		0.07					
		Calcium	N/A	N/A	N/A		165					
		Calcium, Dissolved	N/A	N/A	N/A		121					
		Chromium	1.76	N/A	0.1		0.01					
		Cobalt	0.0906	N/A	N/A		0.01					
		Copper	0.0557	N/A	1.3		0.03					
		Iron	0.449	N/A	N/A		26.9					
		Iron, Dissolved	0.449	N/A	N/A		28.1					
		Lead	0.015	N/A	0.015		0.003U					
		Magnesium	N/A	N/A	N/A		36.1					
		Magnesium, Dissolved	N/A	N/A	N/A		33.8					
		Manganese	0.035	N/A	N/A		4.53					
		Manganese, Dissolved	0.035	N/A	N/A		3.75					
		Molybdenum	0.00753	N/A	N/A		0.04U					
		Nickel	0.0301	N/A	N/A		0.102					
		Nickel, Dissolved	0.0301	N/A	N/A		0.05U					
		Phosphorous	0.00003	N/A	N/A		0.18					
		Phosphorous, Dissolved	0.00003	N/A	N/A		0.12					
		Potassium	N/A	N/A	N/A		12.3					
		Potassium, Dissolved	N/A	N/A	N/A		13					
		Selenium	0.00754	N/A	0.05		0.005U					
		Silicon	N/A	N/A	N/A		14.4					
		Sodium	N/A	N/A	N/A		44					
		Sodium, Dissolved	N/A	N/A	N/A		40.3					
		Uranium	0.000906	N/A	0.03		0.6					
		Uranium, Dissolved	0.000906	N/A	0.03		0.51					
		Vanadium	0.00925	N/A	N/A		0.01					
		Zinc	0.45	N/A	N/A		0.047					
	27-32	Metals (mg/L)										
		Aluminum	1.49	N/A	N/A	0.2U		2.24	1.26			
		Aluminum, Dissolved	1.49	N/A	N/A	0.2U						
		Arsenic	0.000035	N/A	0.01		0.00112	0.00271				
		Barium	0.104	N/A	2	0.214	0.145	0.226				
		Barium, Dissolved	0.104	N/A	2	0.17	0.143	0.203				
		Beryllium	0.00264	N/A	0.004	0.015U	0.005U	0.005U				
		Boron	0.136	N/A	N/A		2U	2U				
		Calcium	N/A	N/A	N/A	49.1	32.6	30.5				
		Calcium, Dissolved	N/A	N/A	N/A	45.1	0.025U	0.2U				
		Chromium	1.76	N/A	0.1	0.197	0.025U	0.00918				
		Cobalt	0.0906	N/A	N/A	0.045U	0.025U	0.00918				
		Copper	0.0557	N/A	1.3	0.031	0.05U	0.0219				
		Iron	0.449	N/A	N/A	11.3	5.81	11.6				
		Iron, Dissolved	0.449	N/A	N/A	1.29						
		Lead	0.015	N/A	0.015		0.005U	0.00549				
		Magnesium	N/A	N/A	N/A	13.5	12.9	12.4				
		Magnesium, Dissolved	N/A	N/A	N/A	12.3						
	27-32	Radionuclides (pCi/L)										
		Uranium-238	N/A	N/A	N/A		0.163					
		Potassium-40	N/A	N/A	N/A		254					
		Radon-222	0.866	N/A	N/A		351					
		Technetium-99	14	N/A	900		47.8					
		Thorium-228	0.129	N/A	N/A		0.165					
		Thorium-230	0.424	N/A	N/A		0.594U					
		Thorium-232	0.382	N/A	N/A		0.365U					
		Thorium-234	N/A	N/A	N/A		156U					
		Uranium	N/A	N/A	N/A		2170					
		Uranium-234	0.546	N/A	N/A		840					
		Uranium-235	0.538	N/A	N/A		53.8					
		Uranium-238	0.443	N/A	N/A		1270					
		Volatiles (mg/L)										
		2-Butanone	0.0868	N/A	N/A		0.005U					
		Acetone	0.0275	N/A	N/A		0.01U					
		Benzene	0.000385	N/A	0.005		0.005					
		Chloroform	0.000287	N/A	N/A		0.005U					
		Ethylbenzene	0.00468	N/A	0.7		0.005U					
		Methylene chloride	0.00426	N/A	0.005		0.001U					
		Total Xylene	0.0653	N/A	10		0.015U					
		Trichloroethene	0.0016	N/A	0.005		0.001					
	27-32	Metals (mg/L)										
		Manganese	0.035	N/A	N/A		0.035	0.074	1.02			
		Manganese, Dissolved	0.035	N/A	N/A		0.035	0.193	0.00165			
		Molybdenum	0.00753	N/A	N/A		0.00753	0.0124	0.0108			
		Nickel	0.0301	N/A	N/A		0.0301	0.442				
		Nickel, Dissolved	0.0301	N/A	N/A		0.0301	0.08				
		Phosphorous, Dissolved	0.00003	N/A	N/A		0.00003	0.587	0.999			
		Potassium	N/A	N/A	N/A		N/A	10.5U				
		Potassium, Dissolved	N/A	N/A	N/A		N/A	10.5U				
		Selenium	0.00754	N/A	0.05	0.005U	0.00754	0.01U	0.01U			
		Sodium	N/A	N/A	N/A	96.3	N/A	144	123			
		Sodium, Dissolved	N/A	N/A	N/A	87.3	N/A					
		Total Metals	N/A	N/A	N/A	5U	N/A					
		Uranium	0.000906	N/A	0.03	0.00522	0.000906	0.027	0.00127			
		Uranium, Dissolved	0.000906	N/A	0.03	0.002	0.000906	0.00134	0.001U			
		Vanadium	0.00925	N/A	N/A		0.00925	0.025U	0.2U			
		Zinc	0.45	N/A	N/A	0.078	0.45	0.0376	0.2U			

Table 4.51. SWMU 145 Locations of Groundwater Contaminants (Continued)

Hydro Unit	Depth Interval	Analysis	Child Resident NAL	Bkgd	MCL	DG-014	MW179	MW181	MW220	MW221	MW222	MW223	MW224	MW263	MW264	MW265	MW266	MW267	MW276	MW277	MW369	MW370	MW384	MW385	MW387	MW388	MW391	MW392	MW394	MW395	MW397					
	50-52	Metals (mg/L)																																		
		Aluminum	1.49	2.189	N/A																0.927															
		Antimony	0.000564	0.06	0.006																0.005U															
		Arsenic	0.000035	0.005	0.01																0.0138															
		Arsenic, Dissolved	0.000035	0.005	0.01																0.0128															
		Barium	0.104	0.235	2																0.486															
		Barium, Dissolved	0.104	0.2	2																0.462															
		Beryllium	0.00264	0.004	0.004																0.005U															
		Boron	0.136	N/A	N/A																2U															
		Cadmium	0.000661	0.01	0.005																0.005U															
		Calcium	N/A	41.238	N/A																29.8															
		Chromium	1.76	0.144	0.1																0.025U															
		Chromium, Dissolved	1.76	0.05	0.1																0.1U															
		Cobalt	0.0906	0.045	N/A																0.0794															
		Copper	0.0557	0.036	1.3																0.05U															
		Iron	0.449	5.03	N/A																21.5															
		Lead	0.015	0.129	0.015																0.005U															
		Magnesium	N/A	16.262	N/A																12															
		Manganese	0.035	0.119	N/A																0.867															
		Mercury	0.000444	0.0002	0.002																0.0002U															
		Molybdenum	0.00753	0.05	N/A																0.025U															
		Nickel	0.0301	0.682	N/A																0.0168															
		Potassium	N/A	5.195	N/A																2.21															
		Selenium	0.00754	0.005	0.05																0.00571															
		Silver	0.0075	0.011	N/A																0.025U															
		Sodium	N/A	59.45	N/A																67.9															
		Uranium	0.000906	0.002	0.03																0.001U															
		Uranium, Dissolved	0.000906	0.002	0.03																0.001U															
		Vanadium	0.000925	0.134	N/A																0.027															
		Zinc	0.45	0.054	N/A																0.0239															
		Pesticides/PCBs (mg/L)																																		
		Heptachlor	0.0000114	N/A	0.0004																0.000059															
		PCB, Total	0.0000793	N/A	0.0005																0.00115															
		PCB-1016	0.0000468	N/A	0.0005																0.000605															
		PCB-1242	0.000123	N/A	0.0005																0.00115															
		PCB-1260	0.0000428	N/A	0.0005																0.001U															
		Radionuclides (pCi/L)																																		
		Iodine-131	N/A	N/A	N/A																30U															
		Radium	N/A	N/A	N/A																0.781U															
		Radium-224	N/A	N/A	N/A																1.74U															
		Radium-226	0.1	0.6	5																0.184															
		Radium-228	N/A	N/A	N/A																7.45															
		Strontium-90	0.522	N/A	N/A																18.2U															
		Technetium-99	14	22.3	900																53.1															
		Thorium-230	0.424	1.1	N/A																1.36U															
		Thorium-234	N/A	N/A	N/A																483U															
		Semivolatile (mg/L)																																		
		1,2-Dichlorobenzene	0.0166	N/A	0.6																0.005U															
		1,4-Dichlorobenzene	0.000578	N/A	0.075																0.005U															
		3-Methylphenol	0.0725	N/A	N/A																0.01U															
		Bis(2-chloroethoxy)methane	N/A	N/A	N/A																0.01U															
		Bis(2-ethylhexyl)phthalate	0.00312	N/A	0.006																0.002															
		Butyl benzyl phthalate	0.259	N/A	N/A																0.01U															
		Chrysene	0.00132	N/A	N/A																0.01U															
		Diethyl phthalate	1.2	N/A	N/A																0.01U															
		Dimethyl phthalate	15.1	N/A	N/A																0.001															
		Di-n-butyl phthalate	0.129	N/A	N/A																0.01U															

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Table 4.51. SWMU 145 Locations of Groundwater Contaminants (Continued)

Hydro Unit	Depth Interval	Analysis	MCL	Bkgd	Child Resident NAL	DG-014	MW179	MW181	MW220	MW221	MW222	MW223	MW224	MW263	MW264	MW265	MW266	MW267	MW276	MW277	MW369	MW370	MW384	MW385	MW387	MW388	MW391	MW392	MW394	MW395	MW397
	50-52	Volatiles (mg/L)																													
		1,1,1-Trichloroethane	0.2	N/A	0.0335	0.001U	0.2														0.005U										
		1,2-Dichloroethane	N/A	N/A	0.00247		N/A														0.002U										
		1,3,5-Trimethylbenzene	N/A	N/A	0.000567		N/A														0.001U										
		1,3-Dimethylbenzene	10	N/A	0.439		10														0.005U										
		1,4-Dioxane	N/A	N/A	0.00479		N/A														0.2U										
		2-Butanone	N/A	N/A	0.0868		N/A														0.05U										
		Acetone	N/A	N/A	0.0275		N/A														5.8										
		Benzene	0.005	N/A	0.000385		0.005														0.005U										
		Bromomethane	N/A	N/A	0.000391		N/A														0.01U										
		Carbon disulfide	N/A	N/A	0.0457		N/A														0.005U										
		Chlorobenzene	0.1	N/A	0.00466		0.1														0.005U										
		Chloroform	N/A	N/A	0.000287		N/A														0.001										
		Chloromethane	N/A	N/A	0.00167		N/A														0.01U										
		cis-1,2-Dichloroethene	0.07	N/A	0.00273		0.07														0.005U										
		Dichlorodifluoromethane	N/A	N/A	0.018		N/A														0.01U										
		Ethanol	N/A	N/A	N/A		N/A														5U										
		Ethylbenzene	0.7	N/A	0.00468		0.7														0.005U										
		m,p-Xylene	10	N/A	N/A		10														0.002U										
		Methylene chloride	0.005	N/A	0.00426		0.005														0.01U										
		Tetrachloroethene	0.005	N/A	0.000582		0.005														0.005U										
		Toluene	1	N/A	0.0338		1														0.005U										
		Total Xylene	10	N/A	0.0653		10														0.015U										
		trans-1,2-Dichloroethene	0.1	N/A	0.00548		0.1														0.005U										
		trans-1,3-Dichloropropene	N/A	N/A	N/A		N/A														0.005U										
		Trichloroethene	0.005	N/A	0.0016		0.005														0.016										
		Trichlorofluoromethane	N/A	N/A	0.0577		N/A														0.005U										
	55-57	Metals (mg/L)																													
		Aluminum	N/A	2.189	1.49		0.2	11.9							5.2																
		Aluminum, Dissolved	N/A	0.311	1.49		0.2	0.2							0.2																
		Antimony	0.006	0.06	0.000564		0.005U	0.006U							0.005U																
		Antimony, Dissolved	0.006	0.06	0.000564		0.011	0.006U							0.008																
		Arsenic	0.01	0.005	0.000035		0.01	0.0026																							
		Barium	2	0.235	0.104		0.03	0.461																							
		Barium, Dissolved	2	0.2	0.104		0.0307	0.176																							
		Beryllium	0.004	0.004	0.00264		0.002	0.002							0.002																
		Beryllium, Dissolved	0.004	0.004	0.00264		0.002	0.002							0.002																
		Boron	N/A	N/A	0.136		1.54	0.03							0.229																
		Boron, Dissolved	N/A	N/A	N/A		1.61	0.03							0.17																
		Cadmium	0.005	0.01	0.000661		0.005U	0.0007							0.005U																
		Calcium	N/A	41.238	N/A		80	33.2							47.4																
		Calcium, Dissolved	N/A	38.166	N/A		64.4	34.3							35.3																
		Cerium	N/A	N/A	N/A		0.08	0.08							0.08																
		Cerium, Dissolved	N/A	N/A	N/A		0.08	0.08							0.08																
		Chromium	0.1	0.144	1.76		0.06	3.18							0.03																
		Chromium, Dissolved	0.1	0.05	1.76		0.03	0.198							0.03																
		Cobalt	N/A	0.045	0.0906		0.01	0.026							0.01																
		Cobalt, Dissolved	N/A	0.045	0.0906		0.01	0.01							0.01																
		Copper	1.3	0.036	0.0557		0.03	0.03							0.03																
		Copper, Dissolved	1.3	0.02	0.0557		0.03	0.03							0.03																
		Gallium	N/A	N/A	N/A		0.09	0.09							0.09																
		Gallium, Dissolved	N/A	N/A	N/A		0.09	0.09							0.09																
		Iron	N/A	5.03	0.449		2.25	46.2							5.8																
		Iron, Dissolved	N/A	2.67	0.449		0.3	0.3							0.3																
		Lead	0.015	0.129	0.0302		0.005U	0.0092							0.005																
		Lithium	N/A	N/A	N/A		0.08	0.08							0.08																
		Lithium, Dissolved	N/A	N/A	N/A		0.08	0.08							0.08																
		Magnesium	N/A	16.262	N/A		24.9	13.2							16.4																
		Magnesium, Dissolved	N/A	16.215	N/A		23.5	12.7							11.5																

RGA

Table 4.51. SWMU 145 Locations of Groundwater Contaminants (Continued)

Hydro Unit	Depth Interval	Analysis	MCL	Bkgd	Child Resident NAL	DG-014	MW179	MW181	MW220	MW221	MW222	MW223	MW224	MW263	MW264	MW265	MW266	MW267	MW276	MW277	MW369	MW370	MW384	MW385	MW387	MW388	MW391	MW392	MW394	MW395	MW397				
55-57		Volatiles (mg/L)																																	
		1,1,1-Trichloroethane	0.2	N/A	0.0335	0.005U	0.005U	0.005U								0.005U									0.005U										
		1,3-Dimethylbenzene	10	N/A	0.439												0.005U								0.005U										
		1,4-Dioxane	N/A	N/A	0.00479																				0.0006										
		2-Butanone	N/A	N/A	0.0868	0.004	0.003										0.01U								0.05U										
		Acetone	N/A	N/A	0.0275	0.007	0.016										0.01U								0.004										
		Benzene	N/A	0.005	0.000385	0.005U	0.00055										0.005U								0.005U										
		Bromomethane	N/A	N/A	0.000391	0.005U	0.00055										0.005U								0.01U										
		Carbon disulfide	N/A	N/A	0.0457	0.005U	0.005U										0.005U								0.005U										
		Chlorobenzene	N/A	0.1	0.00466	0.001	0.005U										0.005U								0.005U										
		Chloroform	N/A	N/A	0.000287	0.0005U	0.00076										0.00061								0.005U										
		Chloromethane	N/A	N/A	0.00167	0.00057	0.00018										0.00009								0.01U										
		cis-1,2-Dichloroethene	N/A	0.07	0.00273	0.00012	0.005U										0.00014								0.005U										
		Dichlorodifluoromethane	N/A	N/A	0.018	0.005U	0.005U										0.005U								0.01U										
		Ethanol	N/A	N/A	N/A	0.25U	0.25U										0.25U								5U										
		Ethylbenzene	N/A	0.7	0.00468	0.001	0.005U										0.005U								0.005U										
		m,p-Xylene	N/A	10	N/A	N/A	0.00046	0.0001U									0.0001U								0.005U										
		Methylene chloride	N/A	0.005	0.00426	0.01U	0.00055										0.00053								0.005U										
		Tetrachloroethene	N/A	0.005	0.000582	0.002	0.005U										0.00077								0.005U										
		Toluene	N/A	1	0.0338	0.001	0.00023										0.0006								0.005U										
		Total Xylene	N/A	10	0.0653	0.003	0.015U										0.015U								0.015U										
		trans-1,2-Dichloroethene	N/A	0.1	0.00548	0.005U	0.005U										0.005U								0.005U										
		trans-1,3-Dichloropropene	N/A	N/A	N/A	N/A	0.005U										0.005U								0.005U										
		Trichloroethene	N/A	0.005	0.0016	0.009	0.001										0.008								0.007										
		Trichlorofluoromethane	N/A	N/A	0.0577	0.005U	0.005U										0.0003								0.005U										
		58-60	RGA	Metals (mg/L)																															
				Aluminum	N/A	2.189	1.49											0.2								0.42									
Aluminum, Dissolved	N/A			0.311	1.49											0.2								0.005U											
Antimony	0.006			0.06	0.000564											0.006U								0.005U											
Antimony, Dissolved	0.006			0.06	0.000564											0.006U								0.005U											
Arsenic	0.01			0.005	0.000035												0.01							0.01											
Arsenic, Dissolved	0.01			0.005	0.000035												0.01							0.01											
Barium	2			0.235	0.104												0.081							0.47											
Barium, Dissolved	2			0.2	0.104											0.082								0.272											
Beryllium	0.004			0.004	0.00264											0.002								0.02U											
Beryllium, Dissolved	0.004			0.004	0.00264											0.002								0.02U											
Boron	N/A			N/A	0.136											0.81								2U											
Boron, Dissolved	N/A			N/A	N/A											0.73								2U											
Cadmium	0.005			0.01	0.000661											0.005U								0.005U											
Calcium	N/A			41.238	N/A											41.3								31											
Calcium, Dissolved	N/A			38.166	N/A											39.8								31											
Cerium	N/A			N/A	N/A											0.08								2.24											
Cerium, Dissolved	N/A			N/A	N/A											0.08								2.24											
Chromium	0.1			0.144	1.76											3.44								2.24											
Chromium, Dissolved	0.1			0.05	1.76											0.03								0.05U											
Cobalt	N/A			0.045	0.0906											0.01								0.028											
Cobalt, Dissolved	N/A			0.045	0.0906											0.01								0.028											
Copper	1.3			0.036	0.0557											0.032								0.102											
Copper, Dissolved	1.3			0.02	0.0557											0.032								0.102											
Gallium	N/A			N/A	N/A											0.09																			
Gallium, Dissolved	N/A			N/A	N/A											0.09																			
Iron	5.03			0.267	0.449											9.12								20.2											
Iron, Dissolved	N/A	0.267	0.449											9.12								20.2													
Lead	0.015	0.129	0.015											0.3								0.003U													
Lead, Dissolved	0.015	0.129	0.015											0.3								0.003U													
Lithium	N/A	N/A	0.0302											0.08																					
Lithium, Dissolved	N/A	N/A	N/A											0.08																					
Magnesium	N/A	16.262	N/A											16.8								12.1													
Magnesium, Dissolved	N/A	16.215	N/A											16.1								12.1													

Table 4.51. SWMU 145 Locations of Groundwater Contaminants (Continued)

Hydro Unit	Depth Interval	Analysis	MCL	Bkgd	Child Resident NAL	DG-014	MW179	MW181	MW220	MW221	MW222	MW223	MW224	MW263	MW264	MW265	MW266	MW267	MW276	MW277	MW369	MW370	MW384	MW385	MW387	MW388	MW391	MW392	MW394	MW395	MW397					
58-60		Volatiles (mg/L)																																		
		1,1,1-Trichloroethane	N/A	0.2	0.0335															0.005U				0.005U												
		1,2-Dichloroethane	N/A	N/A	0.00247															0.01U																
		1,3-Dimethylbenzene	N/A	10	0.439																															
		1,4-Dioxane	N/A	N/A	0.00479																0.15U															
		2-Butanone	N/A	N/A	0.0868																0.01U															
		Acetone	N/A	N/A	0.0275																0.01U															
		Benzene	N/A	0.005	0.000385																0.005U															
		Bromomethane	N/A	N/A	0.000391																0.005U															
		Carbon disulfide	N/A	N/A	0.0457																0.005U															
		Chlorobenzene	N/A	0.1	0.00466																0.005U															
		Chloroform	N/A	N/A	0.000287																0.005U															
		Chloromethane	N/A	N/A	0.00167																0.005U															
		cis-1,2-Dichloroethene	N/A	N/A	0.00273																0.005U															
		Dichlorodifluoromethane	N/A	N/A	0.018																0.005U															
		Ethanol	N/A	N/A	N/A																0.25U															
		Ethylbenzene	N/A	N/A	0.00468																0.005U															
		m,p-Xylene	N/A	10	N/A																0.005U															
		Methylene chloride	N/A	0.005	0.00426																0.01U															
		Tetrachloroethene	N/A	0.005	0.000582																0.005U															
		Toluene	N/A	1	0.0338																0.005U															
		Total Xylene	N/A	10	0.0653																0.015U															
		trans-1,2-Dichloroethene	N/A	0.1	0.00548																0.005U															
		trans-1,3-Dichloropropene	N/A	N/A	N/A																0.005U															
		Trichloroethene	N/A	0.005	0.0016																0.029															
		Trichlorofluoromethane	N/A	N/A	0.0577																0.005U															
		65-68		Metals (mg/L)																																
				Aluminum	N/A	2.189	1.49																													
				Antimony	N/A	0.06	0.000564																													
				Arsenic	N/A	0.01	0.000035																													
				Barium	N/A	2	0.104																													
				Barium, Dissolved	N/A	2	0.104																													
				Beryllium	N/A	0.004	0.00264																													
Boron	N/A			N/A	0.136																															
Cadmium	N/A			0.005	0.000661																															
Calcium	N/A			41.238	N/A																															
Chromium	N/A			0.1	1.76																															
Chromium, Dissolved	N/A			0.1	1.76																															
Cobalt	N/A			N/A	0.0906																															
Copper	N/A			1.3	0.0557																															
Iron	N/A			5.03	0.449																															
Lead	N/A			0.015	0.015																															
Magnesium	N/A			N/A	16.262																															
Manganese	N/A			N/A	0.035																															
Mercury	N/A			0.002	0.000444																															
Molybdenum	N/A			N/A	0.00753																															
Nickel	N/A			N/A	0.0301																															
Potassium	N/A			N/A	5.195																															
Selenium	N/A			0.05	0.00754																															
Silver	N/A			N/A	0.0075																															
Sodium	N/A			N/A	59.45																															
Uranium	N/A			0.03	0.000906																															
Uranium, Dissolved	N/A			0.03	0.000906																															
Vanadium	N/A			N/A	0.00925																															
Zinc	N/A			N/A	0.45																															

RGA

Table 4.51. SWMU 145 Locations of Groundwater Contaminants (Continued)

Hydro Unit	Depth Interval	Analysis	Child Resident NAL	Blkgd	MCL	DG-014	MW179	MW181	MW220	MW221	MW222	MW223	MW224	MW263	MW264	MW265	MW266	MW267	MW276	MW277	MW369	MW370	MW384	MW385	MW387	MW388	MW391	MW392	MW394	MW395	MW397						
65-68		Pesticides/PCBs (mg/L)																																			
		Heptachlor	0.0000114	N/A	0.0004																							0.00005U									
		PCB, Total	0.0000793	N/A	0.0005																							0.00018U									
		PCB-1016	0.0000468	N/A	0.0005																							0.00017U									
		PCB-1242	0.000123	N/A	0.0005																							0.00013U									
		PCB-1260	0.0000428	N/A	0.0005																							0.00009U									
		Radionuclides (pCi/L)																																			
		Iodine-131	N/A	N/A	N/A	N/A																							41.4U								
		Radium	N/A	N/A	N/A	N/A																							0.791U								
		Radium-224	N/A	N/A	N/A	N/A																							0.91U								
		Radium-226	0.1	0.6	5																							0.674									
		Radium-228	N/A	N/A	N/A	N/A																							3.14								
		Strontium-90	0.522	N/A	N/A	N/A																							17.5U								
		Technetium-99	14	22.3	900	40																						28.7									
		Thorium-230	0.424	1.1	N/A																								1.77U								
		Thorium-232	0.382	N/A	N/A	N/A																							0.468								
		Thorium-234	N/A	N/A	N/A	N/A																							585								
		Semivolatiles (mg/L)																																			
		1,2-Dichlorobenzene	0.0166	N/A	N/A	0.6	0.01U																						0.005U								
		1,4-Dichlorobenzene	0.000578	N/A	N/A	0.075	0.01U																						0.005U								
		3-Methylphenol	0.0725	N/A	N/A	N/A																							0.01U								
		Bis(2-chloroethoxy)methane	N/A	N/A	N/A	N/A	0.01U																						0.01U								
		Bis(2-ethylhexyl)phthalate	0.00312	N/A	N/A	0.006	0.01U																						0.005								
		Butyl benzyl phthalate	0.259	N/A	N/A	N/A																							0.01U								
		Chrysene	0.00132	N/A	N/A	N/A	0.01U																						0.01U								
Diethyl phthalate	1.2	N/A	N/A	N/A	0.01U																						0.0005										
Dimethyl phthalate	15.1	N/A	N/A	N/A	0.01U																						0.01U										
Di-n-butyl phthalate	0.129	N/A	N/A	N/A	0.01U																						0.037										
Volatiles (mg/L)																																					
1,1,1-Trichloroethane	0.0335	N/A	N/A	0.2																							0.005U										
1,3-Dimethylbenzene	0.439	N/A	N/A	10																							0.005U										
1,4-Dioxane	0.00479	N/A	N/A	N/A																							0.01U										
2-Butanone	0.0868	N/A	N/A	N/A																							0.05U										
Acetone	0.0275	N/A	N/A	N/A																							0.01U										
Benzene	0.000385	N/A	N/A	0.005																							0.005U										
Bromomethane	0.000391	N/A	N/A	N/A																							0.01U										
Carbon disulfide	0.0457	N/A	N/A	N/A																							0.005U										
Chlorobenzene	0.00466	N/A	N/A	0.1																							0.005U										
Chloroform	0.0000287	N/A	N/A	N/A																							0.005U										
Chloromethane	0.00167	N/A	N/A	N/A																							0.01U										
cis-1,2-Dichloroethene	0.00273	N/A	N/A	0.07	0.001U																						0.005U										
Dichlorodifluoromethane	0.018	N/A	N/A	N/A																							0.01U										
Ethanol	N/A	N/A	N/A	N/A																							5U										
Ethylbenzene	0.00468	N/A	N/A	0.7																							0.005U										
Methylene chloride	0.00426	N/A	N/A	0.005																							0.002										
Tetrachloroethene	0.000582	N/A	N/A	0.005																							0.005U										
Toluene	0.0338	N/A	N/A	1																							0.005U										
Total Xylene	0.0653	N/A	N/A	10																							0.015U										
trans-1,2-Dichloroethene	0.00548	N/A	N/A	0.1	0.001U																						0.005U										
trans-1,3-Dichloropropene	N/A	N/A	N/A	N/A																							0.005U										
Trichloroethene	0.0016	N/A	N/A	0.005	0.001U																						0.019										
Trichlorofluoromethane	0.0577	N/A	N/A	N/A																							0.005U										

Table 4.51. SWMU 145 Locations of Groundwater Contaminants (Continued)

Hydro Unit	Depth Interval	Analysis	Child Resident NAL	Bkgd	MCL	DG-014	MW179	MW181	MW220	MW221	MW222	MW223	MW224	MW263	MW264	MW265	MW266	MW267	MW276	MW277	MW369	MW370	MW384	MW385	MW387	MW388	MW391	MW392	MW394	MW395	MW397					
	69-72	Metals (mg/L)																																		
		Aluminum	1.49	2.189	N/A				3.6							0.2				0.32						2.07										
		Aluminum, Dissolved	1.49	0.311	N/A				0.2							0.2				0.2U						0.005U										
		Antimony	0.000564	0.06	0.006				0.005U							0.01				0.0063						0.005U										
		Antimony, Dissolved	0.000564	0.06	0.006				0.006							0.006U										0.00228										
		Arsenic	0.000035	0.005	0.01				0.00198																	0.00228										
		Barium	0.104	0.235	2				0.605							0.25				0.25						0.208										
		Barium, Dissolved	0.104	0.2	2				0.52							0.229				0.24						0.205										
		Beryllium	0.00264	0.004	0.004				0.002							0.002				0.005U						0.005U										
		Beryllium, Dissolved	0.00264	0.004	0.004				0.002							0.002				0.005U						2U										
		Boron	0.136	N/A	N/A				0.03							0.03				0.03						2U										
		Boron, Dissolved	N/A	N/A	N/A				0.03							0.04				0.01						0.005U										
		Cadmium	0.000661	0.01	0.005				0.0009							0.005U				0.005U						0.005U										
		Calcium	N/A	41.238	N/A				33.9							24.6				27.9						28.7										
		Calcium, Dissolved	N/A	38.166	N/A				23.6							24				27.4																
		Cerium	N/A	N/A	N/A				0.08							0.08																				
		Cerium, Dissolved	N/A	N/A	N/A				0.08							0.08																				
		Chromium	1.76	0.144	0.1				0.719							1.98				0.87						0.0103										
		Chromium, Dissolved	1.76	0.05	0.1				0.04							0.03				0.0011						0.02U										
		Cobalt	0.0906	0.045	N/A				0.161							0.01				0.0105						0.025U										
		Cobalt, Dissolved	0.0906	0.045	N/A				0.01							0.01				0.0055						0.025U										
		Copper	0.0557	0.036	1.3				0.03							0.048				0.085						0.05U										
		Copper, Dissolved	0.0557	0.02	1.3				0.03							0.03				0.006						0.05U										
		Gallium	N/A	N/A	N/A				0.09							0.09																				
		Gallium, Dissolved	N/A	N/A	N/A				0.09							0.09																				
		Iron	0.449	5.03	N/A				11.1							11.9				5.11						7.29										
		Iron, Dissolved	0.449	0.267	N/A				0.3							0.3				0.1U						0.005U										
		Lead	0.015	0.129	0.015				0.00577							0.003U				0.003U						0.005U										
		Lithium	0.0302	N/A	N/A				0.08							0.08																				
		Lithium, Dissolved	N/A	N/A	N/A				0.08							0.08																				
		Magnesium	N/A	16.262	N/A				14.7							9.67				11.6						11.4										
		Magnesium, Dissolved	N/A	16.215	N/A				9.1							9.49				11.3																
		Manganese	0.035	0.119	N/A				2.54							0.204				0.14						0.096										
		Manganese, Dissolved	0.035	0.068	N/A				0.052							0.19				0.05						0.002U										
		Mercury	0.000444	0.0002	0.002				0.0002U							0.0002U				0.0002U						0.0002U										
		Mercury, Dissolved	0.000444	0.0002	0.002				0.0002U							0.0002U				0.0002U						0.0002U										
		Molybdenum	0.00753	0.05	N/A				0.03							0.07				0.04						0.025U										
		Molybdenum, Dissolved	0.00753	0.05	N/A				0.03							0.07				0.04						0.025U										
		Nickel	0.0301	0.682	N/A				1.05							0.53				0.56						0.00568										
		Nickel, Dissolved	0.0301	0.305	N/A				0.204							0.5				0.46						0.00568										
		Potassium	N/A	5.195	N/A				87.5							3				3.23						2.3										
		Potassium, Dissolved	N/A	4.096	N/A				3							3				4.14						0.00942										
		Selenium	0.00754	0.005	0.05				0.0088							0.0029				0.005U						0.00942										
		Silicon	N/A	N/A	N/A				0.03							6.04				7.13																
		Silicon, Dissolved	N/A	N/A	N/A				0.03							6.81				6.45						0.025U										
		Silver	0.0075	0.011	N/A				0.03							0.03				0.05U						0.025U										
		Silver, Dissolved	0.0075	0.06	N/A				0.03							0.03				0.01U						0.025U										
		Sodium	N/A	59.45	N/A				58.2							44.5				33.3						46.7										
		Sodium, Dissolved	N/A	60.433	N/A				36.3							43.4				31.4						50										
		Strontium	0.901	N/A	N/A				0.147							0.09																				
		Strontium, Dissolved	N/A	N/A	N/A				0.139							0.091																				
		Thorium	N/A	N/A	N/A				0.05							0.05				0.05																
		Thorium, Dissolved	N/A	N/A	N/A				0.05							0.05				0.05																
		Titanium	N/A	N/A	N/A				0.06							0.06				0.06																
		Titanium, Dissolved	N/A																																	

Table 4.51. SWMU 145 Locations of Groundwater Contaminants (Continued)

Hydro Unit	Depth Interval	Analysis	MCL	Blgd	Child Resident NAL	DG-014	MW179	MW181	MW220	MW221	MW222	MW223	MW224	MW263	MW264	MW265	MW266	MW267	MW276	MW277	MW369	MW370	MW384	MW385	MW387	MW388	MW391	MW392	MW394	MW395	MW397						
	69-72	Zinc	N/A	0.054	0.45				0.0562							0.105				0.0113		0.0204				0.1U											
		Zinc, Dissolved	N/A	0.049	0.45				0.05							0.05				0.0067																	
		Zirconium	N/A	N/A	N/A				0.02							0.02																					
		Zirconium, Dissolved	N/A	N/A	N/A				0.02							0.02																					
		Pesticides/PCBs (mg/L)																																			
		Heptachlor	0.000114	N/A	0.0004															0.0004U		0.0005U				0.0005U											
		PCB, Total	0.0000793	N/A	0.0005				0.00078											0.0005U		0.000188				0.00018U											
		PCB-1016	0.0000468	N/A	0.0005				0.001U											0.0005U		0.000188				0.00017U											
		PCB-1242	0.000123	N/A	0.0005				0.00078											0.0005U		0.001U				0.00013U											
		PCB-1260	0.0000428	N/A	0.0005				0.001U											0.0011U		0.00017				0.00009U											
		Radionuclides (pCi/L)																																			
		Iodine-131	N/A	N/A	N/A				164													30U				31.5U											
		Radium	N/A	N/A	N/A				0.791U													0.78U				0.791U											
		Radium-224	N/A	N/A	N/A				0.99U													1.15U				0.97U											
		Radium-226	0.1	0.6	5				0.789													0.52				0.567											
		Radium-228	N/A	N/A	N/A				4.56													26.8U				1.25											
		Strontium-90	0.522	N/A	N/A				23.3U													18.2U				17.5U											
		Technetium-99	14	22.3	900				29.5													50.6				228											
		Thorium-230	0.424	1.1	N/A				0.84U													1.35U				0.69U											
		Thorium-234	N/A	N/A	N/A				3.15U													338U				334U											
		Semivolatiles (mg/L)																																			
		1,2-Dichlorobenzene	0.0166	N/A	0.6	0.01U			0.005U													0.005U				0.005U											
		1,4-Dichlorobenzene	0.000578	N/A	0.075	0.01U			0.005U													0.005U				0.005U											
		3-Methylphenol	0.0725	N/A	N/A																	0.01U				0.01U											
		Bis(2-chloroethoxy)methane	N/A	N/A	N/A	0.01U																0.01U				0.01U											
		Bis(2-ethylhexyl)phthalate	0.00312	N/A	0.006	0.01U																0.002				0.002											
		Butyl benzyl phthalate	0.259	N/A	N/A																	0.01U				0.01U											
		Chrysene	0.00132	N/A	N/A	0.01U																0.0006				0.01U											
		Diethyl phthalate	1.2	N/A	N/A	0.01U																0.01U				0.01U											
		Dimethyl phthalate	15.1	N/A	N/A	0.01U																0.01U				0.01U											
		Di-n-butyl phthalate	0.129	N/A	N/A	0.01U																0.01U				0.01U											
		Volatiles (mg/L)																																			
		1,1,1-Trichloroethane	0.0335	N/A	0.2				0.005U								0.004					0.005U				0.005U											
		1,2-Dichloroethane	0.00247	N/A	N/A																	0.001					0.005U										
		1,3-Dimethylbenzene	0.439	N/A	10																	0.005U				0.005U											
		1,4-Dioxane	0.00479	N/A	N/A												0.2U					0.2U				0.2U											
		2-Butanone	0.0868	N/A	N/A												0.004					0.01U				0.05U											
		Acetone	0.0275	N/A	N/A				0.017								0.006					0.01U				0.05U											
		Benzene	0.000385	N/A	0.005				0.005U								0.005U					0.005U				0.005U											
		Bromomethane	0.000391	N/A	N/A				0.005U								0.005U					0.01U				0.01U											
		Carbon disulfide	0.0457	N/A	N/A				0.005U								0.005U					0.005U				0.005U											
		Chlorobenzene	0.00466	N/A	0.1				0.005U								0.005U					0.005U				0.005U											
		Chloroform	0.000287	N/A	N/A				0.005U								0.00049					0.001U				0.005U											
		Chloromethane	0.00167	N/A	N/A				0.000084								0.000072					0.005U				0.01U											
		cis-1,2-Dichloroethane	0.00273	N/A	0.07	0.001U			0.00013							0.00025						0.00553				0.005U											
		Dichlorodifluoromethane	0.018	N/A	N/A				0.005U							0.005U						0.005U				0.01U											
		Ethanol	N/A	N/A	N/A				0.25U							0.25U						5U				5U											
		Ethylbenzene	0.00468	N/A	0.7				0.005U							0.005U						0.005U				0.005U											
		m,p-Xylene	N/A	N/A	10				0.0001U							0.0001U																					
		Methylene chloride	0.00426	N/A	0.005				0.01U							0.000061						0.01U				0.004											
		Tetrachloroethene	0.000582	N/A	0.005				0.005U							0.000061						0.005U				0.005U											
		Toluene	0.0338	N/A	1				0.002							0.00019						0.005U				0.005U											
		Total Xylene	0.0653	N/A	10				0.015U							0.015U						0.015U				0.015U											
		trans-1,2-Dichloroethene	0.00548	N/A	0.1	0.001U			0.005U																												

Table 4.51. SWMU 145 Locations of Groundwater Contaminants (Continued)

Hydro Unit	Depth Interval	Analysis	Child Resident NAL	Bkgd	MCL	DG-014	MW179	MW181	MW220	MW221	MW222	MW223	MW224	MW263	MW264	MW265	MW266	MW267	MW276	MW277	MW369	MW370	MW384	MW385	MW387	MW388	MW391	MW392	MW394	MW395	MW397							
	75-78	Metals (mg/L)																																				
		Aluminum	1.49	2.189	N/A						15.2						0.2																					
		Aluminum, Dissolved	1.49	0.311	N/A						0.2						0.2																					
		Antimony	0.000564	0.06	0.006						0.005U						0.005U																					
		Antimony, Dissolved	0.000564	0.06	0.006						0.006U						0.008																					
		Arsenic	0.000035	0.005	0.01						0.00248						0.08																					
		Barium	0.104	0.235	2						2.25						0.07																					
		Barium, Dissolved	0.104	0.2	2						2.15						0.07																					
		Beryllium	0.00264	0.004	0.004						0.002						0.002																					
		Beryllium, Dissolved	0.00264	0.004	0.004						0.002						0.002																					
		Boron	0.136	N/A	N/A						0.15						1.05																					
		Boron, Dissolved	N/A	N/A	N/A						0.03						0.94																					
		Cadmium	0.000661	0.01	0.005						0.005U						0.005U																					
		Calcium	N/A	41.238	N/A						33.8						59.4																					
		Calcium, Dissolved	N/A	38.166	N/A						21.9						59.4																					
		Cerium	N/A	N/A	N/A						0.08						0.08																					
		Cerium, Dissolved	N/A	N/A	N/A						0.08						0.08																					
		Chromium	1.76	0.144	0.1						0.144						3.04																					
		Chromium, Dissolved	1.76	0.05	0.1						0.03						0.03																					
		Cobalt	0.0906	0.045	N/A						0.574						0.01																					
		Cobalt, Dissolved	0.0906	0.045	N/A						0.01						0.01																					
		Copper	0.0557	0.036	1.3						0.03						0.08																					
		Copper, Dissolved	0.0557	0.02	1.3						0.03						0.03																					
		Gallium	N/A	N/A	N/A						0.09						0.09																					
		Gallium, Dissolved	N/A	N/A	N/A						0.09						0.09																					
		Iron	0.449	5.03	N/A						107						17.6																					
		Iron, Dissolved	0.449	0.267	N/A						0.3						0.3																					
		Lead	0.015	0.129	0.015						0.00723						0.00723																					
		Lithium	0.0302	N/A	N/A						0.08						0.08																					
		Lithium, Dissolved	N/A	N/A	N/A						0.08						0.08																					
		Magnesium	N/A	16.262	N/A						14.7						25.7																					
		Magnesium, Dissolved	N/A	16.215	N/A						9.3						25.7																					
		Manganese	0.035	0.119	N/A						23.2						0.08																					
		Manganese, Dissolved	0.035	0.068	N/A						0.049						0.08																					
		Mercury	0.000444	0.0002	0.002						0.0003						0.0002U																					
		Mercury, Dissolved	0.000444	0.0002	0.002						0.0002U						0.0002U																					
		Molybdenum	0.00753	0.05	N/A						0.03						0.03																					
		Molybdenum, Dissolved	0.00753	0.05	N/A						0.03						0.03																					
		Nickel	0.0301	0.682	N/A						0.883						0.433																					
		Nickel, Dissolved	0.0301	0.305	N/A						0.04						0.04																					
		Potassium	N/A	5.195	N/A						7.97						6.87																					
		Potassium, Dissolved	N/A	4.096	N/A						3						4																					
		Selenium	0.00754	0.005	0.05						0.0107						0.00855																					
		Selenium, Dissolved	0.00754	0.005	0.05						0.0107						0.00855																					
		Silver	0.0075	0.011	N/A						0.03						0.03																					
		Silver, Dissolved	0.0075	0.06	N/A						0.03						0.03																					
		Sodium	N/A	59.45	N/A						72.4						46.6																					
		Sodium, Dissolved	N/A	60.433	N/A						54.3						39.6																					
		Strontium	0.901	N/A	N/A						0.09						0.168																					
		Strontium, Dissolved	N/A	N/A	N/A						0.087						0.167																					
		Thorium	N/A	N/A	N/A						0.05						0.05																					
		Thorium, Dissolved	N/A	N/A	N/A						0.05						0.05																					
		Titanium	N/A	N/A	N/A						0.06						0.06																					
		Titanium, Dissolved	N/A	N/A	N/A						0.06						0.06			</																		

Table 4.51. SWMU 145 Locations of Groundwater Contaminants (Continued)

Hydro Unit	Depth Interval	Analysis	Child Resident NAL	Bkgd	MCL	DG-014	MW179	MW181	MW220	MW221	MW222	MW223	MW224	MW263	MW264	MW265	MW266	MW267	MW369	MW370	MW384	MW385	MW387	MW388	MW391	MW392	MW394	MW395	MW397					
	75-78	Trichloroethene	0.0016	N/A	0.005	0.001U				0.00039	0.000055						0.016	0.00006				0.005U	0.005U											
	82-85	Trichlorofluoromethane	0.0577	N/A	N/A																													
		Metals (mg/L)																																
		Aluminum	1.49	2.189	N/A					0.2		57.2	0.2					0.2																
		Aluminum, Dissolved	1.49	0.311	N/A					0.2		0.2	0.2					0.2													0.737			
		Antimony	0.000564	0.06	0.006					0.005U		0.008	0.005U					0.005U														0.005U		
		Antimony, Dissolved	0.000564	0.06	0.006					0.019		0.006U	0.017					0.006U																
		Arsenic	0.000035	0.005	0.01					0.00218		0.00216	0.00186																				0.00257	
		Barium	0.104	0.235	2					0.304		1.75	0.775					0.246														0.289		
		Barium, Dissolved	0.104	0.2	2					0.291		1.71	0.769					0.239														0.291		
		Beryllium	0.00264	0.004	0.004					0.002		0.002	0.002					0.002														0.005U		
		Beryllium, Dissolved	0.00264	0.004	0.004					0.002		0.002	0.002					0.002														0.005U		
		Boron	0.136	N/A	N/A					0.06		0.16	0.12					0.04														2U		
		Boron, Dissolved	N/A	N/A	N/A					0.07		0.16	0.05					0.15																
		Cadmium	0.000661	0.01	0.005					0.005U		0.005U	0.00111					0.005U														0.005U		
		Calcium	N/A	41.238	N/A					26.7		34.3	34.2					24.6														40.2		
		Calcium, Dissolved	N/A	38.166	N/A					24.9		24.3	27.4					21.3																
		Cerium	N/A	N/A	N/A					0.08		0.08	0.08					0.08																
		Cerium, Dissolved	N/A	N/A	N/A					0.08		0.08	0.08					0.08																
		Chromium	1.76	0.144	0.1					0.277		1.41	0.03					5.4														0.075		
		Chromium, Dissolved	1.76	0.05	0.1					0.03		0.03	0.03					0.03														0.02U		
		Cobalt	0.0906	0.045	N/A					0.0419		0.191	0.306					0.01														0.00151		
		Cobalt, Dissolved	0.0906	0.045	N/A					0.01		0.01	0.01					0.178														0.0281		
		Copper	0.0557	0.036	1.3					0.0505		0.064	0.03					0.03																
		Copper, Dissolved	0.0557	0.02	1.3					0.03		0.03	0.03					0.03																
		Gallium	N/A	N/A	N/A					0.09		0.09	0.09					0.09																
		Gallium, Dissolved	N/A	N/A	N/A					0.09		0.09	0.09					0.09																
		Iron	0.449	5.03	N/A					2.16		117	3.61					21														1.33		
		Iron, Dissolved	0.449	0.267	N/A					0.3		0.3	0.3					0.3																
		Lead	0.015	0.129	0.015					0.005U		0.00145	0.0153					0.005U															0.0167	
		Lithium	0.0302	N/A	N/A					0.08		0.08	0.08					0.08																
		Lithium, Dissolved	N/A	N/A	N/A					0.08		0.08	0.08					0.08																
		Magnesium	N/A	16.262	N/A					11.7		14	15.3					9.2															17.5	
		Magnesium, Dissolved	N/A	16.215	N/A					11		9.9	11.6					8.4																
		Manganese	0.035	0.119	N/A					0.818		36.5	6.32					0.056															0.629	
		Manganese, Dissolved	0.035	0.068	N/A					0.012		0.039	0.038					0.058																
		Mercury	0.000444	0.0002	0.002					0.0002U		0.0002U	0.0004					0.0002U															0.0002U	
		Mercury, Dissolved	0.000444	0.0002	0.002					0.0002U		0.0002U	0.0003					0.0002U																
		Molybdenum	0.00753	0.05	N/A					0.03		0.089	0.03					0.117															0.00609	
		Molybdenum, Dissolved	0.00753	0.05	N/A					0.03		0.03	0.03					0.03																
		Nickel	0.0301	0.682	N/A					0.68		1.48	0.144					0.222															0.029	
		Nickel, Dissolved	0.0301	0.305	N/A					0.05		0.111	0.04					0.04																
		Potassium	N/A	5.195	N/A					9.52		13.9	3					4															1.9	
		Potassium, Dissolved	N/A	4.096	N/A					3		10	3					4																
		Selenium	0.00754	0.005	0.05					0.0118		0.0108	0.00867					0.00685															0.0133	
		Silver	0.0075	0.011	N/A					0.03		0.03	0.03					0.03															0.025U	
		Silver, Dissolved	0.0075	0.06	N/A					0.03		0.03	0.03					0.03																
		Sodium	N/A	59.45	N/A					48.3		66.3	76.1					51.3																53.9
		Sodium, Dissolved	N/A	60.433	N/A					44.7		48.3	71.8					39.6																
		Strontium	0.901	N/A	N/A					0.108		0.125	0.11					0.104																
		Strontium, Dissolved	N/A	N/A	N/A					0.103		0.123	0.109					0.107																
		Thorium	N/A	N/A	N/A					0.05		0.05	0.05					0.05																
		Thorium, Dissolved	N/A	N/A	N/A					0.05		0.05	0.05					0.05																
		Titanium	N/A	N/A	N/A					0.06		0.06	0.06					0.06																
		Titanium, Dissolved	N/A	N/A	N/A					0.06		0.06	0.06					0.06																
		Uranium	0.000906	0.002	0.03					0.001U		0.00384	0.00396																					

Table 4.51. SWMU 145 Locations of Groundwater Contaminants (Continued)

Hydro Unit	Depth Interval	Analysis	MCL	Bkgd	Child Resident NAL	DG-014	MW179	MW181	MW220	MW221	MW222	MW223	MW224	MW263	MW264	MW265	MW266	MW267	MW276	MW277	MW369	MW370	MW384	MW385	MW387	MW388	MW391	MW392	MW394	MW395	MW397
	82-85	Zinc	N/A	0.054	0.45					0.05	0.34	0.05	0.05					0.05												0.1U	
		Zinc, Dissolved	N/A	0.049	0.45					0.05	0.05	0.05	0.05					0.05													
		Zirconium	N/A	N/A	N/A					0.02	0.02	0.02	0.02					0.02													
		Zirconium, Dissolved	N/A	N/A	N/A					0.02	0.02	0.02	0.02					0.02													
		Pesticides/PCBs (mg/L)																													
		Heptachlor	0.000114	N/A	0.000114					0.00787		0.00161	0.000359					0.00017U													0.00005U
		PCB, Total	0.0000793	N/A	0.0000793					0.001U		0.00184	0.000359					0.00017U													0.00018U
		PCB-1016	0.0000468	N/A	0.0000468					0.00787		0.00161	0.00024					0.00016U													0.00013U
		PCB-1242	0.000123	N/A	0.000123					0.001U		0.0001U	0.0001U					0.0001U													0.00009U
		PCB-1260	0.0000428	N/A	0.0000428					0.001U		0.0001U	0.0001U					0.0001U													0.00009U
		Radionuclides (pCi/L)																													
		Iodine-131	N/A	N/A	N/A					499U		118U	101U					47.9U													37.4U
		Radium	N/A	N/A	N/A					0.791U		0.791U	0.791U																		0.79U
		Radium-224	N/A	N/A	N/A					0.97U		0.313	0.86U																		0.87U
		Radium-226	0.1	0.6	0.1					0.101		0.567	0.641					8.47U												0.661	
		Radium-228	N/A	N/A	N/A					29.5U		27U	13.2																		37.9U
		Strontium-90	N/A	N/A	N/A					23.3U		23.3U	23.3U																		17.5U
		Technetium-99	14	22.3	14					30.3		55.5	15																	20.8	
		Thorium-230	0.424	1.1	0.424					0.482		0.46	0.76U																	0.69U	
		Thorium-234	N/A	N/A	N/A					4.290		474	4.490					303U													546
		Semivolatiles (mg/L)																													
		1,2-Dichlorobenzene	0.0166	N/A	0.0166					0.005U		0.005U	0.000057					0.005U													0.005U
		1,4-Dichlorobenzene	0.000578	N/A	0.000578					0.005U		0.005U	0.000062					0.005U													0.005U
		3-Methylphenol	0.0725	N/A	0.0725																										0.01U
		Bis(2-chloroethoxy)methane	N/A	N/A	N/A																										0.01U
		Bis(2-ethylhexyl)phthalate	0.00312	N/A	0.00312																										0.003
		Butyl benzyl phthalate	0.259	N/A	0.259																										0.01U
		Chrysene	0.00132	N/A	0.00132																										0.01U
		Diethyl phthalate	1.2	N/A	1.2																										0.01U
		Dimethyl phthalate	15.1	N/A	15.1																										0.01U
		Di-n-butyl phthalate	0.129	N/A	0.129																										0.01U
		Volatiles (mg/L)																													
		1,1,1-Trichloroethane	0.0335	N/A	0.0335					0.005U		0.005U	0.005U					0.005U													0.005U
		1,3-Dimethylbenzene	0.439	N/A	0.439																										0.005U
		1,4-Dioxane	0.00479	N/A	0.00479																										0.01U
		2-Butanone	0.0868	N/A	0.0868					0.01U		0.004	0.004					0.006													0.05U
		Acetone	0.0275	N/A	0.0275					0.005		0.006	0.17					0.017													0.011
		Benzene	0.000385	N/A	0.000385					0.005U		0.005U	0.005U					0.005U													0.005U
		Bromomethane	0.000391	N/A	0.000391					0.005U		0.005U	0.000062					0.005U													0.01U
		Carbon disulfide	0.0457	N/A	0.0457					0.005U		0.005U	0.005U					0.005U													0.005U
		Chlorobenzene	0.00466	N/A	0.00466					0.005U		0.005U	0.000046					0.005U													0.005U
		Chloroform	0.0000287	N/A	0.0000287					0.000073		0.005U	0.000058					0.00005U													0.005U
		Chloromethane	0.00167	N/A	0.00167					0.005U		0.000094	0.000093					0.000063													0.01U
		cis-1,2-Dichloroethene	0.00273	N/A	0.00273					0.00019		0.005U	0.000078					0.005U													0.005U
		Dichlorodifluoromethane	0.018	N/A	0.018					0.005U		0.005U	0.000074					0.005U													0.01U
		Ethanol	N/A	N/A	N/A					0.25U		0.25U	0.35					0.2													5U
		Ethylbenzene	0.00468	N/A	0.00468					0.005U		0.005U	0.005U					0.005U													0.005U
		m,p-Xylene	N/A	N/A	N/A					0.0001U		0.0001U	0.000055					0.0001U													0.005U
		Methylene chloride	0.00426	N/A	0.00426					0.000085		0.01U	0.000074					0.00005													0.014
		Tetrachloroethene	0.000582	N/A	0.000582					0.005U		0.005U	0.005U					0.005U													0.005U
		Toluene	0.0338	N/A	0.0338					0.005U		0.005U	0.00012					0.000068													0.005U
		Total Xylene	0.0653	N/A	0.0653					0.015U		0.015U	0.015U					0.015U													0.015U
		trans-1,2-Dichloroethene	0.00548	N/A	0.00548					0.005U		0.005U	0.005U					0.005U													0.005U
		trans-1,3-Dichloropropene	N/A	N/A	N/A					0.005U		0.005U	0.00017					0.005U													0.005U
		Trichloroethene	0.0016	N/A	0.0016					0.007		0.001	0.006					0.000065													0.014
		Trichlorofluoromethane	0.0577	N/A	0.0577					0.005U		0.005U	0.005U					0.005U													0.005U

Table 4.51. SWMU 145 Locations of Groundwater Contaminants (Continued)

Hydro Unit	Depth Interval	Analysis	Child Resident NAL	Bkgd	MCL	DG-014	MW179	MW181	MW220	MW221	MW222	MW223	MW224	MW263	MW264	MW265	MW266	MW267	MW276	MW277	MW369	MW370	MW384	MW385	MW387	MW388	MW391	MW392	MW394	MW395	MW397	
91-95		Metals (mg/L)																														
		Aluminum	1.49	2.189	N/A																											1.14
		Antimony	0.000564	0.06	0.006																											0.005U
		Arsenic	0.000035	0.005	0.01																											0.00227
		Barium	0.104	0.235	2																											0.183
		Barium, Dissolved	0.104	0.2	2																											0.169
		Beryllium	0.00264	0.004	0.004																											0.005U
		Boron	0.136	N/A	N/A																											2U
		Cadmium	0.000661	0.01	0.005																											0.005U
		Calcium	N/A	41.238	N/A																											20.8
		Chromium	1.76	0.144	0.1																											0.025U
		Chromium, Dissolved	1.76	0.05	0.1																											0.02U
		Cobalt	0.0906	0.045	N/A																											0.025U
		Copper	0.0557	0.036	1.3																											0.05U
		Iron	0.449	5.03	N/A																											1.58
		Lead	0.015	0.129	0.015																											0.00523
		Magnesium	N/A	16.262	N/A																											10.4
		Manganese	0.035	0.119	N/A																											8.55
		Mercury	0.000444	0.0002	0.002																											0.466
		Molybdenum	0.00753	0.05	N/A																											0.0002U
		Nickel	0.0301	0.682	N/A																											0.00187
		Potassium	N/A	5.195	N/A																											0.0107
		Selenium	0.00754	0.005	0.05																											2.12
		Silver	0.0075	0.011	N/A																											0.00697
		Sodium	N/A	59.45	N/A																											0.025U
		Uranium	0.000906	0.002	0.03																											63.5
		Uranium, Dissolved	0.000906	0.002	0.03																											0.0014
		Vanadium	0.00925	0.134	N/A																											0.001U
		Zinc	0.45	0.054	N/A																											0.025U
																																0.0381
																																0.1U
																																0.00005U
																																0.00018U
																																0.00017U
																																0.00013U
																																0.00009U
																																36.8U
																																0.466
																																0.93U
																																0.576
																																40.2U
																																17.5U
																																25.7
																																0.79U
																																329U
																																0.005U
																																0.005U
																																0.01U
																																0.01U
																																0.006
																																0.01U
																																0.01U
																																0.0006
																																0.01U
																																0.005
																																0.005U
																																0.005U
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																																0.005U

Table 4.51. SWMU 145 Locations of Groundwater Contaminants (Continued)

Hydro Unit	Depth Interval	Analysis	Child Resident NAL	Blgd	MCL	DG-014	MW179	MW181	MW220	MW221	MW222	MW223	MW224	MW263	MW264	MW265	MW266	MW267	MW276	MW277	MW369	MW370	MW384	MW385	MW387	MW388	MW391	MW392	MW394	MW395	MW397								
RGA	91-95	Volatiles (mg/L)																																					
		1,1,1-Trichloroethane	0.0335	N/A	0.2																																		
		1,3-Dimethylbenzene	0.439	N/A	10																																		
		1,4-Dioxane	0.00479	N/A	N/A																																		
		2-Butanone	0.0868	N/A	N/A																																		
		Acetone	0.0275	N/A	N/A																																		
		Benzene	0.000385	N/A	0.005																																		
		Bromomethane	0.000391	N/A	N/A																																		
		Carbon disulfide	0.0457	N/A	N/A																																		
		Chlorobenzene	0.00466	N/A	0.1																																		
		Chloroform	0.000287	N/A	N/A																																		
		Chloromethane	0.00167	N/A	N/A																																		
		cis-1,2-Dichloroethene	0.00273	N/A	0.07																																		
		Dichlorodifluoromethane	0.018	N/A	N/A																																		
		Ethanol	N/A	N/A	N/A																																		
		Ethylbenzene	0.00468	N/A	0.7																																		
		Methylene chloride	0.00426	N/A	0.005																																		
		Tetrachloroethene	0.000582	N/A	0.005																																		
		Toluene	0.0338	N/A	1																																		
		Total Xylene	0.0653	N/A	10																																		
		trans-1,2-Dichloroethene	0.00548	N/A	0.1																																		
		trans-1,3-Dichloropropene	N/A	N/A	N/A																																		
		Trichloroethene	0.0016	N/A	0.005																																		
		Trichlorofluoromethane	0.0577	N/A	N/A																																		

Blank cells indicate interval was not sampled for the specified analysis. Maximum value shown for each depth interval.

N/A = not applicable

Bold indicates result is greater than NAL value.

Italics indicates result is greater than MCL value.

Bold + Italics indicate result is greater than both NAL and MCL values.

Shading indicates result is greater than the background value.

5. FATE AND TRANSPORT

This chapter provides an overview of the fate and transport of the primary analytes for the BGOU. (Appendix E, Section E.3 documents the methods and results of fate and transport modeling performed for the BGOU RI.) The sources modeled are SWMU 2, SWMU 3, SWMU 4, SWMU 5, SWMU 6, SWMU 7, SWMU 30, and SWMU 145. Two pathways were considered in the transport modeling analyses: (1) dissolved-phase transport through the aquifer and (2) vapor transport to a residential basement.

5.1 CONCEPTUAL MODEL

The sources of contamination to the RGA considered in this report are the waste disposal areas in the BGOU SWMUs. Releases from these SWMUs have impacted soils below or adjacent to the source zones and, through vertical infiltration in soil, these sources have the potential to contaminate the groundwater underlying these sources. Subsequently, contaminated groundwater could migrate to the points of exposure (POEs). The potential POEs for the BGOU SWMUs were identified as the SWMU boundary, plant boundary, property boundary, surface seeps at Little Bayou Creek (hereafter referred to as the Little Bayou seeps), and the Ohio River. [Fate and transport modeling assessed the Little Bayou seeps and the Ohio River as the locations to assess risk to the groundwater user through a hypothetical well, consistent with the Risk Methods Document (DOE 2001). Fate and transport modeling also assessed the SWMU boundary.] Not all SWMUs have transport pathways to all of the POEs. For example, SWMU 145 is located outside of the plant boundary and does not contribute to the Little Bayou seeps. SWMUs 3, 6, 7, and 30 were determined to be the only SWMUs contributing to the Little Bayou seeps POE. SWMUs 2, 4, and 5 have POEs at the plant boundary, property boundary, and the Ohio River. Figure 5.1 shows the location of the BGOU SWMUs, plant boundary, property boundary, Little Bayou seeps, Ohio River, and contaminant flow particle tracks from the SWMUs. The uncertainty in the flow paths is discussed in Section 5.5.4.

Contaminant migration could have impacted three HUs underlying the source zones at the BGOU SWMUs. These units, which control the flow of shallow groundwater and contaminant migration, are as follows, in descending depth order:

- UCRS—approximately 60 ft of silt and clay with horizons of sand and gravel;
- RGA—approximately 40 ft of gravel and sand deposits that overlie the McNairy Formation; and
- McNairy Formation—approximately 225 ft of a silty and clayey sand that forms a lower confining unit to the RGA.

Previous work has shown that groundwater flow in the UCRS is primarily vertical to the RGA and then lateral toward the Ohio River and that groundwater flow in the McNairy Formation (both vertical and lateral) is significantly slower than that in the RGA. The primary contaminant pathway for the site-related contaminants is vertical migration through the UCRS followed by lateral migration in the RGA. The RGA discharges to the Ohio River and, for a limited number of SWMUs, to the Little Bayou seeps. Section 3 provides a detailed description of the geology and hydrogeology at PGDP.

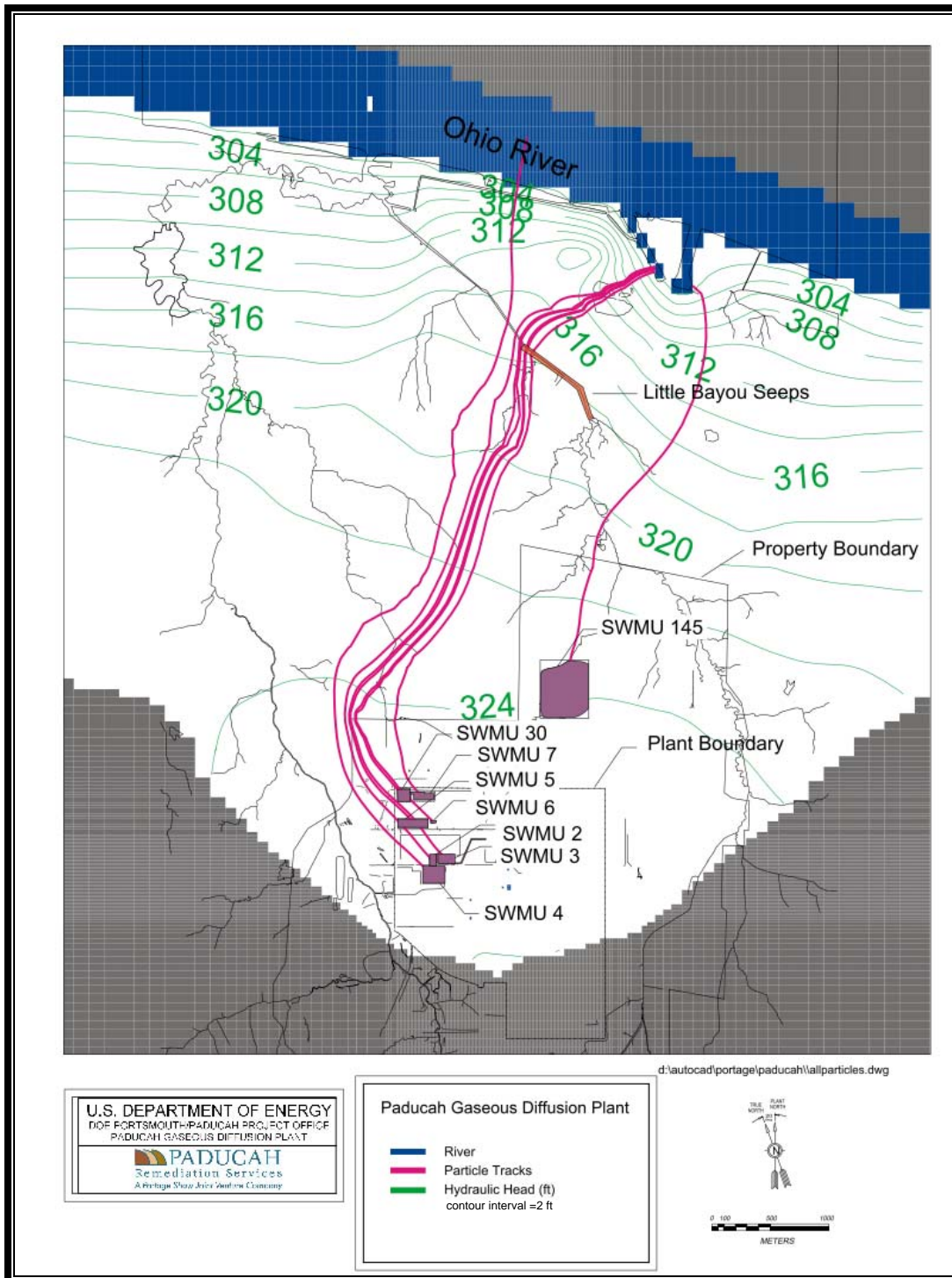


Figure 5.1. Location of the BGOU SWMUs and POEs

5.2 ANALYTE SELECTION AND CHEMICAL PROPERTIES

5.2.1 Analyte Selection

Soil results were screened prior to their use in modeling to determine groundwater analytes for source term development. The following outlines the review and screening process used prior to establishing the analytes for use in modeling:

- *Units of Reported Results.* The units of measure used for analyte classes (i.e., inorganic chemicals, organic compounds, and radionuclides) were assigned consistent units of measure. The units of measure used were mg/kg for inorganic chemicals and organic compounds and pCi/g for radionuclides.
- *Detection Status.* Each result was coded either as a detect or nondetect based on the data qualifier codes present in the database. Results assigned a “U” or “UJ” qualifier were considered nondetects. This coding subsequently was used to calculate the frequency of detection for each constituent. Results for metals and organic compounds were assigned “U” laboratory flags or validation qualifiers based on one of the following: 1) the constituent underwent analysis, but was not detected by the laboratory (laboratory “U” flag); or 2) the constituent was detected, but the measured concentration was indistinguishable from a field, laboratory, or instrument blank (data validation “U” qualifier”). In keeping with MARLAP 2004 and PGDP site-specific data validation guidance, two separate measures were used to define the detectability status of radionuclides: 1) minimum detectable concentration (MDC) and 2) comparison to total propagated 2-sigma counting uncertainties (TPU). Results exceeding the 2-sigma counting uncertainty and the MDC were considered true positive values. Results less than the MDC or the 2-sigma TPU were assigned “U” validation qualifiers.
- *Frequency of Detection.* Those constituents detected in less than 5% of the samples were not retained because their infrequency of detection prevents development of a usable source term.
- *Essential Nutrients.* Results for the seven essential nutrients were removed from the data sets. They are calcium, chloride, iodine, magnesium, potassium, sodium, and phosphorous.
- *Protactinium-234m, potassium-40, and thorium-234.* Results for these isotopes were not retained for source term development. Protactinium-234m was not retained because it is a short-lived daughter of uranium-238 and its presence in the source term is reflected in the uranium-238 results. Potassium-40 and thorium-234 were not retained because the former is naturally occurring and not a site-related contaminant, and the latter is a short-lived isotope not useful in source term development (DOE 2001).

Analytes retained under current conditions are presented for each SWMU in Tables E3.1 through E3.8. The maximum detected soil concentrations (surface to 60 ft bgs) are presented by analyte and by SWMU, with a comparison to the child resident soil screening levels¹ (SSLs) with a dilution attenuation factor (DAF)=1. Those analytes with a maximum concentration greater than their respective SSLs then were compared to soil/sediment child resident NALs. Constituents greater than both the SSL and NAL were retained as analytes for groundwater modeling. The screening values used may be found in Tables A.7 and A.17 of the Risk Methods Document (2001a). The SSLs for protection of groundwater for radionuclides include only ingestion of tap water. Dermal absorbed dose is not applicable to radionuclides per guidance found in EPA’s *Risk Assessment Guidance for Superfund: Volume I - Human Health Evaluation Manual* (EPA2004a). Inhalation exposure though showering and household use is not

¹ SSLs are risk-based soil concentrations considered to be protective of groundwater (DOE 2001).

calculated because the inhalation exposure factor (IEF) is 0 for all the radionuclides with SSLs in Table A.7b of the Risk Methods Document (2001a). There are no SSLs for protection of groundwater for radium-226+D, thorium-228+D, thorium-230, and thorium-232 because RESRAD modeling done to generate those SSLs in Table A.7b of the Risk Methods Document (2001a) shows that the radionuclide does not reach groundwater within 10,000 years precluding receptor uptake.

Exceptions to this are TCE, technetium-99, and uranium isotopes, which were retained in all SWMUs, as they are significant risk contributors or known to be part of the facility's process history.

Following this review and screening process, the analytes retained then were modeled as described below and in Appendix E of the RI Report. Modeled results were used in the BHHRA.

5.2.2 Chemical Properties

Table 5.1 lists the analytes identified for fate and transport assessment along with the parameter values chosen to represent these contaminants in the Seasonal Soil Compartment Model (SESOIL) and Analytical Transient 1-,2-,3- Dimensional (AT123D) models.

Table 5.1. Burial Ground Analytes for the Groundwater Pathway and Properties

Analyte	Mol. Wt. (MW) (g/mol)	Solubility in water (mg/L)	Diffusion in air (cm ² /s)	Diffusion in water (m ² /hr)	Henry's Constant (atm.m3/mol)	Koc (L/kg)	Kd ^a (L/kg)	Half Life (years)
Acenaphthene	154	4.20E+00	4.00E-02	2.77E-06	1.60E-04	4.90E+03	3.9	Infinite
Anthracene	178.24	4.30E-02	3.20E-02	2.79E-06	5.55E-05	2.04E+04	16.3	Infinite
Antimony	121.75	1.00E+07	NA	3.60E-07	NA	NA	45	Infinite
PCB-1248	288	1.70E-02	1.75E-02	2.38E-06	1.60E-04	2.51E+04	20	Infinite
PCB-1254	327	7.00E-02	1.56E-02	1.80E-06	3.40E-04	4.25E+04	34	Infinite
PCB-1260	375.7	2.70E-02	1.38E-02	1.56E-06	7.40E-05	2.07E+05	165.6	Infinite
Arsenic	74.92	1.00E+07	NA	3.60E-07	NA	NA	29	Infinite
Benzo(a)pyrene	252.32	1.62E-03	4.30E-02	3.24E-06	1.13E-06	9.69E+05	772	Infinite
Beryllium	9.01	1.00E+07	NA	3.60E-07	NA	NA	250	Infinite
Cadmium	112.41	1.00E+07	NA	3.60E-07	NA	NA	75	Infinite
cis-1,2-DCE	96.94	3.50E+03	7.00E-02	4.07E-06	4.08E-03	3.55E+01	0.028	Infinite
Cesium-137	137	1.00E+07	NA	3.60E-07	NA	NA	280	30.17
1,1-DCE	97	2.25E+03	9.00E-02	3.74E-06	2.61E-02	6.50E+01	0.013	Infinite
Dibenzo(a,h)-anthracene	278.33	2.50E-03	2.00E-02	1.86E-06	1.47E-08	1.78E+06	1,424	Infinite
Fluoranthene	202.26	2.06E-01	3.00E-02	2.29E-06	1.61E-05	4.91E+04	39.3	Infinite
Fluorene	166	1.90E+00	6.10E-02	2.84E-06	7.70E-05	7.90E+03	6.3	Infinite
Manganese	54.94	1.00E+07	NA	3.60E-07	NA	NA	65	Infinite
Mercury	200.59	6.00E-02	3.07E-02	2.27E-06	2.44E-02	NA	52	Infinite
Molybdenum	95.9	1.00E+07	NA	3.60E-07	NA	NA	10	Infinite
Naphthalene	128.16	3.10E+01	5.90E-02	2.70E-06	4.83E-04	1.19E+03	0.95	Infinite
Nickel	58.69	1.00E+07	NA	3.60E-07	NA	NA	300	Infinite
Plutonium-239	239	1.00E+07	NA	3.60E-07	NA	NA	550	2.41E+04
Pyrene	202.3	1.35E-01	2.72E-02	2.61E-06	1.10E-05	6.80E+04	54.4	Infinite
Selenium	80.98	1.00E+07	NA	3.60E-07	NA	NA	5	Infinite
Technetium-99	99	1.00E+07	NA	3.60E-07	NA	NA	0.2	2.13E+05
Tetrachloroethene	165.8	2.00E+02	7.20E-02	2.95E-06	1.84E-02	2.65E+02	0.053	Infinite
TCE	131	1.10E+03	8.00E-02	3.28E-06	1.03E-02	9.40E+01	0.0752	2.66E+01 ^b
Uranium-234	234	1.00E+07	NA	3.60E-07	NA	NA	66.8	2.44E+05
Uranium-235	235	1.00E+07	NA	3.60E-07	NA	NA	66.8	7.04E+08
Uranium-238	238	1.00E+07	NA	3.60E-07	NA	NA	66.8	4.47E+09
Uranium	238	1.00E+07	NA	3.60E-07	NA	NA	66.8	4.47E+09
Vanadium	50.94	1.00E+07	NA	3.60E-07	NA	NA	1,000	Infinite
Vinyl Chloride	63	2.76E+03	1.10E-01	4.43E-07	2.70E-02	1.88E+01	0.0152	Infinite
Zinc	67.41	1.00E+07	NA	3.60E-07	NA	NA	62	Infinite

^a The Kd of an organic compound depends on the soil's organic content (foc) and compound's organic partition coefficient (Koc). Kd values presented for organic compounds are for UCRS soils (with foc value of 0.08%) only. K_ds used in AT123D are different due to the foc of 0.02% in the RGA.

^b The 26.6 year half-life for TCE is applied to the UCRS only. The TCE half-life is still being researched by a working group and results will be utilized in the FS.

In general, all contaminants were assumed not to degrade in the environment (i.e., infinite half-life), except for radionuclides and TCE. Table 5.1 lists the half-lives assumed in the transport analyses for the analytes.

Although radionuclides behave chemically as metals, the radioactive nuclides undergo spontaneous transformations that involve the emission of particles (alpha and beta particles) and radiant energy (gamma energy). The resulting daughters (i.e., product nuclides) may be radioactive themselves (in which case they too will undergo spontaneous decay) or may be stable nuclides. Natural uranium consists of three primary isotopes: uranium-234, uranium-235, and uranium-238. The decay products of uranium isotopes also are radioactive and form decay chains.

Uranium hexafluoride is the sole raw material used in the enrichment process at PGDP. Some of the uranium feed material that was handled at PGDP has been reclaimed or recycled from reprocessed, spent reactor fuel. The chemical processes by which recycled uranium is purified leave trace amounts of transuranic elements (neptunium and plutonium) and fission products (mainly technetium-99 and cesium-137). Technetium-99 (in the +7 oxidation state) is highly soluble in groundwater and is very mobile (its K_d is similar to that of TCE). The groundwater plumes of TCE and technetium-99 at PGDP, particularly the Northwest Plume, have similarities; however, the technetium-99 plume does not currently exhibit technetium-99 concentrations above the MCL at off-site locations, whereas the TCE plumes do exceed MCLs off-site. Cesium-137 is not highly mobile and does not appear in PGDP plumes. Because cesium-137 has a half-life of 30 years, it is the most likely fission product (except for technetium-99) still to be present at the site.

On an activity basis, the principal radionuclides expected to pass through chemical processing and contaminate the recycled uranium are the transuranics neptunium-237, plutonium-238, plutonium-239, plutonium-240, and americium-241. Characterization studies (DOE 1999c) have shown that these radioisotopes are usually present in activities that are less than 1% of the uranium activity unless treatment processes have collected and concentrated them in sludges or trap material.

An assumption of the modeling for the BGOU RI was TCE degraded in the UCRS with a half-life of 26.6 years, but did not degrade in the RGA.² Although the mechanism is not well understood at PGDP, TCE and its degradation products may be degraded in the environment by various processes including hydrolysis, oxidation/reduction, photolysis, or biodegradation. TCE degradation may result in more toxic degradation products, such as vinyl chloride.

In the degradation of TCE, both aerobic and anaerobic degradation may occur. The anaerobic degradation pathway is as follows:



The anaerobic biodegradation of TCE, which initially forms *cis*-1,2-DCE, occurs under reducing conditions where sulfide- and/or methane-producing conditions exist. Such conditions occur primarily in the presence of other natural or anthropogenic carbon sources. The compounds *cis*- and *trans*-1,2-DCE are indicators of this degradation pathway because neither was used as a pure product at PGDP. The

² The assumption of zero degradation for TCE in the RGA is conservative. An alternative RGA degradation rate for TCE will be selected for use in the FS. The Kentucky Research Consortium for Energy and Environment, with the participation of DOE and its regulatory oversight, is researching TCE attenuation in the RGA at PGDP. PGDP modeling will incorporate these results as they become available. Recent findings (DOE 2007a) indicate the TCE half-life in the Northwest Plume of the RGA ranges from 3.2 to 11.3 years.

presence of *cis*- and *trans*-1,2-DCE also may have been the result of the use of industrial-grade TCE at PGDP. Both *cis*- and *trans*-1,2-DCE may further degrade anaerobically to vinyl chloride, but the rate is slower than the degradation rate of TCE, and the process may require stronger reducing conditions than those required for reduction of TCE. Low-levels of TCE intermediate dechlorination products (produced by anaerobic degradation) are found in RGA groundwater in some on-site locations. These occurrences may be related to degradation of TCE in the UCRS, where anaerobic conditions are known to occur locally.

The RGA is dominantly an aerobic environment. Aerobic biodegradation of TCE may occur under certain conditions. For example, specialized microorganisms have been identified that aerobically degrade some of these solvents in the presence of ammonia, methane, and toluene. In aerobic settings, TCE degrades to epoxides, aldehydes, chlorinated oxides, and ethanols.

Contaminant transport modeling simulates retardation during groundwater transport using indices of water solubility and adsorption to soil. In general, organic chemicals with high solubilities are more mobile in water than those that adsorb more strongly to soils. The following properties dictate an organic chemical's mobility within a specific medium.

- K_{oc} , the soil organic carbon partition coefficient, is a measure of the tendency for organic compounds to be adsorbed to the organic matter of soil and sediments. K_{oc} is expressed as the ratio of the amount of chemical adsorbed, per unit weight of organic carbon, to the chemical concentration in solution at equilibrium.
- K_{ow} , the octanol-water partition coefficient, is an indicator of hydrophobicity (the tendency of a chemical to avoid the aqueous phase) and is correlated with potential adsorption to soils. It also is used to estimate the potential for bioconcentration of chemicals into tissues.
- K_d , the soil/water distribution coefficient, is a measure of the tendency of a chemical to adsorb to soil or sediment particles. For organic compounds, this coefficient is calculated as the product of the K_{oc} value and the fraction of organic carbon in the soils. In general, chemicals with higher K_d values adsorb more strongly to soil/sediment particles and are less mobile than those with lower K_d values.

Release and transport mechanisms for TCE and its degradation products include vertical advective migration through unsaturated soils toward the water table, as well as gravity-driven migration as a DNAPL. The range of K_{oc} values indicates that these chlorinated VOCs are mobile through soils as dissolved constituents and tend not to partition significantly from water to soil.

Inorganic chemicals (i.e., metals) released to the unsaturated soil will be dissolved in soil moisture or absorbed onto soil particles. These dissolved metals are subject to movement with soil water. Aqueous transport mechanisms may result in metal migration through the vadose zone to groundwater. Metals, unlike organic compounds, cannot be degraded; however, metals migration can be attenuated by retardation reactions such as adsorption, surface complexation, and ion-exchange reactions with the soils which they contact. Such reactions are affected by pH, oxidation-reduction conditions, and the type and amount of organic matter, clay, and hydrous oxides present. Some metals, such as arsenic, can be transformed to other oxidation states in soil. Such transformations can affect their mobilities by affecting the way in which they react with soil particles or other solid surfaces by ion exchange, adsorption, precipitation, or complexation.

5.3 GROUNDWATER FATE AND TRANSPORT MODELING

Modeling for the BGOU RI used the Statistical Analysis and Decision Assistance (SADA), SESOIL, and AT123D models, consistent with Tier 3 of the modeling matrix in the PGDP Risk Methods Document (DOE 2001). SADA was used for the definition of the source terms, SESOIL for fate and transport modeling through the UCRS, and AT123D for fate and transport modeling through the RGA to the POEs. In addition to the models used, the MODFLOW/MODPATH models were used along with the previously developed PGDP sitewide groundwater model to establish input parameters for AT123D (i.e., distances to the POEs along flow paths (Figure 5.1), hydraulic gradient, and hydraulic conductivity). These models, along with the fixed parameter values chosen for the analyses (i.e., deterministic analysis), and model implementation are discussed in detail in Appendix E. The fate and transport modeling for the BGOU RI incorporates the sampling results of this RI and more sophisticated geospatial analysis of the source terms than those of previous models for these SWMUs; therefore, these model results differ from those of the previous models.

Modeling predicted the maximum concentration of analytes in groundwater at the boundary of each BGOU SWMU (Table 5.2). Table 5.3 presents the results of the deterministic modeling effort for the BGOU RI for the plant boundary and off-site POEs. Among the analytes, arsenic, technetium-99, and TCE and related VOCs commonly exceeded MCLs. Table 5.4 presents the hazard quotient (HQ) and estimated lifetime cancer risk (ELCR) for each analyte based on the predicted groundwater concentrations at the plant boundary and off-site POEs. The HQs and ELCRs were calculated in accordance with the Risk Methods Document (DOE 2001). Appendix F provides a full description of the risk assessment methodology and calculations. The following discussion summarizes the results for each BGOU SWMU.

Table 5.2. Concentrations of the Analytes in Groundwater at the BGOU SWMU Boundaries Predicted in SESOIL and AT123D Modeling

Analyte	Predicted Maximum Groundwater Concentration (mg/L or pCi/L) ^a	MCL (mg/L or pCi/L)
SWMU 2		
Arsenic	3.54E-02	0.01
<i>cis</i> -1,2-DCE	1.15E+01	0.07
Manganese	7.16E-01	^b
Naphthalene	9.38E-04	^b
PCB-1254	1.54E-03	^b
PCB-1260	8.73E-05	^b
Technetium-99	1.02E+02	900 ^c
TCE	1.48E+00	0.005
Uranium-234	1.58E+00	20 ^d
Uranium-238	1.81E+00	20 ^d
Uranium	9.86E-03	0.03
SWMU 3		
Arsenic	3.29E-02	0.01
Manganese	8.95E-01	^b
Technetium-99	5.560E+03	900 ^c
Uranium-238	1.59E+01	20 ^d
Uranium	4.89E-02	0.03
SWMU 4		
Arsenic	1.77E-02	0.01
<i>cis</i> -1,2-DCE	6.68E-01	0.07
Manganese	5.76E-01	^b
Technetium-99	9.008E+03	900 ^c
TCE	1.18E+00	0.005
Vinyl Chloride	2.61E-02	0.002

Table 5.2. Concentrations of the Analytes in Groundwater at the BGOU SWMU Boundaries Predicted in SESOIL and AT123D Modeling (Continued)

Analyte	Predicted Maximum Groundwater Concentration (mg/L or pCi/L) ^a	MCL (mg/L or pCi/L)
SWMU 5		
Acenaphthene	6.10E-03	b
Arsenic	9.25E-03	0.01
Manganese	1.01E+00	b
Naphthalene	5.55E-03	b
Technetium-99	1.27E+02	900 ^c
Uranium	<i>4.60E-01</i>	0.03
SWMU 6		
Manganese	8.32E-02	b
SWMU 7		
1,1-DCE	<i>8.98E-02</i>	0.07
Arsenic	<i>1.78E-02</i>	0.01
<i>cis</i> -1,2-DCE	2.35E-02	0.07
Manganese	3.32E-01	b
PCB-1254	5.23E-05	b
Technetium-99	<i>9.09E+02</i>	900 ^c
TCE	<i>1.09E-02</i>	0.005
Uranium-234	7.94E+00	20 ^d
Uranium-238	7.59E+00	20 ^d
Uranium	3.46E-03	0.03
Vinyl Chloride	<i>1.35E-02</i>	0.002
SWMU 30		
1,1-DCE	8.18E-05	0.07
Arsenic	<i>1.82E-02</i>	0.01
Manganese	3.78E-01	b
Selenium	1.51E-02	0.05
Technetium-99	2.87E+02	900 ^c
TCE	9.11E-04	0.005
Uranium-234	3.99E+00	20 ^d
Uranium-238	5.91E+00	20 ^d
Uranium	8.40E-03	0.03
SWMU 145		
Antimony	<i>7.99E-02</i>	0.006
Arsenic	<i>6.21E-02</i>	0.01
PCB-1260	1.92E-03	
Technetium-99	<i>1.01E+04</i>	900 ^c
Manganese	8.44E-01	b
Uranium-238	7.67E-02	20 ^d

^a Values in bold, italic font exceed the analytes maximum contaminant level (MCL)

^b MCLs not available for these contaminants

^c Technetium-99 MCL based on a critical organ dose at 4 mrem/yr from drinking water consumption

^d The MCLs for U-234 and U-238 are from Table A.20 of the Risk Methods Document (DOE 2001)

Table 5.3. Concentrations of the Analytes in Groundwater Predicted in SESOIL and AT123D Modeling of the BGOU SWMUs

Analyte	Predicted Maximum Groundwater Concentration ^{a,b}				
	Plant Boundary (mg/L)	Property Boundary (mg/L)	Little Bayou seeps (mg/L)	Ohio River (mg/L)	MCL (mg/L or pCi/L)
SWMU 2					
Arsenic	2.91E-03	8.35E-09	N/A	0.00E+00	0.01
<i>cis</i> -1,2-DCE	1.74E+00	8.58E-01	N/A	3.38E-01	0.07
Manganese	1.86E-05	0.00E+00	N/A	0.00E+00	^c
Naphthalene	1.57E-04	8.27E-05	N/A	3.42E-05	^c
PCB-1248	1.28E-09	0.00E+00	N/A	0.00E+00	^c
PCB-1260	0.00E+00	0.00E+00	N/A	0.00E+00	^c
Technetium-99	1.59E+01	8.06E+00	N/A	3.11E+00	900 ^d
TCE	2.17E-01	1.10E-01	N/A	4.12E-02	0.005
Uranium-234	1.75E-05	0.00E+00	N/A	0.00E+00	20 ^e
Uranium-238	2.03E-05	0.00E+00	N/A	0.00E+00	20 ^e
Uranium	8.33E-08	0.00E+00	N/A	0.00E+00	0.03
SWMU 3					
Arsenic	1.22E-03	0.00E+00	0.00E+00	N/A	0.01
Manganese	4.08E-10	0.00E+00	0.00E+00	N/A	^c
Technetium-99	1.81E+03	1.36E+03	8.04E+02	N/A	900 ^d
Uranium-238	1.59E+01	7.32E-11	0.00E+00	N/A	20 ^e
Uranium	2.27E-13	0.00E+00	0.00E+00	N/A	0.03
SWMU 4					
Arsenic	2.70E-03	4.89E-06	N/A	0.00E+00	0.01
<i>cis</i> -1,2-DCE	1.96E-01	8.94E-02	N/A	3.16E-02	0.07
Manganese	5.01E-03	0.00E+00	N/A	0.00E+00	^c
Technetium-99	2.50E+03	1.20E+03	N/A	3.79E+02	900 ^d
TCE	4.22E-01	2.14E-01	N/A	7.67E-02	0.005
Vinyl Chloride	5.95E-03	2.53E-03	N/A	7.82E-04	0.002
SWMU 5					
Acenaphthene	2.42E-03	1.34E-03	N/A	5.01E-04	NA
Arsenic	1.78E-03	1.27E-04	N/A	0.00E+00	0.01
Manganese	8.69E-02	2.30E-11	N/A	0.00E+00	^c
Naphthalene	9.82E-04	3.72E-04	N/A	1.08E-04	NA
Technetium-99	4.99E+01	2.64E+01	N/A	8.72E+00	900 ^d
Uranium	3.32E-02	4.65E-11	N/A	0.00E+00	0.03
SWMU 6					
Manganese	1.17E-02	2.890E-04	0.00E+00	N/A	^c
SWMU 7					
1,1-DCE	8.24E-02	1.10E-02	4.02E-03	N/A	0.07
Arsenic	1.26E-02	2.35E-03	0.00E+00	N/A	0.01
<i>cis</i> -1,2-DCE	2.15E-02	3.13E-03	1.17E-03	N/A	0.07
Manganese	2.41E-01	1.05E-06	0.00E+00	N/A	^c
PCB-1254	3.09E-05	3.05E-06	1.32E-12	N/A	^c
Technetium-99	8.25E+02	2.70E+02	1.32E+02	N/A	900 ^d
TCE	9.87E-03	1.42E-03	5.06E-04	N/A	0.005
Uranium-234	5.79E+00	5.84E-06	0.00E+00	N/A	20 ^e
Uranium-238	5.58E+00	5.85E-06	0.00E+00	N/A	20 ^e
Uranium	2.53E-03	2.68E-09	0.00E+00	N/A	0.03
Vinyl Chloride	1.24E-02	1.21E-03	4.13E-04	N/A	0.002

Table 5.3. Concentrations of the Analytes in Groundwater Predicted in SESOIL and AT123D Modeling of the BGOU SWMUs (Continued)

Analyte	Predicted Maximum Groundwater Concentration ^{a,b}				
	Plant Boundary (mg/L)	Property Boundary (mg/L)	Little Bayou seeps (mg/L)	Ohio River (mg/L)	MCL (mg/L or pCi/L)
SWMU 30					
1,1-DCE	7.65E-05	6.14E-06	1.86E-06	N/A	0.07
Arsenic	<i>1.21E-02</i>	2.50E-03	0.00E+00	N/A	0.01
Manganese	2.51E-01	2.85E-04	0.00E+00	N/A	^c
Selenium	8.30E-03	9.21E-04	3.15E-04	N/A	0.05
Technetium-99	2.64E+02	7.08E+01	2.92E+01	N/A	900 ^d
TCE	8.60E-04	7.70E-05	2.60E-05	N/A	0.005
Uranium-234	2.75E+00	1.44E-03	0.00E+00	N/A	20 ^e
Uranium-238	4.07E+00	1.98E-03	0.00E+00	N/A	20 ^e
Uranium	4.81E-03	2.41E-06	0.00E+00	N/A	0.03
SWMU 145					
Antimony	N/A	1.51E-06	N/A	0.00E+00	0.006
Arsenic	N/A	1.61E-03	N/A	0.00E+00	0.01
PCB-1260	N/A	0.00E+00	N/A	0.00E+00	^c
Manganese	N/A	0.00E+00	N/A	0.00E+00	^c
Technetium-99	N/A	<i>1.84E+03</i>	N/A	<i>9.65E+02</i>	900 ^d
Uranium-238	N/A	0.00E+00	N/A	0.00E+00	20 ^e

^a Values in bold, italic font exceed the analyte's maximum contaminant level (MCL).

^b Radionuclide concentrations are in pCi/L.

^c MCLs not available for these contaminants.

^d Technetium-99 MCL based on a critical organ dose at 4 mrem/yr from drinking water consumption.

^e The MCLs for U-234 and U-238 are from Table A.20 of the Risk Methods Document (DOE 2001).

N/A = The point of exposure is not applicable to the groundwater pathway for this SWMU.

Table 5.4. Hazard and Cancer Risk Predicted from Maximum Groundwater Concentrations Derived in Modeling of BGOU SWMUs using SESOIL and AT123D^a

Analyte	Plant Boundary		Property Boundary		Little Bayou Seeps		Near Ohio River	
	Hazard Quotient	Cancer Risk	Hazard Quotient	Cancer Risk	Hazard Quotient	Cancer Risk	Hazard Quotient	Cancer Risk
SWMU 2								
Arsenic	0.9	7.7E-05	<0.1	<1.0E-06	N/A	N/A	b	b
<i>cis</i> -1,2-DCE	91.9	b	45.3	b	N/A	N/A	17.9	b
Manganese	<0.1	b	b	b	N/A	N/A	b	b
Naphthalene	0.1	b	<0.1	b	N/A	N/A	<0.1	b
PCB-1248	b	<1.0E-06	b	b	N/A	N/A	b	b
PCB-1260	b	b	b	b	N/A	N/A	b	b
Technetium-99	b	<1.0E-06	b	<1.0E-06	N/A	N/A	b	<1.0E-06
TCE	99.1	6.7E-03	50.3	3.4E-03	N/A	N/A	4.6	1.3E-03
Uranium-234	b	<1.0E-06	b	b	N/A	N/A	b	b
Uranium-238	b	<1.0E-06	b	b	N/A	N/A	b	b
Uranium	0.1	b	b	b	N/A	N/A	b	b
SWMU 3								
Arsenic	0.4	3.2E-05	b	b	b	b	N/A	N/A
Manganese	<0.1	b	b	b	b	b	N/A	N/A
Technetium-99	b	9.9E-05	b	7.5E-05	b	b	N/A	N/A
Uranium-238	b	<1.0E-06	b	b	b	b	N/A	N/A
Uranium	<0.1	b	b	b	b	b	N/A	N/A
SWMU 4								
Arsenic	0.9	7.2E-05	<0.1	<1.0E-06	N/A	N/A	b	b
<i>cis</i> -1,2-DCE	10.4	b	4.7	b	N/A	N/A	0.6	b
Manganese	<0.1	b	b	b	N/A	N/A	b	b
Technetium-99	b	1.4E-04	b	6.6E-05	N/A	N/A	b	2.1E-05
TCE	193	2.0E-02	97.7	6.6E-03	N/A	N/A	32.7	2.4E-03
Vinyl Chloride	0.3	1.9E-04	0.1	7.4E-05	N/A	N/A	<0.1	2.3E-05
SWMU 5								
Arsenic	0.6	4.7E-05	<0.1	3.4E-06	N/A	N/A	b	b
Manganese	0.2	b	<0.1	b	N/A	N/A	b	b
Naphthalene	0.5	b	0.2	b	N/A	N/A	<0.1	b
Technetium-99	b	2.7E-06	b	1.4E-06	N/A	N/A	b	<1.0E-06
Uranium	5.31	b	<0.1	b	N/A	N/A	b	b
SWMU 6								
Manganese	<0.1	b	b	b	b	b	N/A	N/A

Table 5.4. Hazard and Cancer Risk Predicted from Maximum Groundwater Concentrations Derived in Modeling of BGOU SWMUs using SESOIL and AT123D^a (Continued)

Analyte	Plant Boundary		Property Boundary		Little Bayou Seeps		Near Ohio River	
	Hazard Quotient	Cancer Risk	Hazard Quotient	Cancer Risk	Hazard Quotient	Cancer Risk	Hazard Quotient	Cancer Risk
SWMU 7								
1,1-DCE	0.8	1.9E-03	0.1	2.5E-04	<0.1	9.3E-05	N/A	N/A
Arsenic	4.0	3.3E-04 ^b	0.8	6.2E-05 ^b	^b	^b	N/A	N/A
<i>cis</i> -1,2-DCE	1.1	^b	0.2	^b	<0.1	^b	N/A	N/A
Manganese	0.5	^b	<0.1	^b	^b	^b	N/A	N/A
PCB-1254	2.5	4.8E-06	0.2	<1.0E-06	<0.1	<1.0E-06	N/A	N/A
Technetium-99	^b	4.5E-05	^b	1.5E-05	^b	7.3E-06	N/A	N/A
TCE	4.5	3.1E-04	0.6	4.4E-05	0.2	1.6E-05 ^b	N/A	N/A
Uranium-234	^b	8.2E-06	^b	<1.0E-06	^b	^b	N/A	N/A
Uranium-238	^b	9.7E-06 ^b	^b	<1.0E-06 ^b	^b	^b	N/A	N/A
Uranium	0.4	^b	<0.1	^b	^b	^b	N/A	N/A
Vinyl Chloride	0.6	3.6E-04	<0.1	3.6E-05	<0.1	1.2E-05	N/A	N/A
SWMU 30								
1,1-DCE	<0.1	1.8E-06	<0.1	<1.0E-06	<0.1	<1.0E-06	N/A	N/A
Arsenic	3.9	3.2E-04 ^b	0.8	6.2E-05 ^b	^b	^b	N/A	N/A
Manganese	0.5	^b	<0.1	^b	^b	^b	N/A	N/A
Selenium	0.2	^b	<0.1	^b	<0.1	^b	N/A	N/A
Technetium-99	^b	1.4E-05	^b	3.9E-06	^b	1.6E-06	N/A	N/A
TCE	0.4	2.6E-05	<0.1	2.4E-06	<0.1	<1.0E-06 ^b	N/A	N/A
Uranium-234	^b	3.9E-06	^b	<1.0E-06	^b	^b	N/A	N/A
Uranium-238	^b	7.1E-06 ^b	^b	<1.0E-06 ^b	^b	^b	N/A	N/A
Uranium	0.8	^b	<0.1	^b	^b	^b	N/A	N/A
SWMU 145								
Antimony	N/A	N/A	<0.1	^b	N/A	N/A	^b	^b
Arsenic	N/A	N/A	0.5	4.3E-05	N/A	N/A	^b	^b
Technetium-99	N/A	N/A	^b	1.0E-04	N/A	N/A	^b	5.3E-05

^a Contaminants with an HQ greater than 0.1 or a cancer risk greater than 1.00E-06 are considered analytes—all values are rounded to one decimal place.

^b Value not calculated since the groundwater concentrations were predicted as zero at this point of exposure (POE) by AT123D, or the contaminant did not have a reported cancer slope factor or chemical toxicity reference dose (RfD).

N/A = The point of exposure is not applicable to the groundwater pathway for this SWMU

5.3.1 SWMU 2

Analytes retained after the screening process described in Section 5.2.1 are presented for SWMU 2 in Table E3.1. Each analyte was modeled and the resulting peak groundwater concentrations below SWMU 2 shown in Table 5.5 then were compared to the child resident NALs from Table A.18 of the 2001 Risk Methods Document and the provisional groundwater backgrounds shown in the 2001 Risk Method Document in Table A.13. Those analytes with groundwater concentrations below SWMU 2 that exceeded the NALs or background values then were carried through the toxicity and exposure assessments and cancer risk and hazards were calculated for the Rural Resident Groundwater User. Antimony, mercury, nickel, vanadium, PCB-1254, benzo(a)pyrene, plutonium-239, and PCB-1254 were modeled and found not to reach the RGA during the 1,000 year modeling period.

Table 5.5. Screening of Modeled Peak Concentrations in Groundwater for SWMU 2

Analyte	Time (years)	Peak Conc. Below SWMU 2 (mg/L) ^a	Background Groundwater Concentration (mg/L) ^a	Groundwater Child Resident No Action Level (mg/L) ^a	Retain?
Arsenic	9.95E+02	3.54E-02	5.00E-03	3.50E-05	Y
<i>cis</i> -1,2-DCE	1.50E+01	1.15E+01	NA	2.73E-03	Y
Manganese	9.90E+02	7.16E-01	1.19E-01	3.50E-02	Y
Naphthalene	1.40E+02	9.38E-04	NA	2.85E-04	Y
PCB-1248	1.00E+03	1.54E-03	NA	5.74E-02	Y
PCB-1260	1.00E+03	8.73E-05	NA	5.74E-02	Y
TCE	2.00E+01	1.48E+00	NA	1.60E-03	Y
Uranium	1.00E+03	9.86E-03	2.00E-03	9.06E-04	Y
Zinc	9.90E+02	9.83E-03	4.90E-02	4.50E-01	N
Technetium-99	4.00E+01	1.02E+02	2.23E+01	1.40E+01	Y
Uranium-234	9.85E+02	1.58E+00	7.00E-01	5.46E-01	Y
Uranium-238	9.90E+02	1.81E+00	7.00E-01	4.43E-01	Y

^a Units for radionuclides are pCi/L.

NA = not applicable N = No Y = Yes

The groundwater results presented in Table 5.3 for SWMU 2 show that the predicted groundwater concentrations of *cis*-1,2-DCE and TCE exceed their respective MCLs at the plant boundary, property boundary, and Ohio River POEs. All the remaining SWMU 2 analytes are not predicted to exceed their respective MCLs at the POEs. The following summarizes those analytes that exceeded ELCR and HQ risk criteria of 1.0E-06 and 0.1, respectively.

		Plant Boundary	Property Boundary	Ohio River
Arsenic	ELCR	7.7E-05	--	--
	HQ	0.9	--	--
<i>cis</i> -1,2-DCE	ELCR	--	--	--
	HQ	91.9	45.3	17.9
TCE	ELCR	6.7E-03	3.4E-03	1.3E-03
	HQ	99.1	50.3	4.6

-- = does not exceed

All remaining analytes exhibited HQ values less than 0.1 and ELCR values less than 1.0E-06 at all POEs. Figures 5.2 through 5.4 present the predicted concentrations over time of SWMU 2 analytes that exceed a HQ of 0.1 and/or an ELCR of 1.0E-06 (Table 5.4). As shown in these figures, arsenic is predicted to continue rising in concentration at 1,000 years at the plant boundary, will not reach the property boundary or Ohio River in the 1,000 year period. Both *cis*-1,2-DCE and TCE are predicted to exceed their MCLs at all POEs within approximately 100 years and then decline in concentration below the MCLs.

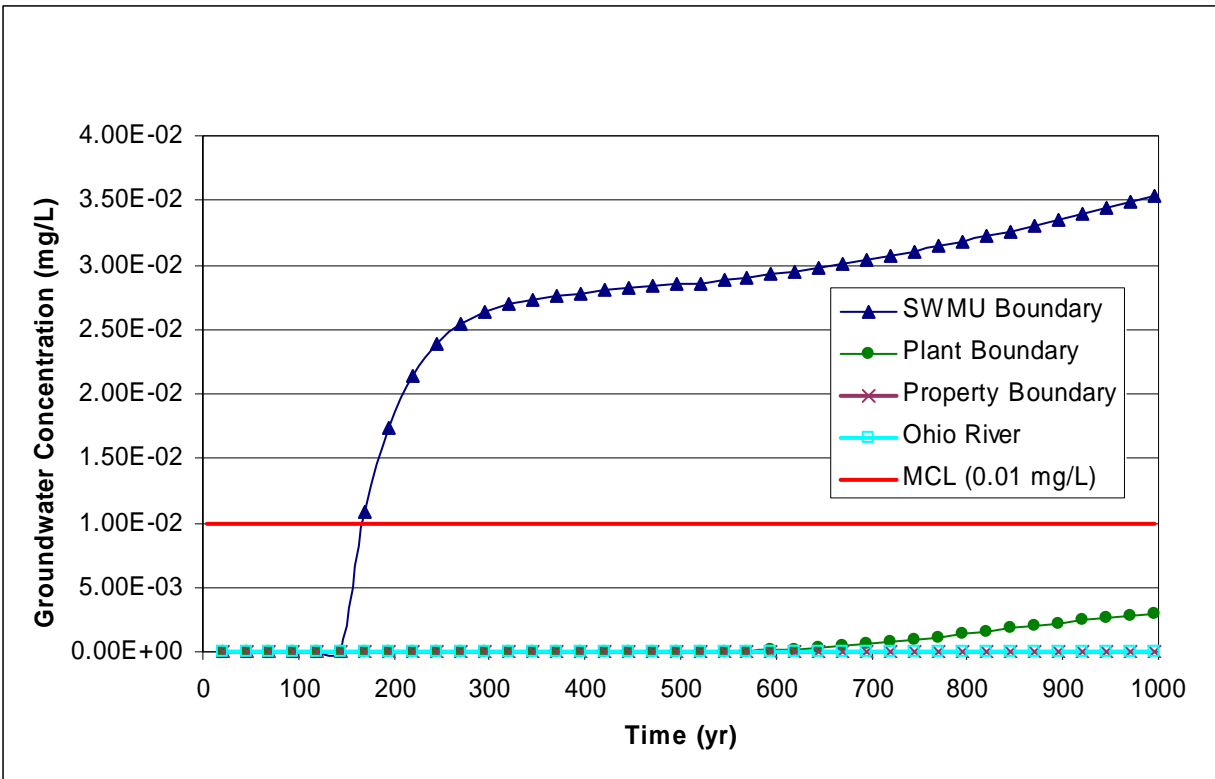


Figure 5.2. Predicted Arsenic Concentration in Groundwater at the POEs Based on Contaminant Leaching from SWMU 2

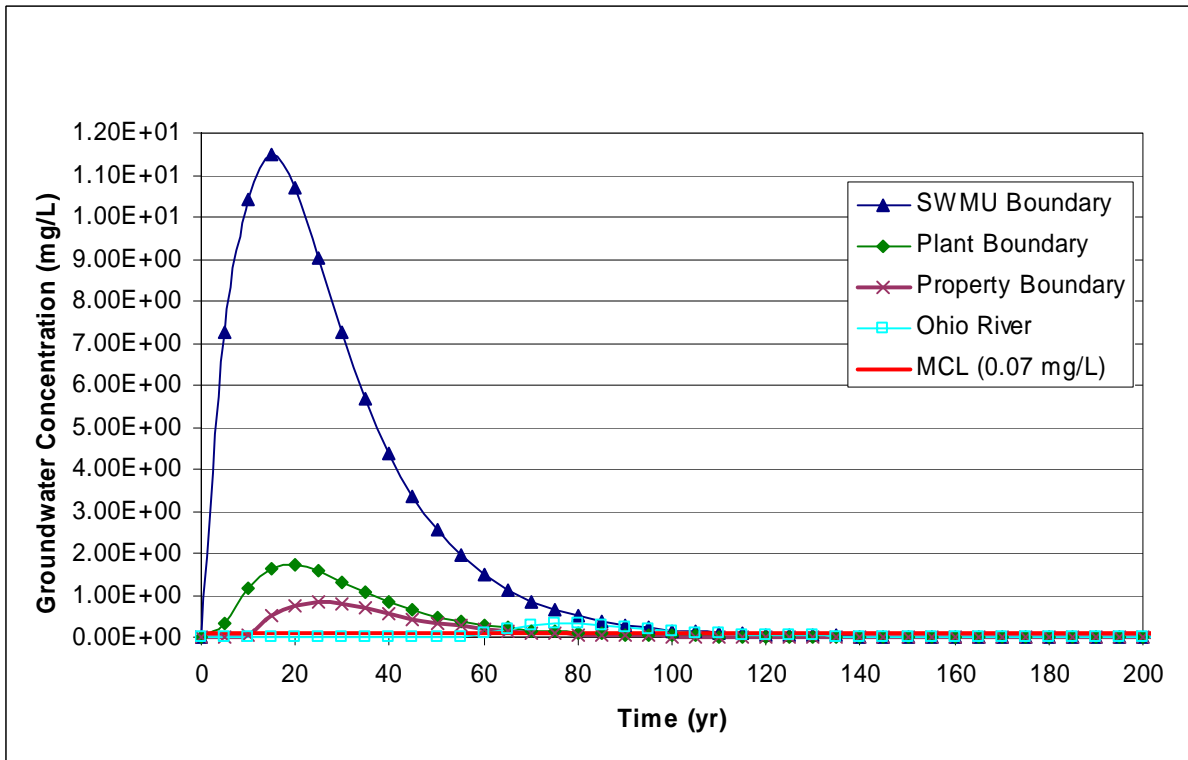


Figure 5.3. Predicted *cis*-1,2-DCE Concentration in Groundwater at the POEs Based on Contaminant Leaching from SWMU 2

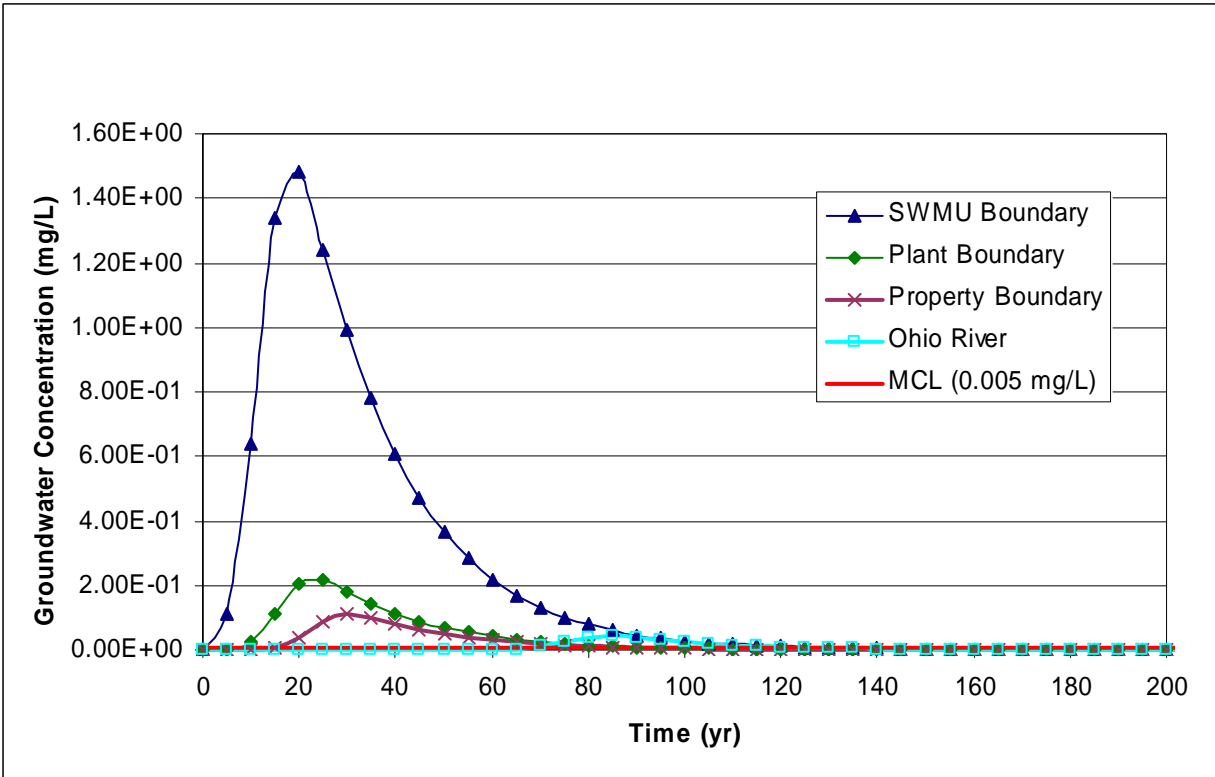


Figure 5.4. Predicted TCE Concentration in Groundwater at the POEs Based on Contaminant Leaching from SWMU 2

5.3.2 SWMU 3

Analytes retained after the screening process described in Section 5.2.1 are presented for SWMU 3 in Table E3.2. Each analyte was modeled and the resulting peak groundwater concentrations below SWMU 3 shown in Table 5.6 then were compared to the child resident NALs from Table A.18 of the 2001 Risk Methods Document and the provisional groundwater backgrounds shown in the 2001 Risk Method Document in Table A.13. Those analytes with groundwater concentrations below SWMU 3 that exceeded the NALs or background values then were carried through the toxicity and exposure assessments and cancer risk and hazards were calculated for the Rural Resident Groundwater User. Molybdenum, nickel and vanadium were modeled and found not to reach the RGA during the 1,000 year modeling period.

Table 5.6. Screening of Modeled Peak Concentrations in Groundwater for SWMU 3

Analyte	Time (years)	Peak Conc. Below SWMU 3 (mg/L) ^a	Background Groundwater Concentration (mg/L) ^a	Groundwater Child Resident No Action Level (mg/L) ^a	Retain?
Arsenic	9.80E+02	3.29E-02	5.00E-03	3.50E-05	Y
Manganese	9.70E+02	8.95E-01	1.19E-01	3.50E-02	Y
Mercury	9.80E+02	9.29E-05	2.00E-04	4.44E-04	N
TCE	3.50E+01	3.45E-04	NA	1.60E-03	N
Uranium	8.75E+02	4.89E-02	2.00E-03	9.06E-04	Y
Zinc	9.55E+02	9.30E-02	4.90E-02	4.50E-01	N
Technetium-99	4.50E+01	5.56E+03	2.23E+01	1.40E+01	Y
Uranium-238	9.90E+02	1.59E+01	7.00E-01	4.43E-01	Y

^a Units for radionuclides are pCi/L; NA = not applicable; N = No; Y = Yes

Screening identified arsenic, manganese, technetium-99, and uranium as analytes for SWMU 3. Fate and transport modeling predicts that technetium-99 will exceed the MCL at the plant and property boundary POEs. The following summarizes those analytes that exceeded ELCR or HQ risk criteria of 1.0E-06 and 0.1, respectively.

		Plant Boundary	Property Boundary	Little Bayou Seeps
Arsenic	ELCR	3.2E-05	--	--
	HQ	0.4	--	--
Technetium-99	ELCR	9.9E-05	7.5E-05	4.4E-05
	HQ	--	--	--

-- = does not exceed

All remaining analytes exhibited HQ values less than 0.1 and ELCR values less than 1.0E-06 at the POEs. Figures 5.5 and 5.6 illustrate the predicted concentrations through time of SWMU 3 analytes that exceed a HQ of 0.1 and/or an ELCR of 1.0E-06 (Table 5.4). As shown in these figures, arsenic is predicted to continue rising in concentration at 1,000 years at the plant boundary at groundwater concentrations less than the MCL, but will not reach the property boundary or Little Bayou Seeps in the 1,000 year period. Technetium-99 is predicted to peak at all POEs within 200 years and at dissolved levels greater than the MCL at the plant and property POEs.

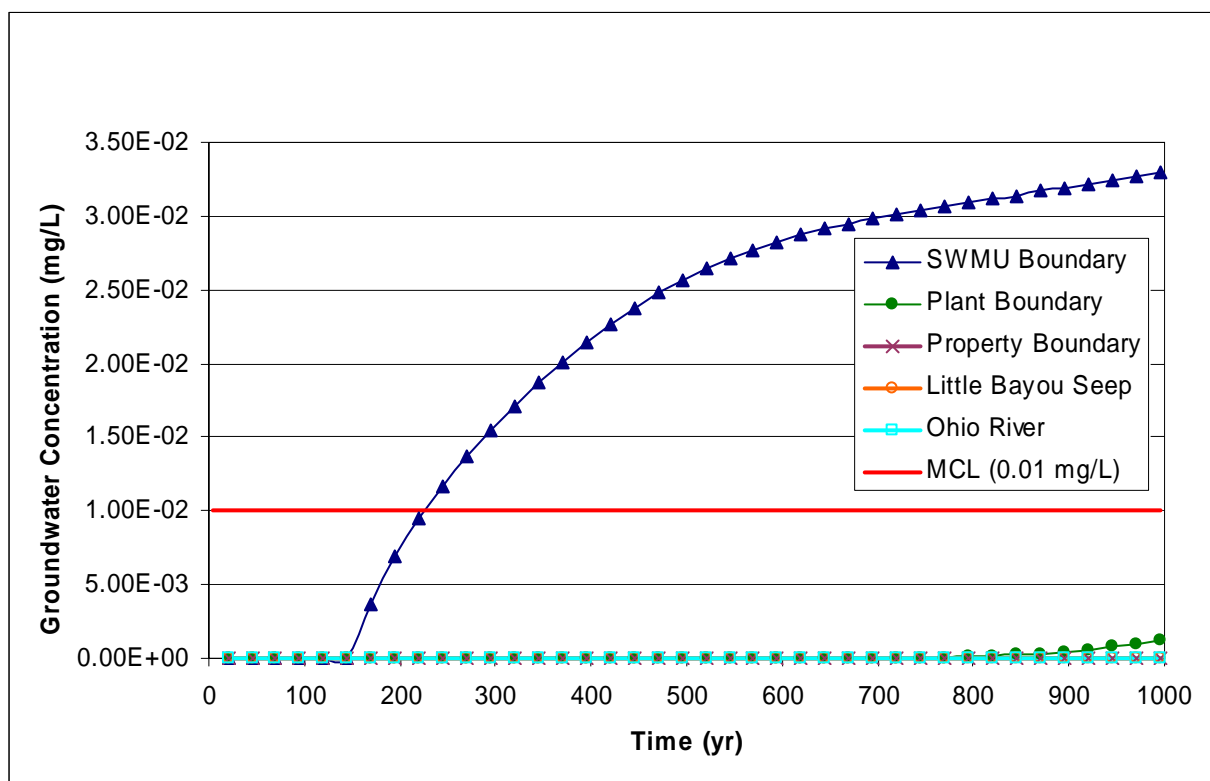


Figure 5.5. Predicted Arsenic Concentrations in Groundwater at the POEs Based on Contaminant Leaching from SWMU 3

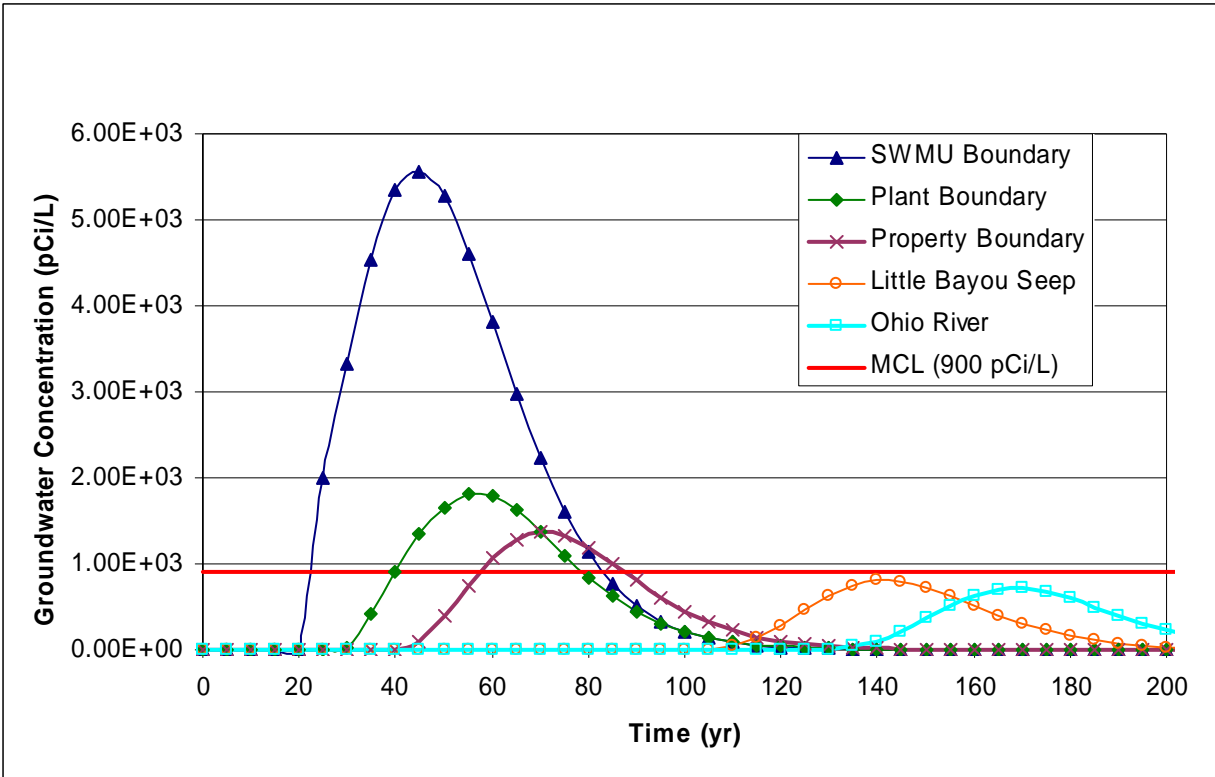


Figure 5.6. Predicted ⁹⁹Tc Activities in Groundwater at the POEs Based on Contaminant Leaching from SWMU 3

5.3.3 SWMU 4

Analytes retained after the screening process described in Section 5.2.1 are presented for SWMU 4 in Table E3.3. Each analyte was modeled and the resulting peak groundwater concentrations below SWMU 4 shown in Table 5.7 then were compared to the child resident NALs from Table A.18 of the 2001 Risk Methods Document and the provisional groundwater backgrounds shown in the 2001 Risk Method Document in Table A.13. Those analytes with groundwater concentrations below SWMU 4 that exceeded the NALs or background values then were carried through the toxicity and exposure assessments and cancer risk and hazards were calculated for the Rural Resident Groundwater User. Antimony, nickel, uranium, vanadium, plutonium-239, uranium-234 and uranium-238 were modeled and found not to reach the RGA during the 1,000 year modeling period.

Table 5.7. Screening of Modeled Peak Concentrations in Groundwater for SWMU 4

Analyte	Time (years)	Peak Conc. Below SWMU 4 (mg/L) ^a	Background Groundwater Concentration (mg/L) ^a	Groundwater Child Resident No Action Level (mg/L) ^a	Retain?
Arsenic	9.90E+02	1.77E-02	5.00E-03	3.50E-05	Y
cis-1,2-DCE	1.50E+01	6.68E-01	NA	2.73E-03	Y
Manganese	1.00E+03	5.76E-01	1.19E-01	3.50E-02	Y
TCE	5.00E+00	1.18E+00	NA	1.60E-03	Y
Zinc	1.00E+03	1.57E-09	4.90E-02	4.50E-01	N
Vinyl Chloride	5.00E+00	2.61E-02	NA	3.50E-05	Y
Technetium-99	5.00E+01	9.00E+03	2.23E+01	1.40E+01	Y

^a Units for radionuclides are pCi/L.

NA = not applicable

N = No

Y = Yes

The groundwater results presented in Table 5.3 for SWMU 4 show that the predicted groundwater concentrations of *cis*-1,2-DCE; technetium-99; TCE, and vinyl chloride will exceed their respective MCLs at the plant boundary and property boundary. TCE also is predicted to exceed the MCL at the Ohio River. The following summarizes those analytes that exceeded ELCR or HQ risk criteria of 1.0E-06 and 0.1, respectively.

		Plant Boundary	Property Boundary	Ohio River
Arsenic	ELCR	7.2E-05	--	--
	HQ	0.9	--	--
<i>cis</i> -1,2-DCE	ELCR	--	--	--
	HQ	10.4	4.7	0.6
Technetium-99	ELCR	1.4E-04	6.6E-05	2.1E-05
	HQ	--	--	--
TCE	ELCR	2.0E-02	6.6E-03	2.4E-03
	HQ	193	97.7	32.7
Vinyl chloride	ELCR	1.9E-04	7.4E-05	2.3E-05
	HQ	0.3	0.1	--

-- = does not exceed

All remaining analytes exhibited HQ values less than 0.1 and ELCR values less than 1.0E-06 at all POEs. Figures 5.7 through 5.11 portray the modeled concentrations over time of SWMU 4 analytes that exceed a HQ of 0.1 and/or an ELCR of 1.0E-06 (Table 5.4). As shown in these figures, the dissolved arsenic concentration is predicted to continue rising at 1,000 years at the plant boundary, but will not reach the property boundary or Ohio River in the 1,000 year period. The chemicals *cis*-1,2-DCE; TCE; vinyl chloride, and technetium-99 are predicted to exceed the MCL at the plant and property boundaries within 100 years. Modeling predicts TCE also will exceed the MCL at the Ohio River within 100 years.

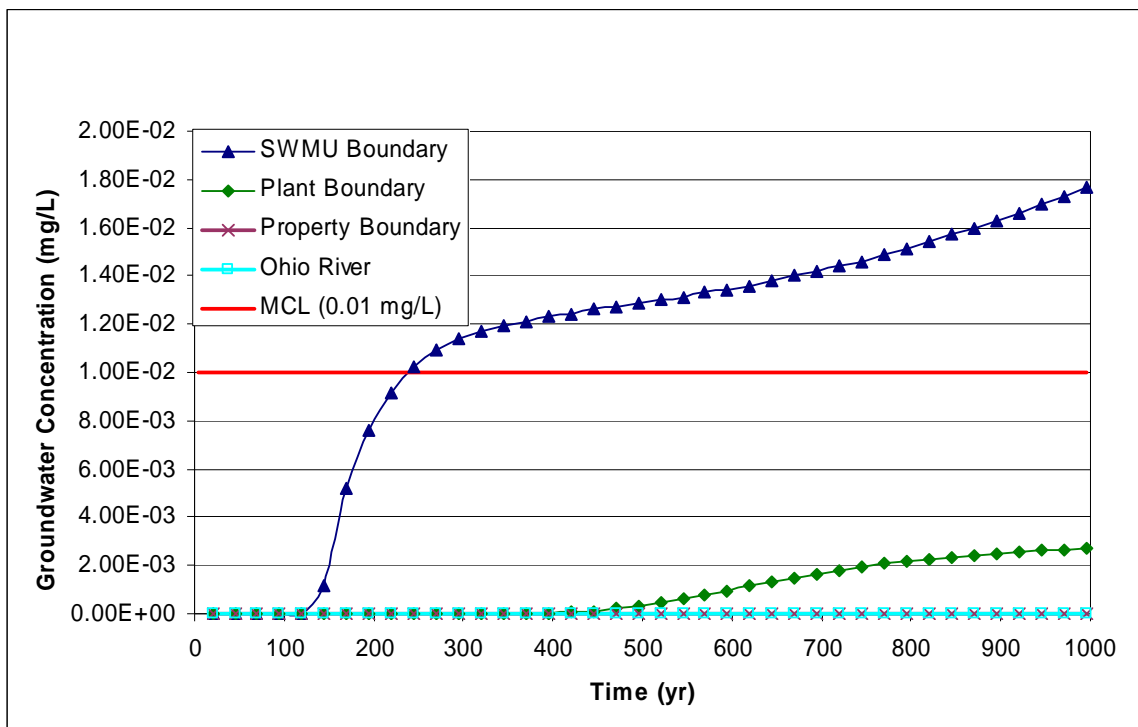


Figure 5.7. Predicted Arsenic Concentrations in Groundwater at the POEs Based on Contaminant Leaching from SWMU 4

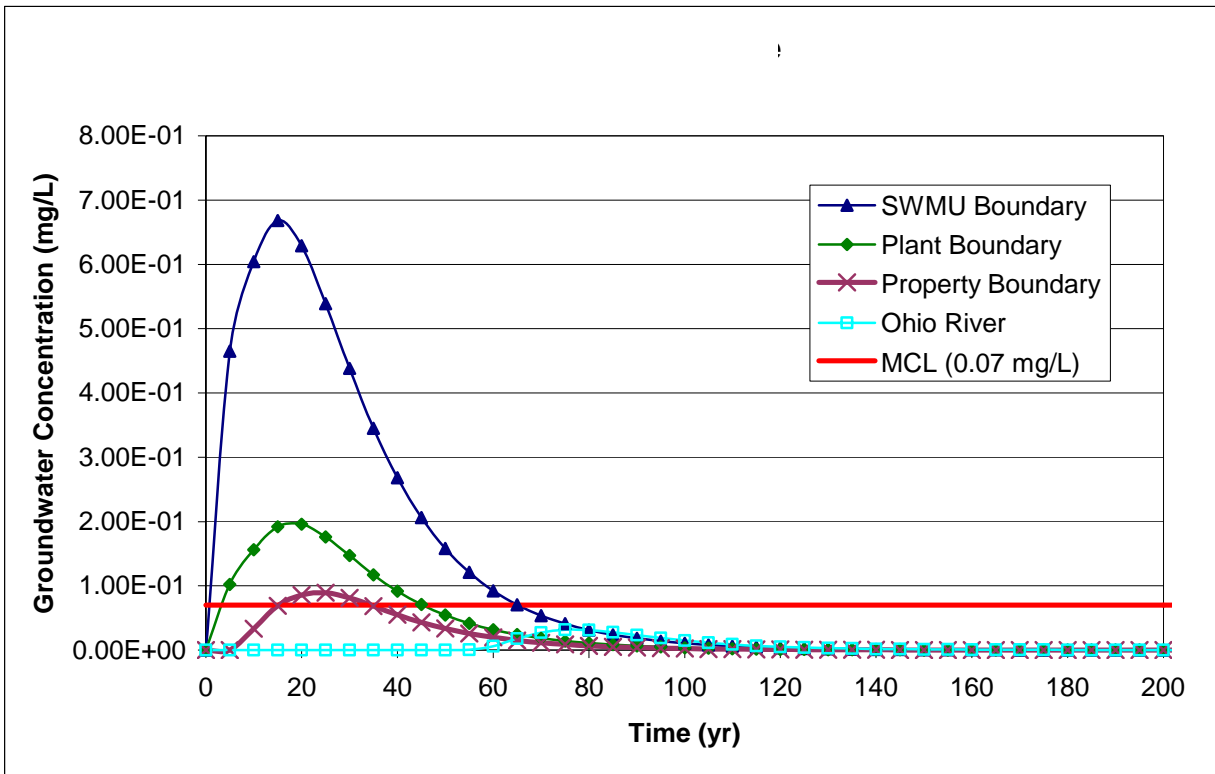


Figure 5.8. Predicted *cis*-1,2-DCE Concentrations in Groundwater at the POEs Based on Contaminant Leaching from SWMU 4

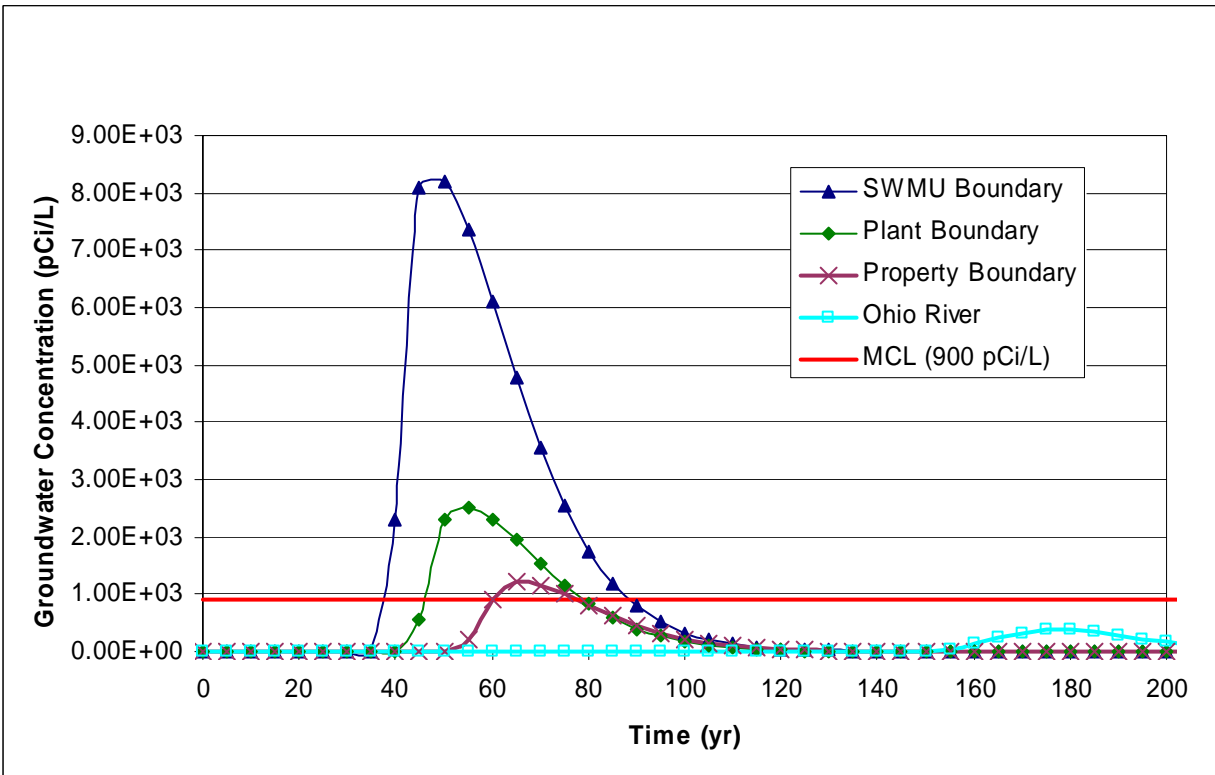


Figure 5.9. Predicted ⁹⁹Tc Activities in Groundwater at the POEs Based on Contaminant Leaching from SWMU 4

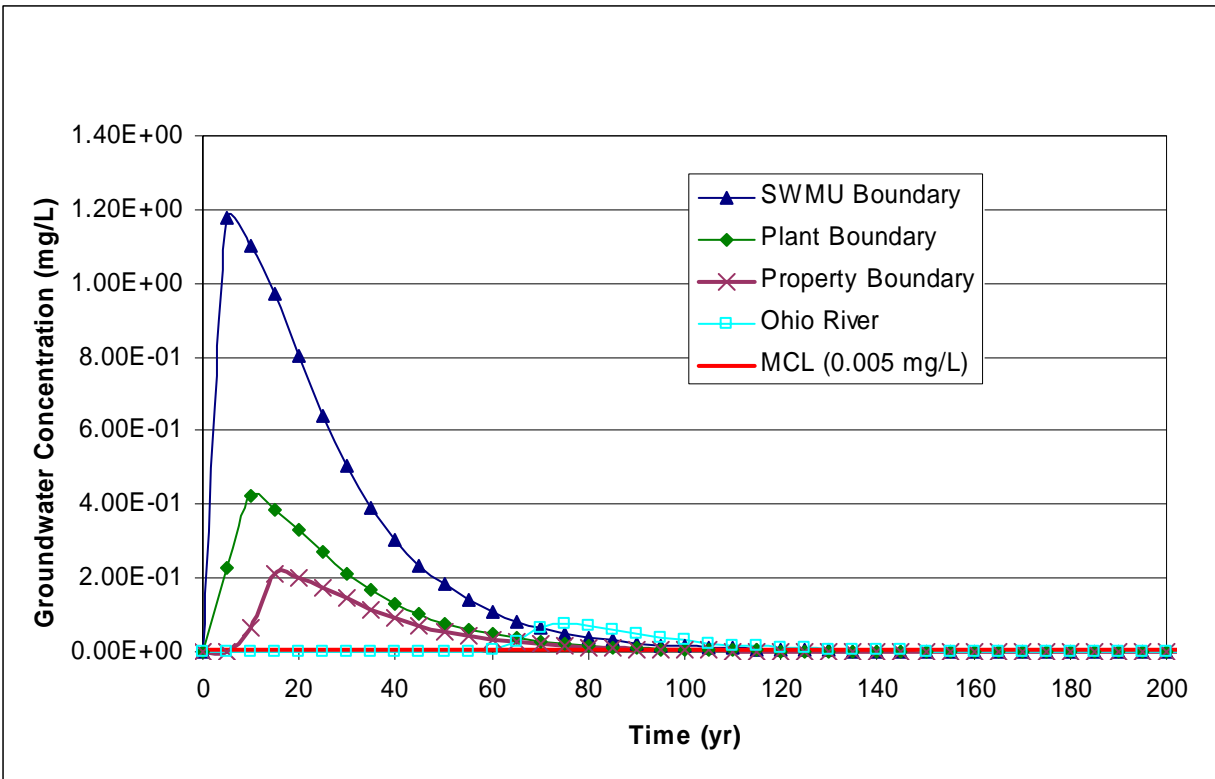


Figure 5.10. Predicted TCE Concentrations in Groundwater at the POEs Based on Contaminant Leaching from SWMU 4

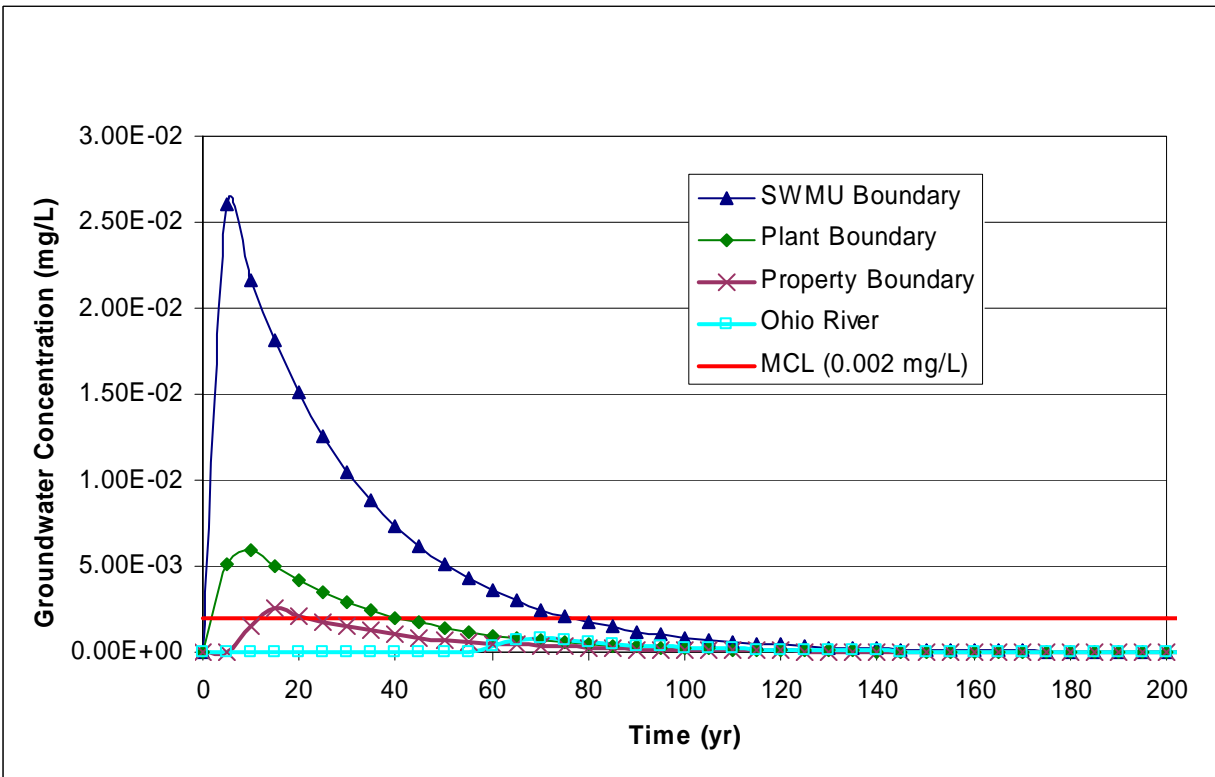


Figure 5.11. Predicted Vinyl Chloride Concentrations in Groundwater at the POEs Based on Contaminant Leaching from SWMU 4

5.3.4 SWMU 5

Analytes retained after the screening process described in Section 5.2.1 are presented for SWMU 5 in Table E3.4. Each analyte was modeled and the resulting peak groundwater concentrations below SWMU 5 shown in Table 5.8 then were compared to the child resident NALs from Table A.18 of the 2001 Risk Methods Document and the provisional groundwater backgrounds shown in the 2001 Risk Method Document in Table A.13. Those analytes with groundwater concentrations below SWMU 5 that exceeded the NALs or background values then were carried through the toxicity and exposure assessments and cancer risk and hazards were calculated for the Rural Resident Groundwater User. Vanadium, benzo(a)pyrene, fluoranthene, PCB-1260, pyrene, cesium-137 and dibenz(a,h,)anthracene were modeled and found not to reach the RGA during the 1,000 year modeling period.

Table 5.8. Screening of Modeled Peak Concentrations in Groundwater for SWMU 5

Analyte	Time (years)	Peak Conc. Below SWMU 5 (mg/L) ^a	Background Groundwater Concentration (mg/L) ^a	Groundwater Child Resident No Action Level (mg/L) ^a	Retain?
Acenaphthene	4.60E+02	6.10E-03	NA	1.36E-02	N
Anthracene	1.25E+02	8.06E-03	NA	7.66E-02	N
Arsenic	1.00E+03	9.25E-03	5.00E-03	3.50E-05	Y
Fluorene	7.20E+02	3.63E-03	NA	9.72E-03	N
Manganese	9.45E+02	1.01E+00	1.19E-01	3.50E-02	Y
Naphthalene	1.30E+02	5.55E-03	NA	2.85E-04	Y
Nickel	1.00E+03	2.01E-03	3.05E-01	3.01E-02	N
Selenium	5.70E+02	1.27E-03	5.00E-03	7.54E-03	N
TCE	1.00E+01	9.91E-04	NA	1.60E-03	N
Zinc	9.40E+02	1.58E-01	4.90E-02	4.50E-01	N
Uranium	9.98E+02	4.60E-01	2.00E-03	9.06E-04	Y
Technetium-99	5.00E+00	1.27E+02	2.23E+01	1.40E+01	Y

^a Units for radionuclides are pCi/L.

NA = not applicable N = No Y = Yes

All SWMU 5 analytes are predicted to be less than their respective MCLs at all POEs, except for uranium at the plant boundary. The following summarizes those analytes that exceeded ELCR or HQ risk criteria of 1.0E-06 and 0.1, respectively.

		Plant Boundary	Property Boundary	Ohio River
Arsenic	ELCR	4.7E-05	3.4E-06	--
	HQ	0.6	--	--
Manganese	ELCR	--	--	--
	HQ	0.2	--	--
Naphthalene	ELCR	--	--	--
	HQ	0.5	0.2	--
Technetium-99	ELCR	2.7E-06	1.4E-06	--
	HQ	--	--	--
Uranium	ELCR	--	--	--
	HQ	5.3	--	--

-- = does not exceed

All remaining analytes exhibited HQ values less than 0.1 and ELCR values less than 1.0E-06 at all POEs. Figures 5.12 through 5.16 illustrate the future predicted concentrations of SWMU 5 analytes that exceed a HQ of 0.1 and/or an ELCR of 1.0E-06 (Table 5.4). As shown in these figures, the dissolved arsenic and manganese concentrations are predicted to continue rising at 1,000 years at the plant boundary.

Manganese will not reach the property boundary or Ohio River in the 1,000 year period. Arsenic begins to increase in concentration at the plant boundary at 1,000 years; however, the concentrations are less than the MCL. Technetium-99 is not predicted to exceed the MCL at the POEs.

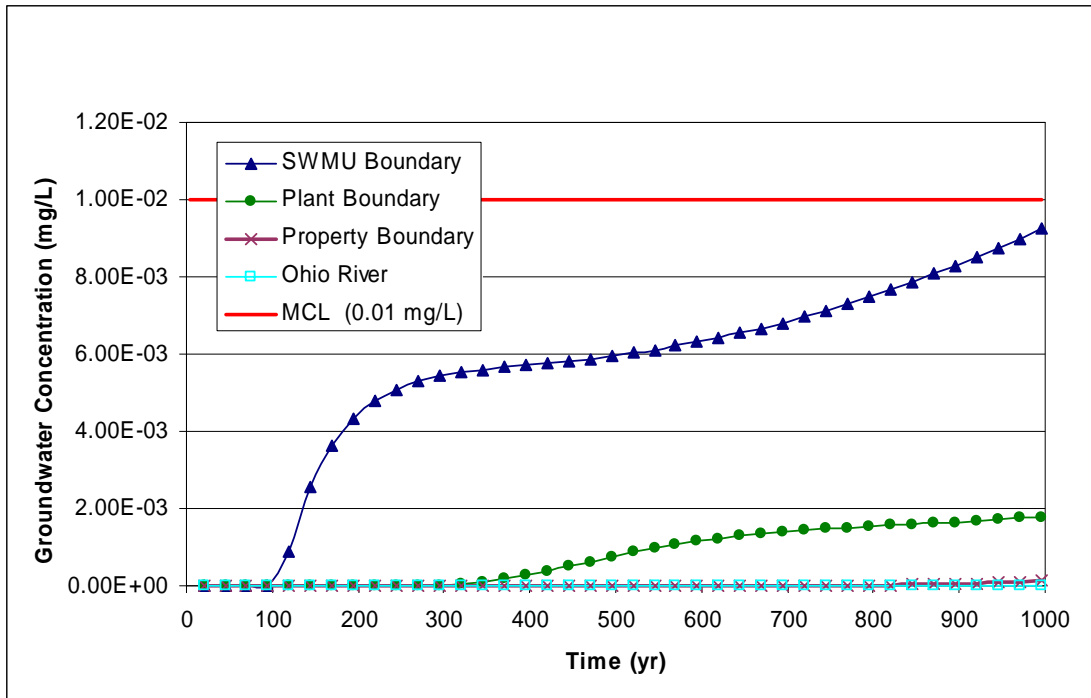


Figure 5.12. Predicted Arsenic Concentrations in Groundwater at the POEs Based on Contaminant Leaching from SWMU 5

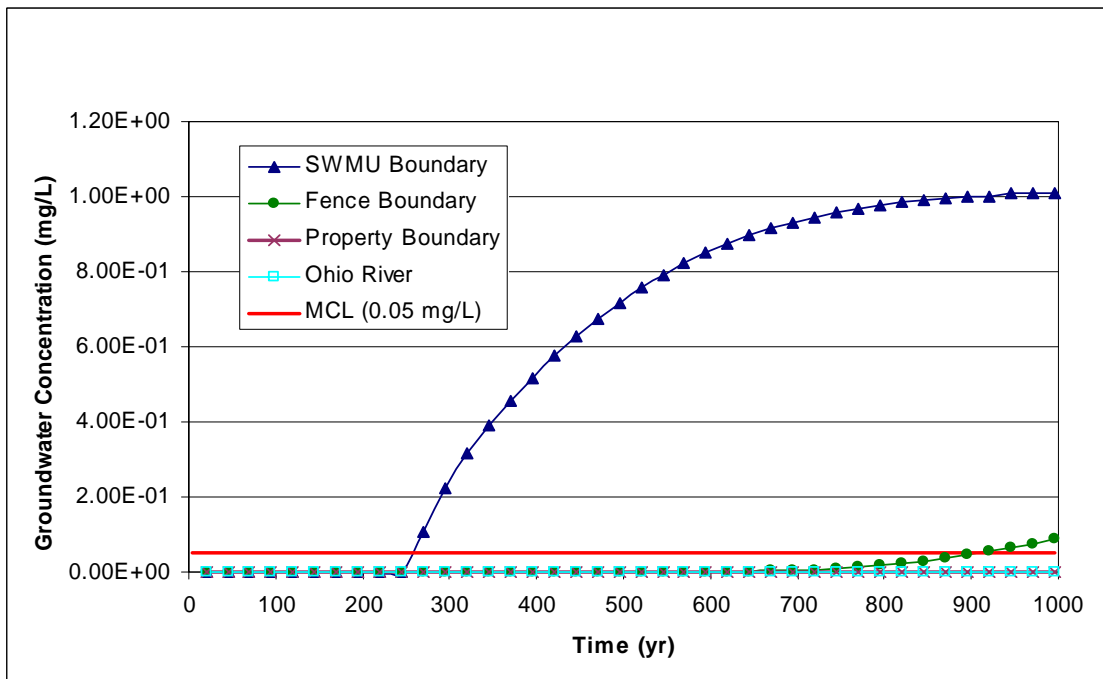


Figure 5.13. Predicted Manganese Concentrations in Groundwater at the POEs Based on Contaminant Leaching from SWMU 5

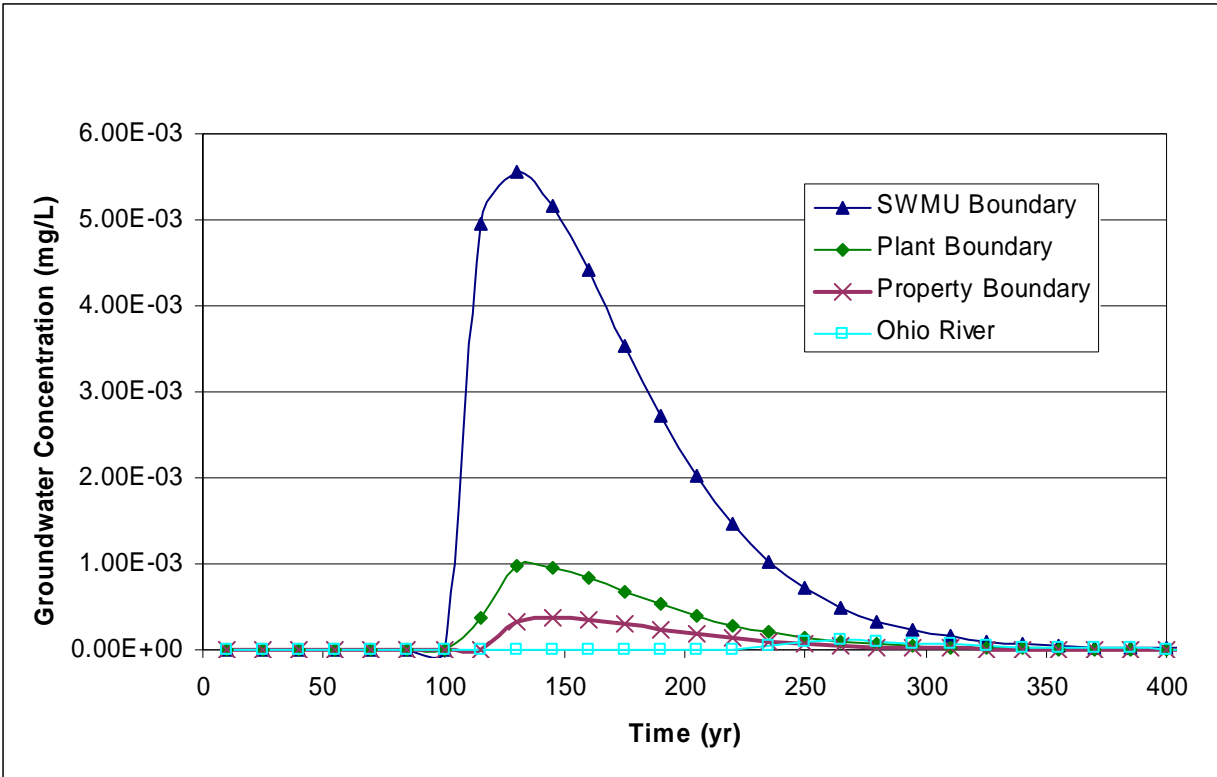


Figure 5.14. Predicted Naphthalene Concentrations in Groundwater at the POEs Based on Contaminant Leaching from SWMU 5

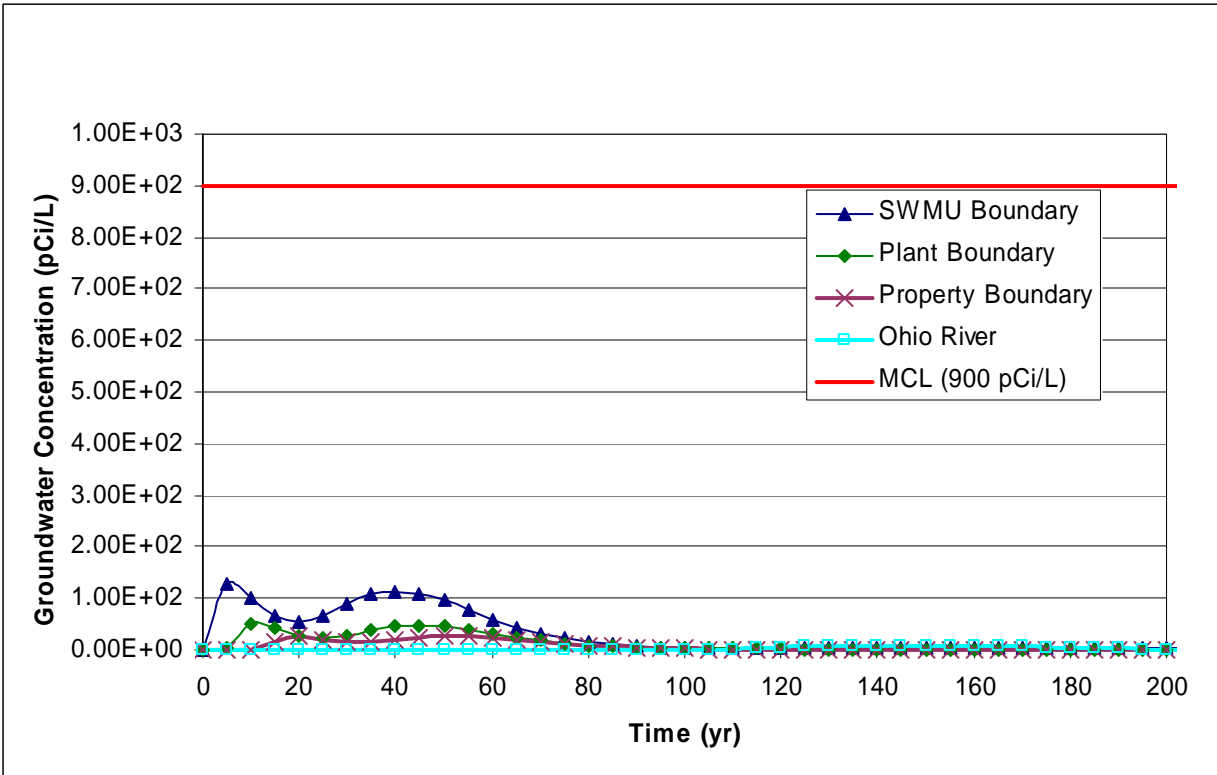


Figure 5.15. Predicted ⁹⁹Tc Activities in Groundwater at the POEs Based on Contaminant Leaching from SWMU 5

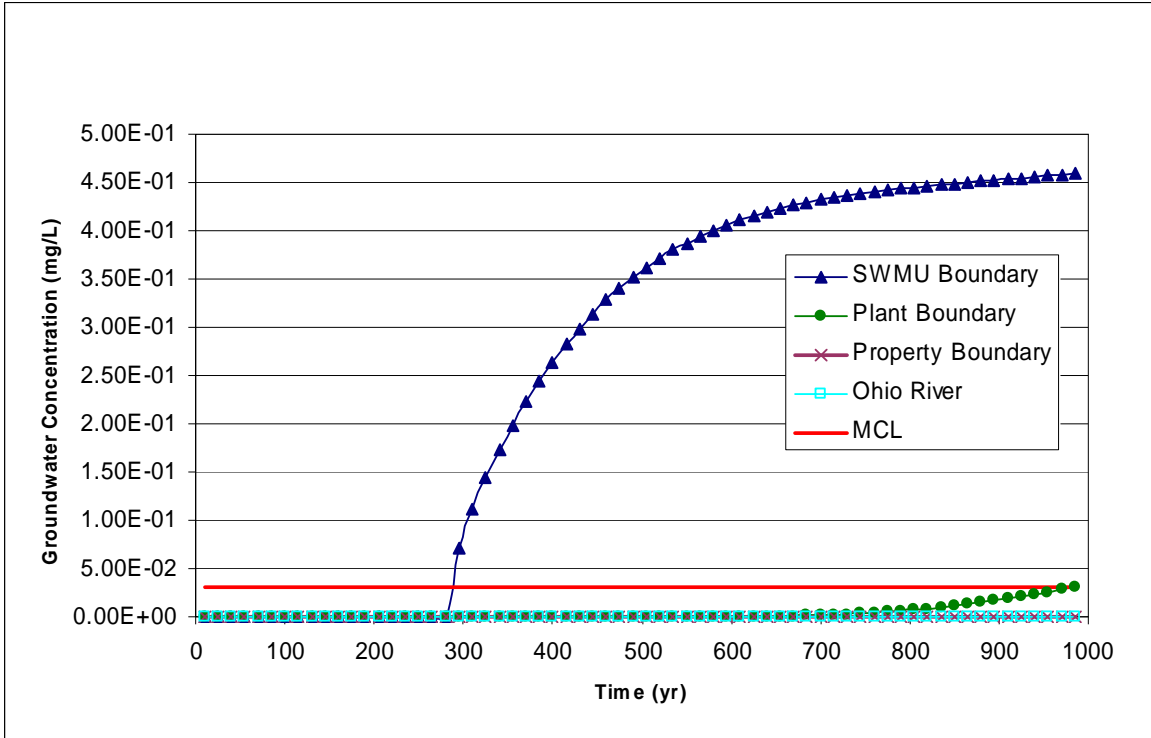


Figure 5.16. Predicted Uranium Activities in Groundwater at the POEs Based on Contaminant Leaching from SWMU 5

5.3.5 SWMU 6

Analytes retained after the screening process described in Section 5.2.1 are presented for SWMU 6 in Table E3.5. Each analyte was modeled and the resulting peak groundwater concentrations below SWMU 6 shown in Table 5.9 then were compared to the child resident NALs from Table A.18 of the 2001 Risk Methods Document and the provisional groundwater backgrounds shown in the 2001 Risk Method Document in Table A.13. Those analytes with groundwater concentrations below SWMU 6 that exceeded the NALs or background values then were carried through the toxicity and exposure assessments and cancer risk and hazards were calculated for the Rural Resident Groundwater User. Beryllium, nickel, vanadium, and Tc-99 were modeled and found not to reach the RGA during the 1,000 year modeling period. Manganese at SWMU 6 exhibited HQ values less than 0.1 at all POEs except at the SWMU boundary, which exhibited an HQ of 0.18. Figure 5.17 illustrates the future predicted concentrations of manganese at SWMU 6 that exceed an HQ of 0.1.

Table 5.9. Screening of Modeled Peak Concentrations in Groundwater for SWMU 6

Analyte	Time (years)	Peak Conc. Below SWMU 6 (mg/L) ^a	Background Groundwater Concentration (mg/L) ^a	Groundwater Child Resident No Action Level (mg/L) ^a	Retain?
Arsenic	9.90E+02	1.92E-03	5.00E-03	3.50E-05	N
Manganese	4.81E+02	8.32E-02	1.19E-01	3.50E-02	Y
TCE	1.10E+01	3.19E-05	NA	1.60E-03	N
Uranium	4.10E+02	1.91E-04	2.00E-03	9.06E-04	N
Zinc	4.45E+02	3.63E-02	4.90E-02	4.50E-01	N

^a Units for radionuclides are pCi/L.
NA = not applicable N = No

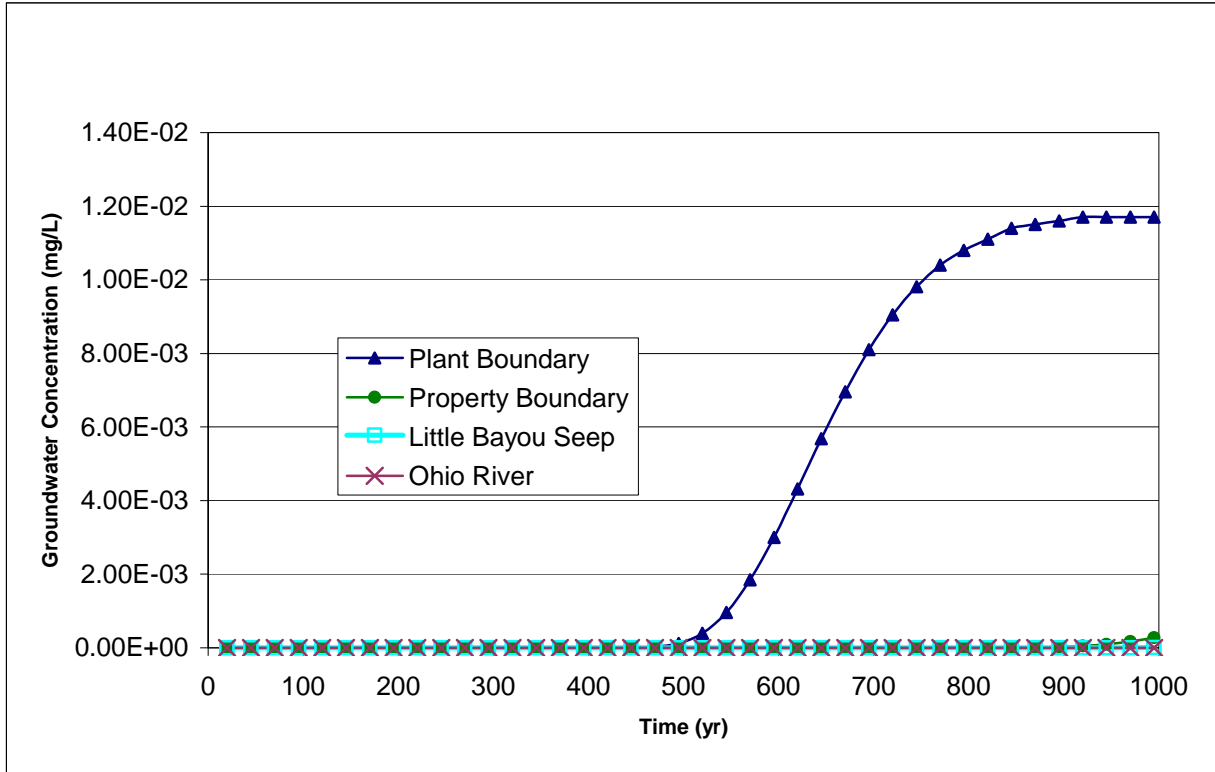


Figure 5.17. Predicted Manganese Activities in Groundwater at the POEs Based on Contaminant Leaching from SWMU 6

5.3.6 SWMU 7

Analytes retained after the screening process described in Section 5.2.1 are presented for SWMU 7 in Table E3.6. Each analyte was modeled and the resulting groundwater concentrations below SWMU 7 shown in Table 5.10 then were compared to the child resident NALs from Table A.18 of the 2001 Risk Methods Document and the provisional groundwater backgrounds shown in the 2001 Risk Method Document in Table A.13. Those analytes with groundwater concentrations below SWMU 7 that exceeded the NALs or background values then were carried through the toxicity and exposure assessments and cancer risk and hazards were calculated for the Rural Resident Groundwater User. Nickel, vanadium, PCB-1260, benzo(a)pyrene, Neptunium-237, and fluoranthene were modeled and found not to reach the RGA during the 1,000 year modeling period.

Table 5.10. Screening of Modeled Peak Concentrations in Groundwater for SWMU 7

Analyte	Time (years)	Peak Conc. Below SWMU 7 (mg/L) ^a	Background Groundwater Concentration (mg/L) ^a	Groundwater Child Resident No Action Level (mg/L) ^a	Retain?
1,1-DCE	9.00E+00	8.98E-02	NA	4.70E-05	Y
Arsenic	1.00E+02	1.78E-02	5.00E-03	3.50E-05	Y
Cadmium	8.55E+02	1.96E-05	1.00E-02	6.61E-04	N
<i>cis</i> -1,2,-DCE	1.00E+01	2.35E-02	NA	2.73E-03	Y
Manganese	8.25E+02	3.32E-01	1.19E-01	3.50E-02	Y
Mercury	9.45E+02	1.01E-05	2.00E-04	4.44E-04	N
PCB-1254	1.00E+03	5.23E-05	NA	1.94E-05	Y
Pyrene	1.00E+03	3.48E-06	NA	1.82E-02	N
Selenium	4.00E+02	1.12E-02	5.00E-03	3.01E-02	N
Tetrachloroethene	1.80E+01	1.40E-04	NA	5.82E-04	N
TCE	4.00E+00	1.09E-02	NA	1.60E-03	Y
Uranium	9.80E+02	3.46E-03	2.00E-03	9.06E-04	Y
Vinyl Chloride	3.00E+00	1.35E-02	NA	3.50E-05	Y
Zinc	7.25E+02	6.73E-02	4.90E-02	4.50E-01	N
Technetium-99	4.00E+01	9.09E+02	2.23E+01	1.40E+01	Y
Uranium-234	9.80E+02	7.94E+00	7.00E-01	5.46E-01	Y
Uranium-235	6.95E+02	8.10E-02	3.00E-01	5.38E-01	N
Uranium-238	9.75E+02	7.59E+00	7.00E-01	4.43E-01	Y

^a Units for radionuclides are pCi/L.

NA = not applicable N = No Y = Yes

The groundwater results presented in Table 5.3 for SWMU 7 show the predicted groundwater concentrations of 1,1-DCE, arsenic, TCE, and vinyl chloride will exceed their respective MCLs at the plant boundary. All SWMU 7 analytes are modeled to be less than their respective MCLs at the property boundary and Little Bayou seeps. The following summarizes those analytes that exceeded ELCR or HQ risk criteria of 1.0E-06 and 0.1, respectively.

		Plant Boundary	Property Boundary	Little Bayou Seeps
1,1-DCE	ELCR	1.9E-03	2.5E-04	9.3E-05
	HQ	0.8	0.1	--
Arsenic	ELCR	3.3E-04	6.2E-05	--
	HQ	4.0	0.8	--
<i>cis</i> -1,2-DCE	ELCR	--	--	--
	HQ	1.1	0.2	--
Manganese	ELCR	--	--	--
	HQ	0.5	--	--
PCB-1254	ELCR	4.8E-06	--	--
	HQ	2.5	0.2	--
Technetium-99	ELCR	4.5E-05	1.5E-05	7.3E-06
	HQ	--	--	--
TCE	ELCR	3.1E-04	4.4E-05	1.6E-05
	HQ	4.5	0.6	0.2
Uranium	ELCR	--	--	--
	HQ	0.4	--	--
Uranium-234	ELCR	8.2E-06	--	--
	HQ	--	--	--
Uranium-238	ELCR	9.6E-06	--	--
	HQ	--	--	--
Vinyl chloride	ELCR	3.6E-04	3.6E-05	1.2E-05
	HQ	0.6	--	--

-- = does not exceed

All remaining analytes exhibited HQ values less than 0.1 and ELCR values less than 1.0E-06 at all POEs. Figures 5.18 through 5.28 portray the predicted concentrations of SWMU 7 analytes that exceed a HQ of 0.1 and/or an ELCR of 1.0E-06 (Table 5.4). Arsenic was modeled and found not to reach the Little Bayou seeps in the 1,000 year modeling period.

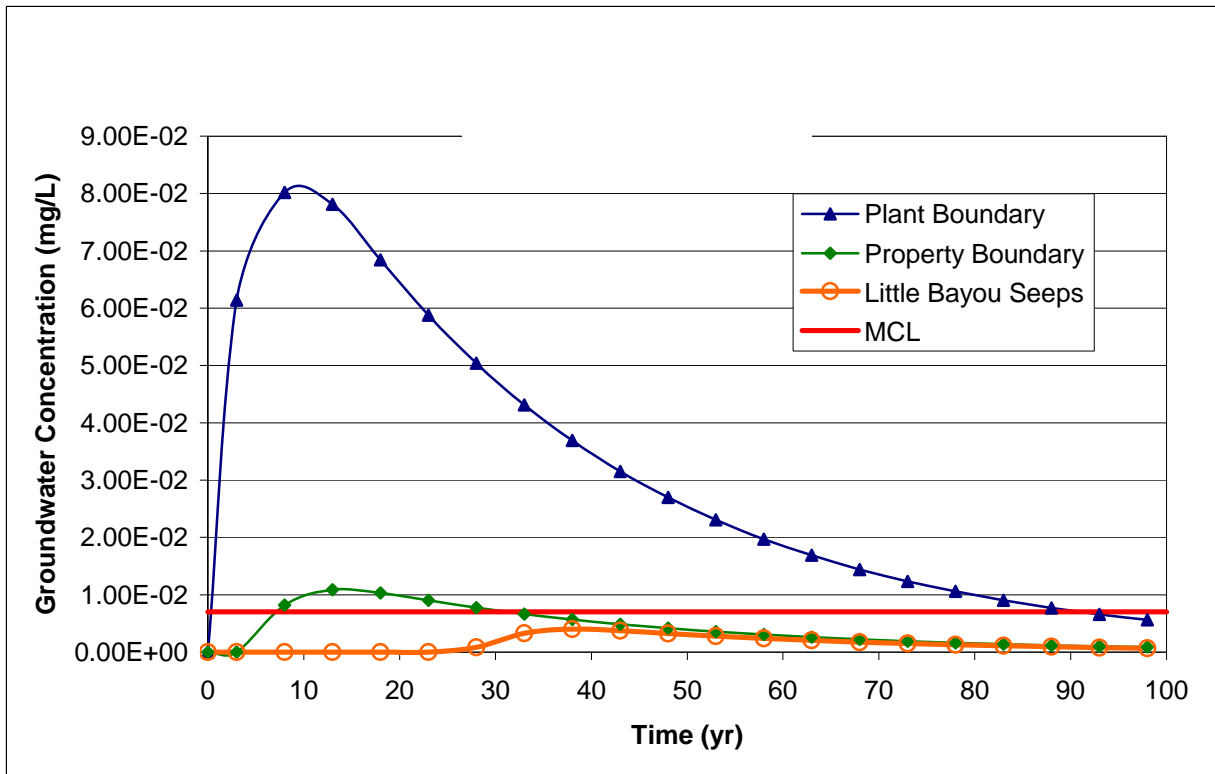


Figure 5.18. Predicted 1,1-DCE Concentrations in Groundwater at the POEs Based on Contaminant Leaching from SWMU 7

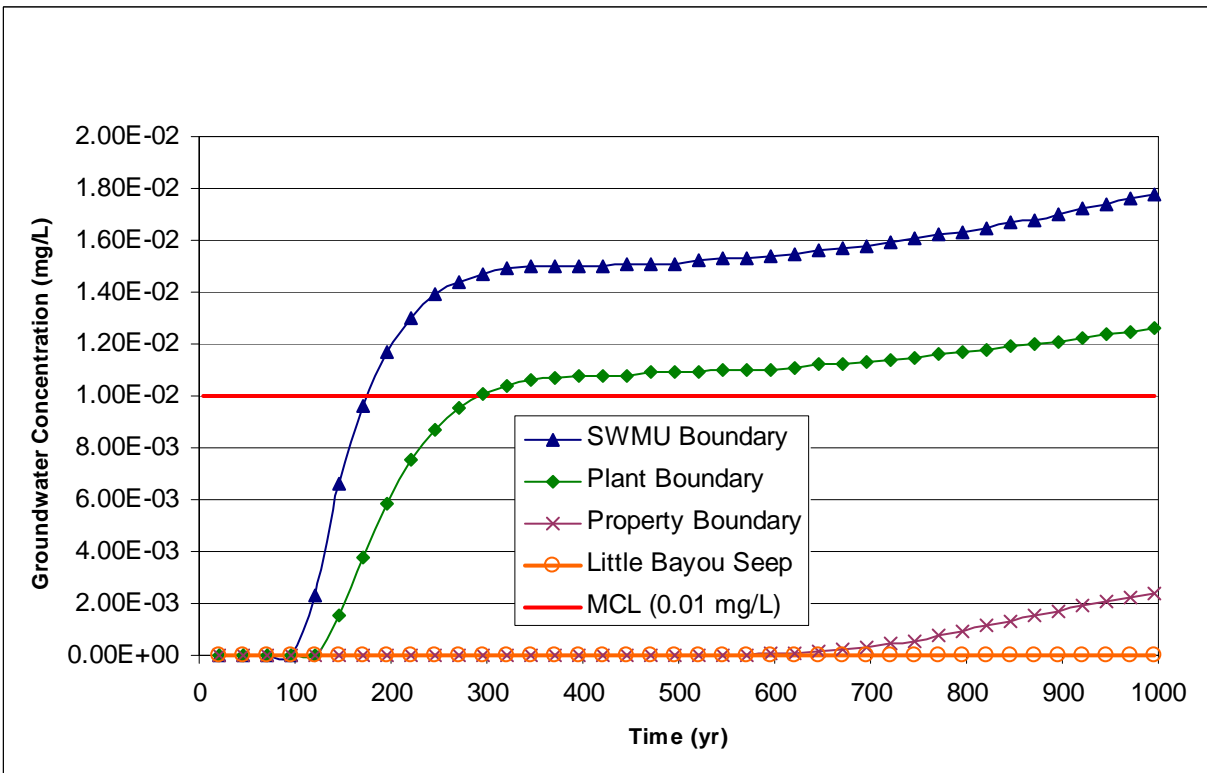


Figure 5.19. Predicted Arsenic Concentrations in Groundwater at the POEs Based on Contaminant Leaching from SWMU 7

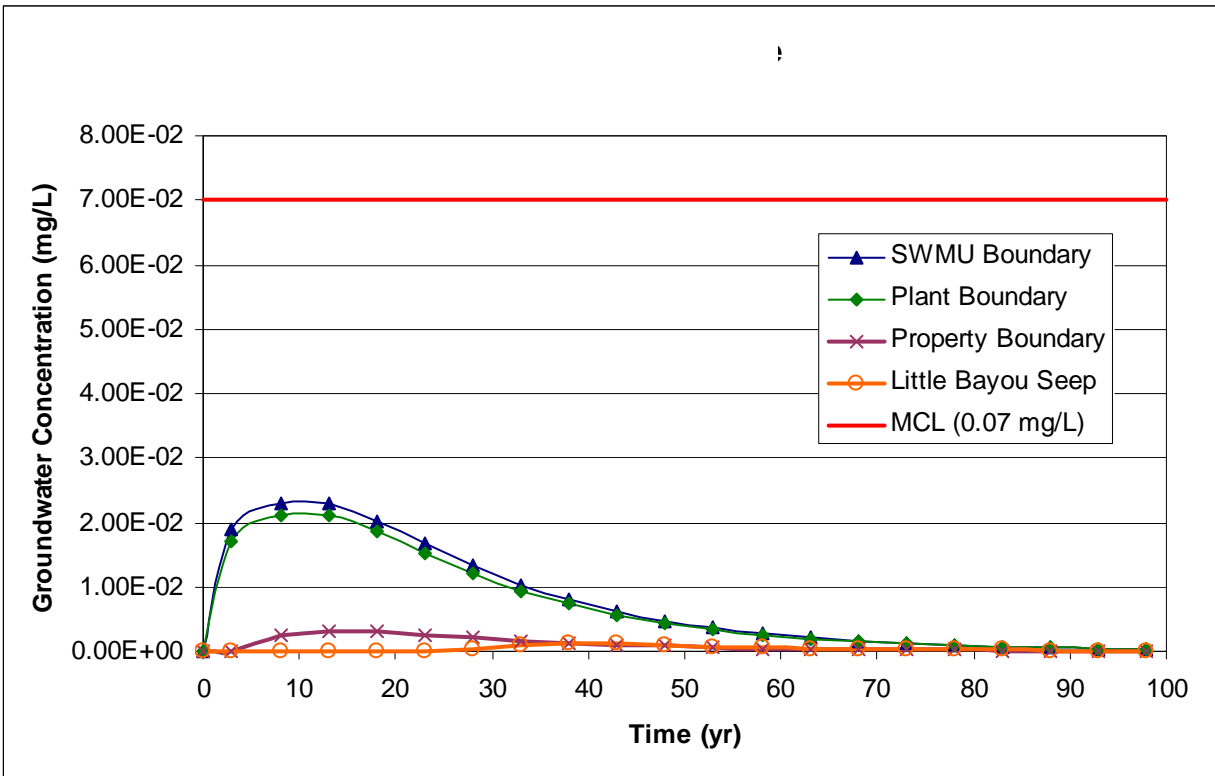


Figure 5.20. Predicted *cis*-1,2-DCE Concentrations in Groundwater at the POEs Based on Contaminant Leaching from SWMU 7

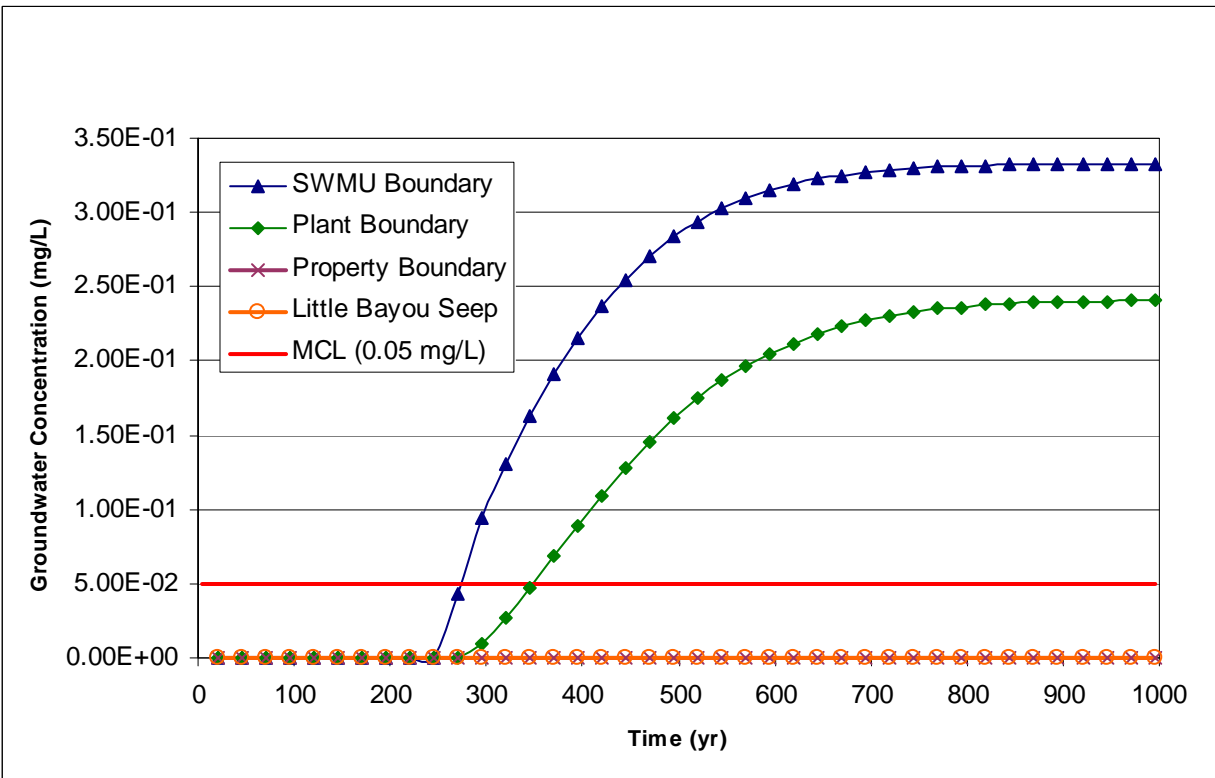


Figure 5.21. Predicted Manganese Concentrations in Groundwater at the POEs Based on Contaminant Leaching from SWMU 7

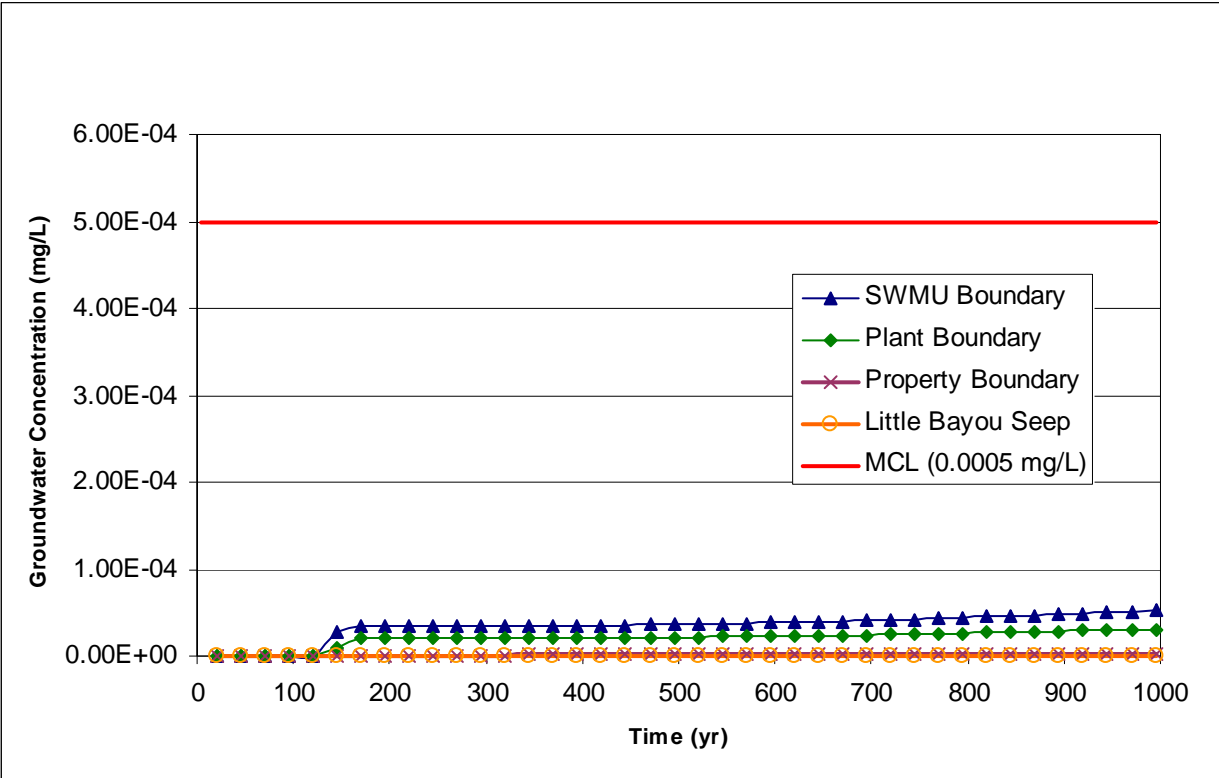


Figure 5.22. Predicted PCB-1254 Concentrations in Groundwater at the POEs Based on Contaminant Leaching from SWMU 7

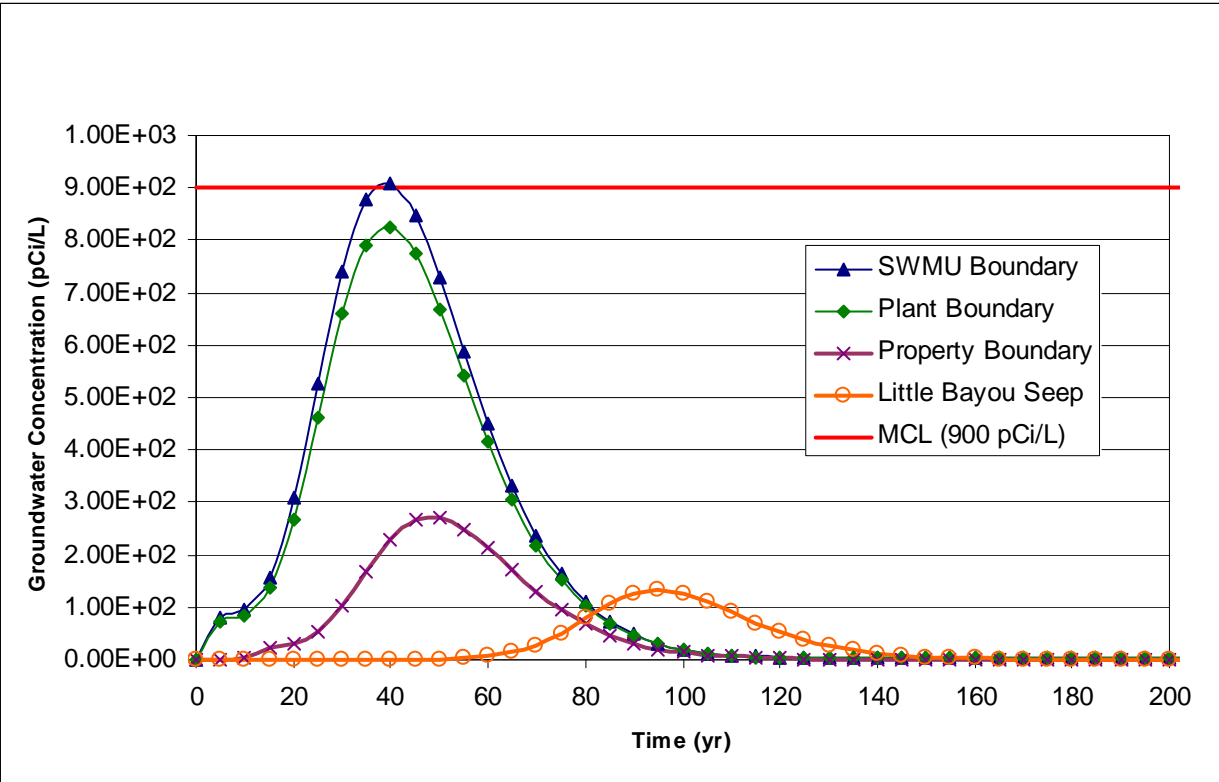


Figure 5.23. Predicted ⁹⁹Tc Activities in Groundwater at the POEs Based on Contaminant Leaching from SWMU 7

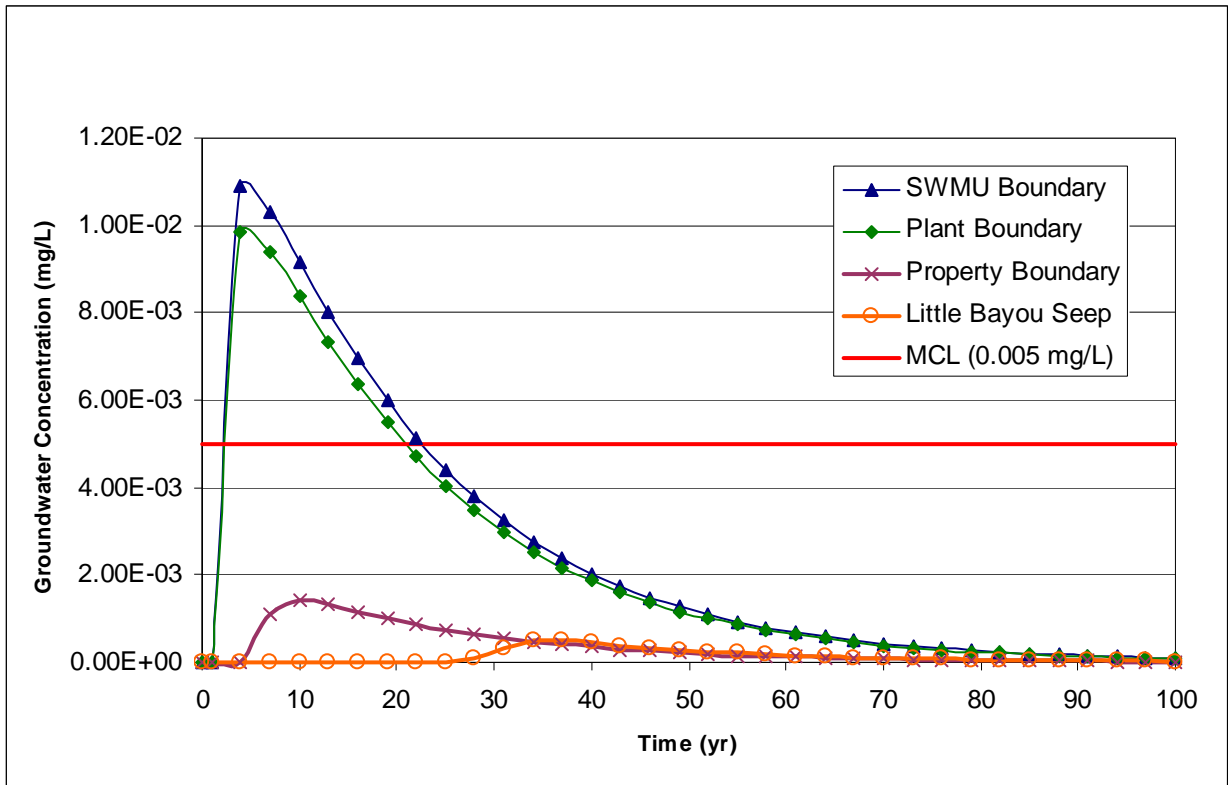


Figure 5.24. Predicted TCE Concentrations in Groundwater at the POEs Based on Contaminant Leaching from SWMU 7

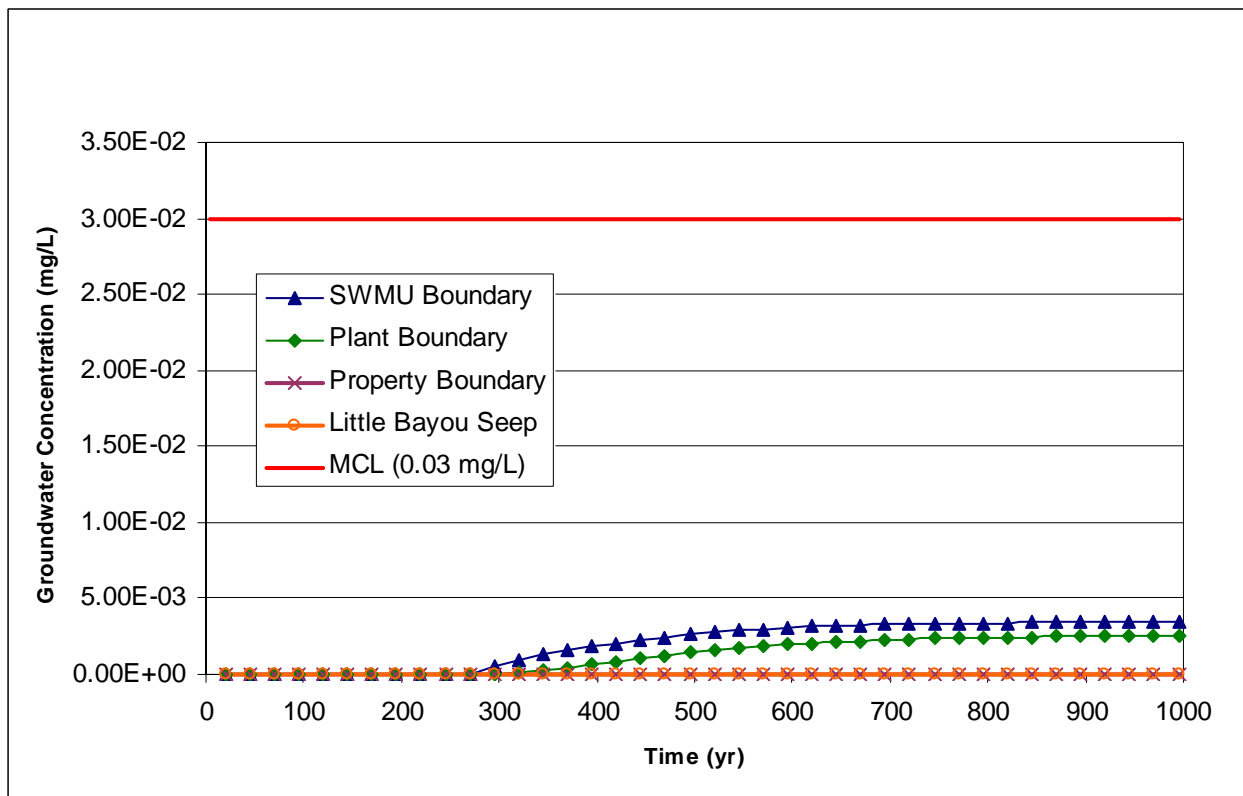


Figure 5.25. Predicted Uranium Concentrations in Groundwater at the POEs Based on Contaminant Leaching from SWMU 7

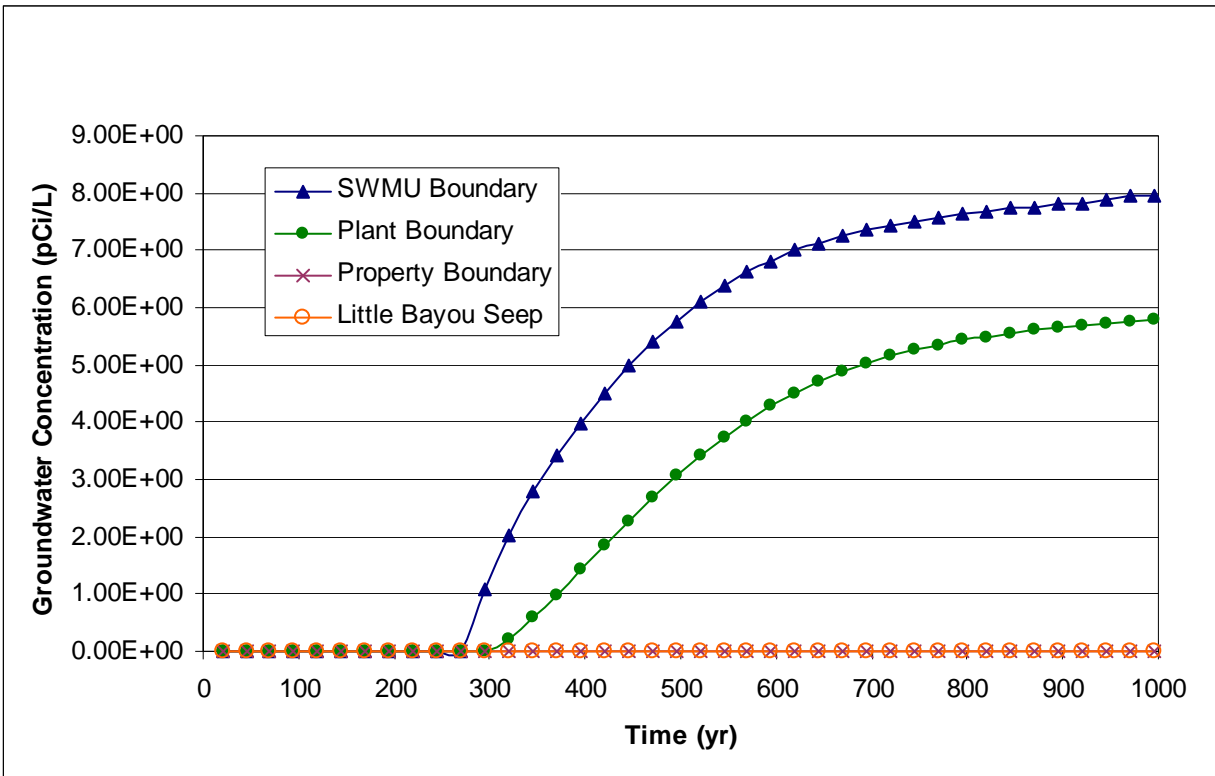


Figure 5.26. Predicted ²³⁴U Activities in Groundwater at the POEs Based on Contaminant Leaching from SWMU 7

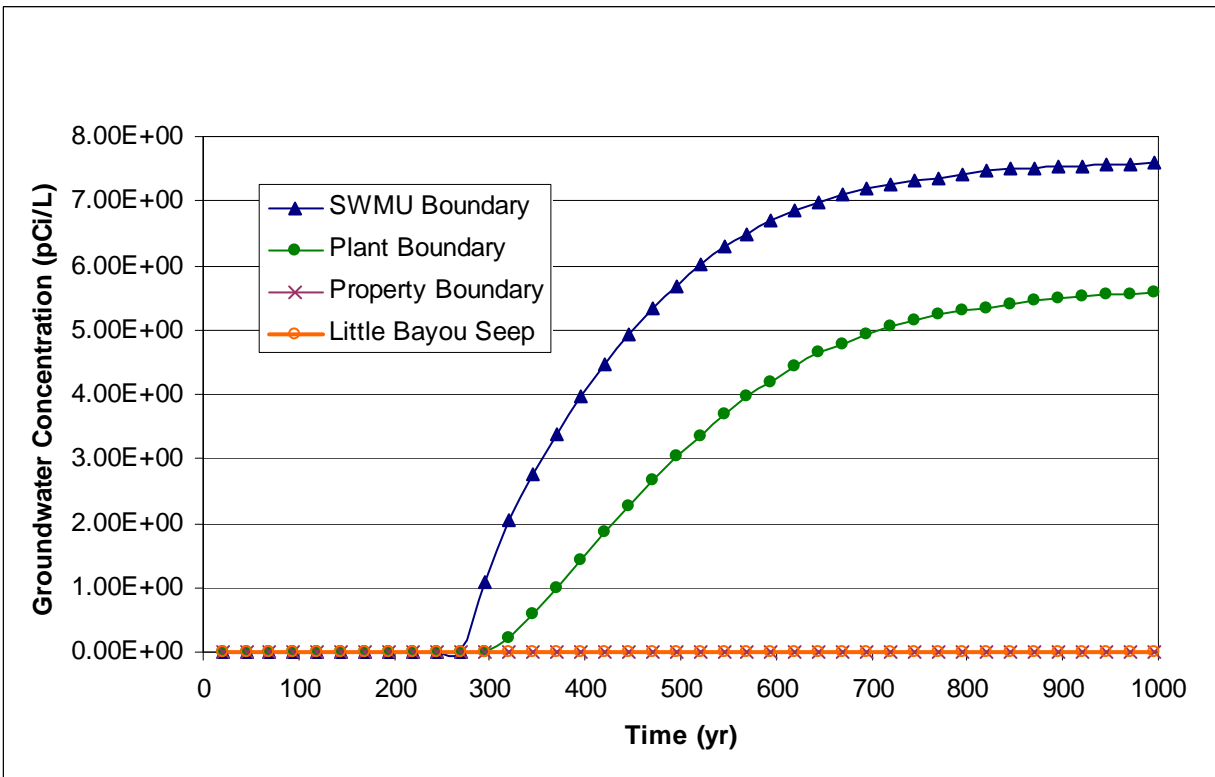


Figure 5.27. Predicted ²³⁸U Activities in Groundwater at the POEs Based on Contaminant Leaching from SWMU 7

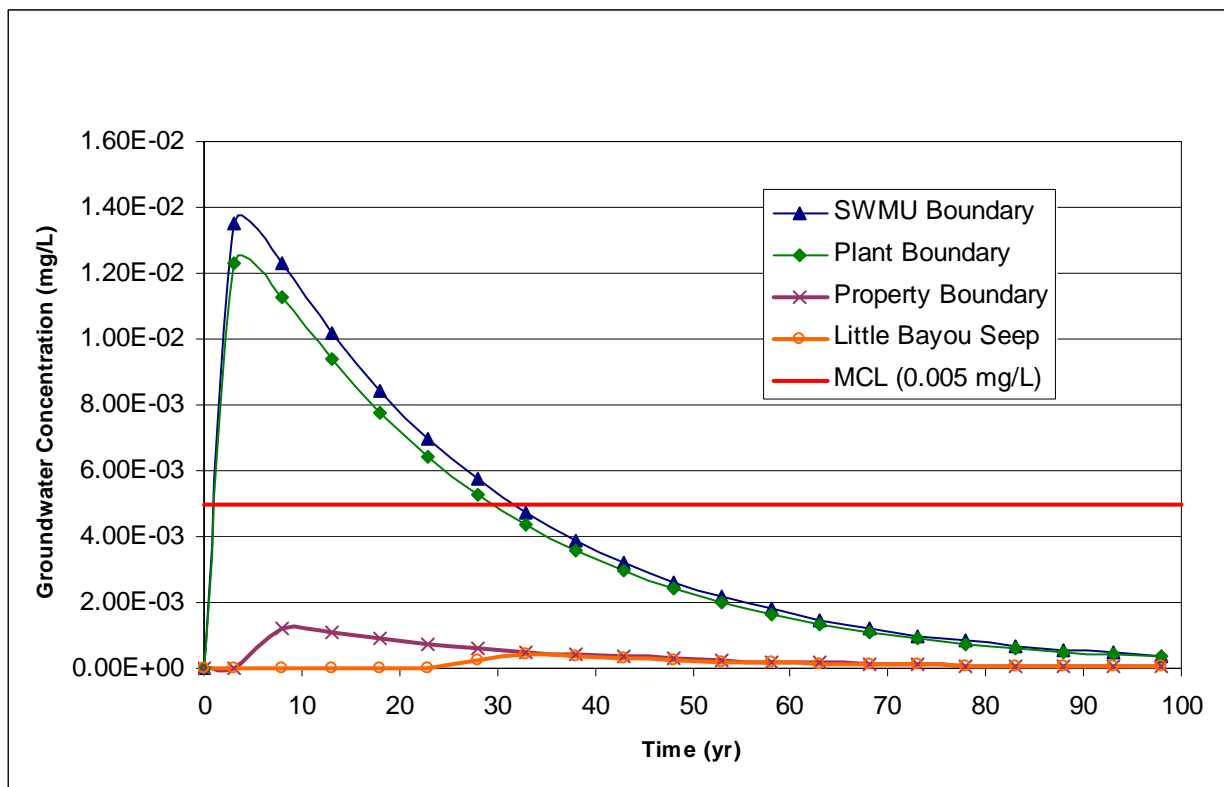


Figure 5.28. Predicted Vinyl Chloride Concentrations in Groundwater at the POEs Based on Contaminant Leaching from SWMU 7

5.3.7 SWMU 30

Analytes retained after the screening process described in Section 5.2.1 are presented for SWMU 30 in Table E3.7. Each analyte was modeled and the resulting peak groundwater concentrations below SWMU 30 shown in Table 5.11 then were compared to the child resident NALs from Table A.18 of the 2001 Risk Methods Document and the provisional groundwater backgrounds shown in the 2001 Risk Method Document in Table A.13. Those analytes with groundwater concentrations below SWMU 30 that exceeded the NALs or background values then were carried through the toxicity and exposure assessments and cancer risk and hazards were calculated for the Rural Resident Groundwater User. Cadmium, nickel, vanadium, benzo(a)pyrene, and dibenz(a,h)anthracene were modeled and found not to reach the RGA during the 1,000 year modeling period.

Table 5.11. Screening of Modeled Peak Concentrations in Groundwater for SWMU 30

Analyte	Time (years)	Peak Conc. Below SWMU 30 (mg/L) ^a	Background Groundwater Concentration (mg/L) ^a	Groundwater Child Resident No Action Level (mg/L) ^a	Retain?
1,1-DCE	2.00E+00	8.18E-05	NA	4.70E-05	Y
Acenaphthene	3.90E+02	2.02E-04	NA	1.36E-02	N
Arsenic	9.90E+02	1.82E-02	5.00E-03	3.50E-05	Y
Fluorene	7.20E+02	1.26E-04	NA	9.72E-03	N
Manganese	7.90E+02	3.78E-01	1.19E-01	3.50E-02	Y
Mercury	1.00E+03	4.41E-06	2.00E-04	4.44E-04	N
Naphthalene	1.35E+02	1.81E-04	NA	2.85E-04	N
PCB-1254	1.00E+03	1.30E-05	NA	1.94E-05	N

**Table 5.11. Screening of Modeled Peak Concentrations in Groundwater for SWMU 30
(Continued)**

Analyte	Time (years)	Peak Conc. Below SWMU 30 (mg/L) ^a	Background Groundwater Concentration (mg/L) ^a	Groundwater Child Resident No Action Level (mg/L) ^a	Retain?
PCB-1260	1.00E+03	5.42E-06	NA	1.94E-05	N
Pyrene	3.30E+02	1.82E-05	NA	1.82E-02	N
Selenium	3.60E+02	1.51E-02	5.00E-03	7.45E-03	Y
TCE	1.30E+01	9.11E-04	NA	5.82E-04	Y
Uranium	5.40E+02	8.40E-03	2.00E-03	9.06E-04	Y
Zinc	9.90E+02	7.77E-02	4.90E-02	4.50E-01	N
Technetium-99	3.70E+01	2.87E+02	2.23E+01	1.40E+01	Y
Uranium-234	7.05E+02	3.99E+00	7.00E-01	5.46E-01	Y
Uranium-235	6.15E+02	1.38E-01	3.00E-01	5.38E-01	N
Uranium-238	6.90E+02	5.91E+00	7.00E-01	4.43E-01	Y

^a Units for radionuclides are pCi/L.

NA = not applicable N = No Y = Yes

The modeled groundwater concentrations of arsenic and TCE exceed their respective MCLs at the plant boundary (Table 5.3). Predicted TCE concentrations also exceed the MCL at the property boundary, and Little Bayou seeps. The following summarizes those analytes that exceeded ELCR or HQ risk criteria of 1.0E-06 and 0.1, respectively.

		Plant Boundary	Property Boundary	Little Bayou Seeps
1,1-DCE	ELCR	1.8E-06	--	--
	HQ	--	--	--
Arsenic	ELCR	3.2E-04	6.2E-05	--
	HQ	3.9	0.8	--
Manganese	ELCR	--	--	--
	HQ	0.5	--	--
Selenium	ELCR	--	--	--
	HQ	0.2	--	--
Technetium-99	ELCR	1.4E-05	3.9E-06	1.6E-06
	HQ	--	--	--
TCE	ELCR	2.6E-05	2.4E-06	---
	HQ	0.4	--	--
Uranium	ELCR	--	--	--
	HQ	0.8	--	--
Uranium-234	ELCR	3.9E-06	--	--
	HQ	--	--	--
Uranium-238	ELCR	7.1E-06	--	--
	HQ	--	--	--

-- = does not exceed

All remaining analytes exhibited HQ values less than 0.1 and ELCR values less than 1.0E-06 at all POEs. Figures 5.29 through 5.37 display the modeled concentrations of SWMU 30 analytes that exceed a HQ of 0.1 and/or an ELCR of 1.0E-06 (Table 5.4). As shown in these figures, the dissolved arsenic and manganese concentrations are predicted to continue rising at 1,000 years at the plant boundary, with arsenic exceeding its MCL. Dissolved arsenic concentrations were less than the MCL at the property

boundary, but dissolved manganese levels have not reached the property boundary or Little Bayou Seeps in the 1,000-year period.

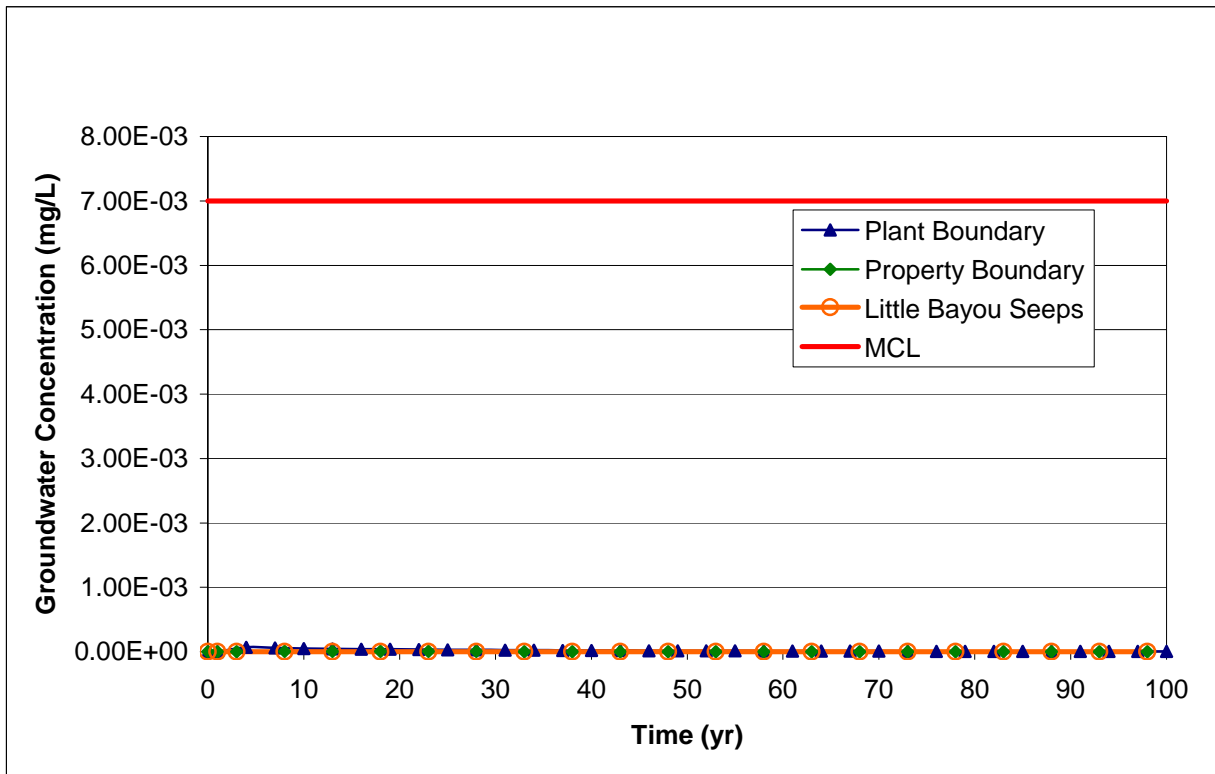


Figure 5.29. Predicted 1,1-DCE Concentrations in Groundwater at the POEs Based on Contaminant Leaching from SWMU 30

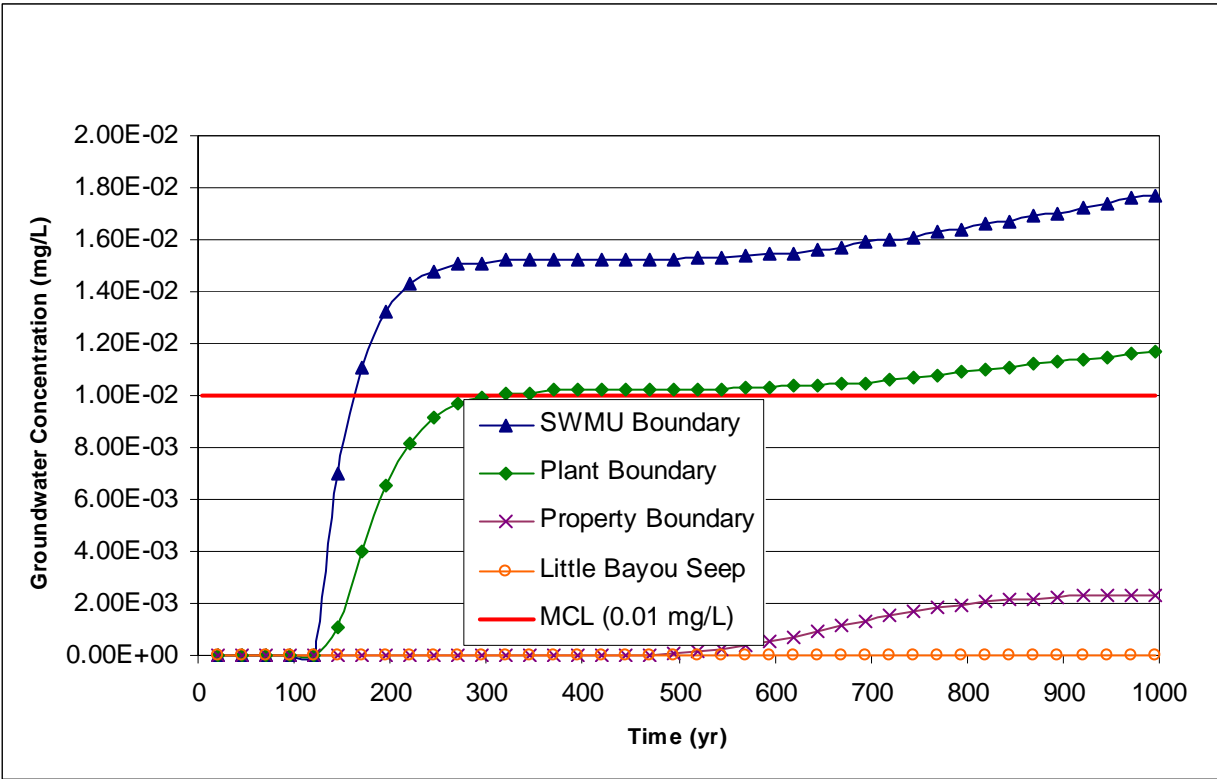


Figure 5.30. Predicted Arsenic Concentrations in Groundwater at the POEs Based on Contaminant Leaching from SWMU 30

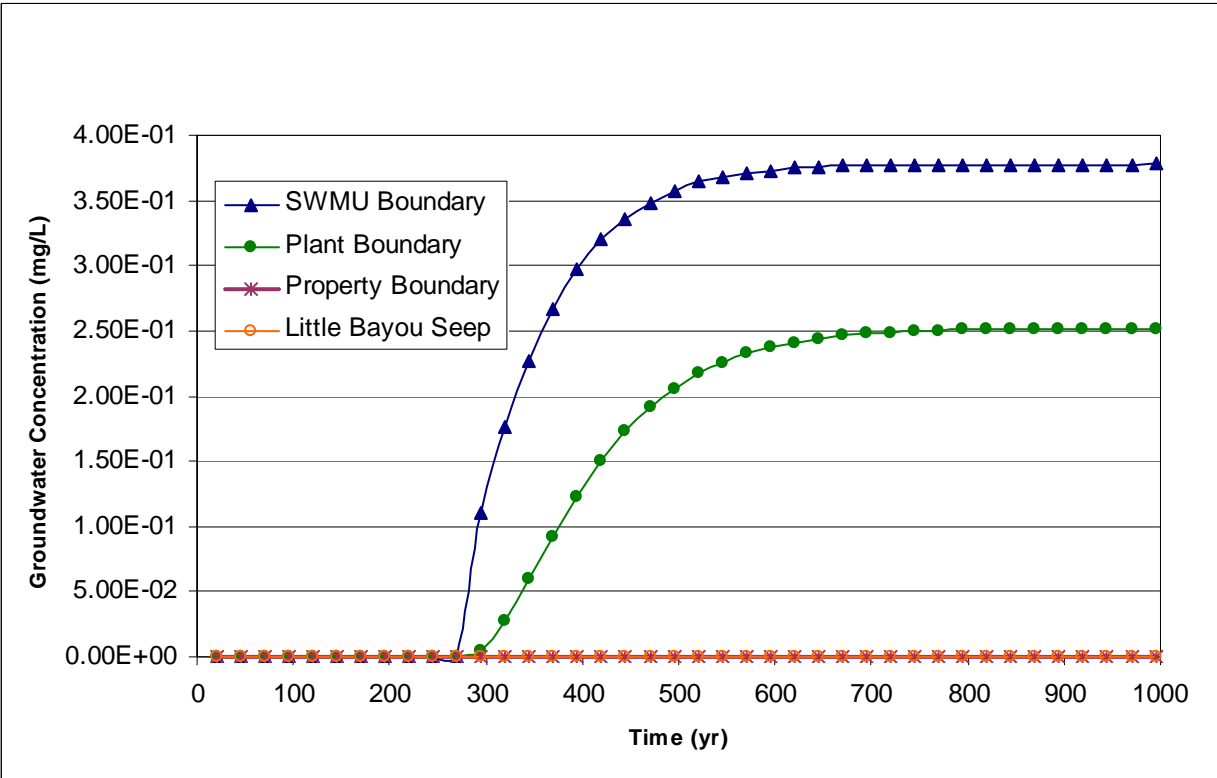


Figure 5.31. Predicted Manganese Concentrations in Groundwater at the POEs Based on Contaminant Leaching from SWMU 30

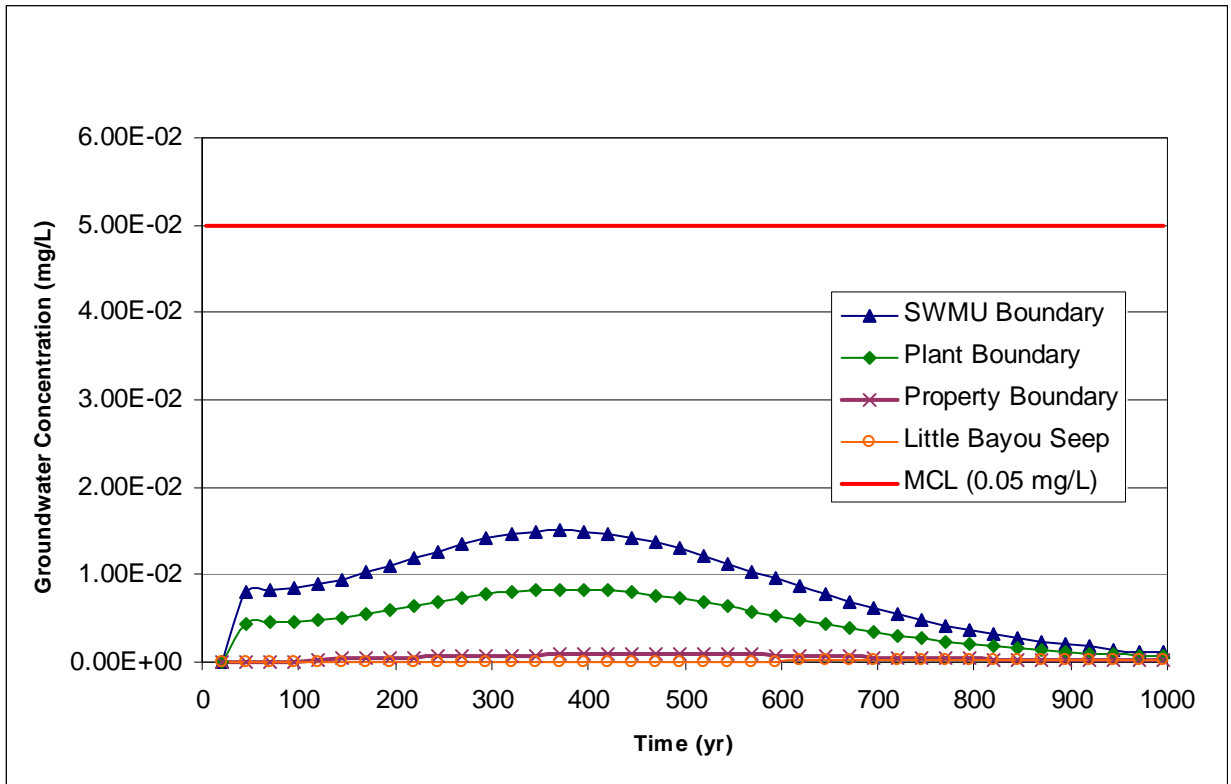


Figure 5.32. Predicted Selenium Concentrations in Groundwater at the POEs Based on Contaminant Leaching from SWMU 30

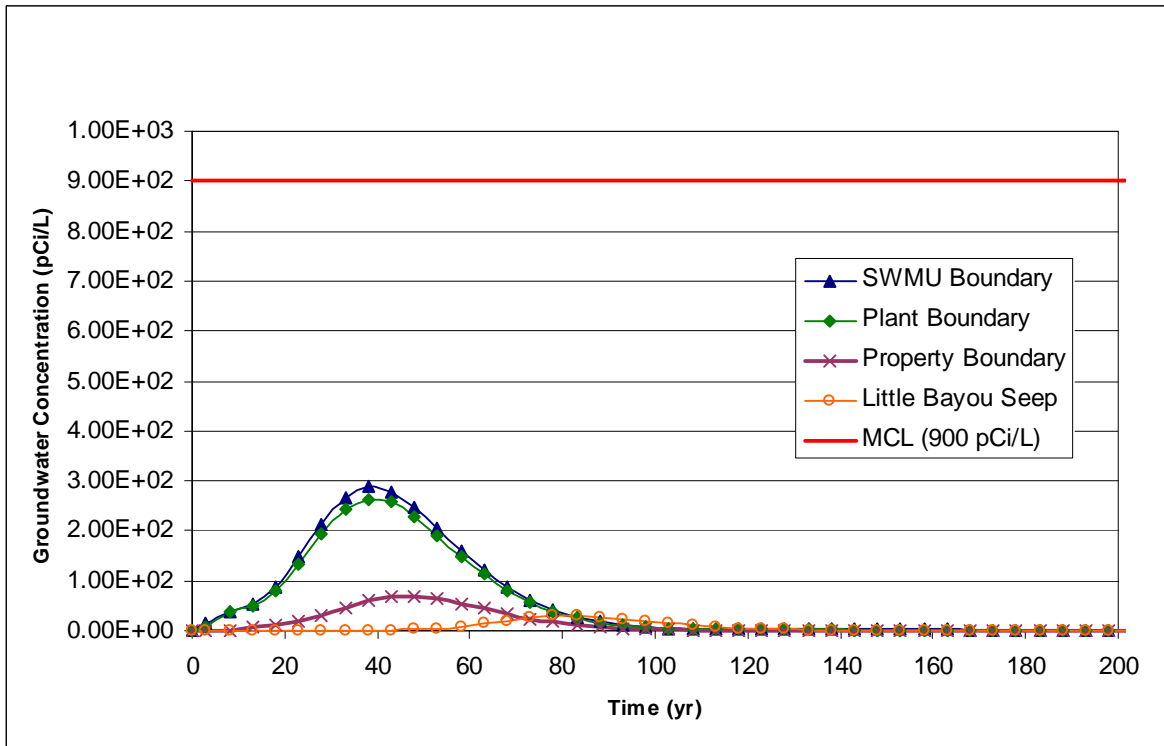


Figure 5.33. Predicted ⁹⁹Tc Activities in Groundwater at the POEs Based on Contaminant Leaching from SWMU 30

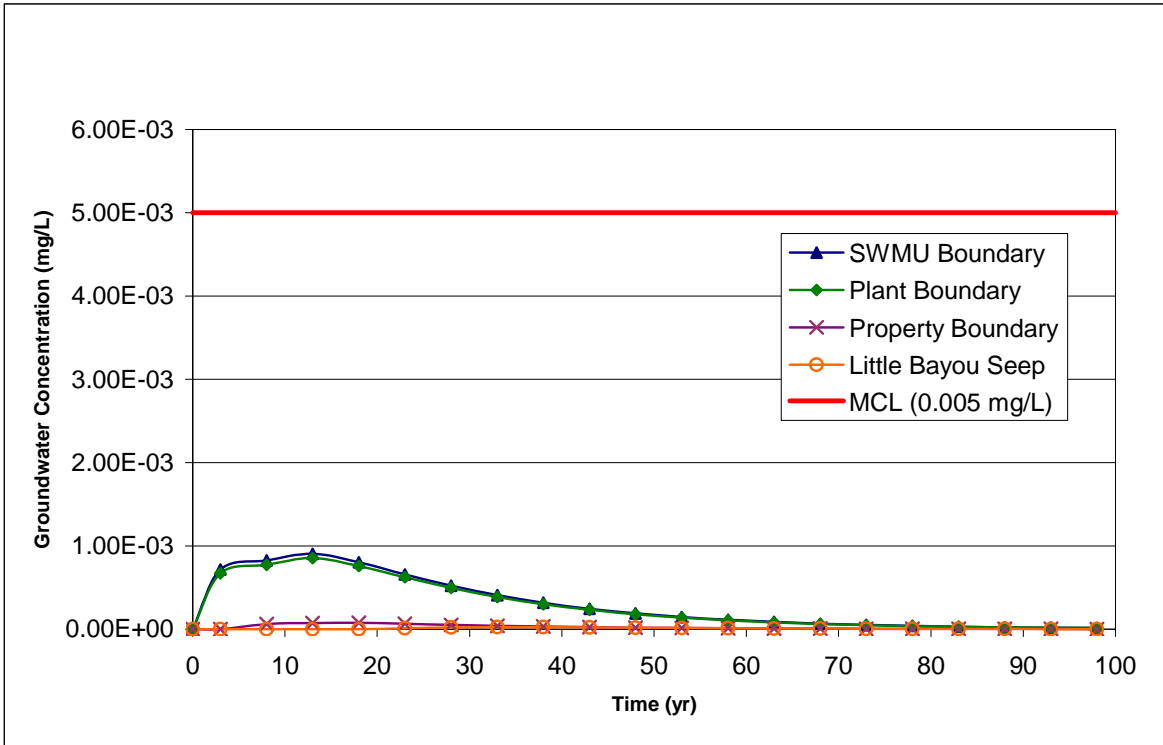


Figure 5.34. Predicted TCE Concentrations in Groundwater at the POEs Based on Contaminant Leaching from SWMU 30

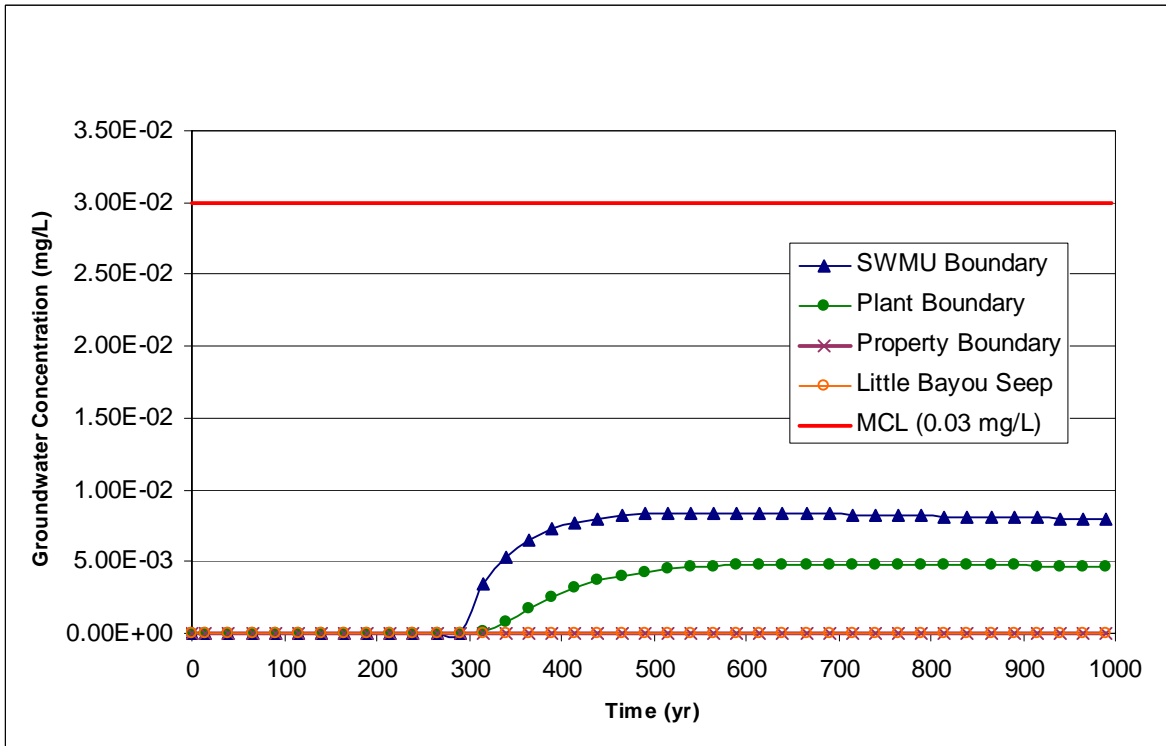


Figure 5.35. Predicted Uranium Concentration in Groundwater at the POEs Based on Contaminant Leaching from SWMU 30

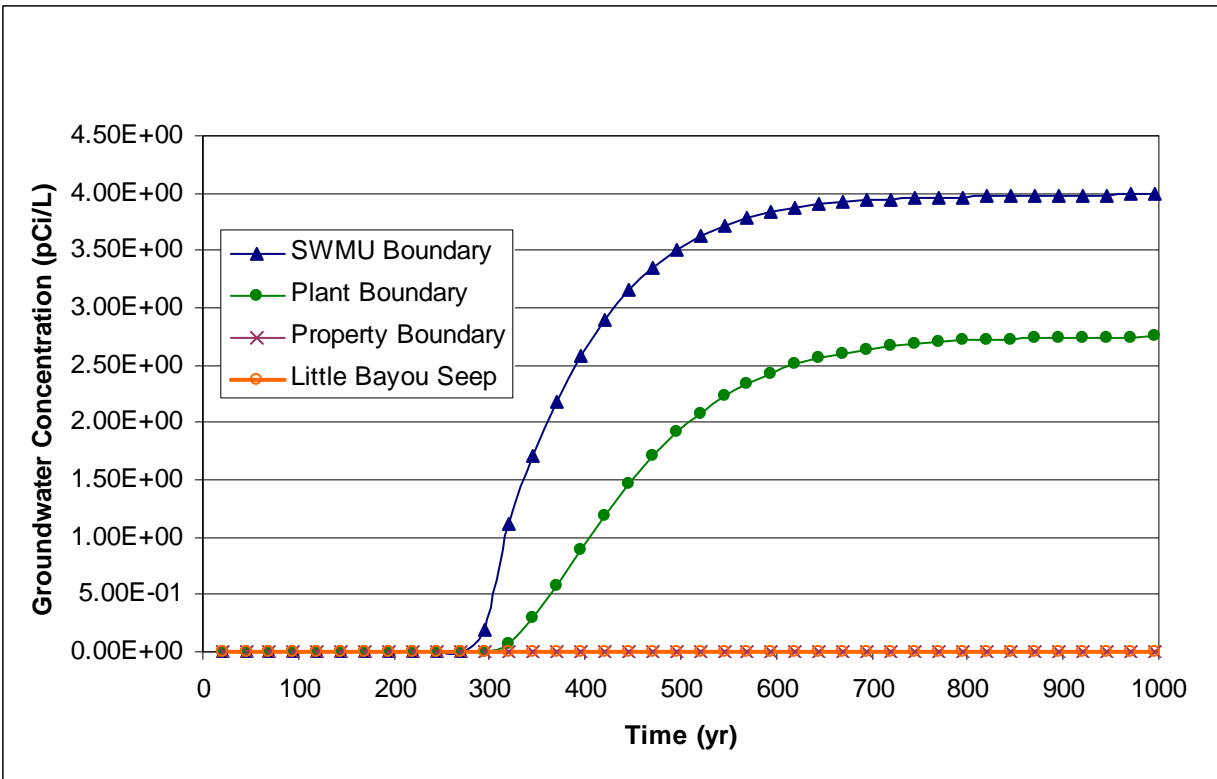


Figure 5.36. Predicted ²³⁴U Activities in Groundwater at the POEs Based on Contaminant Leaching from SWMU 30

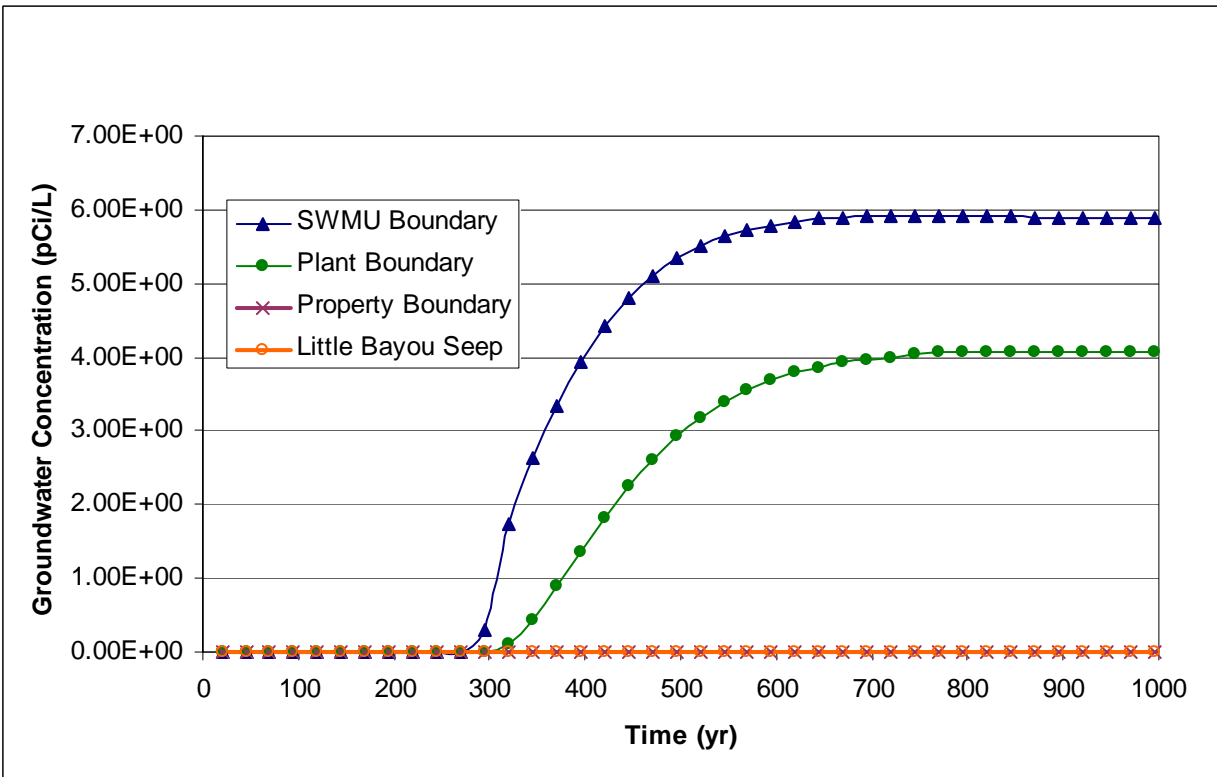


Figure 5.37. Predicted ²³⁸U Activities in Groundwater at the POEs Based on Contaminant Leaching from SWMU 30

5.3.8 SWMU 145

Analytes retained after the screening process described in Section 5.2.1 are presented for SWMU 145 in Table E3.8. Each analyte was modeled and the resulting peak groundwater concentrations below SWMU 145 shown in Table 5.12 then were compared to the child resident NALs from Table A.18 of the 2001 Risk Methods Document and the provisional groundwater backgrounds shown in the 2001 Risk Method Document in Table A.13. Those analytes with groundwater concentrations below SWMU 145 that exceeded the NALs or background values then were carried through the toxicity and exposure assessments and cancer risk and hazards were calculated for the Rural Resident Groundwater User. Vanadium, benzo(a)pyrene, PCB-1254, plutonium-239, uranium-235, and uranium-234 were modeled and found not to reach the RGA during the 1,000 year modeling period.

Table 5.12. Screening of Modeled Peak Concentrations in Groundwater for SWMU 145

Analyte	Time (years)	Peak Conc. Below SWMU 145 (mg/L) ^a	Background Groundwater Concentration (mg/L) ^a	Groundwater Child Resident No Action Level (mg/L) ^a	Retain?
Antimony	1,000	7.99E-02	6.00E-02	5.64E-04	Y
PCB-1260	805	1.92E-03	NA	4.28E-05	Y
Arsenic	1,000	6.21E-02	5.00E-03	3.50E-05	Y
Cadmium	1,000	4.10E-03	1.00E-02	6.61E-04	N
Manganese	1,000	8.44E-01	1.19E-01	3.50E-02	Y
Mercury	850	2.59E-04	2.00E-04	4.44E-04	N
Nickel	1,000	4.14E-03	3.05E-01	3.01E-02	N
Technetium-99	30	1.0106E+04	2.23E+01	1.40E+01	Y
Uranium-238	1,000	2.58E+01	7.00E-01	4.43E-01	Y

^a Units for radionuclides are pCi/L; NA = not applicable; N = No; Y = Yes

The groundwater results presented in Table 5.3 for SWMU 145 show the predicted groundwater concentration of technetium-99 will exceed the MCL at the property boundary and Ohio River. Modeled levels of all remaining SWMU 145 analytes are less than their respective MCLs at the POEs. The following summarizes those analytes that exceeded ELCR or HQ risk criteria of 1.0E-06 and 0.1, respectively.

		Property Boundary	Ohio River
Arsenic	ELCR	4.3E-05	--
	HQ	0.5	--
Technetium-99	ELCR	1.0E-04	5.3E-05
	HQ	--	--

-- = does not exceed

All remaining analytes exhibited HQ values less than 0.1 and ELCR values less than 1.0E-06 at all POEs. Predicted concentrations of SWMU 145 analytes that exceed a HQ of 0.1 and/or an ELCR of 1.0E-06 (Table 5.4) are illustrated in Figures 5.38 and 5.39. As shown in these figures, arsenic is increasing in concentration at the plant boundary at 1,000 years; however, the concentrations are less than the MCL. technetium-99 was predicted to exceed the MCL at the POEs.

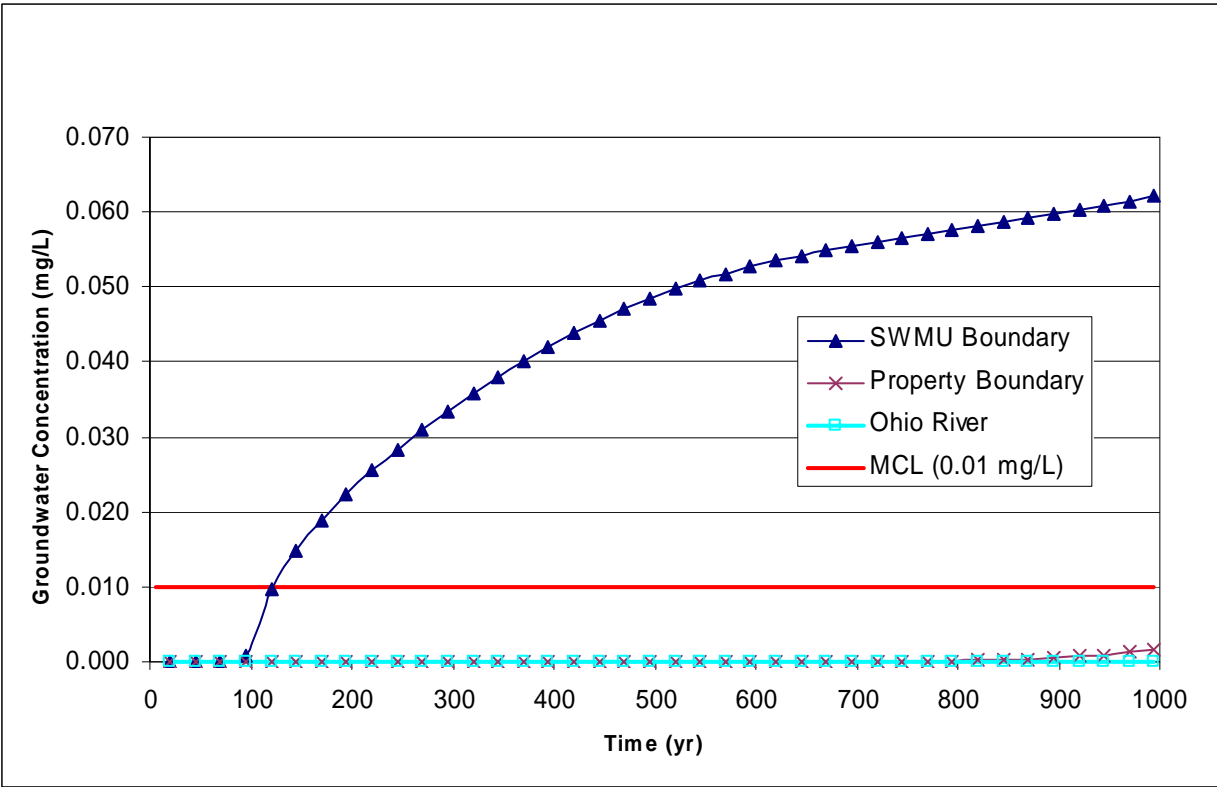


Figure 5.38. Predicted Arsenic Concentrations in Groundwater at the POEs Based on Contaminant Leaching from SWMU 145

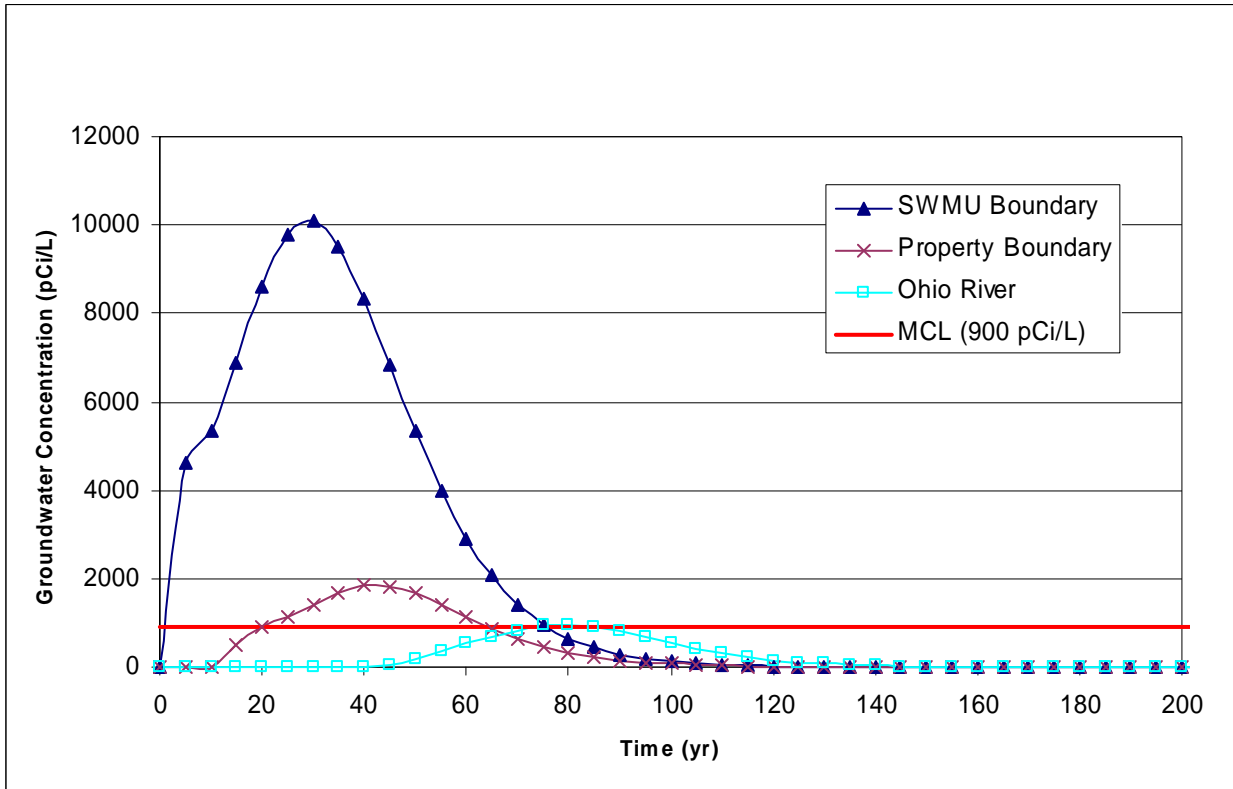


Figure 5.39. Predicted ⁹⁹Tc Activity in Groundwater at the POEs Based on Contaminant Leaching from SWMU 145

5.4 VAPOR TRANSPORT MODELING

The BGOU RI includes vapor transport modeling to evaluate the potential air concentrations in a residential basement for soil and groundwater contamination at the BGOU SWMUs and POEs. Modelers used the Johnson and Ettinger model (1991), coded into spreadsheets by EPA (2004), to assess the potential migration of VOCs into a residential basement (see Appendix E for details of the analysis).

Table 5.13 presents the resulting basement air concentrations, predicted by the model. Table 5.14 summarizes the health and cancer risks calculated in accordance with Appendix A of the Risk Methods Document. The vapor transport modeling for mercury was conservatively based on the metallic form, which has a Henry's Law Constant of $1.07\text{E-}02$ atm-m³/mol. Metallic mercury, with its uniquely high vapor pressure relative to other metals, can enter the atmosphere from the groundwater environment as several different gaseous compounds. The rate of vaporization of mercury and certain of its inorganic compounds decrease in the sequence $\text{Hg} > \text{Hg}_2\text{Cl}_2 > \text{HgCl}_2 > \text{HgS} > \text{HgO}$. The Henry's Law Constant decreases dramatically down the sequence (for example, HgCl_2 has a value of $7.09\text{E-}10$ atm-m³/mol).

The results of the vapor transport modeling (Table 5.14) show that TCE is predicted to have HQ values above 0.1 and/or ELCRs exceeding $1.0\text{E-}06$ for a residential basement exposure above SWMUs 2, 3, 4, 7, and 30. Additional analytes that were found to exceed the HQ value of 0.1 and/or an ELCR value of $1.0\text{E-}06$ within the SWMUs included *cis*-1,2-DCE (SWMUs 2 and 4); vinyl chloride (SWMUs 4 and 7); 1,1-DCE (SWMUs 7 and 30); and mercury (SWMUs 3, 7, 30, and 145).

Derived ELCR values exceeded $1.0\text{E-}06$ for modeled TCE concentrations from groundwater transport at the plant boundary (SWMUs 2 and 4) and property boundary (SWMUs 2 and 4). Modeled vinyl chloride (SWMU 4) and 1,1-DCE (SWMU 7) concentrations also equate to ELCR values greater than $1.0\text{E-}06$ from groundwater transport at the plant boundary.

Table 5.13. Basement Air Concentrations Based on Vapor Transport Modeling Results for Each BGOU SWMU

Source Area	Contaminant	Air concentration (mg/m ³)		
		SWMU Boundary	Plant Boundary	Property Boundary
SWMU 2	TCE	2.81E-02	1.09E-04	5.55E-05
	<i>cis</i> -1,2-DCE	1.95E-01	7.82E-04	3.89E-04
	Naphthalene	2.70E-07	1.56E-08	8.43E-09
SWMU 3	TCE	1.62E-05	8.52E-10	4.47E-10
	Mercury	7.22E-06	1.12E-14	0.00E+00
SWMU 4	TCE	4.90E-03	2.12E-04	1.08E-04
	<i>cis</i> -1,2-DCE	5.76E-03	8.80E-05	4.05E-05
	Vinyl chloride	6.7E-03	1.98E-04	2.55E-06
SWMU 5	TCE	5.41E-06	1.98E-07	9.13E-08
	Acenaphthene	2.04E-07	7.47E-08	4.30E-08
	Fluorene	5.16E-08	2.37E-08	1.27E-08
	Naphthalene	3.80E-06	9.75E-08	3.79E-08
	Pyrene	2.28E-09	NA	NA
SWMU 6	TCE	9.34E-06	3.88E-09	1.92E-09
SWMU 7	TCE	8.63E-05	4.96E-06	7.16E-07
	<i>cis</i> -1,2-DCE	2.13E-04	9.66E-06	1.42E-06
	Vinyl chloride	1.23E-02	1.25E-05	1.22E-06
	1,1-DCE	1.03E-02	6.70E-05	9.03E-06
	Mercury	9.99E-06	2.22E-09	2.41E-12
	Pyrene	7.68E-09	4.93E-12	1.31E-12
	Tetrachloroethene	2.00E-05	6.40E-08	4.70E-09
SWMU 30	TCE	6.75E-05	3.42E-07	2.96E-08
	1,1-DCE	3.36E-05	4.85E-08	3.62E-09
	Acenaphthene	2.77E-08	4.96E-09	9.22E-10
	Fluorene	3.92E-09	NA	NA
	Mercury	1.66E-05	8.91E-1	2.23E-11
	Pyrene	6.56E-10	2.47E-11	6.54E-12
	Naphthalene	3.10E-07	1.90E-08	1.85E-09
SWMU 145	Mercury	1.42E-05	7.95E-08	2.60E-14

Table 5.14. Vapor Hazard Quotients and Risk Based on Vapor Transport Modeling Results for Each BGOU SWMU

Source Area	Contaminant	On-Site			Plant Boundary			Property Boundary		
		HQ	ELCR	HQ	ELCR	HQ	ELCR	HQ	ELCR	
SWMU 2	TCE	3.15E+00	1.84E-03	1.22E-02	7.14E-06	6.22E-03	3.64E-06	0.00E+00	0.00E+00	
	<i>cis</i> -1,2-DCE	2.50E+01	0.00E+00	1.00E-01	0.00E+00	4.99E-02	0.00E+00	0.00E+00	0.00E+00	
	Naphthalene	4.03E-04	0.00E+00	4.99E-06	0.00E+00	1.26E-05	0.00E+00	0.00E+00	0.00E+00	
SWMU 3	TCE	1.82E-03	1.06E-06	9.63E-08	5.68E-11	5.01E-08	2.93E-11	0.00E+00	0.00E+00	
	Mercury	1.08E-01	0.00E+00	1.67E-10	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	
SWMU 4	TCE	5.54E-01	3.23E-04	2.38E-02	1.39E-05	1.21E-02	7.07E-06	0.00E+00	0.00E+00	
	<i>cis</i> -1,2-DCE	7.38E-01	0.00E+00	1.13E-02	0.00E+00	5.19E-03	0.00E+00	0.00E+00	0.00E+00	
	Vinyl Chloride	2.99E-01	4.19E-05	8.85E-03	1.24E-06	1.14E-04	1.60E-08	0.00E+00	0.00E+00	
SWMU 5	TCE	6.06E-04	3.54E-07	2.22E-05	1.30E-08	1.02E-05	5.98E-09	0.00E+00	0.00E+00	
	Acenaphthene	4.37E-06	0.00E+00	1.60E-06	0.00E+00	9.21E-07	0.00E+00	0.00E+00	0.00E+00	
SWMU 6	Fluorene	1.65E-06	0.00E+00	7.57E-07	0.00E+00	4.06E-07	0.00E+00	0.00E+00	0.00E+00	
	Naphthalene	5.67E-03	0.00E+00	1.45E-03	0.00E+00	5.65E-04	0.00E+00	0.00E+00	0.00E+00	
SWMU 7	Pyrene	9.71E-08	0.00E+00	NA	NA	NA	NA	NA	NA	
	TCE	1.05E-03	6.12E-07	4.35E-07	2.54E-10	2.15E-07	1.26E-10	0.00E+00	0.00E+00	
SWMU 30	TCE	9.68E-03	5.65E-06	5.56E-04	3.25E-07	8.03E-05	4.69E-08	0.00E+00	0.00E+00	
	<i>cis</i> -1,2-DCE	2.73E-02	0.00E+00	1.24E-03	0.00E+00	1.82E-04	0.00E+00	0.00E+00	0.00E+00	
	Vinyl Chloride	5.48E-01	7.68E-05	5.59E-04	7.83E-08	5.45E-05	7.64E-09	0.00E+00	0.00E+00	
	1,1-DCE	2.30E-01	3.66E-04	1.50E-03	2.39E-06	2.02E-04	3.21E-07	0.00E+00	0.00E+00	
	Mercury	1.49E-01	0.00E+00	3.31E-05	0.00E+00	3.59E-08	0.00E+00	0.00E+00	0.00E+00	
SWMU 145	Pyrene	3.27E-07	0.00E+00	2.10E-10	0.00E+00	5.58E-11	0.00E+00	0.00E+00	0.00E+00	
	Tetrachloroethene	1.49E-04	8.13E-09	4.78E-07	2.60E-11	3.51E-08	1.91E-12	0.00E+00	0.00E+00	
	TCE	7.57E-03	4.42E-06	3.83E-05	2.24E-08	3.32E-06	1.94E-09	0.00E+00	0.00E+00	
	1,1-DCE	7.52E-04	1.20E-06	1.09E-06	1.73E-09	8.10E-08	1.29E-10	0.00E+00	0.00E+00	
	Acenaphthene	5.93E-07	0.00E+00	1.06E-07	0.00E+00	1.97E-08	0.00E+00	0.00E+00	0.00E+00	
SWMU 145	Fluorene	1.25E-07	0.00E+00	NA	NA	NA	NA	NA	NA	
	Mercury	2.47E-01	0.00E+00	1.33E-05	0.00E+00	3.33E-07	0.00E+00	0.00E+00	0.00E+00	
	Naphthalene	4.62E-04	0.00E+00	2.83E-05	0.00E+00	2.76E-06	0.00E+00	0.00E+00	0.00E+00	
	Pyrene	2.80E-08	0.00E+00	1.05E-09	0.00E+00	2.79E-10	0.00E+00	0.00E+00	0.00E+00	
	Mercury	2.12E-01	0.00E+00	1.19E-03	0.00E+00	3.88E-10	0.00E+00	0.00E+00	0.00E+00	

5.5 FATE AND TRANSPORT UNCERTAINTY

The source inventory, unsaturated zone transport, and saturated zone transport were modeled using the SADA, SESOIL, and AT123D computer codes. The use of these computer codes in the analyses resulted in the use of simplifying assumptions. These assumptions resulted in modeling uncertainties. This section lists the modeling uncertainties and discusses their impacts upon the modeling results. A detailed discussion of the uncertainty in the modeling analyses is provided in Appendix E.

5.5.1 Source Term Development

The analyte screening analysis for the groundwater pathway and modeling is provided in Appendix E, Attachment 3. Very few contaminants detected during the sampling of the BGOU were deleted because of frequency of detection. Many more contaminants were deleted because they were not detected or because the maximum concentration did not exceed the screening criteria. Many organic compounds did not exceed the screening criteria and also could have been screened based on the frequency of detection. Conversely, many contaminants were included in COCs at various SWMUs even though the detection frequency was approximately 1 to 3 percent. These contaminants included vinyl chloride, TCE, 1,1-DCE, naphthalene, and *cis*-1,2-DCE, which were retained at COCs and contributed significantly to overall risk.

Contaminants that were screened based solely on frequency of detection included 1,1,1-trichloroethane, 1,1,2-trichloroethane, 1,2-dichloroethane, dibenzofuran, acenaphthylene, mercury, and silver. The elements or chemicals listed above were screened for frequency at only one or two SWMUs.

The source term was developed using sampling results, geospatial analyses in SADA, and consideration of SESOIL limitations. While the sampling results are appropriate for source identification, SESOIL requires input of the soil concentrations for each layer of interest in the UCRS. Additionally, due to SESOIL's requirement to use the same constant area for each layer, the analyte concentrations of all layers needed to be normalized against the area of the layer with the maximum estimated analyte mass. Geospatial interpolation was used based on the SADA nearest neighbor algorithm to estimate the total mass in each UCRS layer based upon the sampling results.

The techniques in SADA that can be used for source term development are nearest neighbor, inverse distance, and ordinary kriging. The nearest neighbor technique was selected for source zone refinement because it yielded results that were most compatible with the conceptual site model of contaminant release, as described in Attachment 2 to Appendix E.

Each potential analyte source area was discretized using rows and columns with a uniform spacing. Multiple domains with varying depths were used to characterize the analyte source areas vertically in relation to the existing aquifers; therefore, the domain was further discretized into horizontal layers. Analyte results for each domain were compiled, and analyte concentrations in each cell of the domain were predicted using geospatial interpolation (see Appendix E Attachment 2 for details).

The SADA estimated uranium mass in relation to other metals (i.e., vanadium and manganese) appears to be underestimated. The mass of metals, such as vanadium and manganese, also appear to be overestimated using SADA. The SADA interpolation estimates the mass between sample points. This results in an estimated mass of vanadium and manganese in the waste volume based on sample points located outside the waste zone, since these metals tend to be ubiquitous throughout the soils. Likewise, the sample points for uranium outside the waste zone are used to interpolate the mass in the waste zone; however, the transport of uranium from the waste into the surrounding soils is limited due to the sorption of this metal. Since the waste was not sampled, the uranium mass estimates for the waste areas in the

SADA model likely are underestimated due to the limited migration of uranium. The uncertainty in the mass of uranium present in waste will be addressed further during remedial alternatives screening in the feasibility study.

5.5.2 SESOIL and AT123D Transport Uncertainties

SESOIL requires that the same constant area for each layer represented in the model, thus requiring that the analyte concentrations of all layers predicted using SADA be normalized against the area of the layer with the maximum estimated analyte mass. The impact of this normalization was investigated and found that the normalization process has a minor impact on the results (see Appendix E for a detailed discussion).

An additional source of uncertainty in the AT123D modeling runs involves the use of a single hydraulic conductivity and hydraulic gradient. The hydraulic conductivity and gradient are variable from the SWMU locations to the various POEs. The MODPATH model was run to establish the steady-state head distribution in the RGA. MODPATH was used to track flowpaths of particles released from the SWMU location by using the steady-state, head distribution generated by MODFLOW. The distances from the SWMU to the POEs were taken along the flowpaths to determine the distance from the SWMU to the POEs. The hydraulic gradient from the SWMU to the property boundary was estimated using the head difference divided by the distance from the release point to the property boundary POE. The conductivity along the flowpath also was estimated for use in the AT123D model.

Additional uncertainties in the fate and transport analyses include (1) selection of the sorption coefficients (K_d) for uranium in the UCRS, (2) SWMUs on the western side of the plant that may exhibit waste that is below the water table in the UCRS, (3) the fact that SESOIL and AT123D do not consider contaminant transformation such as that for radioactive decay and ingrowth, and (4) the 1,000 year modeling period does not address the fact that ingrowth of progeny from the uranium isotopes and associated risk will, in many cases, continue to increase beyond 1,000 years. These uncertainties are discussed in detail in Appendix E, Section E.3.3, and will be managed in the FS and in future decision and design documents.

5.5.3 Potential Interaction of Sources

The simulations presented in this report for the BGOU SWMUs are based on individual simulations of each SWMU. There is a potential that source plumes from the SWMUs could interact at the POEs. According to the flow paths presented in Figure 5.1, the contaminant plumes from a few of the BGOU SWMUs would interact. The contaminant flow paths from SWMU 6 and SWMU 30 will interact, however, as noted previously, SWMU 6 did not have any groundwater analytes. The contaminant plumes from SWMU 3 and SWMU 5 will interact, and SWMU 2 will interact with a portion of the SWMU 5 contaminant plume. The interaction of the plumes could not be assessed using the SESOIL/AT123D model, since only one SWMU can be discretized in the model for each run.

An evaluation was conducted to ensure that analytes were not eliminated from the groundwater analyses from combined source contributions in the groundwater. The screening evaluation was conducted for the potential interaction of SWMUs 2, 3, and 5. The analysis was based on the conservative summation of the maximum groundwater concentrations below each SWMU provided in Tables 5.5, 5.6, and 5.8. These combined contaminant concentrations are provided in Table E.3.38 and here in Table 5.15. These combined groundwater concentrations then were compared to the child resident NALs from Table A.18 of the 2001 Risk Methods Document and the provisional groundwater backgrounds shown in the 2001 Risk Method Document in Table A.13. The results of the conservative analysis indicate that the selection of analytes evaluated for risk and hazard would not change based on the combination of the source contributions to groundwater from SWMUs 2, 3, and 5.

Table 5.15. Evaluation of Combined Plume Interactions on Selection of Analytes

Analyte	Background Groundwater Concentration (mg/L) ^a	Groundwater Child Resident No Action Level (mg/L) ^a	Peak Conc. Below SWMU 2 (mg/L) ^a	Peak Conc. Below SWMU 3 (mg/L) ^a	Peak Conc. Below SWMU 5 (mg/L) ^a	Sum of Peak Conc. Below SWMUs (mg/L) ^a	Retain as Analyte				
							SWMU 2	SWMU 3	SWMU 5	SUM SWMUs	
Acenaphthene	NA	1.36E-02			6.10E-03	6.10E-03				N	N
Anthracene	NA	7.66E-02			8.06E-03	8.06E-03				N	N
Arsenic	5.00E-03	3.50E-05	3.54E-02	3.29E-02	9.25E-03	7.76E-02	Y	Y	Y	Y	Y
Uranium-234	7.00E-01	5.46E-01	1.58E+00			1.58E+00	Y				Y
Uranium-238	7.00E-01	4.43E-01	1.81E+00	1.59E+01		1.77E+01	Y	Y			Y
Technetium-99	2.23E+01	1.40E+01	1.02E+02	5.56E+03	1.27E+02	5.79E+03	Y	Y	Y	Y	Y
cis-1,2-DCE	NA	2.73E-03	1.15E+01			1.15E+01	Y				Y
Manganese	1.19E-01	3.50E-02	7.16E-01	8.95E-01	1.01E+00	2.62E+00	Y	Y	Y	Y	Y
Mercury	2.00E-04	4.44E-04		9.29E-05		9.29E-05			N		N
Fluorene	NA	9.72E-03			3.63E-03	3.63E-03				N	N
Naphthalene	NA	2.85E-04	9.38E-04		5.55E-03	6.49E-03	Y			Y	Y
Nickel	3.05E-01	3.01E-02			2.01E-03	2.01E-03				N	N
Selenium	5.00E-03	7.54E-03			1.27E-03	1.27E-03				N	N
TCE	NA	1.60E-03	1.48E+00	3.45E-04	9.91E-04	1.48E+00	Y		N	N	Y
Uranium	2.00E-03	9.06E-04	9.86E-03	4.89E-02		5.88E-02	Y	Y	Y	Y	Y
Zinc	4.90E-02	4.50E-01	9.83E-03	9.30E-02	1.58E-01	2.61E-01	N	N	N	N	N

SWMU = solid waste management unit

5.5.4 Location of POEs

The POEs used in the modeling were placed at locations below the SWMU, at the plant boundary, property boundary, Little Bayou seeps, and Ohio River where the greatest contaminant concentrations are expected in the future. By picking locations on the centerline of predicted contaminant plumes as the POEs, the modeling assumed that the hypothetical future resident would pick, by chance, the worst possible location to install a water supply well.

Based on particle tracks taken from the calibrated sitewide numerical flow model developed in MODFLOW for PGDP, SWMUs 2, 4, 5, and 145 were shown not to impact the Little Bayou Seeps. If the SWMUs were to impact the seeps, it has been shown that SWMUs 2 and 145 have modeled groundwater concentrations at the Ohio River that exceed MCLs for several analytes; therefore, the modeled groundwater concentrations at the Little Bayou Seeps also would exceed the MCLs for these analytes. Since there is uncertainty with the flow paths and the seeps radius of influence, the contaminants in SWMU 4 that exceed the MCL at the property boundary POE were evaluated to see if they would exceed the MCL at the Little Bayou seeps. The modeling results indicated that only one contaminant, TCE, exceeded the MCL at the Little Bayou seeps (i.e., 0.083 mg/L); however, TCE also exceeds the MCL at the Ohio River (i.e., 0.077 mg/L) for SWMU 4. Modeling results for SWMU 5 show that the groundwater concentrations at the property boundary do not exceed the MCLs for any analytes modeled; therefore, the groundwater concentrations at the Little Bayou Seeps also would be less than the MCLs for each analyte.

5.5.5 Future Environmental Changes

Several future environmental changes at the PGDP could impact the accuracy of the modeling predictions. These changes include plant shutdown and dam operation on the Ohio River. In a previous modeling effort for a landfill at PGDP, several sensitivity analyses were performed (DOE 2003) to examine the impacts those changes may have on groundwater flow and contaminant transport. The sensitivity analysis of the groundwater travel time due to plant shutdown was studied by varying the recharge over a range of values. The results of the analysis indicated that a decrease in the recharge rate resulted in a monotonic increase in the travel time to the receptor. Thus, chemicals that have short degradation half-lives would show a decrease in concentration due to plant shutdown.

The Olmstead Dam operation is expected to increase the stage (water level) of the Ohio River; therefore, a sensitivity analysis was conducted (DOE 2003) to assess changes in groundwater travel time in relation to dam operation by increasing the river stage between 304.44 ft amsl and 310.04 ft amsl (the baseline river stage is 300.04 ft amsl). The results of the analysis indicated that the travel times in the aquifer changed very little in relation to the Ohio River stage; therefore, the dam operation would have little impact on the results shown in this report.

5.5.6 Burial Cell Waste

Sample data around and beneath the BGOU SWMUs were used to develop a source inventory of contaminants. The premise of this source inventory development is based on the inherent assumption that the contaminants around and beneath the BGOU SWMUs represent the release mass from the Burial Ground disposal cells. The groundwater transport analyses do not model potential future releases directly from the SWMU burial cells.

Waste at several SWMUs was containerized in drums before disposal. Previous inspections of buried drums at PGDP have indicated that the drums were highly corroded. It is considered unlikely that a significant portion of the drummed waste (particularly that buried in steel drums) still is containerized at the BGOU SWMUs due to the length of time the drums have been buried and, thus, susceptible to a corrosive environment. The drums were not modeled in this RI report due to the overall objectives of the RI analyses and uncertainty in the degradation process. Due to the uncertainty in the degradation of the drummed waste, real measured sample data surrounding the SWMUs were used to evaluate the potential risk from the SWMU waste. Test pits at SWMUs 7 and 30, discussed in Sections 4.81 and 4.91, respectively, sampled the waste in the early 1990s and also reported corroded drums. The health-protective approach resulted in all SWMUs with drummed waste exhibiting risk and hazard values that exceeded acceptable levels. All SWMUs, therefore, require screening of remedial alternatives in the BGOU FS.

5.5.7 SWMU 4 RGA TCE Source

The TCE source in SWMU 4 was assessed in this RI based on soil sample results. As discussed in Section 4.5.2, a potential DNAPL zone, less than 200 ft wide, also may be present at the base of the RGA as evidenced by a discrete area with TCE concentrations greater than 10,000 µg/L in the lower RGA immediately downgradient of the SWMU. The volume of soil potentially contaminated with TCE DNAPL at this SWMU is estimated to be approximately 31,480 yd³. This estimate assumes a source area that is 100 ft by 100 ft with a thickness of 85 ft (depth to base of RGA, which is 100 ft minus the estimated depth to base of the waste cell of 15 ft). The DNAPL source term for TCE in the RGA at SWMU 4 was not evaluated in the RI modeling analyses since the RGA concentrations in the lower RGA currently exceed the MCL. The UCRS TCE source concentrations were sufficient to indicate that actions should be taken for SWMU 4. The uncertainties related to source term size and location will be better defined in a remedial design investigation prior to the design and implementation of a remedy.

5.5.8 SWMU 3 UCRS Groundwater Contamination

The groundwater analyses conducted for this RI are based on soil samples obtained from soils surrounding the SWMUs and their subsequent release to the RGA and transport through the RGA. In some instances, water samples from wells in the UCRS indicated additional contaminant concentrations that were not accounted for in the analyses. For example, UCRS wells MW85, MW88, MW91, and MW94 at SWMU 3 indicated elevated levels of TCE. The water data were added to the SWMU 3 TCE soil concentrations and a SADA nearest neighbor interpolation was assessed. The resulting transport analyses indicated that the TCE concentrations were below the MCL.

6. BASELINE HUMAN HEALTH RISK ASSESSMENT

This Baseline Human Health Risk Assessment (BHHRA) utilizes information collected during the recently completed RI of the BGOU SWMUs, in addition to information collected during previous investigations listed here:

- *Remedial Investigation Report for Waste Area Grouping 3 at the Paducah Gaseous Diffusion Plant, Paducah, Kentucky*, DOE/OR/07-1895/V1-V4&D1, U.S. Department of Energy, Paducah, KY (DOE 2000a).
- *Remedial Investigation Report for Solid Waste Management Units 7 and 30 of Waste Area Group 22 at the Paducah Gaseous Diffusion Plant, Paducah, Kentucky*, DOE/OR/07-1604/V1&D2, U.S. Department of Energy, Paducah, KY (DOE 1998c).
- *Remedial Investigation Addendum for Waste Area Grouping 22, Burial Grounds Solid Waste Management Units 2 and 3 at the Paducah Gaseous Diffusion Plant, Paducah, Kentucky*, DOE/OR/07-1141&D2, U.S. Department of Energy, Paducah, KY (DOE 1994).
- *Data Summary and Interpretation Report for Interim Remedial Design at Solid Waste Management Unit 2 of Waste Area Group 22 at the Paducah Gaseous Diffusion Plant, Paducah, Kentucky*, DOE/OR/07-1549&D1, U.S. Department of Energy, Paducah, KY (DOE 1997a).

The purpose of this BHHRA is to characterize the baseline risks posed to human health from contact with contaminants in soil and water at these SWMUs and at locations to which contaminants may migrate. A summary of the data used is presented in Attachment F1 to Appendix F.

Part of Goal 2 for the BGOU RI, as presented in the BGOU Work Plan (DOE 2006a), was to determine if contaminants at the BGOU units are contributing to groundwater contamination; this risk assessment supports that goal by using modeled concentrations of contaminants to the RGA to support the refinement of an assessment of risks to human health and the environment through groundwater. The Work Plan also specified that the RI should include a risk assessment for residential, industrial, and recreational receptors. Risk assessments for each of those scenarios are presented here. The information collected during the RI, the earlier historical data, and the results of this BHHRA will be used to determine if sufficient data are available to evaluate risk and to determine if response actions to reduce risks are needed and, if needed, to screen among response action alternatives.

The methods and presentations used in this BHHRA are consistent with those presented in the Risk Methods Document (DOE 2001). The Risk Methods Document integrates the human health risk assessment guidance from the EPA and the Kentucky Department for Environmental Protection (KDPEP) and incorporates instructions contained in regulatory agency comments on earlier risk assessments performed for PGDP.

6.1 DATA COLLECTION AND EVALUATION

The four previous reports listed above contain risk assessment results for one or more of the burial grounds considered in this RI. The results of these assessments are summarized here, as well as in Section F.1 of Appendix F and the BGOU Work Plan, and are presented in more detail in Attachment F2 to Appendix F. Risks and hazards for soil exposure presented in this BHHRA are taken from these four

previous assessments. For groundwater, these previous assessments were based on measured groundwater concentrations, while this risk assessment used modeled concentrations. Differences can be found in COCs, risk, or hazard level among these previous assessments based on measured concentrations and those resulting from the modeled concentrations presented in this risk assessment. These differences may result from factors such as overestimation by the model or of the source term in the model, or the differences may represent contributions from background or other sources to the measured concentrations in the wells.

6.1.1 Identification of COPCs

Soil COPCs previously were determined in the investigations listed and as outlined in Appendix F, Section F.2. This subsection describes the process used to determine the list of groundwater COPCs used in this BHHRA, including the sources of data and the procedures used to screen the data.

Soil data used to model groundwater concentrations in the BHHRA describing current contaminant concentrations in groundwater at SWMU 2, SWMU 3, SWMU 4, SWMU 5, SWMU 6, SWMU 7, SWMU 30, and SWMU 145 were derived from the recently completed BGOU RI sampling, as well as historical data acquired from the PGDP OREIS database. Methods used to collect and analyze subsurface soil samples used for groundwater modeling as well as the soil data itself were examined to ensure that sampling methods were adequate for determining the nature and extent of contamination and were representative of site conditions. It was determined that samples of the BGOU RI and those selected from the Paducah OREIS database were collected using appropriate methods that were consistent with each project's work plan.

Soil results were screened against a number of criteria prior to their use in groundwater modeling to determine groundwater COPCs for use in the BHHRA. Soil results were used for samples collected from the ground surface to approximately 60 ft bgs. The following outlines the review and screening process used prior to establishing the COPCs for use in modeling:

- *Units of Reported Results.* The units of measure used for analyte classes (i.e., inorganic chemicals, organic compounds, and radionuclides) were assigned consistent units of measure. The units of measure used were mg/kg for inorganic chemicals and organic compounds and pCi/g for radionuclides.
- *Detection Status.* Each result was coded either as a detect or nondetect based on the data qualifier codes present in the database. Results assigned a "U" or "UJ" qualifier were considered nondetects. This coding subsequently was used to calculate the frequency of detection for each constituent.
- *Frequency of Detection.* Those constituents detected in less than 5% of the samples were not considered a COPC.
- *Essential Nutrients.* Results for the seven essential nutrients were removed from the data sets. They are: calcium, chloride, iodine, magnesium, potassium, sodium and phosphorous.
- *Protactinium-234m, potassium-40, and thorium-234.* All results for protactinium-234 were removed from consideration in the BHHRA. Because the uranium-238 toxicity value incorporates the contribution to cancer risk from protactinium-234 and other short-lived progeny, its inclusion would result in a double-counting of the cancer risk had it been retained. Thorium-234 has a half-life so short (24.1 days) that exposure to it is not relevant on the timescale of a risk assessment. Potassium-40 was eliminated because it is ubiquitous.

Analytes retained as COPCs under current conditions are presented for each SWMU in the tables in Appendix F, Section F.2.3.2. The maximum detected soil concentrations (surface to 60 ft bgs) are presented by analyte and by SWMU, with a comparison to the child resident SSLs¹ with a DAF=1. Those analytes with a maximum concentration greater than their respective SSLs were then compared to soil/sediment child resident NALs. Those constituents greater than both the SSL and NAL were retained as COPCs for groundwater modeling. The screening values used may be found in Tables A.7 and A.17 of the Risk Methods Document (DOE 2001). Exceptions to this screening are TCE, technetium-99 and uranium isotopes, which were retained in all SWMUs, as they are significant risk contributors or known to be part of the facility's process history.

Following this review and screening process, the COPCs retained then were modeled as described in Section 5 and Appendix E of the RI Report. Modeled results exceeding both the child resident NALs from Table A.18 and the provisional groundwater backgrounds shown in Table A.13 of the 2001 Risk Method Document then were carried through the toxicity and exposure assessments, and cancer risk and hazards indices were calculated for the Rural Resident Groundwater User.

6.1.2 Points of Exposure for Groundwater COPCs

Groundwater concentrations used in the assessment were modeled to several locations for examination of potential Rural Resident exposure. These were as follows:

- SWMU boundary
- Plant boundary
- Property boundary
- Little Bayou seeps (when particle modeling showed a contribution to the seep), and
- Monitoring well located near the Ohio River

This risk assessment uses the modeled groundwater concentrations at all POEs. A screening of measured concentrations in the groundwater against NALs and action levels is presented in Appendix E of the BGOU Work Plan as ancillary information only. A list of COCs from that screening of measured groundwater is reproduced in Appendix F, Section F.1.5.

6.2 EXPOSURE ASSESSMENT

This section describes the exposure assessment used to determine the pathways of exposure that were considered for the surface and subsurface soil and groundwater at the source units that are part of the BGOU RI. Specifically, the exposure settings of the BGOU are described, the routes of exposure are outlined, and the daily intakes and doses are presented.

6.2.1 Characterization of Exposure Setting

As shown in the physical descriptions presented in Appendix F, Section F.3.2, current land use of all sources investigated during the BGOU RI is industrial. Under current use, because of access restrictions, only plant workers and authorized visitors are allowed access to the source areas. As discussed in the PGDP Site Management Plan (DOE 2009), foreseeable future land use of the area is expected to be industrial as well.

¹ SSLs are risk-based soil concentrations considered to be protective of groundwater (DOE 2001).

At present, both recreational and residential land uses occur in areas surrounding PGDP. Recreational use occurs in the WKWMA. The WKWMA is used primarily for hunting and fishing, but other activities include horseback riding, field trials, hiking, and bird watching. An estimated 5,000 fishermen visit the area annually, according to the Kentucky Department of Fish and Wildlife Resources manager of the WKWMA. Residential use near the plant and in areas to which the groundwater from the BGOU may migrate is rural residential and includes agricultural activities. However, current response actions have eliminated exposure to contaminated groundwater by these rural residents; as part of the WAG 26 Groundwater Removal Action completed August 30, 1994, DOE extended municipal water lines to residences affected by off-site groundwater contamination. More urban residential use occurs in the villages of Heath, Grahamville, and Kevil, which are within 3 miles of DOE property boundaries, but outside of the area projected to be potentially impacted by the BGOU. The closest major urban area is the municipality of Paducah, Kentucky, which has a population of approximately 26,000 and is approximately 10 miles east of PGDP. Other municipalities in the region near PGDP are Cape Girardeau, Missouri, which is approximately 40 miles west of the plant; and the cities of Metropolis and Joppa, Illinois, which are across the Ohio River from PGDP. Total population within a 50-mile radius of the plant is approximately 732,000 people, with about 88,500 people living within 10 miles. The population of McCracken County, in which PGDP lies, is estimated at 65,000 people.

In the area near PGDP and in western Kentucky, in general, the economy historically has been agricultural based; however, industry has increased in recent years. PGDP is a major employer, with approximately 1,400 workers. Another major employer near PGDP is the TVA Shawnee Steam Plant, which employs approximately 260 individuals.

6.2.2 Identification of Exposure Pathways

The following discussions focus on points of potential human contact, types of receptors, and exposure routes that are relevant to exposure to contaminated groundwater and soil evaluated in this and previous BHHRA.

6.2.2.1 Points of Human Contact—Land Use Considerations

The potential BGOU sources are located within a large industrial facility; therefore, the current land use is industrial. Per KDEP and EPA agreement (Risk Methods Document), industrial land use limits the current exposure scenario to an industrial worker (with exposure to the first ft of surface soil) and an excavation worker (with potential exposure to soil in the 0-10 ft bgs depth). The current scenarios do not include any current use of groundwater drawn from the RGA at the sources.

The current land use can be expected to continue in the foreseeable future, and the most plausible future land use of the BGOU sites also is industrial. In the future, the expected exposure frequencies and durations may be higher than duration and frequency of the current exposure. Additionally, use of groundwater drawn from the RGA at the BGOU sources is not expected; however, uses of areas surrounding PGDP indicate that it would be prudent to examine a range of land uses to provide decision makers with estimates of the risk that may be posed to humans under alternate uses, however unlikely. In addition, consideration of a range of land uses is consistent with the approved BGOU RI Work Plan (DOE 2006a). The BHHRA reports the hazards and risks for current and several hypothetical future uses, consistent with regulatory guidance and the approved BGOU RI Work Plan (DOE 2006a). PGDP is an industrial facility and future land use is expected to remain industrial. The future on-site rural resident is not a likely land-use scenario. These factors should be considered in examination of risk information provided in this report. The following future land uses are included in the BHHRA:

- **Future on-site industrial use**—direct contact with surface soil (0 to 1 ft bgs).

- **Future on-site excavation worker**—direct contact with surface and subsurface soil (0 to 10 ft bgs).
- **Future on-site recreational user**—direct contact with surface soils and consumption of game exposed to contaminated surface soil.
- **Future on-site rural resident**—direct contact with surface soil and use of modeled groundwater concentrations from the RGA at source areas, as well as vapor intrusion into a residential basement located above the source.
- **Future off-site rural resident**—use in the home of groundwater drawn from the RGA as well as vapor intrusion into basements at the DOE plant boundary, the DOE property boundary, at Little Bayou seeps (when appropriate) and at the Ohio River.

6.2.2.2 Potential Receptor Populations

The reasonably expected potential receptor population under current and future conditions at the source units are the industrial worker and the excavation worker. Potential receptor populations under future conditions in BGOU areas also include hypothetical recreational and residential exposures. The receptor populations for these scenarios contain age cohorts. For the recreational users, the cohorts include the child (aged 1 to 7), teen (aged 8 to 20), and the adult (older than 21). For rural residents, the cohorts include children (aged 1 to 7) and older individuals (termed adults in this and previous BHHRA). Finally, this and earlier assessments assume that the recreational user is a rural resident who has repeated access to the study area. Recreational users not residing in the study area are not considered separately because nearby residents were determined to be the individuals most likely to take part in recreational activities at PGDP on a continual basis. In addition, the exposure assessment determined that little information useful in remedy selection would be obtained by including a separate visiting recreational user in the assessment. All SWMUs were assessed for these exposure scenarios, with the following exceptions:

- *SWMUs 2 and 3*. SWMUs 2 and 3 were assessed together in the WAG 22 RI. (DOE 1994) Assessment of future residential and excavation worker exposure to on-site soils was not included in the WAG 22 RI and, therefore, not in the current BGOU D2 RI. The risk associated with both these scenarios would be greater than the risk for the industrial worker because industrial worker soil exposure exhibited risks above *de minimis* levels and indicates a need for action; therefore, assessment of the residential and excavation scenarios is not warranted.
- *SWMU 145*. SWMU 145 is partially located beneath SWMUs 9 and 10 and, as a result, limited soil data are available for SWMU 145. Only groundwater exposure pathways were considered in assessing this SWMU. Uncertainty associated with soil exposure at SWMU 145 is addressed in the BGOU FS.

6.2.2.3 Exposure Points and Exposure Routes

Human health risks are assessed by determining POEs and exposure routes. POEs are locations where human receptors can contact contaminated media. Exposure routes are the processes by which human receptors contact contaminated media. The reasons for selecting or not selecting each exposure route for each of the potentially exposed populations are presented in this BHHRA. The exposure routes that were quantitatively assessed in this and previous BHHRAs are listed below. The exposure routes that were quantitatively assessed in this BHHRA using modeled groundwater are highlighted with an asterisk (*).

Further discussion of the rationale behind including these exposure routes in the BHHRA can be found in Appendix F, Section F.3.3.3.

- Ingestion of groundwater as a drinking water source*
- Inhalation of volatile constituents emitted while using groundwater*
- Dermal contact with groundwater while showering*
- Inhalation of vapors released from groundwater into home basements*
- Vapor intrusion into a basement of a residence*
- Incidental ingestion of contaminated soil
- Dermal contact with contaminated soil
- Inhalation of particulates emitted from contaminated soil
- Inhalation of volatile constituents emitted from contaminated soil
- External exposure to ionizing radiation emitted from contaminated soil
- Consumption of game contaminated by consumption of vegetation grown in contaminated soil
- Ingestion of produce

6.2.2.4 Calculation of Exposure Point Concentrations

Groundwater exposure point concentrations (EPCs) used to determine potential future risks for residential use of groundwater at four POEs (i.e., unit boundary, plant boundary, property boundary, and either Ohio River or Little Bayou seeps) were developed from soil data through groundwater modeling. The modeled concentrations in groundwater over time at the four POEs are provided in the figures in Section 5.

6.2.2.5 Calculation of Chronic Daily Intakes

Chronic daily intakes (CDIs), which are calculated for inorganic and organic constituents, and radionuclide intakes, calculated for radionuclides, represent the exposure to a COPC as mass contacted per unit body weight per unit time for the applicable receptor (EPA 1991). CDIs and radionuclide intakes are calculated using the values presented in Tables F.1 through F.4 of Appendix F and are from the 2001 approved version of the Risk Methods Document, except where a footnote indicates that the value is from the draft 2008 revision of the Risk Methods Document. Values in these tables marked as “chemical-specific” were obtained from tables in Appendix B and Appendix D of the draft 2008 Risk Methods Document.

These CDIs derived are presented in Tables F.6 through F.29 of Appendix F. In this presentation, the CDIs used to estimate hazard indices (HIs) (i.e., noncarcinogenic effects) are presented first, and then followed by the values used to estimate ELCRs.

6.3 TOXICITY ASSESSMENT

Many of the toxicological summaries included in Appendix F, section F.4 were obtained from the *Risk Assessment Information System* (RAIS) prepared by the Toxicology and Risk Analysis Section of Oak Ridge National Laboratory for DOE (DOE 2004b). This site also lists toxicity values taken from the EPA’s Integrated Risk Information System (IRIS) database (EPA 2004b), National Center for Environmental Assessment (NCEA), and Health Effects Assessment Summary Tables (HEAST) database (EPA 1998a). This list formed the basis of the toxicity values reported in Appendix F. For those chemicals not profiled in RAIS, a brief summary of information drawn from Agency for Toxic Substances and Disease Registry (ATSDR) or other library research sources is included in this section.

The last paragraph of each profile contains the toxicity values used in this BHHRA. These values also are summarized in Tables 6.1 through 6.4 for groundwater COPCs.

The slope factor for chemicals is defined as a plausible upperbound estimate of the probability of a response (i.e., development of cancer) per unit intake of a chemical over a lifetime (EPA 1989). Slope factors are specific for each chemical and route of exposure.

Toxicity values used in risk calculations also include the chronic RfD, which is used to estimate the potential for systemic toxicity or noncarcinogenic risk. The chronic RfD is defined as an estimate of a daily exposure level for the human population, including sensitive subpopulations, that is likely to be without an appreciable risk of deleterious effects during a lifetime (EPA 1989). RfD values also are specific to the chemical and route of exposure.

Table 6.1. Toxicity Values For Chronic Exposure to Carcinogens Via the Ingestion and Inhalation Exposure Routes

COPC ^a	Class	Oral Slope		Oral Unit Risk ^d	Inhalation Slope Factor ^e	Inhalation		Types of Cancers
		Factor ^b	Factor Source ^c			Slope Factor Source ^c	Unit Risk ^f	
<i>Inorganic Chemicals (Metals)</i>								
Arsenic	A	1.50E+00	a	5.00E-02	1.51E+01	a	4.30E-03	Respiratory system tumors
<i>Organic Compounds</i>								
1,1-DCE	C	6.00E-01	b	1.70E-02	1.75E-01	a	5.00E-05	Kidney, adenocarcinoma
Aroclor-1254	B2	4.00E-01	a		3.50E-01	a	5.71E-01	Liver
Aroclor-1260	B2	4.00E-01	a		3.50E-01	a	5.71E-01	Liver
TCE ^h	C-B2	3.22E-01	c		3.22E-01	c	1.10E-01	Liver and lung cancer
Vinyl Chloride	A	1.50E+00	a	4.20E-02	3.08E-02	a	8.80E-06	Liver, lung, digestive track, and brain tumors
<i>Radionuclides</i>								
	ICRP ^g Lung Class							
Technetium-99	M	2.75E-12	d		1.41E-11	d		Various
Uranium-234	M	7.07E-11	d		1.14E-08	d		Various
Uranium-238	M	8.71E-11	d		9.35E-09	d		Various

Note: Blank cells indicate that data are not available or are not appropriate.

^a All COPCs are listed.

^b The units for the oral slope factors are (mg/kg × day)⁻¹ for nonradionuclides and risk/pCi for radionuclides.

^c Source codes are defined as follows:

a: *Integrated Risk Information System (IRIS)* (EPA 2004)

b: This value has been withdrawn by EPA's National Center for Environmental Assessment (NCEA) and is being reassessed for IRIS. It is acceptable for use in this risk assessment, as no replacement value currently exists (EPA 2000)

c: KDEP

d: *Health Effects Assessment Summary Tables (HEAST)* (EPA 1998b)

^d The units for the oral unit risks are (mg/L)⁻¹.

^e The units for the inhalation slope factors are (mg/kg × day)⁻¹ for nonradionuclides and risk/pCi for radionuclides.

^f The units for inhalation unit risks are m³/μg.

^g ICRP Publication 72 is referenced in the HEAST user's guide (ICRP 1996). Lung class absorption types are defined as follows:

S = slow (particulate)

M = medium (particulate)

F = fast (particulate)

^h Value used is from KDEP (2004) review of TCE slope factors. The slope factors used in previous assessments were 0.052 for the oral slope factor and 0.002 for the inhalation slope factor. This issue is discussed further in the uncertainty section.

Table 6.2. Toxicity Values for Chronic Exposure to Noncarcinogens Via the Ingestion and Inhalation Exposure Routes

COPC ^a	Oral Reference Dose ^b	Oral Reference Dose Source ^c	Inhalation Reference Dose ^d	Inhalation Reference Concentration ^e	Inhalation Reference Concentration Source ^e	RfD basis (vehicle) ^f	Target Organ Critical Effect	Confidence Level ^f	Uncertainty Factor/Modifying Factor ^f
<i>Inorganic Chemicals (Metals)</i>									
Antimony	4.00E-04	a				(O)LOAEL GI		(O)Low	(O)UF=1,000 (O)MF=1
Arsenic	3.00E-04	a				(O)NOAEL Skin /LOAEL		Medium	(O)UF=3 (O)MF=1
Manganese	4.60E-02	a,b	1.43E-05	5.00E-05	a	(I)LOAEL CNS (O)NOAEL		(I)Medium (O)Medium	(I)UF=1,000 (I)MF=1 (O)UF=1 (O)MF=1
Selenium	5.00E-03	a				NOAEL/ LOAEL Lungs (selenosis)		High	(O)UF=3 (O)MF=1
Uranium	6.00E-04	a,d				LOAEL Kidney		NA	(O)UF=100 (O)MF=1
<i>Organic Compounds</i>									
1,1-DCE	5.00E-02	a	5.71E-02	2.00E-01	a	LOAEL	Liver	Medium	(O)UF=1,000 (O)MF=1
1,2-DCE, <i>cis</i> -	1.00E-02	x	9.97E-03	3.49E-02	ex	NOAEL	Blood	Low	(O)UF=3,000 (O)MF=1
Aroclor-1254	2.00E-05	a	1.99E-05		ex	(O)LOAEL	Endocrine System Liver, kidney, CNS	Medium	(O)UF=300 (O)MF=1
TCE	3.00E-04	c	1.14E-02	4.00E-02	c	NA		NA	NA
Vinyl Chloride	3.00E-03	a	2.86E-02	1.00E-01	a	(I)NOAEL/ LOAEL (O)NOAEL /LOAEL	Liver, kidney, CNS	Medium	(I)UF=30 (I)MF=1 (O)UF=3 (I)MF=1
Naphthalene	2.00E-02	a	8.57E-04	3.00E-03	a	(O)NOAEL (I)LOAEL	Decreased body weight Respiratory	(O)Low (I)Medium	(O)UF=3,000 (O)MF=1 (I)UF=3,000 (I)MF=1

Notes: Blank cells indicate that data are not available or are not appropriate. NA=information not readily available at this time; GI=gastrointestinal; CNS=central nervous system

^a All COPCs are listed.

^b The units for the oral reference doses are mg/(kg × day).

^c Source codes are defined as follows:

a: *Integrated Risk Information System (IRIS)* (EPA 2004)

b: IRIS no longer separates manganese values for chronic oral RfDs into water and diet RfDs. IRIS recommends using a modifying factor of 3 to lower the total oral RfD of 1.40E-01 to 4.67E-02. This has been rounded to 4.6E-02 to make the value more conservative

c: EPA National Center for Environmental Assessment (NCEA) (EPA 2001)

d: Also see Soil Screening Guidance for Radionuclides: User's Guide.

ex: Value is extrapolated from the oral reference dose.

x: A provisional value from EPA National Center for Environmental Assessment (NCEA).

^d The units for the inhalation reference doses are mg/(kg × day).

^e The units for the inhalation reference concentrations are mg/m³.

^f O=oral; I=inhalation; UF=uncertainty factor; MF=modifying factor; NA=not available.

Table 6.3. Toxicity Values for Chronic Dermal Contact Exposure to Carcinogens

COPC ^a	Dermal Slope Factor ^b	Dermal Slope Factor Source ^c	GI ABS Factor ^d
<i>Inorganic Chemicals (Metals)</i>			
Arsenic	3.66E+00	a	0.41
<i>Organic Compounds</i>			
1,1-DCE	6.00E-01	b	1.0
Aroclor-1254	4.44E-01	a	0.9
Aroclor-1260	4.44E-01	a	0.9
TCE	2.67E+00	c	0.15
Vinyl Chloride	1.50E+00	a	1

Note: Blank cells indicate that data are not available or are not appropriate.

^a All groundwater COPCs are listed.

^b The units for these dermal dose slope factors are (mg/kg × d)⁻¹ for nonradionuclides. Absorbed cancer slope factors are calculated by dividing the administered cancer slope factor by GI absorption factor; this value is used in the BHHRA to calculate contribution to cancer risk from dermal exposure.

^c Sources for dermal slope factor:

a: *Integrated Risk Information System (IRIS)* (EPA 2004)

b: This value has been withdrawn by EPA's National Center for Environmental Assessment (NCEA) and is being reassessed for IRIS. It is acceptable for use in this risk assessment, as no replacement value currently exists (EPA 2000)

c: KDEP

^d All GI ABS factors from 2008 DRAFT Risk Methods Document (DOE 2008b).

Table 6.4. Toxicity Values for Chronic Exposure to Noncarcinogens Via the Dermal Contact Exposure Route

COPC ^a	Dermal Reference Dose ^b	Administered Reference Dose ^c	GI ABS ^d
<i>Inorganic Chemicals (Metals)</i>			
Antimony	8.00E-06	4.00E-04	0.02
Arsenic	1.23E-04	3.00E-04	0.41
Manganese	1.84E-03	4.60E-02	0.04
Selenium	2.20E-03	5.00E-03	0.44
Uranium ^e	5.10E-04	6.00E-04	0.85
<i>Organic Compounds</i>			
1,1-DCE	5.00E-02	5.00E-02	1
1,2-DCE, <i>cis</i> -	1.00E-02	1.00E-02	1
Aroclor-1254	1.80E-05	2.00E-05	0.9
Naphthalene	1.60E-02	2.00E-02	0.8
TCE	4.50E-05	3.00E-04	0.15
Vinyl Chloride	3.00E-03	3.00E-03	1

Note: Blank cells indicate that data are not available or are not appropriate.

^a All COPCs are listed.

^b The units for the absorbed doses are mg/(kg × day). All dermal reference doses were calculated by multiplying the administered reference dose by the GI absorption factor; this value is used in the BHHRA to calculate contribution to the hazard index from dermal exposure. Based on Risk Assessment Guidance for Superfund Volume I: Human Health Evaluation Manual (Part E, Supplemental Guidance for Dermal Risk Assessment) Final EPA/540/R/99/005 OSWER 9285.7-02EP PB99-963312 July 2004.

^c Administered reference doses are equivalent to the oral reference dose listed in Table 6.2. The units are mg/(kg × day). *Integrated Risk Information System (IRIS)* (EPA 2004)

^d GI absorption factors are from the 2008 DRAFT Risk Methods Document (DOE 2008b) and are unitless.

^e Uranium Source: 40 *CFR* Part 141 (2000).

6.4 RISK CHARACTERIZATION

Risk characterization is the final step in the risk assessment process. In this step, the information from the exposure and toxicity assessments is integrated to quantitatively estimate both carcinogenic health risks and noncarcinogenic hazard potential. For this assessment, risk is defined as both the lifetime probability of excess cancer incidence for carcinogens and the estimate of daily intake exceeding intake that may lead to toxic effects for noncarcinogens. Equations used in the BHHAs to determine the HIs and ELCRs are presented in Sections F.5 and F.5 of Appendix F. These results are summarized in Tables 6.5 through 6.12, as well as in Appendix F, Section F.5.

6.4.1 Risk Characterization for Soil

Results from previous risk assessments were used for the soil risk characterization. Results of previous risk assessments are available for SWMUs 2, 3, 4, 5, 6, 7, and 30 (DOE 1994; DOE 1998c; DOE 2000a). The results for systemic toxicity (HI) and ELCR for soil exposure are discussed below. The percent contribution listed for each soil COC is listed with the same accuracy (i.e., 2% or 2.0%) as the original document from which the value was taken.

6.4.1.1 Future Industrial Worker

Cumulative HIs for the industrial worker were greater than 1 at SWMUs 4, 7, and 30 based on soil exposure.

- At SWMU 4, chromium, iron, and vanadium were the primary drivers contributing 45%, 24%, and 24% to the HI, respectively.
- At SWMU 7, beryllium, chromium, iron, manganese, uranium, and vanadium were the major drivers (> 5%) contributing 9.6%, 13.6%, 20.6%, 10.7%, 13.7%, and 17.7% to the HI, respectively.
- At SWMU 30, aluminum, beryllium, chromium, iron, manganese, uranium, and vanadium were the major drivers contributing 5.1%, 10.8%, 13.5%, 19.8%, 11.3%, 9.0%, and 17.6% to the HI, respectively.

Cumulative ELCRs exceeded 1E-04 for all SWMUs and were greater than 1E-03 at SWMU 7 and SWMU 30 for exposure to soil (SWMU 145 was not evaluated for this scenario). The following summarizes the cumulative risk estimates and major contributors (> 5%) to the ELCR for these SWMUs.

- SWMU 2 cumulative ELCR 1.20E-04; drivers are uranium-235+daughters at 83.9% and uranium-238+daughters at 10.7%.
- SWMU 3 cumulative ELCR 1.20E-04; drivers are uranium-235+daughters at 83.9% and uranium-238+daughters at 10.7%.
- SWMU 4 cumulative ELCR 5.40E-04; the primary driver is beryllium at 97%.
- SWMU 5 cumulative ELCR 4.10E-04; drivers are arsenic at 6%, beryllium at 49%, and Total PAH at 45%.
- SWMU 6 cumulative ELCR 2.40E-04; drivers are beryllium at 90% and Total PAH at 10%.

- SWMU 7 cumulative ELCR 3.90E-03; the primary driver is beryllium at 96%.
- SWMU 30 cumulative ELCR 3.80E-03; the primary driver is beryllium at 96.2%.

6.4.1.2 Future Excavation Worker

Cumulative HIs for the future excavation worker were greater than 1 for SWMUs 4, 5, 6, 7, and 30 based on soil exposure. This exposure scenario was not evaluated for SWMU 2, SWMU 3, and SWMU 145. The following summarizes the cumulative HIs and major contributors to elevated hazards at these SWMUs.

- SWMU 4 cumulative HI 2.61; drivers are aluminum at 8%, chromium at 24%, iron at 24%, manganese 14%, and vanadium 20%.
- SWMU 5 cumulative HI 2.16; drivers are aluminum at 9%, arsenic at 7%, chromium at 18%, iron at 38%, and manganese at 22%.
- SWMU 6 cumulative HI 2.44; drivers are aluminum at 8%, chromium at 15%, iron at 32%, manganese at 15%, and vanadium at 26%.
- SWMU 7 cumulative HI 5.40; drivers are antimony at 11.3%, chromium at 17.6%, iron at 21.3%, manganese at 11%, uranium at 7.5%, and vanadium at 10.9%.
- SWMU 30 cumulative HI 4.50; drivers are antimony at 6.3%, chromium at 10.2%, copper at 7.6%, iron at 19.8%, manganese at 14.3%, uranium at 12.2%, and vanadium at 12.7%.

Cumulative ELCRs exceeded 1E-04 for all SWMUs (except SWMUs 2 and 3) and were greater than 1E-03 at SWMU 4, SWMU 7, and SWMU 30 for exposure to soil. The following summarizes the cumulative risk estimates and major contributors to the ELCR for these SWMUs.

- SWMU 4 cumulative ELCR 2.70E-03; drivers are beryllium at 7% and Total Uranium at 83%.
- SWMU 5 cumulative ELCR 2.90E-04; drivers are arsenic at 8%, beryllium at 62%, and Total PAH at 28%.
- SWMU 6 cumulative ELCR 2.30E-04; drivers are beryllium at 90% and Total PAH at 9%.
- SWMU 7 cumulative ELCR 1.60E-03; drivers are beryllium at 42.2%, uranium-235 at 9.1%, and uranium-238 at 41.3%.
- SWMU 30 cumulative ELCR 1.20E-03; driver is beryllium at 93.7%.

6.4.1.3 Future Recreational Users

Cumulative HIs for the child, teen, and adult recreational users were less than 1 for SWMUs 4, 5, 6, 7, and 30 based on soil exposure. Cumulative ELCRs exceeded 1E-06 for future adult recreational users only at SWMUs 5, 7, and 30 based on consumption of game. This exposure scenario was not evaluated at SWMUs 2, 3, and 145. The following summarizes the cumulative risk estimates and major contributors to the ELCR for these SWMUs.

- SWMU 5 cumulative ELCR 1.0E-05; driver is Total PAH at 96%.
- SWMU 7 cumulative ELCR 1.1E-05; drivers are Aroclor-1260 at 18.6%, benzo(a)pyrene at 9.5%, dibenzo(a,h)anthracene at 42.5%, and uranium-238 at 15.7%.
- SWMU 30 cumulative ELCR 1.5E-05; drivers are Aroclor-1260 at 48.2%, benzo(a)pyrene at 12.9% and dibenzo(a,h)anthracene at 20.8%.

6.4.1.4 Future On-Site Rural Residents

The following summarizes the cumulative HIs and ELCRs observed for each resident cohort.

Hazards—Future Child Residential Exposure to Soil. Cumulative HIs based on direct contact with soil for the child rural resident were greater than 1 for SWMUs 4, 5, 6, 7, and 30. This exposure scenario was not evaluated for SWMUs 2, 3, and 145. The major contributors to elevated hazards are as follows:

- SWMU 4 cumulative HI 98.2: chromium at 24%, iron at 60%, and vanadium at 9%.
- SWMU 5 cumulative HI 46.2: aluminum at 24%, arsenic at 53%, and chromium at 17%.
- SWMU 6 cumulative HI 9.38: beryllium at 8%, chromium at 72%, and nickel at 15%.
- SWMU 7 cumulative HI 370: arsenic at 6.2%, iron at 19.7%, and uranium at 58.4%.
- SWMU 30 cumulative HI 260: arsenic at 7.5%, iron at 22.6%, and uranium at 46.8%.

Hazard—Future Adult Resident Exposure to Soil. Cumulative HIs for the future on-site adult resident were greater than 1 for SWMUs 4, 5, 6, 7, and 30. This exposure scenario was not evaluated for SWMUs 2, 3, and 145. The major contributors to elevated hazards are as follows:

- SWMU 4 cumulative HI 28.4: chromium at 22%, iron at 63%, and vanadium at 8%.
- SWMU 5 cumulative HI 13.9: aluminum at 24%, arsenic at 55%, and chromium at 15%.
- SWMU 6 cumulative HI 2.57: beryllium at 7%, chromium at 70%, nickel at 17%, and zinc at 6%.
- SWMU 7 cumulative HI 110: arsenic at 6.5%, iron at 19.8%, and uranium at 59.5%.
- SWMU 30 cumulative HI 79: arsenic at 7.9%, iron at 22.8%, and uranium at 47.5%.

Risks—Future Adult Residential Exposure to Soil. Cumulative ELCRs exceeding 1E-03 from direct contact with soil was observed for SWMUs 4, 5, 6, 7, and 30. This exposure scenario was not evaluated for SWMUs 2, 3, and 145. Cumulative ELCRs greater than 1E-02 were identified for SWMUs 5, 7, and 30. The major contributors to elevated risks are as follows:

- SWMU 4 cumulative ELCR 4.3E-03: beryllium at 72%, total PCBs at 5%, uranium-234 at 6%, and uranium-238 at 17%.
- SWMU 5 cumulative ELCR 1.0E-02: arsenic at 21%, beryllium at 9%, and Total PAH at 68%.
- SWMU 6 cumulative ELCR 2.4E-03: beryllium at 54% and Total PAH at 46%.
- SWMU 7 cumulative ELCR 3.4E-02: arsenic at 7.3%, beryllium at 65.4%, and uranium-238 at 17.6%.
- SWMU 30 cumulative ELCR 3.2E-02: arsenic at 6.8%, beryllium at 66.7%, and uranium-238 at 11.5%.

6.4.2 Risk Characterization of Vapor Intrusion into Basements from Soil

Characterization of risks from vapor intrusion into basements from soil was conducted as part of the current risk assessment.² To examine potential risks and hazards, vapor intrusion modeling was completed and examined for three POEs: the property boundary, the plant boundary, and at the SWMU. The HQs and ELCRs for the modeled vapor concentrations are presented in Table E.3.34 of Appendix E. Modeled concentrations for the on-site POE showed an HQ greater than 0.1 for vapor intrusion from TCE, *cis*-1,2-DCE, 1,1-DCE, vinyl chloride, or mercury intrusion for the following:

- SWMU 2: TCE, and *cis*-1,2-DCE.
- SWMU 3: mercury.
- SWMU 4: TCE, *cis*-1,2-DCE; and vinyl chloride.
- SWMU 7: 1,1-DCE, mercury; and vinyl chloride.
- SWMU 30: mercury, 1,1-DCE; and TCE.
- SWMU 145: mercury.

ELCRs for the on-site POE were greater than 1E-06 for several SWMUs based on modeled contaminant concentrations. The following summarizes those SWMUs exhibiting elevated risks based on modeled soil concentrations:

- SWMUs 2: TCE.
- SWMU 3: TCE.
- SWMU 4: TCE and vinyl chloride.
- SWMU 7: TCE; vinyl chloride; and 1,1-DCE.
- SWMU 30: TCE and 1,1-DCE.

Vapor intrusion into basements also was modeled at the plant boundary and property boundary. At the plant boundary all HIs were below 0.1. ELCRs were below 1E-06 for all SWMUs except for SWMUs 2, 4, 7, and 30. The following lists the risk driver for each SWMU:

- SWMU 2: TCE.
- SWMU 4: TCE and vinyl chloride.
- SWMU 7: 1,1-DCE.
- SWMU 30: TCE and 1,1-DCE.

At the property boundary all HIs were below 0.1. The ELCR for TCE exceeded 1E-06 risk at the property boundary for TCE at SWMUs 2, 4, and 30. All other risks/hazards were below 1E-06 or an HI of 1 at the property boundary. The quantitative assessment of potential risks and hazards due to exposure to vapor intrusion is summarized in Table 5.14. Table F.73 in Appendix F includes which SWMUs exceeded *de minimis* risk and hazard thresholds for vapor intrusion modeling for the on-site receptor and the receptor at the property boundary.

Characterization of risks and systemic toxicity from vapor intrusion into industrial building basements was conducted as part of the current risk assessment. The vapor intrusion potentially exposes a future industrial worker to vapors from soil. To examine potential risks and hazards, vapor intrusion modeling

² The vapor transport modeling for mercury was conservatively based on the metallic form, which has a Henry's Law Constant of 1.07E-02 atm-m³/mol. The rate of vaporization of mercury and certain of its inorganic compounds decrease in the sequence Hg > Hg₂Cl₂ > HgCl₂ > HgS > HgO. The Henry's Law Constant decreases dramatically down the sequence (for example, HgCl₂ has a value of 7.09E-10 atm-m³/mol).

was completed and examined for one POE. The POE is at the location of the SWMU. Table 6.5 shows the results of the vapor modeling for the industrial worker.

Table 6.5. Future Industrial Worker Exposure to Soil Vapor at SWMU Locations

	HQ	% Contribution to HI	Risk	% Contribution to Risk
SWMU 2				
Trichloroethene	4.82E-01	11.2%	6.33E-04	100.0%
<i>cis</i> -1,2 Dichloroethene	3.83E+00	89.1%	NA	
Naphthalene	6.17E-05	0.0%	NA	
Total	4.3		6.33E-04	
SWMU 3				
Trichloroethene	2.79E-04	100.0%	3.66E-07	100.0%
Total	2.79E-04		3.66E-07	
SWMU 5				
Trichloroethene	9.29E-05	9.7%	3.65E-07	100.0%
Acenaphthene	6.69E-07	0.1%	NA	
Fluorene	2.52E-07	0.0%	NA	
Naphthalene	8.68E-04	90.2%	NA	
Pyrene	1.49E-08	0.0%	NA	
Total	9.62E-04		3.65E-07	
SWMU 6				
Trichloroethene	1.60E-04	100.0%	2.10E-07	100.0%
Total	1.60E-04		2.10E-07	
SWMU 7				
Trichloroethene	1.48E-03	1.7%	5.83E-06	1.3%
<i>cis</i> -1,2 Dichloroethene	4.18E-03	4.7%	NA	
Vinyl chloride	8.39E-02	93.7%	7.92E-05	17.1%
1,1-DCE	3.52E-02	39.3%	3.77E-04	81.6%
Mercury	2.30E-02	25.7%	NA	
Pyrene	5.01E-08	0.0%	NA	
Tetrachloroethene	2.29E-05	0.0%	8.38E-09	0.0%
Total	8.95E-02		4.62E-04	
SWMU 30				
Trichloroethene	1.21E-03	0.1%	1.52E-06	78.7%
1,1-DCE	1.20E-04	0.0%	4.11E-07	21.3%
Acenaphthene	9.45E-08	0.0%	NA	
Fluorene	2.00E-08	0.0%	NA	
Mercury	3.94E-02	2.8%	NA	

Table 6.5. Future Industrial Worker Exposure to Soil Vapor at SWMU Locations (Continued)

	HQ	% Contribution to HI	Risk	% Contribution to Risk
SWMU 30				
Naphthalene	7.37E-05	0.0%	NA	
Pyrene	4.46E-09	0.0%	NA	
	1.4		1.93E-06	
SWMU 145				
Mercury	3.25E-02	100.0%	NA	
Total	3.25E-02			

The vapor intrusion model was used, and the risk and hazard calculations were performed similarly to the residential receptor using the equation shown below except for these changes.

$$Chemical\ Intake\ [mg/(kg\ x\ day)] = \frac{C_{air} \times IR_{air} \times EF \times ED \times ET}{BW \times AT}$$

Parameter	Units	Value
Indoor inhalation rate = IR_{air}	m3/hr	2.5
Exposure frequency = EF	day/year	250
Exposure duration = ED	years	25
Exposure time = ET	hours/day	8

The HI exceed 0.1 at SWMU 2 is due primarily to *cis*-1,2 DCE at a HQ of 4.3. Also, the HI was greater than 0.1 at SWMU 30 due primarily to TCE. Risks exceeded 1E-06 at SWMUs 2 and 30 due to TCE, and risks exceeded 1E-06 at SWMU 7 due to DCE. The vapor intrusion scenario for the future industrial worker is highly uncertain. The industrial worker exposure to contaminated groundwater usually is deemed not to be a plausible scenario because of institutional controls and risk mitigation measures such as supplied drinking water from uncontaminated sources. Similarly, institutional controls and risk mitigation methods for the vapor intrusion scenario to the future industrial worker are likely to be put in place. Buildings likely would not be sited on or near former disposal areas.

6.4.3 Risk Characterization for Residential Use of Groundwater Drawn from the RGA

Risks for residential groundwater use were calculated based on modeled concentrations in the RGA groundwater. The HIs and ELCRs also are presented for each SWMU in Tables F. 41 through F. 64 in Appendix F. Tables F.65 through F.67 show the HIs and ELCRs for the predicted maximum (peak) concentrations for individual contaminants over the 1,000 year time frame of the model. Different contaminants migrate at different rates; therefore, the total HI or ELCR in groundwater at a given time may be less than the sum of the risks of individual peaks. Figures F.2 to F.15 in Appendix F show the total hazard and total risks from the predicted concentrations of all COCs at each time step in the model

for each SWMU except SWMU 6 (which had no modeled COCs). These hazards and risks are calculated using the residential NALs (DOE 2001) for residential use of groundwater.

6.4.3.1 Hazards—Future Resident Exposure to Groundwater at the SWMU Boundary

Cumulative HIs based on exposure to groundwater for the future on-site rural resident were greater than 1 for all of the SWMUs evaluated (SWMU 6 was evaluated as a source of groundwater contamination and it was determined that SWMU was not a source of any contaminant. Specifically, manganese was evaluated, but the manganese concentration in groundwater did not have an HQ greater than 1). The following lists those constituents that contributed to elevated HIs. The major contaminants driving the hazard were ingestion of arsenic and uranium metal, and ingestion and inhalation of TCE and *cis*-1,2 DCE.

The following lists those constituents that contributed to elevated HIs by SWMU for the Child Resident:

- SWMU 2: TCE at 52% and *cis*-1,2-DCE at 47%.
- SWMU 3: arsenic at 52%, uranium at 39%, and manganese at 10%.
- SWMU 4: TCE at 93% and *cis*-1,2-DCE at 6%.
- SWMU 5: uranium at 90%, arsenic at 4%, naphthalene at 3%, and manganese at 3%.
- SWMU 7: arsenic at 30%, TCE at 26%, Aroclor-1254 at 22%, and *cis*-1,2-DCE at 7%.
- SWMU 30: arsenic at 64%, manganese at 9%, uranium at 15%, 1,1-DCE at 5%, and TCE at 5%.
- SWMU 145: antimony at 48% and arsenic at 48%.

6.4.3.2 Risks—Future Residential Exposure to Groundwater at the SWMU Boundary

Cumulative ELCRs exceeding both 1E-06 and 1E-04 from direct exposure to groundwater was observed for all of the SWMUs evaluated (SWMU 6 was not evaluated for carcinogenic risk). The major contaminants (> 5%) driving risk were ingestion of arsenic and TCE.

- SWMU 2: TCE at 98%.
- SWMU 3: arsenic at 72% and technetium-99 at 25%.
- SWMU 4: TCE at 68% and vinyl chloride at 31%.
- SWMU 5: arsenic at 97%.
- SWMU 7: arsenic at 15%, 1,1-DCE at 66%, TCE at 4.1%, and vinyl chloride at 12%.
- SWMU 30: arsenic at 89% and TCE at 5.2%.
- SWMU 145: arsenic at 5% and Aroclor-1260 at 93%.

6.4.4 Risk Characterization for Residential Use of Groundwater at Future Modeled Concentrations on Boundary and River POEs

Risk and hazard estimates for future off-site residential use based on modeled groundwater concentrations were calculated for this BHHRA. The following summarizes the results of the quantitative assessment at the plant boundary, property boundary, and outside the property boundary at either the Little Bayou seeps (applicable to SWMUs 3, 7, and 30) or Ohio River (applicable to SWMUs 2, 4, 5, and 145).

6.4.4.1 Future Residential Exposure to Groundwater—Plant Boundary

SWMU 145 lies outside the plant boundary. Cumulative HIs based on exposure to groundwater at the DOE plant boundary were greater than one for SWMU 2, SWMU 4, SWMU 5, SWMU 7, and SWMU

30. The major contaminants contributing to hazard were TCE, *cis*-1,2-DCE, arsenic, manganese, naphthalene, and PCBs.

The cumulative ELCR was greater than 1E-06 for SWMU 2, SWMU 3, SWMU 4, SWMU 5, SWMU 7, and SWMU 30. The cumulative ELCR was greater than 1E-04 for SWMU 2, SWMU 3, SWMU 4, SWMU 7, and SWMU 30. The major contaminants contributing to risk were TCE, 1,1-DCE, vinyl chloride, technetium-99, and arsenic.

6.4.4.2 Future Residential Exposure to Groundwater – Property Boundary

Cumulative HIs based on exposure to groundwater at the DOE property boundary were greater than 1 for SWMU 2, SWMU 4, and SWMU 7. The major contaminants driving hazard were ingestion of arsenic, TCE, *cis*-1,2-DCE, and PCBs.

Cumulative ELCR exceeded 1E-06 for groundwater exposure for all of the SWMUs, except SWMU 6. Cumulative ELCRs greater than 1E-04 from groundwater use were identified for SWMUs 2, 4, 7, 30, and 145. The major contaminants driving risk were ingestion of arsenic, TCE, 1,1-DCE, vinyl chloride, and technetium-99.

6.4.4.3 Future Residential Exposure to Groundwater–Outside the Property Boundary (Little Bayou seeps or the Ohio River)

Cumulative HIs based on exposure to groundwater for the future off-site rural resident were greater than 1 at the Ohio River for SWMUs 2 and 4. The major contaminants driving hazard were ingestion of TCE and *cis*-1,2-DCE.

Cumulative ELCRs of 1E-06 from groundwater exposure were observed at the Little Bayou seeps for SWMUs 3, 7, and 30, and at the Ohio River for SWMUs 2, 4, and 145. Cumulative ELCRs greater than 1E-04 from groundwater use were identified for SWMU 2, 4, and 7. The contaminants driving risk were ingestion of TCE, 1,1-DCE, vinyl chloride, and technetium-99.

6.4.5 Identification of Land Use Scenarios, Pathways, Media and COCs

This subsection outlines land use scenarios, exposure pathways, media, and COCs for each source area. The results of the previous risk assessments for SWMUs 2, 3, 4, 5, 6, 7, and 30 are used for the risk characterization for soil and are presented in Attachment F2 of this document. Section 6.6 presents the RGOs for each location and land use scenario.

6.4.5.1 Land Use Scenarios of Concern

To determine whether a land use scenario is of concern, quantitative risk and hazard results were compared to risk and hazard benchmarks for each land use scenario. The benchmarks used for this comparison were a) 1 for HI and b) 1×10^{-6} for ELCR. Land use scenarios with total HIs exceeding the benchmark of 1 are deemed land use scenarios of concern for non-cancer hazard. Land use scenarios with a total ELCR exceeding the benchmark of 1×10^{-6} are deemed land use scenarios of concern for cancer risk. These criteria were used in the previous risk assessments for SWMUs 4, 5, 6, 7, and 30 as well (DOE 1998c; DOE 2000). For the risk characterization of soil for SWMUs 2 and 3, land use scenarios of concern were determined by using EPA guidance and policy in effect at the time of the risk assessment (DOE 1994). The following are land uses of concern for BGOU at the SWMUs indicated:

- Industrial: SWMUs 2, 3, 4, 5, 6, 7, and 30.

- Excavation: SWMUs 4, 5, 6, 7, and 30.
- Recreational: SWMUs 5, 7, and 30.
- On-Site Residential: SWMUs 2, 3, 4, 5, 6, 7, 30, and 145.
- Off-Site Residential: SWMUs 2, 3, 4, 5, 7, 30, and 145.

Table F.70 in Appendix F outlines all land use scenarios for all SWMUs that exceed *de minimis* risk or hazard levels.

6.4.5.2 Contaminants of Concern (Soil)

To make a determination about whether contaminants are of concern in soil, quantitative risk and hazard results over all pathways from the previous risk assessments for SWMUs 2, 3, 4, 5, 6, 7, and 30 (DOE 1994; DOE 1998c; DOE 2000a) were compared to risk and hazard benchmarks for land use scenarios of concern. The benchmarks used for this comparison were a) 0.1 for HI and b) 1×10^{-6} for ELCR.

Contaminants with chemical-specific HIs or ELCRs exceeding these benchmarks are deemed COCs. A priority COC is a contaminant whose chemical-specific HI is greater than 1 or whose ELCR is greater than 1×10^{-4} for one or more scenarios (DOE 2008b). The following are priority COCs found in soil at individual SWMUs:

- SWMU 2—none.
- SWMU 3—none.
- SWMU 4—barium, beryllium, cadmium, chromium, iron, nickel, uranium, vanadium, Total dioxins/furans, Total PCBs, uranium-234, and uranium-238.
- SWMU 5—aluminum, arsenic, beryllium, chromium, nickel, Total PAHs, and Total PCBs.
- SWMU 6—beryllium, chromium, nickel, and Total PAHs.
- SWMU 7—aluminum, antimony, arsenic, barium, beryllium, cadmium, chromium, copper, iron, manganese, nickel, uranium, vanadium, benzo(a)pyrene, dibenzo(a,h)anthracene, Aroclor-1254, Aroclor-1260, plutonium-239, uranium-234, uranium-235, uranium-235/236, and uranium-238.
- SWMU 30—aluminum, antimony, arsenic, barium, beryllium, cadmium, chromium, copper, iron, manganese, mercury, nickel, uranium, vanadium, benzo(a)anthracene, benzo(a)pyrene, benzo(b)fluoranthene, dibenzo(a,h)anthracene, indeno(1,2,3-cd)pyrene, Aroclor-1254, Aroclor-1260, uranium-234, uranium-235, uranium-235/236, and uranium-238.

6.4.5.3 Contaminants of Concern (Groundwater—Modeled from Soil)

Similarly for groundwater, to determine whether contaminants are of concern, quantitative risk and hazard results over all pathways were compared to risk and hazard benchmarks for land use scenarios of concern. The benchmarks used for this comparison were a) 0.1 for HI and b) 1×10^{-6} for ELCR.

Contaminants with chemical-specific HIs or ELCRs exceeding these benchmarks were deemed COCs. Priority COCs are contaminants whose chemical-specific HI is greater than 1 or whose ELCR is greater than 1×10^{-4} for one or more scenarios (DOE 2008b). The following presents priority COCs found in groundwater at individual SWMUs:

- SWMU 2—arsenic; Aroclor-1248; manganese; uranium; *cis*-1,2-DCE; and TCE.
- SWMU 3—arsenic; manganese; uranium; and technetium-99.
- SWMU 4—arsenic; manganese; *cis*-1,2-DCE; TCE; vinyl chloride; and technetium-99.
- SWMU 5—arsenic; uranium; manganese; and naphthalene.
- SWMU 6—none.
- SWMU 7—arsenic; 1,1-DCE; *cis*-1,2-DCE; Aroclor-1254; TCE; vinyl chloride.
- SWMU 30—arsenic.
- SWMU 145—antimony; arsenic; manganese; Aroclor-1260; and technetium-99.

“Priority COCs” are identified in this section as an aid to risk managers during decision making. Table F.71 in Appendix F summarizes the COCs for both soil and groundwater.

6.4.6 Pathways of Concern

To determine whether pathways are of concern, the quantitative risks and hazards for each exposure route are summed over all contaminants and compared to benchmarks for land use scenarios of concern. The benchmarks used for this comparison were a) 0.1 for HI and b) 1×10^{-6} for ELCR. For soil, the quantitative risk and hazard results from the previous risk assessments for SWMUs 2, 3, 4, 5, 6, 7, and 30 (DOE 1994; DOE 1998c; DOE 2000a) were used in the comparison. Exposure routes with HIs and ELCRs exceeding these benchmarks are considered pathways of concern (POCs). These POCs are shown by SWMU in Table F.80 of Appendix F. Each of the pathways included in the BHHRA is a POC for at least one SWMU.

6.4.7 Media of Concern

Media of concern are those media that appear in at least one POC. Because they contribute to at least one POC, soil and RGA groundwater are media of concern for all eight SWMUs. Table F.81 in Appendix F provides specific information concerning how each media contributes to risks and hazards for BGOU.

6.4.8 Summary of Risk Characterization

Tables 6.6 through 6.13 present summaries of the risk characterization by location considered in the BHHRA. They present land use scenarios of concern, COCs, and POCs. In addition, each table lists the following:

- Receptor risks for each land use scenario of concern.
- Percent contribution by pathway to the total risk.
- Percent contribution each COC contributes to the total risk.

Table 6.6. Summary of Risk Characterization for SWMU 2

Receptor	Total ELCR ^c	COCs	% Total ELCR	POCs	% Total ELCR	Total HI ^d	COCs	% Total HI	POCs	% Total HI
Current industrial worker/intruder at current concentrations (soil) (from WAG 22 RI Addendum ^b)	1.2E-05	²³⁸ U + daughters ²³⁸ U + daughters	83.8 10.7	External exposure	94.7	6.8E-03	*No COCs	*No COCs	*No COCs	
Future industrial worker at current concentrations (soil) (from WAG 22 RI Addendum ^b)	1.2E-04	Arsenic ²³⁸ U + daughters ²³⁸ U + daughters	2.8 83.9 10.7	Ingestion External exposure	4.7 94.7	7.0E-02	*No COCs		*No COCs	
Future child rural resident at current concentrations (RGA groundwater only)	NA	NA	NA	NA	NA	1.30E+03	Arsenic Manganese Uranium <i>cis</i> -1,2-DCE Naphthalene TCE	0.9 0.1 0.1 46.8 0.0 52.1	Ingestion Dermal Inhalation while showering Household inhalation	46.0 11.7 4.8 37.5

Table 6.6. Summary of Risk Characterization for SWMU 2 (Continued)

Receptor	Total ELCR ^a	COCs	% Total ELCR	POCs	% Total ELCR	Total HI ^a	COCs	% Total HI	POCs	% Total HI
Future adult rural resident at current concentrations (RGA groundwater only)	4.72E-02	Arsenic Aroclor-1248 Aroclor-1268 TCE ⁹⁹ Tc ²³⁴ U ²³⁸ U	2.0 0.4 0.1 97.5 0.0 0.0 0.0	Ingestion Dermal Inhalation while showering Household inhalation	19.8 11.3 7.8 61.0	3.79E+02	Arsenic Manganese Uranium <i>cis</i> -1,2-DCE Naphthalene TCE	0.9 0.1 0.1 36.8 0.0 62.1	Ingestion Dermal Inhalation while showering Household inhalation	45.0 23.9 3.5 27.5
Future child rural resident at modeled concentrations (RGA groundwater drawn at plant boundary)	NA	NA	NA	NA	NA	1.92E+02	Arsenic <i>cis</i> -1,2-DCE Naphthalene TCE	0.5 48 0.1 52	Ingestion Dermal Inhalation while showering Household inhalation	45 12.4 5.4 38
Future adult rural resident at modeled concentrations (RGA groundwater drawn at plant boundary)	6.82E-03	Arsenic TCE	1.1 98.9	Ingestion Dermal Inhalation while showering Household inhalation	19.2 11.1 7.9 61.8	5.08E+01	Arsenic <i>cis</i> -1,2-DCE Naphthalene TCE	0.5 16.2 0.1 83.1	Ingestion Dermal Inhalation while showering Household inhalation	60 32 1 7.2
Future child rural resident at modeled concentrations (RGA groundwater drawn at property boundary)	NA	NA	NA	NA	NA	9.56E+01	<i>cis</i> -1,2-DCE TCE	47.4 52.6	Ingestion Dermal Inhalation while showering Household inhalation	45.4 11.8 4.9 38.0
Future adult rural resident at modeled concentrations (RGA groundwater drawn at property boundary)	3.42E-03	TCE	100	Ingestion Dermal Inhalation while showering Household inhalation	18.3 11.2 8.0 62.5	2.79E+01	<i>cis</i> -1,2-DCE TCE	37.3 62.7	Ingestion Dermal Inhalation while showering Household inhalation	44.4 24.1 3.6 27.9
Future child rural resident at modeled concentrations (RGA groundwater drawn at Ohio River)	NA	NA	NA	NA	NA	2.25E+01	<i>cis</i> -1,2-DCE TCE	79.4 20.5	Ingestion Dermal Inhalation while showering Household inhalation	16.2 18.8 7.4 57.7
Future adult rural resident at modeled concentrations (RGA groundwater drawn at Ohio River)	1.28E-03	TCE	100	Ingestion Dermal Inhalation while showering Household inhalation	18.3 11.2 8.0 62.5	6.7E+00	<i>cis</i> -1,2-DCE TCE	61.2 38.7	Ingestion Dermal Inhalation while showering Household inhalation	15.5 37.7 5.3 41.5

ELCR = excess lifetime cancer risk; HI = hazard index; POC = point of contact; COC = contaminant of concern

Note: NA = ELCR not applicable to child and teen cohorts. ELCR for adult is for lifetime exposure and takes into account exposure as child and teen.

^aNo COCs = there are no COCs or POCs at this SWMU for this endpoint (may apply to ELCR or HI)

^bTotal ELCR and total HI represent total risk or hazard summed across all POCs for all COCs.

^cRemedial Investigation Addendum for Waste Area Grouping 22, Burial Grounds, at the Paducah Gaseous Diffusion Plant, Paducah, Kentucky, DOE/OR/07-1141&D2, September 1994 (DOE 1994), Attachment 2-1 through 2-6. This risk assessment combined SWMU 2 and 3.

Table 6.7. Summary of Risk Characterization for SWMU 3

Receptor	Total ELCR ^a	COCs	% Total ELCR	POCs	% Total ELCR	Total HI ^b	COCs	% Total HI	POCs	% Total HI
Current industrial worker/intruder at current concentrations (soil) (from WAG 22 RI Addendum ^b)	1.2E-05	²³⁵ U + daughters ²³⁸ U + daughters	83.8 10.7	External exposure	94.7	6.8E-03	*No COCs		*No COCs	NE
Future industrial worker at current concentrations (soil) (from WAG 22 RI Addendum ^b)	1.2E-04	Arsenic ²³⁵ U + daughters ²³⁸ U + daughters	2.8 83.9 10.7	Ingestion External exposure	4.7 94.7	7.0E-02	*No COCs		*No COCs	NE
Future child rural resident at current concentrations (RGA groundwater only)	NA	NA	NA	NA	NA	2.03E+01	Arsenic Manganese Uranium	51.9 9.6 38.6	Ingestion Dermal	99.5 0.5
Future adult rural resident at current concentrations (RGA groundwater only)	1.20E-03	Arsenic ⁹⁹ Tc ²³⁸ U	72.4 25.3 2.3	Ingestion Dermal	99.8 0.2	5.83E+00	Arsenic Manganese Uranium	51.7 9.9 38.3	Ingestion Dermal	98.9 1.1
Future child rural resident at modeled concentrations (RGA groundwater drawn at plant boundary)	NA	NA	NA	NA	NA	3.98E-01	Arsenic	100	Ingestion	97.9
Future adult rural resident at modeled concentrations (RGA groundwater drawn at plant boundary)	1.32E-04	Arsenic ⁹⁹ Tc	24.6 75.4	Ingestion	99.9	1.12E-01	Arsenic	100	Ingestion	99.6
Future child rural resident at modeled concentrations (RGA groundwater drawn at property boundary)	NA	NA	NA	NA	NA		*No COCs		*No COCs	
Future adult rural resident at modeled concentrations (RGA groundwater drawn at property boundary)	7.46E-05	⁹⁹ Tc	100	Ingestion	100		*No COCs		*No COCs	

Table 6.7. Summary of Risk Characterization for SWMU 3 (Continued)

Receptor	Total ELCR ^a	COCs	% Total ELCR	POCs	% Total ELCR	Total HI ^b	COCs	% Total HI	POCs	% Total HI
Future child rural resident at modeled concentrations (RGA groundwater drawn at Little Bayou seeps)	NA	NA	NA	NA	NA	*No COCs	*No COCs		*No COCs	
Future adult rural resident at modeled concentrations (RGA groundwater drawn at Little Bayou seeps)	4.41E-05 ^b	⁹⁹ Tc	100.0	Ingestion	100	*No COCs	*No COCs		*No COCs	
Future child rural resident at modeled concentrations (RGA groundwater drawn at Ohio River)		Not a POE for groundwater from this SWMU.								
Future adult rural resident at modeled concentrations (RGA groundwater drawn at Ohio River)		Not a POE for groundwater from this SWMU.								

ELCR = excess lifetime cancer risk; HI = hazard index; POC = point of contact; COC = contaminant of concern
 Note: NA = ELCR not applicable to child and teen cohorts. ELCR for adult is for lifetime exposure and takes into account exposure as child and teen.
 *No COCs = there are no COCs or POCs.

^a Total ELCR and total HI represent total risk or hazard summed across all POCs for all COCs.

^b Remedial Investigation Addendum for Waste Area Grouping 22: Burial Grounds, at the Paducah Gaseous Diffusion Plant, Paducah, Kentucky, DOE/OR-07-1141&D2, September 1994 (DOE 1994), Attachment 2-1 through 2-6. This risk assessment combined SWMU 2 and 3.

Table 6.8. Summary of Risk Characterization for SWMU 4

Receptor	Total ELCR ^a	COCs	% Total ELCR	POCs	% Total ELCR	Total HI ^r	COCs	% Total HI	POCs	% Total HI
Current industrial worker at current concentrations (soil) (WAG 3 RI ^b)	5.4E-04	Beryllium ²³⁸ U	97 2	Dermal External exposure	97 2	3.62E+00	Beryllium Chromium Iron Vanadium Barium	5 45 24 24 2	Dermal	99
Future industrial worker at current concentrations (soil) (WAG 3 RI ^b)	5.4E-04	Beryllium ²³⁸ U	97 2	Dermal External exposure	97 2	3.62E+00	Beryllium Chromium Iron Vanadium Barium	5 45 24 24 2	Dermal	99
Future child rural resident at current concentrations (soil) (WAG 3 RI ^b)	NA	NA	NA	NA	NA	9.82E+01	Barium Beryllium Cadmium Chromium Iron Nickel Vanadium	2 2 2 24 60 2 2 9	Ingestion Dermal Ingestion of vegetables	1 21 78
Future adult rural resident at current concentrations (soil) (WAG 3 RI ^b)	4.3E-03	Beryllium Total PCB ²³⁴ U ²³⁸ U	72 5 6 17	Dermal External exposure Ingestion of vegetables	36 2 61	2.84E+01	Barium Beryllium Cadmium Chromium Iron Nickel Vanadium	2 2 2 22 63 2 2 8	Dermal Ingestion of vegetables	14 85
Future child rural resident at current concentrations (RGA groundwater only)	NA	NA	NA	NA	NA	5.82E+02	Arsenic Manganese <i>cis</i> -1,2-DCE TCE Vinyl Chloride	1.0 0.2 6.1 92.5 0.2	Ingestion Dermal Inhalation while showering Household inhalation	67.2 20.2 1.4 11.2
Future adult rural residents at current concentrations (RGA groundwater only)	5.41E-02	Arsenic TCE Vinyl chloride ⁹⁹ Tc	0.9 67.7 30.5 0.9	Ingestion Dermal Inhalation while showering Household inhalation	15.4 36.7 5.4 42.4	1.98E+02	Arsenic Manganese <i>cis</i> -1,2-DCE TCE Vinyl chloride	0.8 0.2 4.1 94.7 0.2	Ingestion Dermal Inhalation while showering Household inhalation	56.5 35.6 0.9 7.0
Future child rural resident at modeled concentrations (RGA groundwater drawn at plant boundary)	NA	NA	NA	NA	NA	2.04E+02	Arsenic <i>cis</i> -1,2-DCE TCE Vinyl chloride	0.4 4.6 94.4 0.1	Ingestion Dermal Inhalation while showering Household inhalation	67.5 20.6 1.4 10.6

Table 6.8. Summary of Risk Characterization for SWMU 4 (Continued)

Receptor	Total ELCR ^a	COCs	% Total ELCR	POCs	% Total ELCR	Total HI ^d	COCs	% Total HI	POCs	% Total HI
Future adult rural resident at modeled concentrations (RGA groundwater drawn at plant boundary)	2.03E-02	Arsenic TCE Vinyl chloride ⁹⁹ Tc	0.4 98.0 0.9 0.7	Ingestion Dermal Inhalation while showering Household inhalation	13.6 7.2 5.2 74.0	6.97E+01	Arsenic <i>cis</i> -1,2-DCE TCE	0.4 3.0 96.6	Ingestion Dermal Inhalation while showering Household inhalation	56.5 36.1 0.8
Future child rural resident at modeled concentrations (RGA groundwater drawn at property boundary)	NA	NA	NA	NA	NA	1.03E+02	<i>cis</i> -1,2-DCE TCE Vinyl chloride	4.6 95.3 0.1	Ingestion Dermal Inhalation while showering Household inhalation	67.6 20.8 1.3 10.3
Future adult rural resident at modeled concentrations (RGA groundwater drawn at property boundary)	6.79E-03	TCE Vinyl chloride ⁹⁹ Tc	97.9 1.1 1.0	Ingestion Dermal Inhalation while showering Household inhalation	19.8 11.0 7.8 61.3	3.51E+01	<i>cis</i> -1,2-DCE TCE	3.1 96.8	Ingestion Dermal Inhalation while showering Household inhalation	56.4 36.3 0.8 6.4
Future child rural resident at modeled concentrations (RGA groundwater drawn at Ohio River)	NA	NA	NA	NA	NA	3.33E+01	<i>cis</i> -1,2-DCE TCE	1.7 98.2	Ingestion Dermal Inhalation while showering Household inhalation	74.6 22.9 1.4 1.0
Future adult rural resident at modeled concentrations (RGA groundwater drawn at Ohio River)	2.43E-03	TCE Vinyl chloride ⁹⁹ Tc	98.2 0.9 0.9	Ingestion Dermal Inhalation while showering Household inhalation	19.6 11.0 7.9 61.5	1.26E+01	<i>cis</i> -1,2-DCE TCE	3.0 96.9	Ingestion Dermal Inhalation while showering Household inhalation	56.4 36.3 0.8 6.4
Future child recreational user at current concentrations (soil) (WAG 3 RI ^b)	NA	NA	NA	NA	NA	<1	*No COCs		*No COCs	
Future teen recreational user at current concentrations (soil) (WAG 3 RI ^b)	NA	NA	NA	NA	NA	<1	*No COCs		*No COCs	
Future adult recreational user at current concentrations (soil) (WAG 3 RI ^b)	<1.0E-06	*No COCs		*No COCs		<1	*No COCs		*No COCs	
Future excavation worker at current concentrations (soil and waste) (WAG 3 RI ^b)	2.7E-03	Arsenic Beryllium Total dioxins/furans Total PCB ²²⁶ Ra Total uranium ^c ²³⁸ U	1 7 4 2 2 83 1	Ingestion Dermal External exposure	37 10 54	2.61E+00	Aluminum Arsenic Barium Beryllium Cadmium Chromium Iron Manganese Vanadium	8 4 2 2 1 24 24 14 20	Ingestion Dermal	13 87

ELCR = excess lifetime cancer risk; HI = hazard index; POC = point of contact; COC = contaminant of concern; Note: NA = ELCR not applicable to child and teen cohorts. ELCR for adult is for lifetime exposure and takes into account exposure as child and teen. *No COCs = There are no COCs or POCs. ^a Total ELCR and total HI represent total risk or hazard summed across all POCs for all COCs. ^b Remedial Investigation Report for Waste Area Grouping 3 at the Paducah Gaseous Diffusion Plant, Paducah, Kentucky, DOE/OR/07-1895-6DDJ, September 2000 (DOE 2000a), Table 1.55. In this table, lead has been excluded as a COC. ^c Risk associated with total uranium at SWMU 4 was calculated using a total uranium analytical result in pCi/g units and toxicity information for ²³⁸U. Individual isotopes also were included in the risk calculation, resulting in a double-counting of risk due to uranium isotopes. This approach likely accounts for the discrepancy between risk related to total uranium and ²³⁸U.

Table 6.9. Summary of Risk Characterization for SWMU 5

Receptor	Total ELCR ^a	COCs	% Total ELCR	POCs	% Total ELCR	Total HI ^c	COCs	% Total HI	POCs	% Total HI
Current industrial worker at current concentrations (soil) (WAG 3 RI ^b)	4.1E-04	Arsenic Beryllium Total PAH	6 49 45	Ingestion Dermal	2 98	< 1	*No COCs		*No COCs	
Future industrial worker at current concentrations (soil) (WAG 3 RI ^b)	4.1E-04	Arsenic Beryllium Total PAH	6 49 45	Ingestion Dermal	2 98	< 1	*No COCs		*No COCs	
Future child rural resident at current concentrations (soil) (WAG 3 RI ^b)	NA	NA	NA	NA	NA	4.62E+01	Aluminum Arsenic Beryllium Chromium Nickel Zinc	24 53 1 17 3 1	Ingestion Dermal Ingestion of vegetables	1 12 87
Future adult rural resident at current concentrations (soil) (WAG 3 RI ^b)	>1.0E-02*	Arsenic Beryllium Total PAH Total PCB	21 9 68 2	Dermal Ingestion of vegetables	9 90	1.39E+01	Aluminum Arsenic Beryllium Chromium Nickel Zinc	24 55 1 15 3 1	Dermal Ingestion of vegetables	8 92
Future child rural resident at current concentrations (RGA groundwater only)	NA	NA	NA	NA	NA	8.15E+01	Uranium Arsenic Manganese Naphthalene	90.3 3.6 2.6 3.4	Ingestion Dermal Inhalation while showering Household inhalation	96.4 0.2 0.4 3.0
Future adult rural resident at current concentrations (RGA groundwater only)	2.52E-04	Arsenic ⁹⁹ Tc	97.2 2.8	Ingestion Dermal	99.7 0.3	2.31E+01	Uranium Arsenic Manganese Naphthalene	91.0 3.7 2.7 2.6	Ingestion Dermal Inhalation while showering Household inhalation	97.1 0.3 0.3 2.3
Future child rural resident at modeled concentrations (RGA groundwater drawn at plant boundary)	NA	NA	NA	NA	NA	6.56E+00	Uranium Arsenic Manganese Naphthalene	81 9 27.5	Ingestion Household inhalation	92.4 6.6
Future adult rural resident at modeled concentrations (RGA groundwater drawn at plant boundary)	4.99E-05	Arsenic ⁹⁹ Tc	94.5 5.5	Ingestion	99.7	1.84E+00	Uranium Arsenic Naphthalene manganese	82.4 8.9 5.8 2.1	Ingestion Household inhalation	93.9 5.1

Table 6.9. Summary of Risk Characterization for SWMU 5 (Continued)

Receptor	Total ELCR ^a	COCs	% Total ELCR	POCs	% Total ELCR	Total HI ^c	COCs	% Total HI	POCs	% Total HI
Future child rural resident at modeled concentrations (RGA groundwater drawn at property boundary)	NA	NA	NA	NA	NA	2.28E-01	Naphthalene	82.2	Household inhalation	72.0
Future adult rural resident at modeled concentrations (RGA groundwater drawn at property boundary)	4.81E-06	Arsenic ⁹⁹ Tc	69.9 30.1	Ingestion	99.8	<0.1	*No COCs		*No COCs	
Future child rural resident at modeled concentrations (RGA groundwater drawn at Ohio River)	NA	NA	NA	NA	NA		*No COCs		*No COCs	
Future adult rural resident at modeled concentrations (RGA groundwater drawn at Ohio River)		*No COCs		*No COCs			*No COCs		*No COCs	
Future child recreational user at current concentrations (soil) (WAG 3 RI ^b)	NA	NA	NA	NA	NA	<1	*No COCs		*No COCs	
Future teen recreational user at current concentrations (soil) (WAG 3 RI ^b)	NA	NA	NA	NA	NA	<1	*No COCs		*No COCs	
Future adult recreational user at current concentrations (soil) (WAG 3 RI ^b)	1.0E-05	Arsenic Total PAH Total PCB	2 96 2	Ingestion of venison Ingestion of rabbit Ingestion of quail	16 63 21	<1	*No COCs		*No COCs	
Future excavation worker at current concentrations (soil and waste) (WAG 3 RI ^b)	2.9E-04	Arsenic Beryllium Total PAH Total PCB	8 62 28 1	Ingestion Dermal	13 87	2.16E+00	Aluminum Arsenic Barium Beryllium Chromium Iron Manganese	9 7 2 3 18 38 22	Ingestion Dermal	18 82

ELCR = excess lifetime cancer risk; HI = hazard index; POC = point of contact; COC = contaminant of concern
 Note: NA = ELCR not applicable to child and teen cohorts. ELCR for adult is for lifetime exposure and takes into account exposure as child and teen.

*No COCs = There are no COCs or POCs.

* = The ELCR is approximate because the linearized multistage model returns imprecise values at risks > 1.0E-02.

^a Total ELCR and total HI represent total risk or hazard summed across all POCs for all COCs.

^b Remedial Investigation Report for Waste Area Grouping 3 at the Paducah Gaseous Diffusion Plant, Paducah, Kentucky, DOE/OR/07-1895, & D1, September 2000 (DOE 2000a), Table 1.56. In this table, lead has been excluded as a COC.

Table 6.10. Summary of Risk Characterization for SWMU 6

Receptor	Total ELCR ^a	COCs	% Total ELCR	POCs	% Total ELCR	Total HI ^c	COCs	% Total HI	POCs	% Total HI
Current industrial worker at current concentrations (soil) (WAG 3 RI ^b)	2.4E-04	Beryllium Total PAH	90 10	Dermal	99	< 1	*No COCs		*No COCs	
Future industrial worker at current concentrations (soil) (WAG 3 RI ^b)	2.4E-04	Beryllium Total PAH	90 10	Dermal	99	< 1	*No COCs		*No COCs	
Future child rural resident at current concentrations (soil) (WAG 3 RI ^b)	NA	NA	NA	NA	NA	9.38E+00	Beryllium Chromium Nickel Zinc	8 72 15 5	Dermal Ingestion of vegetables	34 65
Future adult rural resident at current concentrations (soil) (WAG 3 RI ^b)	2.4E-03	Beryllium Total PAH	54 46	Dermal Ingestion of vegetables	30 69	2.57E+00	Beryllium Chromium Nickel Zinc	7 70 17 6	Dermal Ingestion of vegetables	24 75
Future child rural resident at current concentrations (RGA groundwater only)		*No COCs		*No COCs		1.77E-01	Manganese	100	Ingestion of water Dermal	97.9 2.1
Future adult rural resident at current concentrations (RGA groundwater only)		*No COCs		*No COCs		5.18E-02	Manganese	100	Ingestion of water Dermal	95.7 4.3
Future child rural resident at modeled concentrations (RGA groundwater drawn at plant boundary)		*No COCs		*No COCs			*No COCs		*No COCs	
Future adult rural resident at modeled concentrations (RGA groundwater drawn at plant boundary)		*No COCs		*No COCs			*No COCs		*No COCs	
Future child rural resident at modeled concentrations (RGA groundwater drawn at property boundary)		*No COCs		*No COCs			*No COCs		*No COCs	
Future adult rural resident at modeled concentrations (RGA groundwater drawn at property boundary)		*No COCs		*No COCs			*No COCs		*No COCs	
Future child rural resident at modeled concentrations (RGA groundwater drawn at Ohio River)		*No COCs		*No COCs			*No COCs		*No COCs	
Future adult rural resident at modeled concentrations (RGA groundwater drawn at Ohio River)		*No COCs		*No COCs			*No COCs		*No COCs	

Table 6.10. Summary of Risk Characterization for SWMU 6 (Continued)

Receptor	Total ELCR ^a	COCs	% Total ELCR	POCs	% Total ELCR	Total HI ^c	COCs	% Total HI	POCs	% Total HI
Future child recreational user at current concentrations (soil) (WAG 3 RI ^b)	NA	NA	NA	NA	NA	< 1	*No COCs		*No COCs	
Future teen recreational user at current concentrations (soil) (WAG 3 RI ^b)	NA	NA	NA	NA	NA	< 1	*No COCs		*No COCs	
Future adult recreational user at current concentrations (soil) (WAG 3 RI ^b)	< 1.0E-06	*No COCs		*No COCs		< 1	*No COCs		*No COCs	
Future excavation worker at current concentrations (soil and waste) (WAG 3 RI ^b)	2.3E-04	Beryllium Total PAH	90 9	Ingestion Dermal	5 95	2.44E+00	Aluminum Barium Beryllium Chromium Iron Manganese Vanadium	8 2 3 15 32 15 26	Ingestion Dermal	12 88

ELCR = excess lifetime cancer risk; HI = hazard index; POC = point of contact; COC = contaminant of concern
 Note: NA = ELCR not applicable to child and teen cohorts. ELCR for adult is for lifetime exposure and takes into account exposure as child and teen.
 * = There are no COCs or POCs.

^a Total ELCR and total HI represent total risk or hazard summed across all POCs for all COCs.

^b Remedial Investigation Report for Waste Area Grouping 3 at the Paducah Gaseous Diffusion Plant, Paducah, Kentucky, DOE/OR/07-1895&D1, September 2000 (DOE 2000a), Table 1.57. In this table, lead has been excluded as a COC.

Table 6.11. Summary of Risk Characterization for SWMU 7

Receptor	Total ELCR ^a	COCs	% Total ELCR	POCs	% Total ELCR	Total HI ^a	COCs	% Total HI	POCs	% Total HI
Current industrial worker at current concentrations (soil) (from WAG 22 RI ^b)	3.8E-03	Arsenic	0.6	Ingestion	0.5	5.0E+00	Aluminum	4.1	Ingestion	3.6
		Beryllium	97.6	Dermal	97.4		Antimony	4.4	Dermal	96.4
Future industrial worker at current concentrations (soil) (from WAG 22 RI ^b)	3.9E-03	Benzo(a)anthracene	<0.1	External exposure	2.5	5.0E+00	Arsenic	2.6	Ingestion Dermal	3.6 96.4
		Benzo(a)pyrene	0.3	Ingestion Dermal External exposure	0.5 97.4 2.5		Beryllium	9.6		
		Benzo(b)fluoranthene	<0.1				Chromium	13.6		
		Dibenzo(a,h)anthracene	0.4				Iron	20.6		
		Indeno(1,2,3-cd)pyrene	0.1				Manganese	10.7		
		²³⁷ Np	<0.1				Uranium	13.7		
		²³⁴ U	<0.1				Vanadium	17.7		
		²³⁵ U	0.2				Aluminum Antimony Arsenic Beryllium Chromium Iron Manganese Uranium Vanadium	4.1 4.4 2.6 9.6 13.6 20.6 10.7 13.7 17.7		
		^{235/236} U	0.3							
		²³⁸ U	2.1							
		Arsenic	0.6							
		Beryllium	96.0							
		Benzo(a)anthracene	<0.1							
		Benzo(a)pyrene	0.3							
Benzo(b)fluoranthene	<0.1									
Dibenzo(a,h)anthracene	0.4									
Indeno(1,2,3-cd)pyrene	0.1									
²³⁷ Np	<0.1									
²³⁴ U	<0.1									
²³⁵ U	0.2									
^{235/236} U	0.3									
²³⁸ U	2.1									
Future child rural resident at current concentrations (soil) (from WAG 22 RI ^b)	NA	NA	NA	NA	3.7E+02	Aluminum Antimony Arsenic Barium Beryllium Cadmium Chromium Cobalt Copper Iron Manganese Nickel Uranium Vanadium Zinc Aroclor-1254	2.7 0.9 6.2 0.3 1.3 0.8 2.7 0.1 0.3 19.7 1.9 0.4 58.4 2.4 0.2 1.7	Ingestion Dermal Ingestion of vegetables from soil	1.4 7.7 90.9	

Table 6.11. Summary of Risk Characterization for SWMU 7 (Continued)

Receptor	Total ELCR ^a	COCs	% Total ELCR	POCs	% Total ELCR	Total HI ^c	COCs	% Total HI	POCs	% Total HI
Future adult rural residents at current concentrations (soil) (from WAG 22 RI ^b)	3.4E-02	Arsenic Beryllium Aroclor-1254 Aroclor-1260 Benzo(a)anthracene Benzo(a)pyrene Benzo(b)fluoranthene Benzo(k)fluoranthene Dibenzo(a,h)anthracene Indeno(1,2,3-cd)pyrene ²³⁷ Np ²³⁹ Pu ²³⁴ U ²³⁵ U ^{235/236} U ²³⁸ U	7.3 65.4 0.2 0.4 0.2 1.7 0.2 <0.1 1.9 0.3 0.2 0.4 3.3 0.3 0.5 17.6	Ingestion Dermal External exposure Ingestion of vegetables from soil	0.5 33.0 1.9 64.6	1.1E+02	Aluminum Antimony Arsenic Barium Beryllium Cadmium Chromium Copper Iron Manganese Nickel Uranium Vanadium Zinc Aroclor-1254	2.7 0.8 6.5 0.3 1.1 0.8 2.3 0.3 19.8 1.6 0.4 59.5 2.0 0.2 1.7	Ingestion Dermal Ingestion of vegetables from soil	0.5 5.0 94.6
Future child rural residents at current concentrations (RGA groundwater only)	NA	NA	NA	NA	NA	1.89E+01	Arsenic Manganese Uranium 1,1-DCE <i>cis</i> -1,2-DCE Aroclor-1254 TCE Vinyl chloride	30.2 3.7 2.9 4.5 6.6 22.3 26.4 3.4	Ingestion Dermal contact Inhalation while showering Inhalation household use	60.9 21.0 2.0 16.0
Future adult rural residents at current concentrations (RGA groundwater only)	3.13E-03	Arsenic 1,1-DCE Total PCBs TCE Vinyl chloride ⁹⁹ Tc ²³⁴ U ²³⁸ U	15.1 66.4 0.2 4.1 11.9 1.6 0.4 0.4	Ingestion Dermal contact Inhalation while showering Inhalation during household use	61.2 3.7 4.9 30.3	6.39E+00	Arsenic Manganese Uranium 1,1-DCE <i>cis</i> -1,2-DCE Total PCBs TCE Vinyl chloride	25.5 3.2 2.5 3.1 4.5 31.4 27.1 2.7	Ingestion Dermal contact Inhalation while showering Inhalation household use	51.4 37.2 1.3 10.1

Table 6.11. Summary of Risk Characterization for SWMU 7 (Continued)

Receptor	Total ELCR ^c	COCs	% Total ELCR	POCs	% Total ELCR	Total HI ^r	COCs	% Total HI	POCs	% Total HI
Future child rural resident at modeled concentrations (RGA groundwater drawn at plant boundary)	NA	NA	NA	NA	1.45E+01	Arsenic Manganese Uranium 1,1-DCE <i>cis</i> -1,2-DCE Total PCBs TCE Vinyl chloride	27.9 3.6 2.8 5.4 7.9 17.2 31.2 4.1	Ingestion Dermal contact Inhalation while showering Inhalation household use	62.3 18.7 2.2 16.9	
Future adult rural resident at modeled concentrations (RGA groundwater drawn at plant boundary)	2.98E-03	Arsenic 1,1-DCE Total PCBs TCE Vinyl chloride ^{99m} Tc ²³⁴ U ²³⁸ U	11.2 63.9 0.2 10.3 12.3 1.5 0.3 0.3	Ingestion Dermal contact Inhalation while showering Inhalation during household use	55.4 3.4 4.7 36.5	4.78E+00	Arsenic Manganese Uranium 1,1-DCE <i>cis</i> -1,2-DCE Total PCBs TCE Vinyl chloride	24.2 3.1 2.4 3.8 5.5 24.8 32.9 3.3	Ingestion of groundwater Dermal contact Inhalation household use	53.8 33.8 11.0
Future child rural resident at modeled concentrations (RGA groundwater drawn at property boundary)	NA	NA	NA	NA	1.97E+00	Arsenic 1,1-DCE <i>cis</i> -1,2-DCE Total PCBs TCE	38.1 5.3 8.4 12.4 32.9	Ingestion Dermal contact Inhalation household use	66.3 15.8 15.9	
Future adult rural resident at modeled concentrations (RGA groundwater drawn at property boundary)	4.11E-04	Arsenic 1,1-DCE TCE Vinyl chloride ^{99m} Tc	15.1 61.8 10.7 8.7 3.6	Ingestion Dermal contact Inhalation while showering Inhalation during household use	56.7 3.2 4.5 35.5	6.36E-01	Arsenic Total PCBs TCE	33.9 18.4 35.5	Ingestion Dermal contact	58.8 29.3
Future child rural resident at modeled concentrations (RGA groundwater drawn at Little Bayou seeps)	NA	NA	NA	NA	3.373E-01	TCE	61.0	Ingestion Inhalation household use	52.5 30.0	
Future adult rural resident at modeled concentrations (RGA groundwater drawn at Little Bayou seeps)	1.28E-04	1,1-DCE TCE Vinyl chloride ^{99m} Tc	72.6 12.3 9.5 5.7	Ingestion Dermal contact Inhalation while showering Inhalation during household use	49.6 3.6 5.3 41.4	1.15E-01	*No COCs	*No COCs		

Table 6.11. Summary of Risk Characterization for SWMU 7 (Continued)

Receptor	Total ELCR ^a	COCs	% Total ELCR	POCs	% Total ELCR	Total HI ^c	COCs	% Total HI	POCs	% Total HI
Future child rural resident at modeled concentrations (RGA groundwater drawn at Ohio River)		Not a POE for groundwater from this SWMU.								
Future adult rural resident at modeled concentrations (RGA groundwater drawn at Ohio River)		Not a POE for groundwater from this SWMU.								
Future child recreational user at current concentrations (from WAG 22 RI ^b)	NA	NA	NA	NA	NA	7.3E-02	*No COCs		*No COCs	
Future teen recreational user at current concentrations (from WAG 22 RI ^b)	NA	NA	NA	NA	NA	6.4E-02	*No COCs		*No COCs	
Future adult recreational user at current concentrations (from WAG 22 RI ^b)	1.1E-05	Arochlor-1260 Benzo(a)pyrene Dibenzo(a,h)anthracene ²³⁸ U	18.6 9.5 42.5 15.7	Ingestion of deer Ingestion of rabbit Ingestion of quail	10.0 70.9 21.8	7.5E-02	*No COCs		*No COCs	
Future excavation worker at current concentrations (soil) (from WAG 22 RI ^b)	1.6E-03	Arsenic Beryllium Benzo(a)pyrene Dibenzo(a,h)anthracene ²³⁷ Np ²³⁹ Pu ²³⁴ U ²³⁵ U ^{235/236} U ²³⁸ U	1.8 42.2 0.1 1.7 0.4 0.5 3.4 9.1 0.4 41.3	Ingestion Dermal External exposure	25.6 43.8 32.5	5.4E+00	Aluminum Antimony Arsenic Chromium Copper Iron Manganese Nickel Uranium Vanadium	5.0 11.3 3.4 17.6 2.9 21.3 11.0 3.9 7.5 10.9	Ingestion Dermal	18.4 81.5

ELCR = excess lifetime cancer risk; HI = hazard index; POC = point of contact; COC = contaminant of concern

Note: NA = ELCR not applicable to child and teen cohorts. ELCR for adult is for lifetime exposure and takes into account exposure as child and teen.

*No COCs = There are no COCs or POCs.

^a Total ELCR and total HI represent total risk or hazard summed across all POCs for all COCs.

^b Remedial Investigation Report for Solid Waste Management Units 7 and 30 of Waste Area Group 22 at the Paducah Gaseous Diffusion Plant, Paducah, Kentucky, DOE/OR/07-1604/V1&D2, January 1998 (DOE 1998a), Tables 1.59 through 1.68, excluding lead as a COC.

Table 6.12. Summary of Risk Characterization for SWMU 30

Receptor	Total ELCR ^a	COCs	% Total ELCR	POCs	% Total ELCR	Total HI ^c	COCs	% Total HI	POCs	% Total HI	
Current industrial worker at current concentrations (soil) (from WAG 22 RI ^b)	3.7E-03	Arsenic	0.5	Ingestion	0.5	4.4E+00	Aluminum	5.1	Ingestion	5.1	
		Beryllium	97.5	Dermal	97.3		Antimony	3.7	Dermal	3.7	
		Aroclor-1260	0.1	External exposure	1.7		Arsenic	2.7			2.9
		Benzo(a)anthracene	0.1				Beryllium	10.8			97.1
		Benzo(a)pyrene	0.8				Cadmium	3.5			
		Benzo(b)fluoranthene	0.1				Chromium	13.5			
		Dibenzo(a,h)anthracene	0.3				Iron	19.8			
		Indeno(1,2,3-cd)pyrene	0.1				Manganese	11.3			
		²³⁷ Np	<0.1				Uranium	9.0			
		²³⁴ U	<0.1				Vanadium	17.6			
		²³⁵ U	0.2								
		^{235/236} U	0.3								
		²³⁸ U	1.4								
Future industrial worker at current concentrations (soil) (from WAG 22 RI ^b)	3.8E-03	Arsenic	0.5	Ingestion	0.5	4.4E+00	Aluminum	5.1	Ingestion	5.1	
		Beryllium	96.2	Dermal	97.8		Antimony	3.7	Dermal	3.7	2.9
		Aroclor-1260	0.1	External exposure	1.7		Arsenic	2.7			97.1
		Benzo(a)anthracene	0.1				Beryllium	10.8			
		Benzo(a)pyrene	0.8				Cadmium	3.5			
		Benzo(b)fluoranthene	0.1				Chromium	13.5			
		Dibenzo(a,h)anthracene	0.3				Iron	19.8			
		Indeno(1,2,3-cd)pyrene	0.1				Manganese	11.3			
		²³⁷ Np	<0.1				Uranium	9.0			
		³⁴ U	<0.1				Vanadium	17.6			
		³⁵ U	0.2								
		^{235/236} U	0.3								
		²³⁸ U	1.4								
Future child rural resident at current concentrations (soil) (from WAG 22 RI ^b)	NA					2.6E+02	Aluminum	4.1	Ingestion	4.1	
							Antimony	0.9	Dermal	0.9	1.3
							Arsenic	7.5	Ingestion of vegetables from soil	7.5	9.4
							Barium	0.6			89.3
							Beryllium	1.8			
							Cadmium	2.2			
							Chromium	3.2			
							Copper	0.6			
							Iron	22.6			
							Manganese	2.5			
							Mercury	0.7			
							Nickel	0.8			
							Uranium	46.8			
					Vanadium	3.0					
					Zinc	0.2					
					Aroclor-1254	2.6					

Table 6.12. Summary of Risk Characterization for SWMU 30 (Continued)

Receptor	Total ELCR ^d	COCs	% Total ELCR	POCs	% Total ELCR	Total HI ^e	COCs	% Total HI	POCs	% Total HI
Future adult rural resident at current concentrations (soil) (from WAG 22 RI ^b)	3.2E-02	Arsenic Beryllium Aroclor-1254 Aroclor-1260 Benzo(a)anthracene Benzo(a)pyrene Benzo(b)fluoranthene Benzo(k)fluoranthene bis(2-ethylhexyl)phthalate Chrysene Dibenzo(a,h)anthracene Indeno(1,2,3-cd)pyrene ²³⁷ Np ²³⁴ U ²³⁵ U ^{235,236} U ²³⁸ U	6.8 66.7 0.2 1.8 0.4 4.4 0.5 <0.1 <0.1 <0.1 1.7 0.4 0.2 4.5 0.3 0.6 11.5	Ingestion Dermal External exposure Ingestion of vegetables from soil	0.5 35.4 1.3 62.8	7.9E+01	Aluminum Antimony Arsenic Barium Beryllium Cadmium Chromium Copper Iron Manganese Mercury Nickel Uranium Vanadium Zinc Aroclor-1254	4.1 0.8 7.9 0.6 1.5 2.2 2.9 0.6 22.8 2.1 0.7 0.9 47.5 2.4 0.2 2.7	Ingestion Dermal Ingestion of vegetables from soil	0.5 6.1 93.4
Future child rural resident at current concentrations (RGA groundwater only)	NA	NA	NA	NA	NA	9.14E+00	Arsenic Manganese Selenium Uranium 1,1-DCE TCE	63.8 8.8 3.2 14.7 5 4.6	Ingestion Dermal contact Inhalation while showering Inhalation household use	93.3 1.3 0.6 4.7
Future adult rural resident at current concentrations (RGA groundwater only)	5.44E-04	Arsenic 1,1-DCE TCE ⁹⁹ Tc ²³⁴ U ²³⁸ U	88.6 0.3 5.2 2.9 1 1.3	Ingestion Dermal contact Inhalation while showering Inhalation household use	95.3 0.9 0.4 3.4	3.31E+00	Arsenic Manganese Selenium Uranium 1,1-DCE TCE	50.5 7.1 2.5 11.6 23.9 4.4	Ingestion Dermal contact Inhalation while showering Inhalation household use	88.8 9.8 0.2 1.2
Future child rural resident at modeled concentrations (RGA groundwater drawn at plant boundary)	NA	NA	NA	NA	NA	6.14E+00	Arsenic Manganese Selenium Uranium 1,1-DCE TCE	63.1 8.7 2.6 12.5 0.1 12.9	Ingestion of groundwater Dermal contact Inhalation while showering Inhalation household use	91.1 1.7 0.1 7.1
Future adult rural resident at modeled concentrations (RGA groundwater drawn at plant boundary)	3.75E-04	Arsenic 1,1-DCE TCE ⁹⁹ Tc ²³⁴ U ²³⁸ U	85.6 0.5 7.1 3.9 1 1.9	Ingestion Dermal contact Inhalation while showering Inhalation household use	93.6 1.1 0.6 4.7	2.10E+00	Arsenic Manganese Selenium Uranium 1,1-DCE TCE	52.9 7.4 2.2 10.5 0.4 26.6	Ingestion Dermal contact Inhalation while showering Inhalation household use	76.1 3 0 20.8

Table 6.12. Summary of Risk Characterization for SWMU 30 (Continued)

Receptor	Total ELCR ^a	COCs	% Total ELCR	POCs	% Total ELCR	Total HI ^r	COCs	% Total HI	POCs	% Total HI
Future child rural resident at modeled concentrations (RGA groundwater drawn at property boundary)	NA	NA	NA	NA	NA	8.40E-01	Arsenic Selenium 1,1-DCE TCE Manganese	89.2 2.1 0.1 8.5 0.1	Ingestion Dermal contact Inhalation while showering Inhalation household use	94.2 1.1 0 4.6
Future adult rural resident at modeled concentrations (RGA groundwater drawn at property boundary)	6.85E-05	Arsenic 1,1-DCE TCE Technetium-99	90.6 0.2 3.5 5.7	Ingestion Dermal contact Inhalation while showering Inhalation household use	96.7 0.7 0.3 2.3	2.76E-01	Arsenic Selenium 1,1-DCE TCE Manganese	77.9 1.8 0.3 19.9 0.1	Ingestion Dermal contact Inhalation while showering Inhalation household use	82 2 1.8 14.2
Future child rural resident at modeled concentrations (RGA groundwater drawn at Little Bayou seeps)	NA	NA	NA	NA	NA	3.02E-02	Selenium 1,1-DCE TCE	20 0.6 79.3	Ingestion Dermal contact Inhalation household use	47.5 8.6 0.4 43.5
Future adult rural resident at modeled concentrations (RGA groundwater drawn at Little Bayou seeps)	2.45E-06	1,1-DCE TCE ⁹⁹ Tc	1.8 32.9 65.3	Ingestion Dermal contact Inhalation while showering Inhalation household use	72.1 3.7 2.7 21.4	9.17E-03	Selenium 1,1-DCE TCE	18.9 2.3 78.8	Ingestion Dermal contact Inhalation while showering Inhalation household use	44.7 17 18.3 20
Future child rural resident at modeled concentrations (RGA groundwater drawn at Ohio River)		Not a POE for groundwater from this SWMU								
Future adult rural resident at modeled concentrations (RGA groundwater drawn at Ohio River)		Not a POE for groundwater from this SWMU								
Future child recreational user at current concentrations (from WAG 22 RI ^b)	NA	NA	NA	NA	NA	4.2E-02	*No COCs		*No COCs	
Future teen recreational user at current concentrations (from WAG 22 RI ^b)	NA	NA	NA	NA	NA	3.8E-02	*No COCs		*No COCs	

Table 6.12. Summary of Risk Characterization for SWMU 30 (Continued)

Receptor	Total ELCR ^a	COCs	% Total ELCR	POCs	% Total ELCR	Total HI ^c	COCs	% Total HI	POCs	% Total HI
Future adult recreational user at current concentrations (from WAG 22 RI ^b)	1.5E-05	Aroclor-1260	48.2	Ingestion of deer	8.7	4.3E-02	*No COCs		*No COCs	
		Benzo(a)pyrene	12.9	Ingestion of rabbit	80.0					
		Dibenzo(a,h)anthracene	20.8	Ingestion of quail	11.3					
Future excavation worker at current concentrations (soil) (from WAG 22 RI ^b)	1.2E-03	Arsenic	1.9	Ingestion	6.3	4.5E+00	Aluminum Antimony Arsenic Beryllium Cadmium Chromium Copper Iron Manganese Uranium Vanadium	4.6	Ingestion Dermal	26.4
		Beryllium	93.7	Dermal	91.7					
		Aroclor-1248	0.1	External exposure	3.3					
		Benzo(a)anthracene	0.1							
		Benzo(a)pyrene	0.8							
		Benzo(b)fluoranthene	0.1							
		Dibenzo(a,h)anthracene	0.4							
		Indeno(1,2,3-cd)pyrene	0.1							
		²³⁷ Np	0.3							
		²³⁹ Pu	0.2							
		²³⁴ U	0.8							
		²³⁵ U	0.1							
		^{235/236} U	0.8							
²³⁸ U	0.6									

ELCR = excess lifetime cancer risk; HI = hazard index; POC = point of contact; COC = contaminant of concern

Note: NA = ELCR not applicable to child and teen cohorts. ELCR for adult is for lifetime exposure and takes into account exposure as child and teen.

^a Total ELCR and total HI represent total risk or hazard summed across all POCs for all COCs.

^b Remedial Investigation Report for Solid Waste Management Units 7 and 30 of Waste Area Group 22 at the Paducah Gaseous Diffusion Plant, Paducah, Kentucky; DOE/OR-07-1604/V1&D2, January 1998 (DOE 1998a), Tables 1.59 through 1.68, excluding lead as a COC.

Table 6.13. Summary of Risk Characterization for SWMU 145

Receptor	Total ELCRa	COCs	% Total ELCR	POCs	% Total ELCR	Total HI ^a	COCs	% Total HI	POCs	% Total HI
Future child rural resident at current concentrations (RGA groundwater only)	NA	NA	NA	NA	NA	4.17E+01	Antimony Arsenic Manganese	48.0 47.7 4.3	Ingestion Dermal contact	97.8 2.2
Future adult rural resident at current concentrations (RGA groundwater only)	3.27E-02	Arsenic ⁹⁹ Tc Aroclor-1260	5.1 1.7 93.2	Ingestion Dermal contact	6.9 93.1	1.22E+01	Antimony Arsenic Manganese	49.0 46.7 4.3	Ingestion Dermal contact	95.5 4.5
Future child rural resident at modeled concentrations (RGA groundwater drawn at property boundary)	NA	NA	NA	NA	NA	5.16E-01	Arsenic	99.9	Ingestion	99.8
Future adult rural resident at modeled concentrations (RGA groundwater drawn at property boundary)	1.44E-04	Arsenic ⁹⁹ Tc	29.7 70.3	Ingestion	99.9	1.48E-01	Arsenic	99.9	Ingestion	99.6
Future child rural resident at modeled concentrations (RGA groundwater drawn at Ohio River)	NA	NA	NA	NA	NA		*No COCs		*No COCs	
Future adult rural resident at modeled concentrations (RGA groundwater drawn at Ohio River)	5.29E-05	⁹⁹ Tc	100.0	Ingestion	100		*No COCs		*No COCs	
Future child recreational user at current concentrations (soil)	NE	NE	NE	NE	NE	NE	NE	NE	NE	NE

Table 6.13. Summary of Risk Characterization for SWMU 145 (Continued)

Receptor	Total ELCR ^a	COCs	% Total ELCR	POCs	% Total ELCR	Total HI ^c	COCs	% Total HI	POCs	% Total HI
Future teen recreational user at current concentrations (soil)	NE	NE	NE	NE	NE	NE	NE	NE	NE	NE
Future adult recreational user at current concentrations (soil)	NE	NE	NE	NE	NE	NE	NE	NE	NE	NE
Future excavation worker at current concentrations	NE	NE	NE	NE	NE	NE	NE	NE	NE	NE

ELCR = excess lifetime cancer risk; HI = hazard index; POC = point of contact; COC = contaminant of concern

Note: NA = ELCR not applicable to child and teen cohorts. ELCR for adult is for lifetime exposure and takes into account exposure as child and teen.

NE = not evaluated; land use scenario was not assessed because surface soil was not assessed for this SWMU.

^a Total ELCR and total HI represent total risk or hazard summed across all POCs for all COCs.

6.4.9 Uncertainty in the Risk Assessment

Risk and hazard estimates could vary if different assumptions were used in deriving the risk estimates or if better information were available for some parameters. No uncertainties were estimated to have a large effect on the risk characterization, and only the following were estimated to have a moderate effect:

- Exclusion of some potential biota (produce and fish) for future receptors,
- Migration of groundwater to off-site receptors,
- Calculation of toxicity values for chemicals (particularly TCE), and
- Updates to toxicity values.

Uncertainty on toxicity factors plays a major role in this risk assessment. Because the RfD for lead was in question even when the previous soil assessments were written, the results were calculated without lead in those previous documents, and those results were summarized in this assessment. At the time the WAG 22 and WAG 3 RI reports were developed, beryllium still was evaluated as a carcinogen through the oral route of exposure. Since the completion of those BHHRAs, the oral cancer slope factor for beryllium has been withdrawn from IRIS, and there has been an agreement not to use this withdrawn value for risk assessments performed in EPA Region 4. At several SWMUs, beryllium was a significant contributor to the total cancer risk from soil exposure; generally, beryllium accounted for greater than 90% of the risk to the industrial worker and greater than 65% of the risk to the resident, with nearly all the risk due to the oral exposure route. While the inhalation pathway for beryllium exposure is valid, it minutely contributed to the total risk due to beryllium exposure. For example, SWMU 7 exhibited the highest beryllium inhalation risk for a residential user of any SWMU at 5.59E-09, while the total oral ingestion risk was 3.22E-02. The inhalation risk accounted for 0.00002% of the total risk. While beryllium would be a COPC, it would be screened from evaluation as a COC because the highest risk from any SWMU was three orders of magnitude less than 1 E-06.

When beryllium is removed from consideration as a carcinogen, the total ELCR becomes much lower at those SWMUs for which it is a COC. The total risk including and excluding the contribution from beryllium is shown below by SWMU and receptor.

	Receptor	Risk Including Beryllium	Risk Excluding Beryllium
SWMU 4	Future industrial worker	5.4E-04	1.6E-05
SWMU 4	Adult Resident	4.3E-03	1.2E-03
SWMU 5	Future industrial worker	4.1E-04	2.1E-04
SWMU 5	Adult Resident	1.0E-02	9.1E-03
SWMU 6	Future industrial worker	2.4E-04	2.4E-05
SWMU 6	Adult Resident	2.4E-03	1.1E-03
SWMU 7	Future industrial worker	3.9E-03	1.6E-04
SWMU 7	Adult Resident	3.4E-02	1.2E-02
SWMU 30	Future industrial worker	3.8E-03	1.4E-04
SWMU 30	Adult Resident	3.2E-02	1.1E-02

For SWMUs 4 and 6, removal of the contribution of beryllium to the ELCR reduces the total ELCR to within the EPA risk range of 1E-04 to 1E-06 for the industrial worker scenario.

Summary of Historical Data Quality Assessment

A summary of uncertainties and any data evaluation or DQA is included in Appendix F as part of the summary of uncertainties. The wording of the historical uncertainty assessment has not been changed but some text and tables have been removed to be succinct. Every effort was made to accurately report the uncertainty sections of the historical investigations in summary form. The DQA for each of the SWMUs was performed as part of each SWMU SI, RI, or investigation. This BHHRA does not include a further analysis of this data.

6.5 BHHRA OBSERVATIONS AND CONCLUSIONS

This section summarizes the results of this and previous BHHRAs and draws conclusions from the results. This section also includes a series of observations in which the results of the BHHRAs are combined with the uncertainties in the risk assessment.

Appendix F provides observations regarding the results and uncertainties of this and previous risk assessments in detail. This section provides a summary of the results of those observations. The discussion focuses on the individual exposure scenarios examined for the assessment.

6.5.1 Future Industrial Worker

SWMUs 4, 7, and 30 hazard levels exceed 1 for industrial worker exposure to soil, with chromium, iron, and vanadium serving as the primary hazard drivers for elevated HIs. All SWMUs (SWMU 145 was not evaluated for this scenario) exceed risk levels of 1E-04 for industrial worker exposure to soil, with uranium-235+daughters, uranium-238+daughters, and beryllium serving as the primary risk drivers. Other COCs contributing to elevated risks include Total PAH and arsenic. SWMUs 2, 3, 5, 7, and 30 exceed risk levels of 1E-04 for industrial worker exposure to soil, if beryllium is not included in the cumulative ELCR for each of the SWMUs.

6.5.2 Future Excavation Worker

SWMUs 4, 5, 6, 7, and 30 exceed a hazard level of 1 for excavation worker exposure to soil (SWMUs 4, 5 and 6 were evaluated for exposure to soil and waste), with aluminum, antimony, chromium, iron, manganese, uranium, and vanadium serving as the primary hazard driver for elevated HIs. Other COCs contributing to hazards include arsenic and copper. SWMUs 4, 5, 6, 7, and 30 exceed the risk level of 1E-04 for excavation worker exposure to soil, with beryllium, uranium, Total PAH, and uranium-238 serving as the primary risk drivers. Other COCs contributing to elevated risks include arsenic and uranium-235. SWMUs 4, 5, 7, and 145 exceed the risk level of 1E-04 for excavation worker exposure to soil when beryllium is not included in the cumulative ELCR. SWMUs 4, 5 and 6 included exposure to soil and waste which was included in this BHHRA but referred to as one media type, soil.

6.5.3 Future On-Site Residents (Groundwater)

For residential groundwater use at the SWMU boundary, ELCR was greater than 1E-04 and HI was greater than 1 for all SWMUs except SWMU 6. The primary risk drivers are TCE, arsenic, vinyl chloride, 1,1-DCE, and technetium-99.

6.5.4 Future Off-Site Residents (Groundwater)

SWMUs 2, 4, 5, and 7 exceed a hazard level of 1 for off-site residential exposure to groundwater at the PGDP plant boundary. SWMUs 2, 4, and 7 exceed a hazard level of 1 at the property boundary. SWMUs 2, and 4 exceed a hazard level of 1 at the Ohio River (or seeps). The primary drivers for hazard are arsenic, TCE, *cis*-1,2-DCE, and 1,1-DCE. SWMUs 2, 3, 4, and 7 at the plant boundary, SWMUs 2, 4, 7, 30, and 145 at the property boundary, and SWMUs 2, 4, and 7 at the Ohio River (or seeps) exceed a risk level of 1E-04 for off-site residential exposure to groundwater. The primary risk drivers are TCE, 1,1-DCE, and technetium-99.

6.6 REMEDIAL GOAL OPTIONS

RGOs are presented in Table 6.14 for soil for the industrial worker, excavation worker, and residential user scenarios and in Table 6.15 for the residential groundwater user. RGOs were calculated for each COC from the modeled groundwater concentrations considering residential use of groundwater at each source and at the property boundary POE. When calculating the HI-based RGOs, the more conservative child-based values are reported. In addition, for comparison to the RGOs, the MCL for each COC is presented. Note, MCLs are not clean-up criteria, though they may be ARARs. The National Contingency Plan (NCP) notes that clean-up criteria different from MCLs may be required if multiple contaminants are present or if contaminants may reach a receptor through exposure routes different from those considered in the development of MCLs. Risks for use of contaminated groundwater must be presented in addition to a simple screen against MCLs so that risk managers can make appropriate decisions. The calculation used to derive groundwater RGOs can be found in Section 8.1 of Appendix F. The soil RGOs were calculated from the NALs (DOE 2008b) for all COCs from the previous risk assessments for SWMUs 2, 3, 4, 5, 6, 7, and 30 (DOE 1994; DOE 1998a; DOE 2000a).

Table 6.14. RGOs for Soil COCs of the BGOU SWMU5

COC ^A	Noncancer			RGO ^B at			RGO at			RGO at			Units
	Cancer NAL	NAL	HI=0.1	HI=1	HI=3	1 x 10 ⁻⁶	1 x 10 ⁻⁵	1 x 10 ⁻⁴	1 x 10 ⁻⁵	1 x 10 ⁻⁴	1 x 10 ⁻⁴		
Residential User Soil Exposure													
Aluminum		9.69E+02	9.69E+01	9.69E+02	2.91E+03							mg/kg	
Antimony		8.69E-02	8.69E-03	8.69E-02	2.61E-01							mg/kg	
Arsenic	1.44E-01	1.16E+00	1.16E-01	1.16E+00	3.48E+00	1.44E-01	1.44E+00	1.44E+01				mg/kg	
Barium		1.40E+02	1.40E+01	1.40E+02	4.20E+02							mg/kg	
Beryllium and compounds	1.19E-03	2.20E-01	2.20E-02	2.20E-01	6.60E-01	1.19E-03	1.19E-02	1.19E-01				mg/kg	
Cadmium	2.00E+00	3.26E+00	3.26E-01	3.26E+00	9.78E+00	2.00E+00	2.00E+01	2.00E+02				mg/kg	
Chromium	1.10E+02	8.32E+01	8.32E+00	8.32E+01	2.50E+02	1.10E+02	1.10E+03	1.10E+04				mg/kg	
Cobalt	4.69E+02	6.95E+01	6.95E+00	6.95E+01	2.09E+02	4.69E+02	4.69E+03	4.69E+04				mg/kg	
Copper		9.39E+01	9.39E+00	9.39E+01	2.82E+02							mg/kg	
Iron		4.14E+02	4.14E+01	4.14E+02	1.24E+03							mg/kg	
Manganese		5.60E+01	5.60E+00	5.60E+01	1.68E+02							mg/kg	
Nickel	5.06E+03	4.35E+01	4.35E+00	4.35E+01	1.31E+02	5.06E+03	5.06E+04	5.06E+05				mg/kg	
Uranium		2.57E+00	2.57E-01	2.57E+00	7.71E+00							mg/kg	
Vanadium		7.71E-01	7.71E-02	7.71E-01	2.31E+00							mg/kg	
Zinc		5.21E+02	5.21E+01	5.21E+02	1.56E+03							mg/kg	
Aroclor 1260	6.08E-02					6.08E-02	6.08E-01	6.08E+00				mg/kg	
Benz[a]anthracene	7.48E-02					7.48E-02	7.48E-01	7.48E+00				mg/kg	
Benzo[a]pyrene	7.48E-03					7.48E-03	7.48E-02	7.48E-01				mg/kg	
Benzo[b]fluoranthene	7.48E-02					7.48E-02	7.48E-01	7.48E+00				mg/kg	
Dibenz[a,h]anthracene	7.48E-03					7.48E-03	7.48E-02	7.48E-01				mg/kg	
Total Dioxins/Furans	6.78E-07					6.78E-07	6.78E-06	6.78E-05				mg/kg	
Indeno[1,2,3-cd]pyrene	7.48E-02					7.48E-02	7.48E-01	7.48E+00				mg/kg	
Total PCBs	5.78E-02					5.78E-02	5.78E-01	5.78E+00				mg/kg	
Total PAHs	7.48E-03					7.48E-03	7.48E-02	7.48E-01				mg/kg	
Neptunium-237+D	8.39E-02					8.39E-02	8.39E-01	8.39E+00				pCi/g	
Plutonium-239*	3.15E+00					3.15E+00	3.15E+01	3.15E+02				pCi/g	
Radium-226+D	7.94E-03					7.94E-03	7.94E-02	7.94E-01				pCi/g	
Uranium-234	5.47E+00					5.47E+00	5.47E+01	5.47E+02				pCi/g	
Uranium-235+D	1.22E-01					1.22E-01	1.22E+00	1.22E+01				pCi/g	
Uranium-238+D	5.17E-01					5.17E-01	5.17E+00	5.17E+01				pCi/g	

Table 6.14. RGOs for Soil COCs of the BGOU SWMUs (Continued)

COC ^A	Cancer NAL		Noncancer NAL		RGO ^B at HI=0.1		RGO at HI=1		RGO at HI=3		RGO at ELCR=1 x 10 ⁻⁶		RGO at ELCR=1 x 10 ⁻⁵		RGO at ELCR=1 x 10 ⁻⁴		Units
	Cancer NAL	Noncancer NAL	HI=0.1	HI=1	HI=3	1 x 10 ⁻⁶	1 x 10 ⁻⁵	1 x 10 ⁻⁴	1 x 10 ⁻⁶	1 x 10 ⁻⁵	1 x 10 ⁻⁴	1 x 10 ⁻⁶	1 x 10 ⁻⁵	1 x 10 ⁻⁴			
Industrial Worker Soil Exposure																	
Aluminum		4.22E+03	4.22E+02	4.22E+03	1.27E+04												mg/kg
Antimony		3.46E-01	3.46E-02	3.46E-01	1.04E+00												mg/kg
Arsenic	4.84E-01	7.78E+00	7.78E-01	7.78E+00	2.33E+01	4.84E-01	4.84E+00	4.84E+01	4.84E-01	4.84E+00	4.84E+01	4.84E-01	4.84E+00	4.84E+01	4.84E+01	4.84E+01	mg/kg
Barium		5.92E+02	5.92E+01	5.92E+02	1.78E+03												mg/kg
Beryllium and compounds	2.83E-03	8.68E-01	8.68E-02	8.68E-01	2.60E+00	2.83E-03	2.83E-02	2.83E-01	2.83E-03	2.83E-02	2.83E-01	2.83E-03	2.83E-02	2.83E-01	2.83E-01	2.83E-01	mg/kg
Cadmium	1.49E+01	1.97E+01	1.97E+00	1.97E+01	5.91E+01	1.49E+01	1.49E+02	1.49E+03	1.49E+01	1.49E+02	1.49E+03	1.49E+01	1.49E+02	1.49E+03	1.49E+03	1.49E+03	mg/kg
Chromium	2.11E+02	3.26E+02	3.26E+01	3.26E+02	9.78E+02	2.11E+02	2.11E+03	2.11E+04	2.11E+02	2.11E+03	2.11E+04	2.11E+02	2.11E+03	2.11E+04	2.11E+04	2.11E+04	mg/kg
Cobalt	9.05E+02	4.48E+02	4.48E+01	4.48E+02	1.34E+03	9.05E+02	9.05E+03	9.05E+04	9.05E+02	9.05E+03	9.05E+04	9.05E+02	9.05E+03	9.05E+04	9.05E+04	9.05E+04	mg/kg
Copper		4.91E+02	4.91E+01	4.91E+02	1.47E+03												mg/kg
Iron		1.90E+03	1.90E+02	1.90E+03	5.70E+03												mg/kg
Manganese		2.29E+02	2.29E+01	2.29E+02	6.87E+02												mg/kg
Nickel	9.75E+03	2.22E+02	2.22E+01	2.22E+02	6.66E+02	9.75E+03	9.75E+04	9.75E+05	9.75E+03	9.75E+04	9.75E+05	9.75E+03	9.75E+04	9.75E+05	9.75E+05	9.75E+05	mg/kg
Uranium		1.88E+01	1.88E+00	1.88E+01	5.64E+01												mg/kg
Vanadium		3.04E+00	3.04E-01	3.04E+00	9.12E+00												mg/kg
Zinc		2.50E+03	2.50E+02	2.50E+03	7.50E+03												mg/kg
Aroclor 1260	1.75E-01					1.75E-01	1.75E+00	1.75E+01	1.75E-01	1.75E+00	1.75E+01	1.75E-01	1.75E+00	1.75E+01	1.75E+01	1.75E+01	mg/kg
Benz[a]anthracene	1.94E-01					1.94E-01	1.94E+00	1.94E+01	1.94E-01	1.94E+00	1.94E+01	1.94E-01	1.94E+00	1.94E+01	1.94E+01	1.94E+01	mg/kg
Benzo[a]pyrene	1.94E-02					1.94E-02	1.94E-01	1.94E+00	1.94E-02	1.94E-01	1.94E+00	1.94E-02	1.94E-01	1.94E+00	1.94E+00	1.94E+00	mg/kg
Benzo[b]fluoranthene	1.94E-01					1.94E-01	1.94E+00	1.94E+01	1.94E-01	1.94E+00	1.94E+01	1.94E-01	1.94E+00	1.94E+01	1.94E+01	1.94E+01	mg/kg
Dibenz[a,h]anthracene	1.94E-02					1.94E-02	1.94E-01	1.94E+00	1.94E-02	1.94E-01	1.94E+00	1.94E-02	1.94E-01	1.94E+00	1.94E+00	1.94E+00	mg/kg
Total Dioxins/Furans	1.89E-06					1.89E-06	1.89E-05	1.89E-04	1.89E-06	1.89E-05	1.89E-04	1.89E-06	1.89E-05	1.89E-04	1.89E-04	1.89E-04	mg/kg
Indeno[1,2,3-cd]pyrene	1.94E-01					1.94E-01	1.94E+00	1.94E+01	1.94E-01	1.94E+00	1.94E+01	1.94E-01	1.94E+00	1.94E+01	1.94E+01	1.94E+01	mg/kg
Total PCBs	1.63E-01					1.63E-01	1.63E+00	1.63E+01	1.63E-01	1.63E+00	1.63E+01	1.63E-01	1.63E+00	1.63E+01	1.63E+01	1.63E+01	mg/kg
Total PAHs	1.94E-02					1.94E-02	1.94E-01	1.94E+00	1.94E-02	1.94E-01	1.94E+00	1.94E-02	1.94E-01	1.94E+00	1.94E+00	1.94E+00	mg/kg
Neptunium-237+D	2.71E-01					2.71E-01	2.71E+00	2.71E+01	2.71E-01	2.71E+00	2.71E+01	2.71E-01	2.71E+00	2.71E+01	2.71E+01	2.71E+01	pCi/g
Plutonium-239*	1.07E+01					1.07E+01	1.07E+02	1.07E+03	1.07E+01	1.07E+02	1.07E+03	1.07E+01	1.07E+02	1.07E+03	1.07E+03	1.07E+03	pCi/g
Radium-226+D	2.56E-02					2.56E-02	2.56E-01	2.56E+00	2.56E-02	2.56E-01	2.56E+00	2.56E-02	2.56E-01	2.56E+00	2.56E+00	2.56E+00	pCi/g
Uranium-234	1.89E+01					1.89E+01	1.89E+02	1.89E+03	1.89E+01	1.89E+02	1.89E+03	1.89E+01	1.89E+02	1.89E+03	1.89E+03	1.89E+03	pCi/g
Uranium-235+D	3.95E-01					3.95E-01	3.95E+00	3.95E+01	3.95E-01	3.95E+00	3.95E+01	3.95E-01	3.95E+00	3.95E+01	3.95E+01	3.95E+01	pCi/g
Uranium-238+D	1.70E+00					1.70E+00	1.70E+01	1.70E+02	1.70E+00	1.70E+01	1.70E+02	1.70E+00	1.70E+01	1.70E+02	1.70E+02	1.70E+02	pCi/g

Table 6.14. RGOs for Soil COCs of the BGOU SWMUs (Continued)

Excavation Worker Soil Exposure	COC ^A		Noncancer			RGO at			RGO at			RGO at			Units
	Cancer NAL	NAL	HI=0.1	HI=1	HI=3	ELCR=	ELCR=	ELCR=	ELCR=	ELCR=	ELCR=	ELCR=	ELCR=		
Aluminum		4.84E+03	4.84E+02	4.84E+03	1.45E+04									mg/kg	
Antimony		4.52E-01	4.52E-02	4.52E-01	1.36E+00									mg/kg	
Arsenic	3.13E-01	5.03E+00	5.03E-01	5.03E+00	1.51E+01	3.13E-01	3.13E+00	3.13E+00	3.13E+00	3.13E+01				mg/kg	
Barium		7.11E+02	7.11E+01	7.11E+02	2.13E+03									mg/kg	
Beryllium and compounds		1.15E+00	1.15E-01	1.15E+00	3.45E+00	3.83E-03	3.83E-02	3.83E-02	3.83E-01	3.83E-01				mg/kg	
Cadmium	2.12E+00	1.45E+01	1.45E+00	1.45E+01	4.35E+01	2.12E+00	2.12E+01	2.12E+01	2.12E+02	2.12E+02				mg/kg	
Chromium	2.85E+02	4.36E+02	4.36E+01	4.36E+02	1.31E+03	2.85E+02	2.85E+03	2.85E+03	2.85E+04	2.85E+04				mg/kg	
Cobalt	1.22E+03	3.11E+02	3.11E+01	3.11E+02	9.33E+02	1.22E+03	1.22E+04	1.22E+04	1.22E+05	1.22E+05				mg/kg	
Copper		4.37E+02	4.37E+01	4.37E+02	1.31E+03									mg/kg	
Iron		2.02E+03	2.02E+02	2.02E+03	6.06E+03									mg/kg	
Manganese		2.90E+02	2.90E+01	2.90E+02	8.70E+02									mg/kg	
Nickel	1.32E+04	2.05E+02	2.05E+01	2.05E+02	6.15E+02	1.32E+04	1.32E+05	1.32E+05	1.32E+06	1.32E+06				mg/kg	
Uranium		1.10E+01	1.10E+00	1.10E+01	3.30E+01									mg/kg	
Vanadium		4.03E+00	4.03E-01	4.03E+00	1.21E+01									mg/kg	
Zinc		2.50E+03	2.50E+02	2.50E+03	7.50E+03									mg/kg	
Aroclor 1260	1.55E-01					1.55E-01	1.55E+00	1.55E+00	1.55E+01	1.55E+01				mg/kg	
Benzo[a]anthracene	2.16E-01					2.16E-01	2.16E+00	2.16E+00	2.16E+01	2.16E+01				mg/kg	
Benzo[a]pyrene	2.16E-02					2.16E-02	2.16E-01	2.16E-01	2.16E+00	2.16E+00				mg/kg	
Benzo[b]fluoranthene	2.16E-01					2.16E-01	2.16E+00	2.16E+00	2.16E+01	2.16E+01				mg/kg	
Dibenz[a,h]anthracene	2.16E-02					2.16E-02	2.16E-01	2.16E-01	2.16E+00	2.16E+00				mg/kg	
Total Dioxins/Furans	1.79E-06					1.79E-06	1.79E-05	1.79E-05	1.79E-04	1.79E-04				mg/kg	
Indeno[1,2,3-cd]pyrene	2.16E-01					2.16E-01	2.16E+00	2.16E+00	2.16E+01	2.16E+01				mg/kg	
Total PCBs	1.48E-01					1.48E-01	1.48E+00	1.48E+00	1.48E+01	1.48E+01				mg/kg	
Total PAHs	2.16E-02					2.16E-02	2.16E-01	2.16E-01	2.16E+00	2.16E+00				mg/kg	
Neptunium-237+D	3.27E-01					3.27E-01	3.27E+00	3.27E+00	3.27E+01	3.27E+01				pCi/g	
Plutonium-239*	1.62E+00					1.62E+00	1.62E+01	1.62E+01	1.62E+02	1.62E+02				pCi/g	
Radium-226+D	3.30E-02					3.30E-02	3.30E-01	3.30E-01	3.30E+00	3.30E+00				pCi/g	
Uranium-234	2.83E+00					2.83E+00	2.83E+01	2.83E+01	2.83E+02	2.83E+02				pCi/g	
Uranium-235+D	4.55E-01					4.55E-01	4.55E+00	4.55E+00	4.55E+01	4.55E+01				pCi/g	
Uranium-238+D	1.17E+00					1.17E+00	1.17E+01	1.17E+01	1.17E+02	1.17E+02				pCi/g	

^A COC = contaminant of concern

^B RGO = remedial goal option. RGOs for soil for both HI and ELCR are calculated from the 2008 draft NALs (DOE 2008)

Table 6.15. RGOs for Groundwater COCs of the BGOU SWMUs

Residential User Groundwater Exposure												
COC ^A	EPC ^B	SWMU ^C	ELCR at EPC	HI at EPC	RGO ^D at HI=0.1	RGO at HI=1	RGO at HI=3	RGO at ELCR=1 x 10 ⁻⁶	RGO at ELCR=1 x 10 ⁻⁵	RGO at ELCR=1 x 10 ⁻⁴	MCL	mg/L
Antimony	7.99E-02	145	5.97E+00	1.34E-03	1.34E-02	1.34E-01	3.76E-05	3.76E-04	3.76E-03	0.006	0.006	mg/L
Arsenic	6.21E-02	145	1.99E+01	3.12E-04	3.12E-03	9.36E-03	3.76E-05	3.76E-04	3.76E-03	0.010	0.010	mg/L
Manganese	1.01E+00	5	2.15E+00	4.70E-02	4.70E-01	1.41E+00	---	---	---	---	---	mg/L
Selenium	1.51E-02	30	2.90E-01	5.21E-03	5.21E-02	1.56E-01	0.05	0.05	0.05	0.05	0.05	mg/L
Uranium	4.89E-02	3	7.82E+00	604	603	1.88E-02	0.03	0.03	0.03	0.03	0.03	mg/L
Aroclor-1254	5.23E-05	7	7.09E-06	1.25E-06	1.25E-05	3.74E-05	7.38E-06	7.38E-05	7.38E-04	0.0005	0.0005	mg/L
1,1-DCE	8.98E-02	7	8.51E-01	1.06E-02	1.06E-01	3.17E-01	4.32E-05	4.32E-04	4.32E-03	---	---	mg/L
cis-1,2-DCE	1.15E+01	2	6.07E+02	1.89E-03	1.89E-02	5.68E-02	0.07	0.07	0.07	0.07	0.07	mg/L
Naphthalene	5.55E-03	5	2.80E+00	1.98E-04	1.98E-03	5.95E-03	---	---	---	---	---	mg/L
TCE	1.18E+00	4	5.39E+02	2.19E-04	2.19E-03	6.57E-03	3.22E-05	3.22E-04	3.22E-03	0.005	0.005	mg/L
Vinyl Chloride	2.61E-02	4	1.21E+00	2.16E-03	2.16E-02	6.47E-02	1.58E-06	1.58E-05	1.58E-04	0.002	0.002	mg/L
Technetium-99	1.01E+04	145	5.54E-04	1.82E+01	1.82E+02	1.82E+03	900 ^E	900 ^E	900 ^E	900 ^E	900 ^E	pCi/L
Uranium-234	7.94E+00	7	1.11E-05	7.12E-01	7.12E+00	7.12E+01	20 ^F	20 ^F	20 ^F	20 ^F	20 ^F	pCi/L
Uranium-238	1.59E+01	3	2.76E-05	5.76E-01	5.76E+00	5.76E+01	20 ^F	20 ^F	20 ^F	20 ^F	20 ^F	pCi/L

^A COC = contaminant of concern
^B EPC = exposure point concentration; represents maximum EPC value for all SWMUs where constituent was a COC for the applicable scenario
^C SWMU = the SWMU associated with the maximum EPC value
^D RGO = remedial goal option
^E converted from MCL of 4 mrem/yr dose (DOE 2001)
^F converted from MCL for total uranium of 0.03 mg/L (DOE 2001)

6.7 SCREENING ECOLOGICAL RISK ASSESSMENT

Appendix G provides summaries of the results of ERAs previously completed for SWMUs 2, 4, 5, 6, 7, and 30 within the BGOU of PGDP (Figure G.1). SWMUs 3 and 145 are not included because SWMU 3 is covered by a RCRA cap and SWMU 145 is sited on 44 acres that now lie beneath the C-746-S&T Landfills. For most of the SWMUs, no new surface data have been collected since the previous risk assessments were performed. In addition, the soils at these units are outside the scope of the BGOU RI as defined in the approved work plan (DOE 2006a); therefore, a new quantitative risk assessment was not performed for soils. A summary of the results of the comparison in previous assessments of the site data to the ecological screening levels is provided in Table 6.16. This table lists the number of COPCs in each suite retained for each site and the medium for further consideration. This table shows that a number of inorganic analytes detected above background values and detected organic analytes were retained. Radionuclides were eliminated as COPCs for all sites except for SWMUs 7 and 30.

Table 6.16. Summary of Suite of COPCs Retained in Surface Soil

Area	Media	Metal	Rad	Pesticide/PCB	SVOC	VOC
SWMU 2	Soil	6	----	----	----	----
SWMU 3	Soil	NE	NE	NE	NE	NE
SWMU 4	Soil	5	----	1	----	----
SWMU 5	Soil	5	----	1	3	----
SWMU 6	Soil	2	----	----	1	----
SWMU 7	Soil	19	Total*	1	----	----
SWMU 30	Soil	17	Total*	1	----	----
SWMU 145	Soil	NE	NE	NE	NE	NE

----: no COPCs

NE: SWMU did not undergo an ecological evaluation.

*Radionuclide risk was assessed based on a total dose benchmark for all radionuclides.

SVOC=semivolatile organic compound

VOC=volatile organic compound

Each of the sites evaluated in the ERAs summarized in this section retained a number of COPCs as COPCs. Some metals at concentrations above background were retained as COPCs for ecological risk at each SWMU. Total PCBs were retained as COPCs for all SWMUs except SWMU 6. This is based on direct risk from soil as well as risks to some wildlife receptors from bioaccumulation through the food chain. The only other COPCs retained are three SVOCs (fluorene, phenanthrene, and di-n-butyl phthalate) at SWMU 5 and di-n-butyl phthalate at SWMU 6.

The current plan is to conduct further ecological risk assessment in future activities. In the absence of these activities, the benchmarks used in the screenings presented here and in the no further action levels the PGDP Ecological Risk Methods Document (DOE 2001) and the 2008 draft revision of the same document will be used to develop ecologically-based RGOs.

7. SUMMARY AND CONCLUSIONS

This chapter summarizes and presents conclusions about the nature and extent of contamination, fate and transport, and risk assessment at the eight burial grounds evaluated during this RI. The conclusions are drawn from known site conditions, historical knowledge of the burial grounds, and geological and environmental sampling data collected from the burial areas.

7.1 OVERVIEW

The PGDP SMP (DOE 2009) focuses environmental restoration activities into five strategic initiatives, as follows:

- BGOU Strategic Initiative,
- Decontamination and Decommissioning (D&D) OU Strategic Initiative,
- Groundwater OU Strategic Initiative,
- Soils OU Strategic Initiative, and
- Surface Water OU Strategic Initiative.

These initiatives include a series of prioritized response actions, ongoing site characterization activities to support future response action decisions, and D&D of the operating gaseous diffusion plant once it ceases operation. After completion of these activities, the Comprehensive Site OU evaluation will be conducted, with implementation of additional actions, as needed, to ensure long-term protectiveness.

7.2 RECOMMENDED REMEDIAL ACTION OBJECTIVES

General site cleanup objectives have been developed that serve as guiding principles for creating more detailed remedial action objectives (RAOs) to focus OUs on site-specific problems. A primary objective for the BGOU is to contribute to the protection of off-site residents by addressing sources of groundwater contamination. Based on the current and reasonably anticipated future land use, on-site industrial workers, recreational users, and off-site residents are the primary human receptors having the greatest potential for exposure to site contamination originating from PGDP. The primary pathways of exposure are (1) the groundwater pathway for off-site residents; (2) the surface water pathway (i.e., surface water and sediments) for recreational users (assumed to be primarily local residents); and (3) direct contact with waste, soil, and sediment for industrial workers. The following are the preliminary BGOU RAOs.

- Contribute to protection of current and future residential receptors from exposure to contaminated groundwater by addressing sources of groundwater contamination.
- Protect industrial workers from exposure to waste and contaminated soils.
- Treat or remove principal threat wastes wherever practicable, consistent with 40 *CFR* § 300.430 (a)(iii)(A).

These preliminary RAOs were developed to guide the evaluation of data in this RI. More specific and detailed RAOs, including SWMU-specific RAOs, will be developed to support remedy screening and evaluation in the FS.

The selected response actions for each OU must meet the Comprehensive Environmental Response, Compensation, and Liability Act (CERCLA) threshold criteria: 1) be protective of human health and the environment and 2) attain Applicable or Relevant and Appropriate Requirements (ARARs) [or provide grounds for invoking a waiver under CERCLA 121(d)(4)]. The NCP defines protectiveness in terms of risk-based levels and states that acceptable health-based exposure levels for known or suspected carcinogens are concentration levels that represent an excess upper-bound lifetime cancer risk between 10^{-4} to 10^{-6} . The NCP requires the 10^{-6} risk level be used as the point of departure for determining remediation goals for alternatives when ARARs are not available or are not sufficiently protective because of the presence of multiple contaminants at a site or multiple pathways of exposure. For systemic toxicants, EPA guidance defines a HI less than 1 as an acceptable health-based exposure level.

A summary of potential location-specific ARARs that have been identified for the BGOU is included as Appendix H of this document.

7.3 REMEDIAL INVESTIGATION SCOPE

The scope of the BGOU Strategic Initiative includes an RI, BHHRA, FS, remedy selection, and implementation of actions, as necessary, for protection of human health and the environment. This BGOU RI addresses eight SWMUs containing burial grounds and landfills at PGDP: seven (SWMUs 2, 3, 4, 5, 6, 7, and 30) are located within the main PGDP secure area; and one (SWMU 145) is located within a controlled access area to the north. Two other SWMUs in the BGOU are the C-746-S (SWMU 9) and C-746-T (SWMU 10) Landfills, which are closed landfills that were not included in this RI.

<u>SWMU</u>	<u>Facility</u>
2	C-749 Uranium Burial Ground
3	C-404 Low-Level Radioactive Waste Burial Ground
4	C-747 Contaminated Burial Yard and C-748-B Burial Area
5	C-746-F Burial Yard
6	C-747-B Burial Ground
7	C-747-A Burial Ground
30	C-747-A Burn Area
145	Area P

Ditches essentially bound each of the BGOU SWMUs to facilitate surface drainage. The nature and extent of contamination within these ditches is within the scope of the Surface Water OU Strategic Initiative.

The BGOU RI/FS Work Plan identified four primary goals for this RI and for the follow-up FS (DOE 2006a). Table 7.1 summarizes these goals and references sections of the RI report (where applicable) that address these goals. Both the historical data and new data collected during this RI were of sufficient quality to address the data needs identified during the DQO process.

The potential areas of buried metal within the C-746-P and C-746-P1 Scrap Yards (SWMU 13) identified during this BGOU RI field investigation will be characterized with a Field Sampling Plan addendum to the BGOU RI/FS Work Plan and follow-on field investigation. The results will be discussed with the FFA parties and, if further action is necessary, a path forward will be determined.

Table 7.1. Goals Identified for the BGOU RI

GOAL 1: CHARACTERIZE NATURE OF SOURCE ZONE

- 1-1: What are the suspected contaminants?
- 1-2: What are the plant processes that could have contributed to the contamination? When and over what duration did releases occur?
- 1-3: What are the concentrations and activities at the source?
- 1-4: What is the area and volume of the source zone?
- 1-5: What are the chemical and physical properties of associated material at the source areas?

GOAL 2: DEFINE EXTENT OF SOURCE ZONE AND CONTAMINATION IN SOIL AND OTHER SECONDARY SOURCES AT ALL UNITS

- 2-1: What are the past, current, and potential future migratory paths?
- 2-2: What are the past, current, and potential future release mechanisms?
- 2-3: What are the contaminant concentrations or activity gradients?
- 2-4: What is the vertical and lateral extent of contamination?
- 2-5: What is the relationship of the UCRS gradient to the source, to surface water bodies, and to the RGA?

GOAL 3: DETERMINE SURFACE AND SUBSURFACE TRANSPORT MECHANISMS AND PATHWAYS

- 3-1: What are the contaminant migration trends?
- 3-2: To what area is the dissolved-phase plume migrating?
- 3-3: What are the effects of underground utilities and plant operations on migration pathways including ditches?
- 3-4: What is the role of the UCRS in contaminant transport?
- 3-5: What are the physical and chemical properties of the formations and subsurface matrices?

GOAL 4: SUPPORT EVALUATION OF REMEDIAL ALTERNATIVES

- 4-1: What are the possible remedial technologies applicable for this unit?
 - 4-2: What are the physical and chemical properties of media to be remediated?
 - 4-3: Are cultural impediments present?
 - 4-4: What is the extent of contamination (geologic limitations presented by the source zone or secondary contamination source)?
 - 4-5: What would be the impact of action on and by other sources?
 - 4-6: What would the impact of an action at the source be on the integrator units?
 - 4-7: What are stakeholders' perceptions of contamination at or migrating from source zone or secondary contamination sources?
-

Table 7.1. Goals Identified for the BGOU RI (Continued)

GOAL 1: Characterize Nature of Source Zone

		SWMU	Location in Text
		SWMU 2	Section 4.3
1-1	What are the suspected contaminants?	SWMU 3	Section 4.4
		SWMU 4	Section 4.5
&		SWMU 5	Section 4.6
		SWMU 6	Section 4.7
1-3	What are the concentrations and activities at the source?	SWMU 7	Section 4.8
		SWMU 30	Section 4.9
		SWMU 145	Section 4.10
		SWMU 2	Section 1.3.1.2
1-2	What are the plant processes that could have contributed to the contamination?	SWMU 3	Section 1.3.2.2
		SWMU 4	Section 1.3.3.2
	When and over what duration did releases occur?	SWMU 5	Section 1.3.4.2
&		SWMU 6	Section 1.3.5.2
		SWMU 7	Section 1.3.6.2
1-5	What are the chemical and physical properties of associated material at the source areas?	SWMU 30	Section 1.3.7.2
		SWMU 145	Section 1.3.8.2
		SWMU 2	Section 1.3.1.1 and Appendix E, Table E.3.3
		SWMU 3	Section 1.3.2.1 and Appendix E, Table E.3.7
		SWMU 4	Section 1.3.3.1 and Appendix E, Table E.3.11
		SWMU 5	Section 1.3.4.1 and Appendix E, Table E.3.15
1-4	What is the area and volume of the source zone?	SWMU 6	Section 1.3.5.1 and Appendix E, Table E.3.19
		SWMU 7	Section 1.3.6.1 and Appendix E, Table E.3.21
		SWMU 30	Section 1.3.7.1 and Appendix E, Table E.3.25
		SWMU 145	Section 1.3.8.1 and Appendix E, Table E.3.29

Table 7.1. Goals Identified for the BGOU RI (Continued)

GOAL 2: Define Extent of Source Zone and Contamination in Soil and Other Secondary Sources at All Units

		SWMU	Location in Text
2-1	What are the past, current, and potential future migratory paths?	All SWMUs	Section 5.1 and Figure 5.1
2-2	What are the past, current, and potential future release mechanisms?	All SWMUs	Section 4
2-3	What are the contaminant concentrations or activity gradients?	SWMU 2	Section 4.3
		SWMU 3	Section 4.4
		SWMU 4	Section 4.5
		SWMU 5	Section 4.6
		SWMU 6	Section 4.7
		SWMU 7	Section 4.8
		SWMU 30 SWMU 145	Section 4.9 Section 4.10
2-4	What is the vertical and lateral extent of contamination?	SWMU 2	Table 4.8, Appendix D, SWMU 2
		SWMU 3	Table 4.14, Appendix D, SWMU 3
		SWMU 4	Table 4.20, Appendix D, SWMU 4
		SWMU 5	Table 4.25, Appendix D, SWMU 5
		SWMU 6	Table 4.30, Appendix D, SWMU 6
		SWMU 7	Table 4.35, Appendix D, SWMU 7
2-5	What is the relationship of the UCRS gradient to the source, to surface water bodies, and to the RGA?	SWMU 30	Table 4.41, Appendix D, SWMU 30
		SWMU 145	Table 4.47, Appendix D, SWMU 145
		SWMU 2	Section 3.9.3.1
		SWMU 3	Section 3.9.3.1
		SWMU 4	Section 3.9.3.2
		SWMU 5	Section 3.9.3.3
		SWMU 6	Section 3.9.3.3
SWMU 7	Section 3.9.3.4		
SWMU 30	Section 3.9.3.4		
SWMU 145	Section 3.9.3.5		

Table 7.1. Goals Identified for the BGOU RI (Continued)

GOAL 3: Determine Surface and Subsurface Transport Mechanisms and Pathways

		SWMU	Location in Text
		SWMU 2	Section 5.3.1
3-1	What are the contaminant migration trends?	SWMU 3	Section 5.3.2
		SWMU 4	Section 5.3.3
&		SWMU 5	Section 5.3.4
		SWMU 6	Section 5.3.5
3-2	To what area is the dissolved-phase plume migrating?	SWMU 7	Section 5.3.6
		SWMU 30	Section 5.3.7
		SWMU 145	Section 5.3.8
3-3	What are the effects of underground utilities and plant operations on migration pathways including ditches?	All SWMUs	Section 3.9.2
		SWMU 2	Section 4.3.2
		SWMU 3	Section 4.4.2
		SWMU 4	Section 4.5.2
3-4	What is the role of the UCRS in contaminant transport?	SWMU 5	Section 4.6.2
		SWMU 6	Section 4.7.2
		SWMU 7	Section 4.8.2
		SWMU 30	Section 4.9.2
		SWMU 145	Section 4.10.2
3-5	What are the physical and chemical properties of the formations and subsurface matrices?	All SWMUs	Section 3.9.3

Table 7.1. Goals Identified for the BGOU RI (Continued)

GOAL 4: Support Evaluation of Remedial Alternatives

		SWMU	Location in Text
4-1	What are the possible remedial technologies applicable for this unit?	All SWMUs	Tables 7.9 and 7.10, To be evaluated in FS
4-2	What are the physical and chemical properties of media to be remediated?	All SWMUs	Sections 1.3 and 3.9.3
4-3	Are cultural impediments present?	All SWMUs	Section 3.9.1. To be further evaluated in FS
4-4	What is the extent of contamination (geologic limitations presented by the source zone or secondary contamination source)?	SWMU 2	Section 3.9.3.1
		SWMU 3	Section 3.9.3.1
		SWMU 4	Section 3.9.3.2
		SWMU 5	Section 3.9.3.3
		SWMU 6	Section 3.9.3.3
		SWMU 7	Section 3.9.3.4
		SWMU 30 SWMU 145	Section 3.9.3.4 Section 3.9.3.5
4-5	What would be the impact of action on and by other sources?	All SWMUs	To be evaluated in FS
4-6	What would the impact of an action at the source on the integrator units?	All SWMUs	To be evaluated in FS
4-7	What are stakeholders' perceptions of contamination at or migrating from source zone or secondary contamination sources?	All SWMUs	To be evaluated in FS

7.4 NATURE AND EXTENT OF CONTAMINATION

Materials that were disposed of in each of the SWMUs of the BGOU contained hazardous substances. The conceptual model applicable to all of the BGOU SWMUs is that releases from these SWMUs have impacted soils below or adjacent to the source zones and, through vertical infiltration in the soil, have the potential to contaminate the groundwater underlying these sources. Analysis of soil and groundwater from the area of each SWMU documents the presence of metals, organic compounds, and radionuclides above screening levels. Section 4 summarizes the characterization of the area of these SWMUs, as part of the BGOU RI and previous investigation efforts.

Principal threat waste (PTW) is defined by EPA as “source materials considered to be highly toxic or highly mobile that generally cannot be reliably contained or would present a significant risk to human health or the environment should exposure occur” (EPA 1991). EPA also recognizes that “although no threshold level of risk has been established to identify principal threat waste, a general rule of thumb is to

consider as a principal threat those source materials with toxicity and mobility characteristics that combine to pose a potential risk several orders of magnitude greater than the risk level that is acceptable for the current or reasonably anticipated future land use, given realistic exposure scenarios” (EPA 1997).

For the BGOU SWMUs, the TCE DNAPL at SWMUs 4, 7, and 30 and uranium at SWMUs 2 and 3 potentially are PTW. Dissolved contaminant trends in the RGA indicate that SWMU 4 and the adjoining areas of SWMUs 7 and 30 could contain TCE sources as DNAPL. The mobility and toxicity of DNAPLs make them PTW.

The uranium at SWMU 2 presents risk greater than 1E-03 under some hypothetical exposure scenarios. Some forms of the buried uranium could be considered potential PTW where toxicity and mobility combine to pose such a risk to human health. These hypothetical exposure scenarios assume a direct contact exposure to buried waste (DOE 1997a). The uranium metal present at SWMUs 2 and 3 likely is not mobile due to its insolubility in water. The Data Summary and Interpretation Report (DOE 1997a) concluded that only some forms of uranium present may be mobile (e.g., uranyl fluoride at SWMU 2). Uncertainties concerning the risks associated with the toxicity and mobility of the uranium will be considered further during alternative evaluation in the FS.

Several analytes in subsurface soils (soils deeper than 1 ft) were frequently detected above screening criteria used to identify contaminants for the assessment of nature and extent. Table 7.2 indicates the frequency of detection above the excavation worker NALs (DOE 2001). All inorganic and radiological values in the table also correspond to at least one detection above background. As shown in Table 7.2, aluminum, iron, manganese, and vanadium are the most prevalent of the frequently detected contaminants in subsurface soils, often detected above screening criteria in more than 50% of samples representative of the nature and extent of contamination in six of the eight BGOU SWMUs. Uranium-238 was detected above screening criteria at all SWMUs except SWMU 6.

**Table 7.2. Frequency of Subsurface Soil Analytes Detected above No Action Levels
(for Analytes Detected above Background)**

Analyte	SWMU 2	SWMU 3	SWMU 4	SWMU 5	SWMU 6	SWMU 7	SWMU 30	SWMU 145
<i>Inorganics</i>								
Aluminum			94%	83%	79%	61%	88%	84%
Antimony		8%						33%
Arsenic	97%	90%	10%			72%		61%
Barium	3%		<1%	2%				2%
Beryllium	3%		5%	14%	17%	1%	8%	8%
Iron	100%		100%	97%	98%	94%	100%	
Manganese	86%		73%	75%	91%	64%	72%	
Thallium	24%							
Uranium	12%	10%				1%		13%
Vanadium	93%		99%	100%	98%		68%	
<i>Organics</i>								
Total PCB	4%		7%			3%	4%	11%
Total PAH							4%	
TCE	2%		3%					
1,1-DCE						3%		
Cis-1,2-DCE	3%							
Vinyl Chloride	3%		1%			1%		
<i>Radionuclides</i>								
Uranium-234	2%	3%	57%			11%	14%	19%
Uranium-235			<1%					5%

Table 7.2. Frequency of Subsurface Soil Analytes Detected above No Action Levels (Continued)

Analyte	SWMU 2	SWMU 3	SWMU 4	SWMU 5	SWMU 6	SWMU 7	SWMU 30	SWMU 145
Uranium-235/236	2%					38%	20%	
Uranium-238	21%	15%	61%	21%		29%	21%	26%
Technetium-99			1%					8%
Cesium-137		3%	1%					11%
Thorium-228								85%
Thorium-230			50%			3%		9%
Thorium-232								3%
Americium-241								2%

The "no action levels" are for the Excavation Worker scenario (DOE 2001).

Background values are from the Risk Methods Document (DOE 2001).

Blanks represent analytes that (1) did not exceed NAL or (2) exceeded no action level (NAL), but not background.

Metals are the most common of the frequently detected (for groundwater, frequently detected refers to inorganic or radiological analytes detected in 25% or more samples above screening criteria; all organic detections are listed in the tables) contaminants in both UCRS and RGA groundwater samples (Tables 7.3 and 7.4). Iron and manganese are the predominant contaminants in the UCRS. Iron is less prevalent in the RGA. In both Table 7.3 and 7.4, all organic analytes are listed if they were detected.

Table 7.3. Metals and Radionuclides Frequently Detected above Screening Levels and All Organic Compounds Detected in UCRS Groundwater

Source Area	Metals	Organic Compounds	Radionuclides
SWMU 2	Beryllium, Iron, Manganese, Uranium, Vanadium	1,1-DCE; TCE: <i>cis</i> -1,2-DCE; Vinyl chloride	²³⁴ U, ²³⁸ U
SWMU 3	Arsenic, Iron, Manganese, Molybdenum	TCE	⁹⁹ Tc, ²³⁴ U, ²³⁸ U
SWMU 4	Arsenic, Beryllium, Cadmium, Chromium, Copper, Iron, Lead, Manganese, Nickel, Vanadium, Zinc	<i>cis</i> -1,2-DCE; TCE; 1,1-DCE; Vinyl chloride; <i>trans</i> -1,2-DCE; Naphthalene; PCB	⁹⁹ Tc
SWMU 5	Arsenic, Beryllium, Chromium, Copper, Iron, Lead, Manganese, Molybdenum, Nickel, Vanadium, Zinc	TCE; Pyrene	--
SWMU 6	Arsenic, Chromium, Iron, Lead, Manganese, Molybdenum, Nickel, Uranium, Vanadium, Zinc	PCB-1016	⁹⁹ Tc, ²³⁴ U, ²³⁸ U
SWMU 7	Arsenic, Beryllium, Cadmium, Chromium, Iron, Lead, Manganese, Molybdenum, Nickel, Vanadium, Zinc	<i>cis</i> -1,2-DCE; TCE; Vinyl chloride ; 1,1-DCE ; Benzene ; Naphthalene	²²² Rn, ²³⁴ U, ²³⁸ U
SWMU 30	Arsenic, Cadmium, Iron, Lead, Manganese, Molybdenum, Nickel, Uranium, Vanadium	TCE ; Vinyl chloride; Benzene; Naphthalene; PCB-1260	²³⁴ U, ²³⁸ U
SWMU 145	Arsenic, Iron, Manganese, Uranium	Chloroform	²²² Rn, ²³⁴ U, ²³⁸ U

-- = none

DCE = dichloroethene

²²²Rn = radon-222

TCE = trichloroethene

⁹⁹Tc = technetium-99

²³⁴U = uranium-234

²³⁸U = uranium-238

PCB = polychlorinated biphenyl

Table 7.4. Metals and Radionuclides Frequently Detected above Screening Levels and All Organic Compounds Detected in RGA Groundwater

Source Areas	Metals	Organic Compounds	Radionuclides
SWMU 2	Arsenic, Beryllium, Cadmium, Iron, Manganese, Vanadium	1,1-DCE; TCE; <i>cis</i> -1,2-DCE; Chloroform	²³⁴ U, ²³⁸ U
SWMU 3	Arsenic, Iron, Manganese	TCE; 1,1-DCE; Chloroform	--
SWMU 4	Arsenic, Manganese, Iron, Lead	1,1-DCE; Carbon Tetrachloride; Chloroform; <i>cis</i> -1,2-DCE; TCE; Vinyl Chloride	--
SWMU 5	Iron, Lead, Manganese	TCE	--
SWMU 6	Arsenic, Iron, Lead, Manganese	TCE	--
SWMU 7	Arsenic, Beryllium, Cadmium, Iron, Lead, Manganese, Molybdenum, Nickel, Vanadium, Zinc	TCE; <i>cis</i> -1,2-DCE; Vinyl chloride; Chloroform; Carbon tetrachloride	⁹⁹ Tc, ²³⁴ U, ²³⁸ U
SWMU 30	Arsenic, Iron, Manganese	TCE, Tetrachloroethene; Chloroform	²²² Rn, ⁹⁹ Tc
SWMU 145	Arsenic, Iron, Manganese, Nickel, Vanadium	TCE; Chloroform; PCB	--

-- = none

DCE = dichloroethene

²²²Rn = radon-222

TCE = trichloroethene

⁹⁹Tc = technetium-99

²³⁴U = uranium-234

²³⁸U = uranium-238

PCB = polychlorinated biphenyl

7.5 FATE AND TRANSPORT

Modeling assessed fate and transport of contaminants for two pathways: (1) dissolved-phase transport through the aquifer and (2) vapor transport to a residential basement. Section 5 and Appendix E document the fate and transport modeling applied to the BGOU RI.

Previous work has shown that the primary pathway for groundwater flow and the site-related contaminants is predominantly vertical migration through the UCRS, followed by lateral migration in the RGA. Contaminated groundwater could migrate to the POEs. The POEs evaluated were at the SWMU, at the plant boundary, at the property boundary, and near the Ohio River (either at the Little Bayou Creek seeps or at the Ohio River, depending on modeled groundwater flow paths beginning at each SWMU). Not all SWMUs have transport pathways to all of the POEs. For example, SWMU 145 is located outside of the plant boundary and does not contribute to the Little Bayou seeps. SWMUs 3, 6, 7, and 30 were determined to be the only SWMUs contributing to the Little Bayou seeps POE. While there is some uncertainty related to modeling in predicting whether a SWMU would contribute to the Little Bayou seeps, this uncertainty should not affect or drive remedial decisions. Table 7.5 identifies analytes that were modeled to exceed MCLs at the POEs.

Table 7.5. Analytes Predicted to Exceed Maximum Contaminant Levels at the Points of Exposure

Source Area	Contaminant	SWMU Boundary	Plant Boundary	Property Boundary	Little Bayou seeps	Ohio River
SWMU 2	Arsenic	Yes ^a	No ^b	No	N/A ^c	No
	<i>cis</i> -1,2-DCE	Yes	Yes	Yes	N/A	Yes
	TCE	Yes	Yes	Yes	N/A	Yes
SWMU 3	Arsenic	Yes	No	No	No	N/A
	⁹⁹ Tc	Yes	Yes	Yes	No	N/A
	Uranium	Yes	No	No	No	N/A
SWMU 4	Arsenic	Yes	No	No	N/A	No
	<i>cis</i> -1,2-DCE	Yes	Yes	Yes	N/A	No
	⁹⁹ Tc	Yes	Yes	Yes	N/A	No
	TCE	Yes	Yes	Yes	N/A	Yes
	Vinyl Chloride	Yes	Yes	Yes	N/A	No
SWMU 5	No analytes predicted to exceed MCLs at POEs					
SWMU 6	No analytes predicted to exceed MCLs at POEs					
SWMU 7	1,1-DCE	Yes	Yes	No	No	N/A
	Arsenic	Yes	Yes	No	No	N/A
	⁹⁹ Tc	Yes	No	No	No	N/A
	TCE	Yes	Yes	No	No	N/A
	Vinyl Chloride	Yes	Yes	No	No	N/A
SWMU 30	Arsenic	Yes	Yes	No	No	N/A
SWMU 145	Antimony	Yes	N/A	No	N/A	No
	Arsenic	Yes	N/A	No	N/A	No
	⁹⁹ Tc	Yes	N/A	Yes	N/A	Yes

^a Yes = The modeled analyte concentration exceeds its MCL

^b No = The modeled analyte concentration does not exceed its MCL

^c N/A = The POE does not apply to the SWMU

DCE = dichloroethene

MCL = maximum contaminant level

POE = point of exposure

SWMU = solid waste management unit

TCE = trichloroethene

⁹⁹Tc = technetium-99

Vapor transport modeling assessed contaminant concentrations in a hypothetical residential basement at the SWMU and in hypothetical residential basements at the POEs. (Appendix E, Section E.3.2 documents the vapor transport modeling performed for the BGOU RI.) The resident scenario provides bounding risks and hazards for the vapor pathways when compared to the industrial worker exposure scenario. Hence, the industrial worker exposure scenario was not evaluated in the vapor modeling analysis. Table 7.6 summarizes the results of vapor transport modeling. At some POEs, the excess lifetime cancer risk (ELCR) or hazard posed by hypothetical exposure to the modeled air concentration exceeded 1E-06 or 0.1, respectively. Currently there are no buildings, or pipelines connected to buildings, located over the contaminated material at the BGOU SWMUs. The existing buildings at PGDP are slated to be decontaminated and demolished after the plant closes; therefore, the on-site industrial worker scenario was not evaluated. There are no plans to construct future facilities on the BGOU SWMUs.

Table 7.6. Analytes with Basement Air Concentrations of Concern Based on Vapor Transport Modeling Results at the Points of Exposure

Source Area	Contaminant	SWMU Boundary	Plant Boundary	Property Boundary
SWMU 2	TCE	Yes ^a	Yes	Yes
	<i>cis</i> -1,2-DCE	Yes	No ^b	No
SWMU 3	TCE	Yes	No	No
	Mercury ^c	Yes	No	No
SWMU 4	TCE	Yes	Yes	Yes
	Vinyl Chloride	Yes	Yes	No
	<i>cis</i> -1,2-DCE	Yes	No	No
SWMU 5	No analytes with air concentrations of concern			
SWMU 6	No analytes with air concentrations of concern			
SWMU 7	TCE	Yes	No	No
	Vinyl Chloride	Yes	No	No
	1,1-DCE	Yes	Yes	No
	Mercury	Yes	No	No
SWMU 30	TCE	Yes	No	No
	1,1-DCE	Yes	No	No
	Mercury	Yes	No	No
SWMU 145	Mercury	Yes	No	No

DCE = dichloroethene

SWMU = solid waste management unit

TCE = trichloroethene

^a Yes = Modeled air concentration equals or exceeds 1.0E-06 excess lifetime cancer risk (ELCR) or 0.1 hazard quotient (HQ)

^b No = Modeled air concentration is less than 1.0E-06 ELCR or 0.1 HQ

^c The vapor transport modeling for mercury was conservatively based on the metallic form, which has a Henry's Law Constant of 1.07E-02 atm-m³/mol. The rate of vaporization of mercury and certain of its inorganic compounds decrease in the sequence Hg > Hg₂Cl₂ > HgCl₂ > HgS > HgO. The Henry's Law Constant decreases dramatically down the sequence (for example, HgCl₂ has a value of 7.09E-10 atm-m³/mol).

Vapor transport modeling conducted with the Johnson and Ettinger model (1991), coded into spreadsheets by EPA (2004).

7.6 RISK ASSESSMENT

PGDP is an industrial facility. The reasonably anticipated future use of the area within the current plant boundary is expected to remain industrial. This expectation should be considered when using the risk information provided in the report to support risk management decision making.

For soil, results from previous risk assessments were used. The BGOU Work Plan did not call for additional surface (0-1 ft bgs) or subsurface (0-10 ft bgs) soil sampling at most SWMUs. The risk for the on-site resident for soil exceeds 1E-04 and the HI is greater than 1 at all SWMUs except for SWMU 2, 3, and SWMU 145 (which were not evaluated for soil exposure for this scenario). The contaminants that are risk drivers for soil are aluminum, arsenic, beryllium, chromium, iron, nickel, Total PAHs, uranium (as a metal), uranium-234, uranium-238, vanadium, and zinc.

Residential use of groundwater was evaluated at the SWMU boundary, plant boundary, property boundary, and Ohio River (or seeps) for all SWMUs except SWMU 6 (SWMU 6 had no groundwater COPCs) and SWMU 145 (SWMU 145 was not evaluated at the plant boundary since it lies outside that boundary). At the SWMU boundary, risks and hazards from groundwater use for all evaluated SWMUs exceeded 1E-04 risk and exceeded an HI of 1. The major contaminants driving the groundwater risks and hazards at SWMU boundary point of exposure are arsenic (at SWMUs 3, 5, 7, and 145); antimony (at SWMU 145); Aroclor-1260 (at SWMU 145); *cis*-1,2-DCE (at SWMUs 2 and 7); 1-1-DCE (at SWMUs 7 and 30); manganese (at SWMUs 3 and 5); naphthalene (at SWMU 5); Aroclor-1254 (at SWMU 7); TCE (at SWMUs 2, 4, 7, and 30); technetium-99 (at SWMU 3); uranium (at SWMU 3); and vinyl chloride (at

SWMUs 4 and 7). At the plant boundary, risks and hazards from groundwater for SWMUs 2, 3, 4, 5, 7, and 30 exceeded 1E-04 risk or exceeded an HI of 1. At the property boundary, risks and hazards from groundwater for SWMUs 2, 4, 7, 30, and 145 exceeded 1E-04 risk or exceeded an HI of 1. At the Ohio River (or seeps), risks and hazards from groundwater for SWMUs 2, 4, 7, and 30 exceeded 1E-04 risk or exceeded an HI of 1. The major contaminants driving the groundwater risks and hazards at the property boundary and Ohio River (or Little Bayou Creek seeps) POEs are arsenic, *cis*-1,2-DCE, 1,1-DCE, TCE, technetium-99, and vinyl chloride. While the migration of contamination from the potential TCE DNAPL zones at SWMU 4 and SWMUs 7 and 30 were not modeled due to uncertainties in source term development, a qualitative analysis completed considering results from previous studies done for the PGDP (e.g., C-400 DNAPL source) indicates that TCE migration from these sources would have resulted in potential risks exceeding 1E-04 at all POEs.

For exposure to soil, at least one of the on-site receptor scenarios (industrial worker, excavation worker, or recreational user), all SWMUs (except SWMU 145 where these scenarios were not assessed) have an ELCR $\geq 1.0E-06$. For at least one of these scenarios, SWMUs 4, 5, 6, 7, and 30 have HIs > 1 . Soil exposures to industrial or excavation workers are more relevant to the potential future uses of the site.

For the excavation worker who is exposed to both surface soil and subsurface soil (soil from 0 to 10 ft bgs), HIs were greater than one at SWMUs 4, 5, 6, 7, and 30. Risks for the excavation worker exceeded 1E-04 at SWMUs 4, 5, 6, 7, and 30. The risk/hazard drivers for the excavation worker scenario were arsenic, beryllium, Total PAHs, uranium, uranium-235, and uranium-238.

Given the reasonably anticipated future industrial use of the areas within the plant boundary, the most likely future and current scenario is the industrial worker. The ELCR for the future industrial worker scenario exceeded 1E-04 at SWMUs 2, 3, 4, 5, 6, 7, and 30 primarily due to risk from arsenic, beryllium, Total PAHs, uranium-235, and uranium-238. The HI exceeds 1 for the industrial worker at SWMUs 4, 7, and 30; aluminum, beryllium, chromium, iron, manganese, uranium, and vanadium are the hazard drivers. Risks for the current industrial worker (at 16 days per year of exposure) were less than those for the future industrial worker; risks for the current industrial worker exceeded 1E-04 at SWMUs 4, 5, 6, 7, and 30.

Table 7.7 presents a summary of the dominant exposure pathways and COCs for each SWMU for exposure to subsurface soil and groundwater.

The inclusion of beryllium as a risk driver is a result of incorporating the historical risk assessments. At the time those risk assessments were developed, beryllium still was evaluated as a carcinogen. Since then, the oral cancer slope factor for beryllium has been withdrawn and no longer is used for PGDP risk assessments. As a result, the total ELCR becomes much lower at those SWMUs where beryllium is a contaminant of concern. For SWMUs 4 and 6, removal of the contribution of beryllium to the ELCR reduces the total ELCR to within the EPA risk range for the industrial worker scenario.

Table 7.7. Exposure Routes and Exposure Pathways and COCs Associated with Dominant Risk for Each SWMU

Source Area	HI	ELCR
SWMU 2	<ul style="list-style-type: none"> Ingestion of groundwater and household inhalation of vapors (TCE; <i>cis</i>-1,2-DCE) 	<ul style="list-style-type: none"> Household inhalation of vapors (TCE) Ingestion of groundwater (TCE) External exposure to subsurface soil (uranium-235, uranium-238)
SWMU 3	<ul style="list-style-type: none"> Ingestion of groundwater (arsenic, uranium) 	<ul style="list-style-type: none"> Ingestion of groundwater (arsenic, technetium-99) External exposure to subsurface soil (uranium-235, uranium-238)
SWMU 4	<ul style="list-style-type: none"> Ingestion of groundwater (TCE) Dermal exposure to soil (chromium, iron) 	<ul style="list-style-type: none"> Household inhalation of vapors and dermal exposure (TCE, vinyl chloride) Dermal exposure to subsurface soil (beryllium)
SWMU 5	<ul style="list-style-type: none"> Ingestion of RGA groundwater (arsenic, naphthalene) Ingestion of vegetables (arsenic, aluminum) 	<ul style="list-style-type: none"> Ingestion of RGA groundwater (arsenic)
SWMU 6	<ul style="list-style-type: none"> Ingestion of subsurface soil (chromium) Dermal exposure to soil (chromium) 	<ul style="list-style-type: none"> Dermal exposure to subsurface soil (PAHs, beryllium) Ingestion of vegetables (PAHs, beryllium)
SWMU 7	<ul style="list-style-type: none"> Ingestion of RGA groundwater (TCE, arsenic, Total PCBs) Ingestion of vegetables (iron, uranium) Dermal exposure to soil (vanadium, iron, uranium) 	<ul style="list-style-type: none"> Household inhalation of vapors and ingestion of RGA groundwater (1,1-DCE) Dermal exposure and ingestion of vegetables (beryllium, uranium-238)
SWMU 30	<ul style="list-style-type: none"> Ingestion of RGA groundwater (TCE) Ingestion of subsurface soil (uranium) Dermal exposure to soil (vanadium, iron) 	<ul style="list-style-type: none"> Household inhalation of vapors (TCE) Ingestion of vegetables (beryllium, uranium-238)
SWMU 145	<ul style="list-style-type: none"> Ingestion of RGA groundwater (antimony, arsenic) 	<ul style="list-style-type: none"> Ingestion of RGA groundwater (Aroclor-1260)

DCE = dichloroethene
PCB = polychlorinated biphenyl
TCE = trichloroethene

PAH = polycyclic aromatic hydrocarbon
RGA = Regional Gravel Aquifer

The BGOU RI includes a summary of previous ecological risk assessments for SWMUs 2, 4, 5, 6, 7, and 30. Neither SWMU 3 nor SWMU 145 has been assessed for ecological risk. SWMU 3 is covered by a RCRA cap, and SWMU 145 is located beneath the C-746-S & -T Landfills, which also are covered by caps. Comparison of site characterization data against No Action screening levels determined that all of the SWMUs have metals and organic compounds (in surface soil) that are COPCs for ecological risk to the environment, while SWMUs 7 and 30 have a radionuclide COPC (in surface soil).

7.7 UNCERTAINTIES/ASSUMPTIONS

The BGOU Work Plan identified data gaps on a SWMU-by-SWMU basis that were necessary to be filled in order to move forward with the FS. The Work Plan was implemented to reduce any remaining uncertainties from previous investigations regarding the nature of the source zone, extent of the source zone and secondary sources, surface and subsurface transport mechanisms, and to support evaluation of remedial technologies in the FS.

Nature of the Source Zone

A key project assumption for the upcoming FS is that the available historical documentation and soil and groundwater characterization data is sufficient relative to waste characteristics, to chemical and physical properties, and to waste volume estimates to evaluate general response actions, to screen technology types, and to conduct detailed alternative analysis for the BGOU. The potential impact of source zone uncertainties on alternatives analysis will be further documented and analyzed in the FS. While the BGOU RI field investigation sampled directly beneath the waste units using angled borings, it remains possible that the buried waste contains hazards or constituents that current sample results do not characterize (historical disposal records and waste manifests are incomplete). Additional uncertainty exists for SWMU 5 because wastes from other facilities were buried on-site through a “Work for Others” program. These waste streams may be unrelated to the typical materials used in gaseous diffusion. A related uncertainty is that the field investigation was unable to sample to the middle of a few of the larger SWMUs (SWMUs 5 and 145, particularly); therefore, there are some uncertainties in the nature and extent of the contaminant source that will need to be managed during the decision-making process.

Aerial photographs were reviewed to help with locating the extent of burial pits or past activities associated with the BGOU SWMUs; however, there is some uncertainty with this approach since aerial photographs are for sporadic time periods and there were several years where photographs are not available to view to aid with delineating the extent of each burial ground.

Many of the SWMUs have been investigated previously. The BGOU RI uses a combination of historical and current sample results of soil and groundwater from the area of each SWMU. The results of previous investigations and RI sampling document and confirm the presence of metals, organic compounds, and radionuclides in the BGOU burial grounds. The associated samples were collected and analyzed over several previous and continuing investigations, as well as the BGOU RI, using several methods. QA/QC practices at PGDP, now and previously, limit the uncertainty associated with the sampling and analysis process. Nevertheless, changes have occurred to analytical methods that limit the strict comparison of data (e.g., laboratory reporting limits have varied over time). In some cases, analytical method detection limits are above screening criteria, such as the Excavation Worker No Action Level; these data are of limited value to the RI. One potential change with large impact to the interpretation of data is the preservation of groundwater samples. Both filtered and unfiltered samples were collected in the field as “unfiltered,” with one of the fractions (the sample preserved for unfiltered analyses) being acidified in the field. The other sample was filtered and acidified in the laboratory. The unfiltered analyses represent the “whole water” chemistry (including both the dissolved and suspended fractions). These analyses may be significantly biased by turbid samples derived from temporary borings and likely would lead to an overestimation of soluble transportable contaminants. Groundwater metals and radionuclide analyses would be most affected.

To minimize the potential for “age” to bias the analysis of the data, the historical sample analyses used in the BGOU RI are limited to groundwater samples collected in January 1995 and later and soil samples collected in June 1996 and later. This criterion, which was established during scoping for the BGOU RI and is documented in Section 5 of the BGOU RI Work Plan, maximizes the number of historical sample

analyses available to the RI, while providing a reasonable assurance of the comparability of the data. There are limited monitoring wells in close proximity to many of the SWMUs, but temporary borings provide a snapshot of the conditions where groundwater samples could be obtained. Permanent MWs would reduce the uncertainties associated with seasonal variations in water levels and contaminant trends.

Where changing conditions are present, the earlier data will be less representative of current conditions. Factors in the UCRS that might cause significant change include the degradation of organic wastes or the episodic breach of a waste container. An additional factor that might impact conditions in the RGA is an advancing contaminant plume from an upgradient source. The increasing TCE level in the RGA at SWMUs 2 and 3 is an example of conditions in the RGA being impacted by an upgradient source.

The potential for acidic leachate at each SWMU is uncertain due to the lack of disposal records. SWMUs with the greatest potential for acidic leachate are SWMU 6 (exhaust fans with perchloric acid) and SWMU 4 (records of chemicals buried are incomplete). It should be noted that angled borings beneath SWMU 6 found no evidence of acidic leachate, either from subsurface metal concentrations or groundwater pH. There are no pH measurements from shallow groundwater at SWMU 4 to allow an evaluation of the uncertainty related to acidic leachate. The potential for acid leaching at the SWMUs will be evaluated further relative to the importance of acid leaching in screening, and detailed analysis of alternatives in the FS.

For SWMUs 2 and 6, where the last disposal occurred in 1977 (32 years ago) and 1976 (33 years ago), respectively, it is reasonable to assume most, if not all, drums have failed (an Oak Ridge National Laboratory researcher estimated that failure of steel drums would be expected to occur within 18 to 36 years). At SWMU 5, where the last disposal occurred 22 years ago, it is reasonable to assume some drums still may be intact. For SWMUs 4, 7, 30, and 145, it can be assumed that drums likely are breached since they were dumped rather than having been carefully stacked. Because all drummed waste was assumed to have been released to the environment during disposal or through degradation, samples from soils surrounding the buried wastes were used to evaluate potential contaminant migration and risks associated with the SWMUs. This approach resulted in the inclusion of SWMUs with drummed waste exhibiting risk and hazard values that exceeded acceptable levels; therefore, though the integrity of buried drums is an uncertainty, the overall objectives of the RI analysis were met. The risk assessment concluded that these uncertainties, related to the source zone, were not estimated to have a large effect on the risk characterization and does not affect the need for taking action.

The BGOU FS may identify the need for remedial design support or additional source delineation after final selection of media-specific remedial goals and remedial alternatives. In addition, monitoring of the source zones will be evaluated in the FS.

Extent of the source zone and secondary sources

Secondary sources of groundwater contamination that are derived from the BGOU burial grounds, such as the potential DNAPL source zones beneath SWMU 4 and SWMU 7, are within the scope of the BGOU for evaluation and remedial action. The evidence for UCRS DNAPL presence is documented in previous investigations (DOE 2007a; DOE 1998a) and discussed in the RI. Collection of UCRS groundwater samples was attempted from 32 angled borings in order to detect releases or secondary sources that may be related to the SWMUs. Of the 32 attempts, 17 boring locations provided enough groundwater to collect a sample. Assessment of the secondary source in the UCRS at SWMUs 7 and 30 was based on both historical and newly generated data, while the assessment of the secondary source in both the UCRS and RGA at SWMU 4 is based primarily on historical data. Because the UCRS water samples supplement only the characterization of the BGOU SWMUs (the analysis of subsurface soil samples is the primary measure that supports the assessment of nature and extent and risk) and secondary sources, the lack of

UCRS water samples from all soil borings does not limit the planned assessment of the SWMUs. There is also potential for a TCE DNAPL source at SWMU 2 based on historical disposal records; however, neither the subsurface soil nor shallow groundwater data at SWMU 2 indicate a DNAPL source. The volumetric extent of secondary source contamination has been approximated and constitutes a project assumption for evaluation of the alternatives. The impact of source volume or DNAPL uncertainties will be evaluated and further discussed in the FS.

There remains some uncertainty with regard to the boundaries of the burial pits. Geophysical surveys have not been completed across the entire area of all SWMUs (SWMU 6 had equipment in the way when it was geophysically surveyed). There also were single-point geophysical anomalies identified (i.e., SWMU 30), the nature of which has not been identified. These uncertainties will need to be managed in the FS.

The nature and extent of contaminants in surface soil at selected SWMUs (e.g., SWMU 145) is not well characterized. This uncertainty will need to be managed in the FS.

Surface and subsurface transport mechanisms

Previous work has shown that the primary pathway for groundwater flow and the site-related contaminants is vertical migration through the UCRS, followed by lateral migration in the RGA. Contaminated groundwater could migrate to the POEs identified for the BGOU SWMUs at the plant boundary, property boundary, surface seeps at Little Bayou Creek, and near the Ohio River. Not all SWMUs have transport pathways to all of the POEs. For example, SWMU 145 is located outside of the plant boundary and does not contribute to the Little Bayou seeps. SWMUs 3, 6, 7, and 30 were determined to be the only SWMUs contributing to the Little Bayou seeps POE. SWMUs 2, 4, and 5 are potential contributors to the Little Bayou Creek seeps, but the numerical model used to define flowpaths indicated releases from those SWMUs would go to the Ohio River. While there is some uncertainty related to modeling in predicting whether a SWMU would contribute to the Little Bayou seeps or the Ohio River, the model results indicate this uncertainty has an insignificant effect on the modeled contaminant concentrations and should not affect or drive remedial decisions.

The location of the water table varies in burial ground SWMUs. Most of the buried waste at SWMU 2 is saturated. The westward slope of the water table below SWMU 2 indicates that the water table must be equally shallow beneath SWMU 3. Because SWMU 3 is an aboveground facility with a RCRA multi-layered cap, all but the base of the landfill wastes likely are unsaturated. The stratigraphy of SWMU 4 is comparable to that of SWMUs 2 and 3. It appears that the hydrogeologic setting is similar, and the water table likely extends up into the waste burial pits. At SWMUs 5 and 6, even the shallowest wastes (with top near 365 ft amsl), likely are buried below the water table (at an elevation of approximately 367 ft amsl on the north side of SWMU 5). The SWMUs 7 and 30 RI (DOE 1998a) determined that a shallow water table exists approximately 5 ft bgs (Figure 3.22) and within the burial cells. UCRS piezometer and well measurements documented a strong downward gradient within the area. The elevation of the water table remains poorly documented at SWMU 145. Some buried waste at SWMU 145 likely is saturated.

There are water pipelines and electrical conduits in the proximity of some of the SWMUs. These conduits could act as pathways for contaminant transport. Because they were not specifically evaluated in the RI, or in past RIs, uncertainty remains as to any contaminant transport along these pipelines and conduits. Due to the strong downward hydraulic gradient in the UCRS and the fact that the pipelines are not located below the waste cells, transport along these type of features should be limited.

Uncertainty does exist with regard to the dissolved oxygen in the UCRS at SWMUs 4 and 6 due to a lack of data. The majority of dissolved oxygen measurements from UCRS wells range from near zero to four

mg/L and oxidation/reduction potential commonly ranges from -100 to 300 microVolts, with the majority of measurements greater than zero. Line plots in Figure 3.9 of the RI further demonstrate trends of dissolved oxygen (517 measurements) and oxidation/reduction potential (136 measurements) in the UCRS at the BGOU SWMUs. Due to the relative abundance of measurements for most SWMUs that demonstrate that the cumulative trend is likely to be representative of conditions at each SWMU, the oxidation/reduction potential in the UCRS at SWMUs 4 and 6 will be assumed to be similar to that in the UCRS at other BGOU SWMUs. The impact of this assumption will be evaluated further in the FS. If determined necessary to support implementation of a remedial alternative, dissolved oxygen could be measured as part of a remedial design support investigation for SWMUs 4 and 6.

Another potential pathway that exists, likely at SWMUs 7 and 30, but also possibly at other BGOU SWMUs, is lateral seepage from the burial pits into nearby ditches. The SWMU 7 and 30 RI Report (DOE 1998a) reported that water was observed emanating from the slope of a nearby ditch following a heavy rainfall. It is uncertain whether the seepage was derived from the burial pits. The RI report concluded that uranium activity concentrations in the ditch sediments suggest SWMUs 7 and 30 are contributing to contamination in the ditch, but the uranium isotope activity ratios in surface water in the ditch argued against waste burial pit waters as contributors to surface water contamination. The increased radiological activity is best explained by surface erosion carrying soil-bound radionuclides to the ditch. Likewise, some discharge of shallow groundwater in the ditch south of SWMU 2 has been observed, but the report was unclear as to the contribution of contamination to the ditch (the report concluded that contaminant migration to Outfall 015 and Bayou Creek is unlikely to exceed preliminary remediation goals) (DOE 1997a). This uncertainty will need to be managed in the FS.

Several uncertainties identified in this section will need to be managed in the FS. These include the following:

- Uncertainties related to risks associated with the mobility of uranium (the FS will manage this uncertainty by evaluating appropriate technologies for SWMUs where uranium is a primary contaminant);
- Uncertainties concerning the extent of source zones (burial areas) and unidentified single-point geophysical anomalies and the impact on alternative analyses (the FS will use existing knowledge and manage the uncertainties regarding the volume requiring removal or treatment);
- Uncertainties regarding the potential for acidic leachate, oxidation/reduction conditions, and degree of waste saturation (the FS will manage these uncertainties by evaluating robust technologies that are not sensitive to these types of uncertainties);
- Uncertainties regarding the extent and volume of secondary source zones (TCE DNAPL) (the FS will manage uncertainties regarding the extent and volume of these sources for comparison);
- Uncertainties related to limited groundwater monitoring around the BGOU SWMUs (the FS will manage this uncertainty by incorporating additional groundwater monitoring where appropriate at SWMUs where effectiveness monitoring is needed or where waste is left in place).
- Uncertainties related to the potential for releases from burial areas to impact adjacent surface water ditches (the FS will manage these uncertainties by recommending additional shallow groundwater monitoring during remedial design).

- Uncertainties related to the nature and extent of contaminants in surface soil at selected SWMUs, primarily SWMU 145. (The FS will manage this uncertainty by evaluating remedial alternatives that would address this uncertainty.)

7.8 CONCLUSIONS

Table 7.8 summarizes the decision rules of the BGOU Strategic Initiative (DOE 2006a). For each SWMU of the BGOU, risk levels associated with contamination at the SWMUs and associated with groundwater contamination derived from all of the SWMUs meet the criteria of the decision rules to progress to evaluate actions that will mitigate risk and to achieve ARARs; to seek an ARAR waiver in accordance with EPA guidance; or to propose alternative standards.

The following are the major contaminant distribution findings for sources investigated in the BGOU RI.

- Environmental media, specifically subsurface soil and groundwater, have been impacted by releases of contaminants at all of the BGOU SWMUs.
- TCE trends in the RGA indicate that TCE DNAPL potentially is present at SWMU 4 and in the vicinity of the shared border between SWMUs 7 and 30. (See Sections 3.9.4, 4.5.2, and 4.8.2.) Concentrations of TCE at SWMU 4 suggest this potential TCE DNAPL may be present both in the waste cells and underlying soils of the UCRS and in the matrix of the RGA. TCE trends at SWMUs 7 and 30 indicate that the potential TCE DNAPL source is likely constrained to the UCRS soils.
- The BHHRA completed as part of the BGOU RI indicates that ELCRs greater than the upper end of EPA's acceptable risk range (i.e., 1E-04) and HIs greater than 1 exist at all SWMUs; therefore, an FS is appropriate for impacted media at each SWMU. The metals arsenic, beryllium, and uranium; the organic compounds Total PAHs and Total PCBs; and the radionuclides uranium-235 and uranium-238 are common contaminants that present the dominant risks from exposure to surface and subsurface soil. The major contaminants driving the groundwater risks at the on-site POEs are arsenic, Aroclor-1260, 1,1-DCE, TCE, technetium-99, and vinyl chloride.
- Migration of contaminants through groundwater from all but SWMU 6 to locations at the SWMU boundary, the plant boundary, property boundary, and near the Ohio River, also posed greater than *de minimis* risks to a hypothetical residential groundwater user. Arsenic, TCE, 1,1-DCE, technetium-99, and vinyl chloride are the primary risk drivers.
- The Screening Ecological Risk Assessment retained a number of COPCs, primarily metals, at each of the sites. Each SWMU requires further ecological evaluation.

Table 7.8. Decision Rules for the BGOU Strategic Initiative

GOAL	DECISION RULE		<i>Then statement</i>
	<i>If statement</i>		
Nature of Contamination	1a	If the concentration of analytes found in the source zone could result in a cumulative excess lifetime cancer risk greater than 1×10^{-6} or a cumulative Hazard Index greater than 1 through contact with contaminated media, or if the concentration of analytes in the source zone could result in detrimental impacts to nonhuman receptors through contact with contaminated media as indicated by exceeding ecological screening criteria, and if the concentrations of analytes in the source zone are greater than those expected to occur naturally in the environment,	then evaluate actions that will mitigate risk; otherwise pursue a “no further action” decision (see D1b and D1c).
	1b	If concentrations of analytes found in the source zone exceed ARARs,	then evaluate actions that will bring contamination within the source zone into compliance with ARARs; seek an ARAR waiver (such as technical impracticability, inconsistent application of state standards, interim measure, greater risk to human health and the environment, equivalent standard of performance) in accordance with EPA guidance; or propose/obtain alternative standards.
	1c	If contaminants found at the site are known to transform or degrade into chemicals that could lead to increased risks to human health or the environment or into chemicals for which there are ARARs, and if the concentrations of these contaminants could result in risks greater than those defined in D1a or concentrations greater than ARARs,	then evaluate actions that will mitigate potential future risk or obtain compliance with ARARs; seek an ARAR waiver (such as technical impracticability, inconsistent application of state standards, interim measure, greater risk to human health and the environment, equivalent standard of performance) in accordance with EPA guidance; or propose/obtain alternative standards.
Extent of Contamination	2a	If secondary contamination sources are found, and if the concentration of analytes within the secondary contamination source is found to potentially result in a cumulative excess lifetime cancer risk greater than 1×10^{-6} or a cumulative Hazard Index greater than 1 through contact with contaminated media at the unit, and if the concentrations of analytes are greater than those expected to occur naturally in the environment,	then evaluate actions that will mitigate risk; otherwise, do not consider secondary contamination sources when making remedial decisions for the unit.

Table 7.8. Decision Rules for the BGOU Strategic Initiative (Continued)

GOAL	DECISION RULE		
	<i>If statement</i>	<i>Then statement</i>	
Fate and Transport	3a	<p>If contaminants are found in the source zone, or if secondary contamination sources are found, and if these contaminants are found to be migrating or may migrate from the source zone or from secondary contamination sources at concentrations that may potentially result in a cumulative excess lifetime cancer risk greater than 1×10^{-6} or a cumulative Hazard Index greater than 1 through use of contaminated media at downgradient points of exposure, and the concentrations of analytes are greater than those expected to occur naturally in the environment,</p>	<p>then evaluate actions that will mitigate risk; otherwise, do not consider risk posed by migratory pathways when evaluating remedial alternatives for the unit (see D3b).</p>
	3b	<p>If contaminants are found in the source zone, or if secondary contamination sources are found, and if these contaminants are found to be migrating or may migrate from the source zone or from the secondary contamination source at concentrations that exceed ARARs,</p>	<p>then evaluate actions that will bring migratory concentrations into compliance with ARARs; seek an ARAR waiver (such as technical impracticability, inconsistent application of state standards, interim measure, greater risk to human health and the environment, equivalent standard of performance) in accordance with EPA guidance; or propose/obtain alternative standards; otherwise, do not consider ARARs when examining migratory pathways during the evaluation of remedial actions (see D3a).</p>
Risk Assessment	4a	<p>If Decision D1a, D1b, D1c, D2a, D3a, or D3b indicate that response actions are needed,</p>	<p>then evaluate response actions to mitigate risk in the source zone.</p>

7.8.1 Recommendations for Future Work

Based on results in this RI Report, an FS will be conducted for each of the SWMUs in the BGOU. A listing of potential process options is detailed in Tables 7.9 and 7.10 and is consistent with data collection objectives in the work plan.

Table 7.9. Potential Process Options for Primary Sources (Waste and Vadose Soils)

Soil	
Institutional Controls	<ul style="list-style-type: none"> • Land-use restrictions • Environmental media monitoring
Containment	<ul style="list-style-type: none"> • Low-permeability capping • Constructed barriers • Dust and vapor suppression • Erosion control • Retro-fitted liners • Surface water control
Recovery or Removal	<ul style="list-style-type: none"> • Excavation/storage • Excavation/disposal
Treatment	<ul style="list-style-type: none"> • <i>In situ</i> grouting • Freezing

Table 7.10. Potential Process Options for Secondary Sources (DNAPL)

Groundwater	
Institutional Controls	<ul style="list-style-type: none"> • Land-use restrictions • Environmental media monitoring
Containment	<ul style="list-style-type: none"> • Constructed barriers • Hydraulic containment • Retro-fitted liners • Subsurface drainage
Recovery or Removal	<ul style="list-style-type: none"> • Extraction/storage • Extraction/disposal • Electrical resistivity heating • Dynamic underground stripping • Soil vapor extraction
<i>Ex Situ</i> Treatment	<ul style="list-style-type: none"> • Coagulation/flocculation • Freeze crystallization • Gravity separation • Media filtration • Membrane separation
<i>In Situ</i> Treatment	<ul style="list-style-type: none"> • Neutralization • <i>In situ</i> neutralization • Reactive walls • Phytoremediation

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APPENDIX A

TECHNICAL MEMORANDUM FOR FIELD ACTIVITIES

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ACRONYMS

AOC	area of contamination
BGOU	Burial Grounds Operable Unit
bgs	below ground surface
CERCLA	Comprehensive Environmental Response, Compensation, and Liability Act
<i>CFR</i>	<i>Code of Federal Regulations</i>
DOE	U.S. Department of Energy
DPT	direct push technology
EM	electromagnetometer
EPA	U.S. Environmental Protection Agency
ES&H	Environment, Safety, and Health
HSA	hollow-stem auger
NSDD	North-South Diversion Ditch
Paducah OREIS	Paducah Oak Ridge Environmental Information System
PCB	polychlorinated biphenyl
PEMS	Project Environmental Measurements System
PGDP	Paducah Gaseous Diffusion Plant
pH	negative logarithm of the hydrogen-ion concentration
PPE	personal protective equipment
QC	quality control
RCRA	Resource Conservation and Recovery Act
RGA	Regional Gravel Aquifer
RI	remedial investigation
SWMU	solid waste management unit
UCRS	Upper Continental Recharge System
VOC	volatile organic compound
WAG	Waste Area Group

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A.1. INTRODUCTION

The purpose of this memorandum is to provide certain technical details regarding field activities pertaining to the Burial Grounds Operable Unit (BGOU) Remedial Investigation (RI). A brief summary of project objectives is provided below; a more thorough discussion is contained in the body of the report.

The BGOU is one of six operable units located within the Paducah Gaseous Diffusion Plant (PGDP). This operable unit consists of contamination associated with PGDP's landfills and burial grounds. Burial grounds addressed by this RI include the following solid waste management units (SWMUs):

SWMU 2	C-749 Uranium Burial Ground
SWMU 3	C-404 Low-Level Radioactive Waste Burial Ground
SWMU 4	C-747 Contaminated Burial Yard and C-748-B Burial Area
SWMU 5	C-746-F Burial Yard
SWMU 6	C-747-B Burial Ground
SWMUs 7 and 30	C-747-A Burial Ground and Burn Area
SWMU 145	C-746-P Construction/Demolition Debris Disposal and Spoils Area (including the residential/inert borrow area and old North-South Diversion Ditch (NSDD) disposal trench)

The primary focus of this RI was to collect field and analytical data necessary to determine the nature and extent of any soil and groundwater contamination originating from, and immediately under, the burial cells; support the completion of a baseline human health risk assessment and a screening-level ecological risk assessment; and evaluate appropriate remedial alternatives (if necessary) at each of the SWMUs. The RI had the following four specific objectives:

- Characterize Nature of Source Zone—Characterize the nature of contaminant source materials by using existing data and, if required, by collecting additional data;
- Define Extent of Source Zone and Contamination in Soil and Other Secondary Sources at All Units—Define the nature, extent (vertical and lateral), and magnitude of contamination in soils, sediments, surface water, and groundwater by using existing data and, if required, by collecting additional data; determine the presence, general location (if practicable), and magnitude of any dense nonaqueous-phase liquid zones;
- Determine Surface and Subsurface Transport Mechanisms and Pathways—Gather existing quality data and, if necessary, collect additional adequate-quality data to analyze contaminant transport mechanisms, evaluate risk, and support a Feasibility Study; and
- Support Evaluation of Remedial Technologies—Determine if the existing data are sufficient to evaluate alternatives that will reduce risk to human health and the environment and/or control the migration of contaminants off-site.

The following table presents various procedures and work instructions that were used to complete the fieldwork conducted as part of the BGOU RI.

Table A.1. Procedures Used in the RI of the BGOU

Work Instructions or Procedures Required for Fieldwork and Sampling Activities
Archival of Environmental Data Within the Environmental Restoration Program
Chain-of-Custody
Cleaning and Decontaminating Sample Containers and Sampling Equipment
Data Entry
Data Management Coordination
Data Validation
Environmental Radiological Screening
Equipment Decontamination
Field Measurement Procedures: pH, Temperature, and Conductivity, and Dissolved Oxygen
Field Quality Control
Filter Pack and Screen Selection for Wells and Piezometers
Groundwater Sampling Procedures: Water Level Measurements
Identification and Management of Waste Not From a Radioactive Material Management Area
Labeling, Packaging, and Shipping of Environmental Field Samples
Lithologic Logging
Monitoring Well Development
Monitoring Well Installation
Monitoring Well Purging and Groundwater Sampling
Off-Site Decontamination Pad Operating Procedures
On-Site Handling and Disposal of Waste Materials
Opening Containerized Waste
Paducah Contractor Records Management Program
Pumping Liquid Wastes Into Tankers
Quality Assured Data
Sampling of Containerized Wastes
Use of Field Logbooks
Well and Temporary Boring Abandonment

The existing data for SWMU 4 was determined to be sufficient to evaluate the nature and extent of contamination and provide data from under the burial cells; therefore, no additional samples were collected for this SWMU. Borings were collected from under some of the burial cells for SWMUs 5 and 6 in a previous investigation; however, not all cells were evaluated. For SWMU 5, additional borings were collected from cells not previously targeted. For SWMU 6, physical constraints limited access to the area during previous investigations. Equipment had been removed subsequent to the Waste Area Group (WAG) 3 RI from the area, and it was possible to collect samples and evaluate those cells during this RI.

Activities addressed in this technical memorandum (Appendix A) are discussed in the following chapters:

- Chapter 2—Sampling Strategy
- Chapter 3—Surveying
- Chapter 4—Sampling Procedures
- Chapter 5—Field Decontamination
- Chapter 6—Waste Management
- Chapter 7—Environment, Safety, and Health
- Chapter 8—Fieldwork Documentation
- Chapter 9—Deviation from Planned Sample Locations

A.2. SAMPLING STRATEGY

The field sampling strategy used for the RI consisted of intrusive media sampling (surface and subsurface soil, and groundwater). The investigation activities used standard industry practices that were consistent with U.S. Environmental Protection Agency (EPA) procedures and protocols. Sampling activities at the burial grounds focused on the soils and groundwater beneath the burial pits down to a depth of 60 ft bgs (below ground surface). Surface and subsurface soils adjacent to but not beneath the burial pits were not part of this investigation. These will be evaluated through the Soils Operable Unit. Likewise, the Regional Gravel Aquifer (RGA) was not part of this investigation. It will be evaluated through the Groundwater Operable Unit. Borings adjacent to the NSDD were advanced to a depth of 15 ft bgs to evaluate impacts from the pipeline that once discharged leachate from SWMU 3 into the NSDD.

A.2.1 SOIL/SEDIMENT SAMPLING

The drilling technology for the angled borings and the six shallow vertical borings was a track-mounted rig capable of both direct push technology (DPT) and hollow-stem auger (HSA) drilling. This track-mounted drill rig utilized push rods to advance a soil sample tube with an acetate liner to collect undisturbed soil samples. If refusal was met using the push rods, auger flights then were drilled over the push rods to advance the borings to the sample depth. The deep vertical borings were completed with a larger HSA drill rig capable of reaching a depth of 100 ft bgs. For the deeper samples, drillers advanced the hollow stem augers to near the sample depth and then pushed the DPT sample tube through the hollow-stem augers to the sample depth to collect the soil samples.

The BGOU RI Work Plan directed the use of angled borings to sample from beneath the burial cells. Per the RI Work Plan, the field crew did not collect soil samples at or near the surface in the angled borings because these borings were begun at a specified distance away from the burial cells, outside the influence of the buried waste. Table A.2 summarizes the common soil sample depths.

Table A.2. BGOU RI Soil Sample Depths

Target Vertical Depth (ft)*		Actual Drilled Length (ft) at 45° Angle		Actual Vertical Depth (ft)	
Top	Bottom	Top	Bottom	Top	Bottom
8	10	10	15	7	11
13	15	15	20	11	14
28	30	40	45	28	32
43	45	60	65	42	46
58	60	80	85	57	60

*Specified in the BGOU RI Work Plan

Drilling and sampling difficulties necessitated slight adjustments to sampling depths in some instances. Collection of duplicate samples, likewise, required a longer sample interval to accommodate the increased sample volume. In addition, the sampling depths of the two lower intervals in boreholes 145-106 and 145-107 were adjusted to accommodate a locally thinner Upper Continental Recharge System (UCRS). Cross sections and tables in Section 4 and the boring logs in Appendix B document the depth of each soil sample.

The BGOU RI included sampling from both shallow and deep vertical borings. Ten shallow borings were installed along a former drainage ditch that connected the C-404 Landfill and the NSDD. The field crew collected samples from these borings at the surface and from 1 to 5 ft, 5 to 10 ft, and 10 to 15 ft. Three deep borings were installed within SWMU 7. With the exception of the surface soil sample in boring 007-011, the field crew collected soil samples in these borings from depths of 0 to 1 ft, 3 to 5 ft, 8 to 10 ft, 13 to 15 ft, 28 to 30 ft, 43 to 45 ft, and 58 to 60 ft. (The 0-1 ft depth interval in boring 007-011 consisted of gravel road base which was not amenable to laboratory analysis).

The field crew sampled the soil borings in accordance with U.S. Department of Energy (DOE) Prime Contractor-approved procedures, consistent with *Environmental Investigation Standard Operating Procedures and Quality Assurance Manual*, EPA Region 4, November 2001. As soon as the drill crew recovered the acetate liner containing the soil sample, the soil core was placed in the sample preparation area. A health and safety officer and radiation control officer scanned the acetate sleeve and the ends of the soil core for volatile organic compounds and radiation before releasing the core to the sample crew. Once the soil core in acetate sleeve was cleared, the sample crew opened the acetate sleeve with a utility knife and, once again, a health and safety officer and radiation control officer scanned the sample for contamination. The field scans of the acetate liner and soil core rarely identified contamination. When contamination was found, the health and safety officer and radiation control officer directed the field crew in any additional personal protective equipment (PPE) requirements and appropriate handling precautions.

Immediately upon approval from the health and safety officer and radiation control officer for the field crew to sample the soil core, the field crew collected the samples for volatile organic compound (VOC) analysis by filling two 2 ounce, wide-mouth, sample bottles with soil, ensuring that no air space was present, and securely sealing the filled bottles. At the same time, the project geologist examined soil core samples for lithologic description. After the collection of the VOC samples and the description of the lithology were complete, the remaining soil was placed in a clean bowl and mixed thoroughly. Samplers placed the resulting soil mixture in the appropriate sample jars for analysis.

A.2.2 GROUNDWATER SAMPLING

Groundwater samples were collected from multiple discrete depths within the UCRS and RGA using temporary borings at various locations. The RI field crew collected water samples in the UCRS where the temporary soil borings intersected water-producing zones. Water sampling in the RGA began at the top of the RGA (approximately 60 ft bgs) and continued at 10 ft intervals to the base of the RGA (approximately 100 ft bgs). This strategy resulted in a total of up to six water samples collected from the borings, depending on the presence of water-bearing zones in the UCRS and the thickness of the RGA at a boring's location. The drilling and sampling process allowed collection of discrete-depth water samples with minimum vertical cross-contamination.

A.3. SURVEYING

Two types of surveying were performed for the BGOU RI. They were civil surveying to locate sampling points and geophysical surveying to locate areas of potential burial.

A.3.1 CIVIL SURVEYING

As the field crew performed the BGOU RI sampling, they marked the boring locations using flagging and wooden stakes. Entries in project logbooks and on field maps further documented the sample locations. Brass markers were incorporated as part of pad installation for any monitoring wells. The BGOU RI included surveying of sampling locations upon completion of the RI field activities. This survey work was performed by or under responsible charge of a Professional Land Surveyor registered in the Commonwealth of Kentucky, locating each sample point with its horizontal and vertical position using the PGDP coordinate system for horizontal control. Additionally, the survey identified the State Plane Coordinates for each sample location using the U.S. Coast and Geodetic Survey North American Datum of 1983. The datum for vertical control was the U.S. Coast and Geodetic Survey North American Vertical Datum of 1988. Accuracy for this work was that of a Class 1 First Order survey.

Project personnel entered the coordinates into the Paducah Project Environmental Measurements System (PEMS), and the coordinate locations were transferred with the station's ready-to-load file to the Paducah Oak Ridge Environmental Information System (Paducah OREIS).

A.3.2 GEOPHYSICAL SURVEYING

The BGOU RI performed nonintrusive data collection (surface geophysics) for several of the SWMUs in order to better delineate areas of subsurface burial. Because these SWMUs consist of one or more burial pits of various depths that are filled with a heterogeneous collection of wastes and backfill soils, the BGOU represented a difficult target for geophysical characterization. Magnetic properties of the metal drums and buried metal scrap offered the best contrast with the native soils for imaging.

A Geonics electromagnetometer (EM)-61 metal locator pulled on a trailer was used to perform the geophysics. The EM-61 detects buried metal with very good resolution utilizing time-domain electromagnetic phenomena to detect buried metal up to 16 ft below the surface (http://www.geologyuk.com/geophysics/tech_summary/07_Technical%20summary%20sheet_EM61.pdf). Generally, readings from the EM-61 were taken along continuous lines spaced 5 ft apart in both north-south and east-west orientations (see Figure A.1). Locations of anomalies were pin flagged in the field and recorded on a PC600 Data Logger. The pin-flagged locations were mapped and transferred to digital maps. Depths of the anomalies were taken from the data recorded and also transferred to the maps.

Geophysical surveys of SWMUs 7 and 30 and 145 delineated the burial pits exact location and burial extent prior to sampling activities. The survey grid covered an area that extended at least 10 ft beyond the currently identified burial pit edges. The survey grid for SWMU 145 was set to 5 ft by 10 ft (5-ft line spacing in the east-west direction and 10-ft line spacing in the north-south direction; see Figure A.2) due to the size of the SWMU.

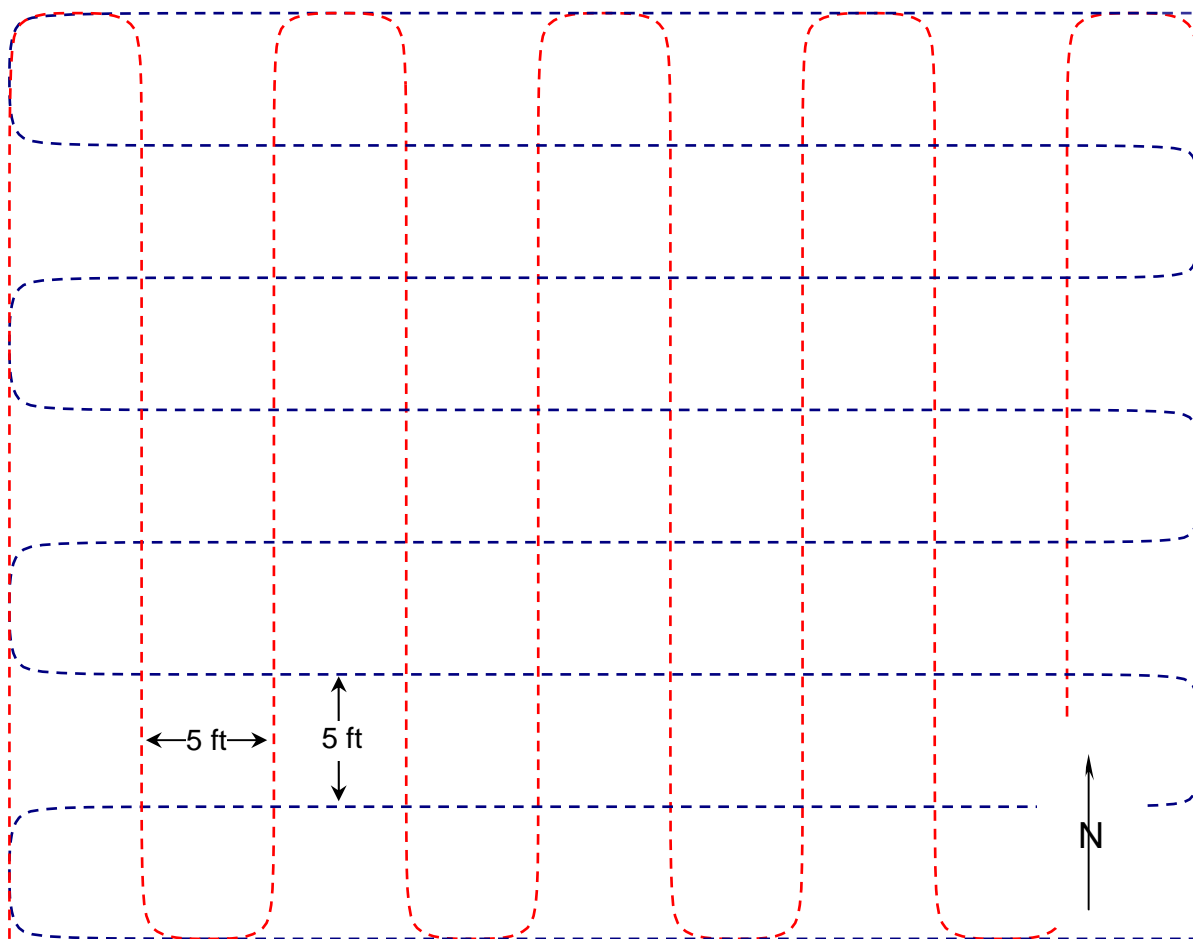


Figure A.1. Example EM-61 Survey Grid for BGOU RI

Portions of SWMUs 2, 3, 5, and 6 were surveyed to delineate accurately burial pits prior to drilling. For these SWMUs, grid spacing was not consistent along continuous lines since the results of these surveys were specifically targeted to aid in placement of the planned drilling. Additionally, geophysical surveys were used at SWMU 2 to delineate the exact location of a buried electrical conduit within the burial yard. A portion of SWMU 5 was surveyed to delineate the location of an abandoned waterline at the north of the burial yard.

Following an employee interview, geophysical surveys were performed in three areas within SWMU 13 utilizing the same methodology that was applied for SWMUs 7 and 30 and 145 (i.e., trailer-mounted EM-61 pulled along a grid of continuous lines spaced 5 ft apart). The EM-61 survey identified metal throughout the three areas of interest beginning at a depth of 2 ft bgs. This metal likely is small scrap material related to previous activities that followed removal of the scrap piles (spreading and covering small amounts of metal as part of yard cleanup). The EM-61 survey detected an anomaly in the third area at the western end of the SWMU, beginning at approximately 4 to 8 ft in depth.

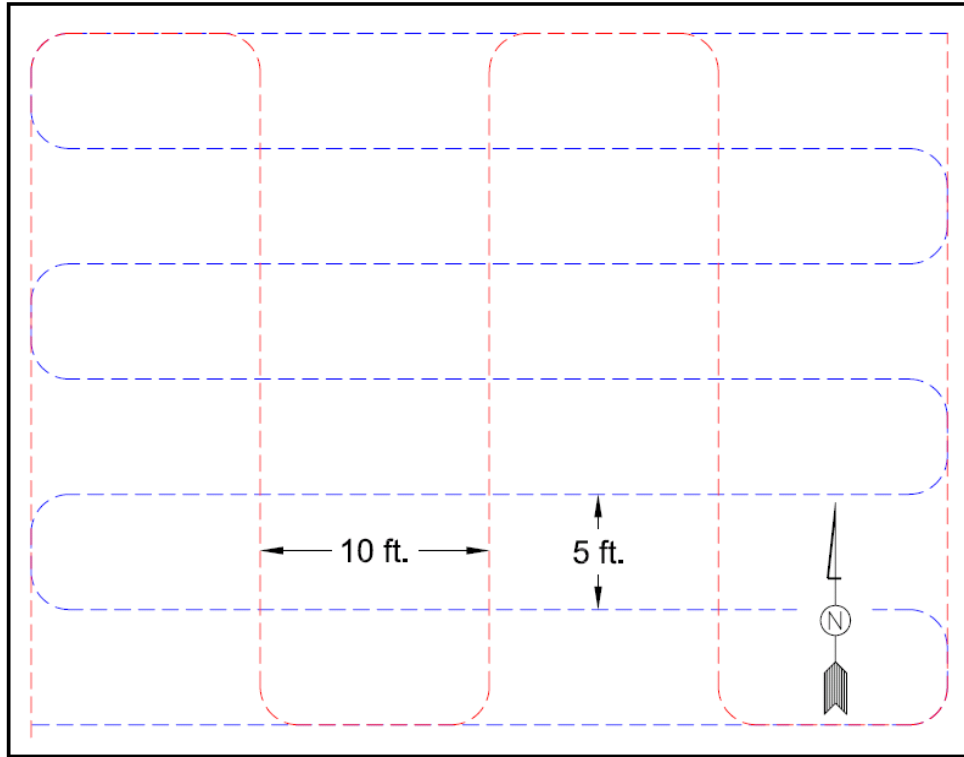


Figure A.2. Example EM-61 Survey Grid for BGOU RI at SWMU 145

A.4. SAMPLING PROCEDURES

During the sampling event, three types of samples—soil/sediment, groundwater, and field quality control (QC)—were collected and submitted for analysis. The sampling team varied between two and three members. Prior to initiation of field sampling, all sample team members completed general and project-specific training.

The sampling team collected, stored, and shipped the samples according to preestablished QC protocols and approved project procedures, which were consistent with EPA Region 4 sampling methodologies. Sample container, preservation, and holding time requirements were in accordance with the EPA Engineering Support Branch Standard Operating Procedures.

Samples collected for this project were assigned unique sample identifiers that were recorded on the sample labels and chain-of-custody forms. Sample team crew members directly affixed labels to the sample containers that included the following information:

- Station name,
- Sample identification number,
- Sample matrix,
- Sample type,
- Type or types of analysis required,
- Date and time of collection,
- Sampler name,

- Sample preservation (if required), and
- Destination laboratory.

The sampling team wore proper PPE during sampling. PPE consisted of, in part, company-issued clothing, safety glasses, and latex gloves. Sampling in radiological contamination areas sometimes necessitated modifications of the PPE requirements (as prescribed in work permits and directed by the project's Health Physics technician).

A.4.1 SOIL/SEDIMENT SAMPLES

The field crew sampled the soil borings in accordance with DOE Prime Contractor-approved procedures, consistent with *Environmental Investigation Standard Operating Procedures and Quality Assurance Manual*, EPA Region 4, November 2001, collecting soil for VOC analysis, followed by samples for lithologic description as soon as the acetate sleeve was cut open. After the description of the lithology was complete, the remaining soil was placed in a clean stainless steel bowl and mixed thoroughly using a stainless steel spoon to homogenize the soil taken from the sample interval before sampling for other analyses. Since round bowls were used for sample preparation, adequate mixing was achieved by stirring the material in a circular fashion, reversing direction, and occasionally turning the material over.

Sample team members filled the sample containers and ensured that each lid was securely tightened. The sample container then was placed in a cooler with an ice pack to maintain a preservation temperature of 4 degrees Celsius. Crew members recorded all pertinent information in the sampling logbook.

A.4.2 GROUNDWATER SAMPLES

Where the temporary soil borings intersected water-bearing units, the BGOU RI collected a groundwater sample from the UCRS and multiple discrete depths in the RGA. This RI (fieldwork performed in 2007) resulted in a total of 30 groundwater samples. The first step in collecting both UCRS and RGA groundwater sample was to purge the drill pipe and the disturbed soil in the vicinity of the open pipe. The field crew used bladder pumps to purge the boring and to collect the water samples.

Since sampling took place immediately after drilling ceased, there was no stagnant water to remove from the boring and, therefore, no predetermined minimum purge volume. The sample crew collected the water sample in both the UCRS and RGA only after the measure of select geochemical parameters [i.e., acidity as reported as the negative logarithm of the hydrogen-ion concentration (pH), specific conductivity, and temperature] stabilized within the purge water (signifying that the discharging water was representative of groundwater quality). The geochemical parameters were considered stabilized when the following criteria were met:

- At least three measurements taken three minutes apart have consistent readings for temperature, conductivity, and pH;
- Temperature measurements agree within 1 °C;
- Conductivity measurements agree within 10%; and
- pH measurements agree within 0.5 units.

When the geochemical parameters stabilized, the sampling crew adjusted the flow rate of the pump for sampling. Groundwater samples were collected in accordance with SWMU-specific sampling plans (*Work Plan for the Burial Grounds Operable Unit Remedial Investigation/Feasibility Study at Paducah Gaseous Diffusion Plant, Paducah, Kentucky, DOE/OR/07-2179&D2/R1*). (The sampling plans were specific as to the analytes for each SWMU and the horizons, UCRS and RGA, to be sampled.) All samples required multiple analyses. With the exception of a duplicate groundwater sample in boring 007-007, each groundwater interval that was sampled yielded sufficient volume to fill sample bottles for all analyses. The sampling crew collected the field parameters of groundwater temperature, pH, and specific conductance during each sampling event.

After sampling was completed, the sample crew removed the tubing and pump from the boring. The pump and tubing was decontaminated in accordance with DOE Prime Contractor-approved procedures prior to its next use.

A.4.3 FIELD QC SAMPLES

To ensure reliability of the analytical data and to meet the data quality objectives for the project, the following QC sample types were obtained during sample collection:

- Trip Blanks—Analysis of trip blanks documented the occurrence of cross contamination by VOCs during sample handling and shipping. The sample crew prepared trip blanks by filling VOC vials with deionized water before collection of the field samples. These trip blanks accompanied the filled sample bottles in ice chests in the field and during shipment and through interim storage in secured refrigerators until laboratory analysis. The trip blanks were analyzed for VOCs only.
- Field Blanks—Field blanks served as a check for potential airborne environmental contamination at the sample site. For the field blanks, the sample crew typically filled sample bottles with deionized water in the project's sample staging area and transported the bottles to the field sample station, where they were opened during the sampling process. Field blanks also were used as a reagent blank, as needed. The BGOU RI required field blanks at a frequency of one in 20 samples (5%) for each sample matrix.
- Field Duplicate Samples—Field duplicate samples determined the sampling variance. The sampling crew collected one duplicate for every ten samples (10%), per matrix. The field duplicate was analyzed for the same set of analytical parameters as the sample it duplicated.
- Equipment Blanks or Rinsate Samples—Equipment blanks provided a measure of the decontamination process effectiveness and were used as reagent blanks, as needed. These equipment blanks were required only when nondisposable equipment was being used. The equipment blanks consisted of deionized water passed through or over decontaminated sampling equipment and analyzed for the same parameters as the samples collected with the equipment. Equipment blanks were collected at a frequency of one for every 20 samples (5%).

In addition to the QC samples that were collected for laboratory analysis, temperature blanks accompanied the soil and groundwater samples in the transport coolers to document proper preservation of the samples. All transport coolers contained temperature blanks.

A.5. FIELD DECONTAMINATION

The field decontamination procedure, *Decontamination of Sampling Equipment and Devices*, PRS-ENM-2702, determined the decontamination activities for the stainless steel spoons and bowls used in soil sampling and the pumps and tubing used for groundwater sampling. This procedure, as applied during the RI, is summarized as follows:

- Equipment first was cleaned with tap water and nonphosphate detergent, using a brush if necessary, to remove particulate matter and surface films.
- The equipment then was rinsed thoroughly with tap water, followed by an analyte-free water rinse, and then wiped with an isopropyl alcohol towelett.
- The inside of the pump and tubing was cleaned by purging soap water, followed by tap water and analyte-free water, through the pump and tubing.
- Cleaned sample equipment was allowed to air dry.
- Cleaned equipment was handled only by personnel wearing clean latex gloves to prevent recontamination.
- If cleaned sampling equipment was not reused immediately, it was wrapped in aluminum foil.

Large Equipment Decontamination, PRS-FCD-2701, governed the cleaning of other sampling equipment such as the drill rigs and associated tooling. This procedure provides for the use of high-pressure steam as the primary cleaning agent. Because of its remote location, the BGOU RI drill crew constructed a temporary decontamination pad at SWMU 145 that was used in cleaning the drill rig and tooling. The on-site decontamination facility, C-416, supported cleaning activities for the drill rig and associated tooling during sampling at all other (on-site) BGOU RI locations.

A.6. WASTE MANAGEMENT

The RI work plan included a project-specific waste management plan to provide instruction regarding waste storage and disposition. A variety of wastes were generated during the field investigation, including sample residuals and associated waste derived from sample collection. The waste generated was stored in Comprehensive Environmental Response, Compensation, and Liability Act (CERCLA) waste storage areas within the CERCLA area of contamination (CERCLA AOC) during the characterization period and prior to disposal. Consistent with EPA Policy, the storage of waste within the CERCLA AOC does not trigger Resource Conservation and Recovery Act (RCRA) storage requirements (similarly, movement of waste within a CERCLA AOC does not trigger RCRA disposal requirements). As a best management practice, waste storage areas within the CERCLA AOC were managed in accordance with the substantive RCRA 90-Day storage standards; the 90-Day storage restriction and the requirement to label hazardous waste was not applied to the storage areas.

PPE was considered to fall into the same waste classification as the environmental media with which it came into contact. PPE, plastic, and paper were segregated by classification, collected in plastic bags, and labeled appropriately. These items then were handled as solid waste.

Decontamination water that included small quantities of soil sediments/mud was generated from cleaning the equipment. The water was collected and stored in a polyethylene tank and discharged to the Kentucky Pollutant Discharge Elimination System Outfall 001 after final characterization documented that the stored water met release criteria in the Kentucky Pollutant Discharge Elimination System permit for Outfall 001.

Solid waste was containerized in 55-gal drums, or approved equivalent, that were lined with a thick plastic liner and placed in CERCLA waste storage areas. The amount of free liquid was minimized. Any substantial amount of free liquid is being decanted and placed in an approved container. Drummed soils and other solid wastes have been disposed of in the C-746-U Landfill.

All clean trash (i.e., trash that was not chemically or radiologically contaminated) was segregated according to established guidelines and then collected and disposed of. Examples of clean trash are office paper, aluminum cans, packaging materials, glass bottles not used to store potentially hazardous chemicals, aluminum foil, and food items.

Based on sample analyses, existing data, or process knowledge, the waste was classified into one of the following categories:

- RCRA-listed hazardous waste,
- RCRA-characteristic hazardous waste,
- Polychlorinated biphenyl (PCB) waste,
- Transuranic waste,
- Low-level waste,
- Mixed waste, or
- Nonhazardous waste.

Waste minimization requirements were implemented, as appropriate, and included those established by the 1984 Hazardous and Solid Waste Amendments of RCRA; DOE Orders 5400.1, 5400.3, 435.1; and DOE Prime Contractor's requirements. Requirements specified in the waste management plan regarding waste generation, waste tracking, waste reduction techniques, and the waste reduction program, in general, also were implemented.

To support DOE's commitment to waste reduction, an effort was made during field activities to minimize waste generation as much as possible, largely through ensuring that potentially contaminated wastes were localized and did not come into contact with any clean media (which could create more contaminated waste). Waste minimization also was accomplished through waste segregation, selection of PPE, waste handling (spill control), and the use of alternative treatment standards.

A.7. ENVIRONMENT, SAFETY, AND HEALTH

A project-specific environment, safety, and health (ES&H) plan was included in the approved work plan and was used to provide instruction regarding safety and health of workers, the public, and the environment. The ES&H Plan established the specific applicable standards and practices to be used during execution of the RI to protect the safety and health of workers, the public, and the environment. The document contained information about the sites, potential contaminants and hazards that may be encountered on-site, and hazards inherent in routine procedures. The list of contaminants was site-specific and based on previous investigations. The plan also outlined directly, or by reference, federal and state

standards, pertinent consensus standards, and applicable contract requirements. The ES&H plan was implemented in accordance with 29 *CFR* § 1910.120, Hazardous Waste Operations and Emergency Response. Additional health and safety requirements were incorporated into the ES&H plan for the various field activities through preparation of project-specific activity hazard analyses.

The project team held daily safety and plan of the day meetings at the beginning of each shift. This approach ensured that the planned daily activities were reviewed prior to execution and the potential hazards were identified and discussed with the entire field team. These meetings are documented in the project work package and in the field logbooks.

A.8. FIELDWORK DOCUMENTATION

Field documentation was maintained throughout the BGOU RI in various types of documents and formats, including the field logbooks, sample labels, sample tags, chain-of-custody forms, and field data sheets. The following general guidelines for maintaining field documentation was implemented. Documentation requirements are listed below. Entries were written clearly and legibly using indelible ink.

- Corrections were made by striking through the error with a single line that did not obliterate the original entry. Corrections were dated and initialed.
- Dates and times were recorded using the format “mm/dd/yy” for the date and the military (i.e., 24-hour) clock for the time.
- Zeroes were recorded with a slash (/) to distinguish them from letter Os.
- Blank lines were prohibited. Information was recorded on each line or a blank line was lined out, initialed, and dated.
- No documents were altered, destroyed, or discarded, even if they were illegible or contained inaccuracies that required correction.
- Information blocks on field data forms were completed or a line was drawn through the unused section, and the area was dated and initialed.
- Unused logbook pages were marked with a diagonal line drawn from corner to corner and a signature and date was placed on the line.
- Photocopies of logbooks, field data sheets, and chain-of-custody forms were made and stored in the project file.
- The following information was recorded on the outside of the front cover of each logbook using indelible ink:
 - Project name,
 - Unique logbook name and number,
 - Client and contract number,
 - Task and document control number,

- Activity or site name, and
- Start and completion date of the logbook.

Quality assurance personnel conducted periodic reviews of the data forms and logbooks (including data forms placed in the logbooks) prepared by field personnel to verify the following:

- Accuracy of entries;
- Legibility and clarity of entries;
- Completeness, to ensure that at least the minimum required information was recorded;
- Consistency of information recorded; and
- Signature and date of entries by the designated team member.

A.9. DEVIATION FROM PLANNED SAMPLE LOCATIONS

A.9.1 INTRODUCTION

A Geographic Information System provided sample coordinates from maps of the intended sample locations in the BGOU RI Work Plan. Some of these locations were later adjusted to address additional data regarding the placement of waste. Once these locations were agreed upon, conventional survey methods located the sample coordinates at each SWMU. Table A.3 lists originally planned sample location coordinates and the final sample location coordinates.

A.9.2 DISCUSSION OF DEVIATION FROM COORDINATE LOCATIONS

During the survey and location of the sample boreholes, there were some boreholes that could not be located at the planned coordinates due to steep topography and surface structures, the presence of buried shock sensitive (explosive) waste, and High Radiation Areas. When obstructions or conditions prevented location of a sample at the planned location, the samples locations were offset close to the intended site. This section presents a summary of the samples that were relocated and provides the distance that the samples were offset from the intended coordinates.

At SWMU 5, it was discovered at the time of installation that an angled boring (005-103) at the southeast corner was being drilled in a diagonal orientation to the waste cells and not in a perpendicular orientation, as stated in the text of the work plan. The error was confirmed and the installation was stopped. A new boring was started a few ft away with a perpendicular orientation. The 10 ft and the 15 ft samples collected from the diagonally-oriented boring were discarded. The 30 ft sample was submitted for analysis along with the full set of samples from the new boring oriented perpendicular to the waste cells. Data from the initial boring are included with the summaries in the RI Report with the other data from boring 005-103. This data is available individually within the dataset and is identified by the sample number '005103SA030-2'.

At SWMU 145, continuing detection of PCBs in groundwater samples from the area monitoring wells at the time of the BGOU RI field investigation refocused the sampling strategy. With the participation of the Kentucky Department for Environmental Protection and the consent of EPA, the field crew relocated the SWMU 145 soil borings. The locations remained focused on areas of suspected disposal, as originally sited, but also considered the location of a potential TCE source (which may have facilitated PCB

transport) and the location of the last construction on the C-746-S Landfill, which may have disturbed PCB-contaminated soils.

Table A.3. BGOU RI Sample Locations

Sample Location	Planned		Final		Displacement (ft)
	Easting	Northing	Easting	Northing	
<i>SWMU 2</i>					
002-001	-6,312.96	-924.80	-6,275.65	-1,030.25	111.9
002-002	-6,228.85	-824.49	-6,285.73	-813.89	57.9
<i>SWMU 3</i>					
003-001	-6,071.13	-826.20	-6,072.45	-814.33	11.9
003-002	-5,901.41	-824.49	-5,904.5	-808.99	15.8
003-003	-5,751.48	-823.53	-5,755.25	-802.32	21.5
003-004	-5,654.54	-908.49	-5,634.54	-910.43	20.1
003-005	-5,361.38	-899.92	-5,372.98	-899.56	11.6
003-006	-5,292.80	-725.05	-5,296.92	-723.92	4.3
003-007	-5,212.22	-533.04	-5,210.8	-533.49	1.5
003-008	-5,133.36	-363.32	-5,132.26	-363.78	1.2
003-009	-4,968.78	-363.32	-4,968.32	-363.17	0.5
003-010	-4,804.20	-361.60	-4,804.22	-361.38	0.2
<i>SWMU 5</i>					
005-101	-6,676.72	200.58	-6,615.28	194.07	61.8
005-102	-6,545.21	199.28	-6,352.97	86.61	222.8
005-103	-6,345.99	19.59	-6,350.41	3.87	16.3
<i>SWMU 6</i>					
006-101	-6,253.46	121.60	-6,224.07	156.46	45.6
006-102	-6,288.55	97.72	-6,275.76	95.12	13.1
006-103	-6,234.47	75.00	-6,217.78	68.71	17.8
006-104	-6,180.98	104.34	-6,178.21	93.91	10.8
<i>SWMU 7</i>					
007-001	-6,270.86	913.50	-6,271.04	913	0.5
007-002	-6,342.41	807.58	-6,328.25	859.9	54.2
007-003	-6,557.62	784.20	-6,566.19	834.02	50.6
(offset)			-6,561.94	834.35	50.3
007-004	-6,682.69	788.71	-6,711.86	794.3	29.7
007-005	-6,745.51	759.69	-6,713.91	838.26	84.7
007-006	-6,751.14	884.76	-6,786.05	876.72	35.8
007-007	-6,588.89	883.64	-6,602.83	849.08	37.3
007-008	-6,061.29	924.76	-6,064.71	934.28	10.1
007-009	-6,830.86	990.12	-6,785.37	964.98	52.0
007-010	-6,590.86	988.99	-6,543.57	957.99	56.5
007-011	-6,260.72	810.96	-6,282.4	869.1	62.1
<i>SWMU 30</i>					
030-001	-7,114.24	994.91	-7,187.03	976.62	75.1
030-002	-6,954.24	922.79	-7,035.76	883.46	90.5
030-003	-6,926.07	883.36	-6,939.76	881.58	13.8
030-004	-6,814.52	793.22	-6,800.23	781.03	18.8
<i>SWMU 145</i>					
145-101	-1,832.30	4,329.28	-2,156.26	4,493.87	363.4
145-102	-1,821.86	3,790.02	-1,895.42	4,042.42	262.9
145-103	-2,413.30	3,748.28	-2,465.84	3,406.88	345.4
145-104	-2,618.56	3,716.96	-2,333.16	3,381.42	440.5
145-105	-2,917.76	4,367.55	-2,765.27	3,183.99	1,193.3
145-106	-2,343.72	4,565.85	-3,088.28	3,545.83	1,262.9
145-107	-2,044.52	4,510.19	-2,809.44	4,138.49	850.4

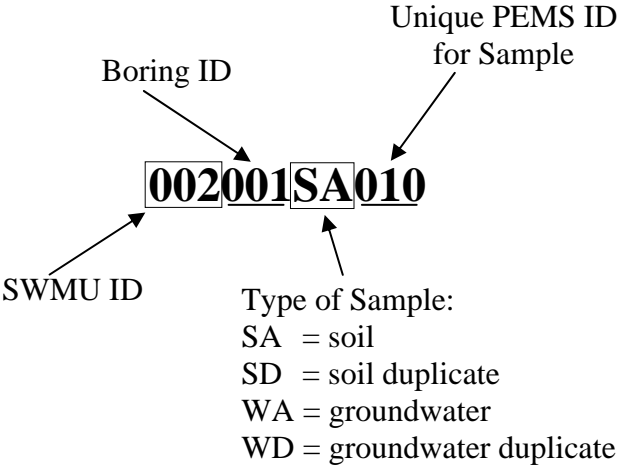
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APPENDIX B

**LITHOLOGIC LOGS AND WELL CONSTRUCTION DIAGRAMS,
GROUNDWATER STABILIZATION LOGS, AND WELL
DEVELOPMENT LOGS**

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SAMPLE ID LEGEND



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SWMU 2

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LITHOLOGIC LOG			BORING/WELL ID 002-001-ASB		Page 1 of 1
Facility: Paducah Gaseous Diffusion Plant, Paducah, KY			Site: SWMU- 002		
Project: BGOU RI			Client: USDOE/PRS		
Drilling Contractor: Chase Environmental, LLC			Driller: Jeff Brownfield		
Start Time/Date: 14:30/4-4-07			End Time/Date: 11:15/4-5-07		
Borehole Diameter: 6.25"			Drilling Method: Direct Push Through Augers		
Sampling Method: DT-21 Dual Tube / 2.25" ID Augers			Drill Rig: Geoprobe® 6620DT		
Total Depth (Vertical): 60 feet			Angle: 45°		
Logged By: Mark Gartner			Protective Level: Modified Level D		
Coordinates: E -6275.65 N -1030.259			Direction (plant grid): North		
Depth (ft)	SAMPLE		LITHOLOGIC DESCRIPTION	GRAPHIC LOG	COMMENTS
	INTERVAL	ID			
0	0-7	N/A	Core not retrieved	N/A	N/A
5	7-11	002001SA010	CLAY, light brown, 7.5YR 6/4, firm, moist, with orange streaks		14:44
10	11-14	002001SA015	CLAY, gray, 7.5YR 6/1, firm, moist, mottled, with gray streaks		14:52
15	14-28	N/A	Core not retrieved	N/A	N/A
20	28-32	002001SA030	CLAY, brown, 7.5YR 4/2, soft, moist, mottled, with black streaks: 5% chert fragments 1/8" across		16:30 Additional sample was required. Collected a second sample liner.
25	32-42	N/A	Core not retrieved	N/A	N/A
30	42-46	002001SA045	SAND, medium to coarse grained, poorly sorted with angular to rounded chert approximately 1/8" to 1/2" across		10:36
35	46-57	N/A	Core not retrieved	N/A	N/A
40	57-60	002001SA060	SAND, fine to coarse grained, poorly sorted		11:15
Total Vertical Depth is 60 feet. Total Linear Depth is 85 feet.			LEGEND: CLAY		
			SAND		

LITHOLOGIC LOG			BORING/WELL ID 002-002-ASB		Page 1 of 1
Facility: Paducah Gaseous Diffusion Plant, Paducah, KY			Site: SWMU- 002		
Project: BGOU RI			Client: USDOE/PRS		
Drilling Contractor: Chase Environmental, LLC			Driller: Jeff Brownfield		
Start Time/Date: 14:38/4-3-07			End Time/Date: 10:40/4-4-07		
Borehole Diameter: 6.25"			Drilling Method: Direct Push Through Augers		
Sampling Method: DT-21 Dual Tube / 2.25" ID Augers			Drill Rig: Geoprobe® 6620DT		
Total Depth (Vertical): 60 feet			Angle: 45°		
Logged By: Mark Gartner			Protective Level: Modified Level D		
Coordinates: E -6285.732 N -813.893			Direction (plant grid): South		
Depth (ft)	SAMPLE		LITHOLOGIC DESCRIPTION	GRAPHIC LOG	COMMENTS
	INTERVAL	ID			
0	0-7	N/A	Core not retrieved	N/A	N/A
5	7-11	002002SA010	CLAY, light brown, 7.5YR 6/4, firm, moist, with gray and orange streaks		14:44
10	11-14	002002SA015	CLAY, brown, 7.5YR 5/4, firm, moist, mottled, with gray and orange streaks		14:49
15					
20	14-28	N/A	Core not retrieved	N/A	N/A
25					
30	28-32	002002SA030 002002SD030	CLAY, strong brown, 7.5YR 5/6, firm, moist, with gray streaks: 5% sand		15:19
35	32-42	N/A	Core not retrieved	N/A	N/A
40					
45	42-46	002002SA045	CLAY, strong brown, 7.5YR 5/8, hard, dry, mottled with black streaks		9:00 / 4/4/07
50	46-57	N/A	Core not retrieved	N/A	N/A
55	57-60	002002SA060	SAND, medium grained, dark red: 10% clay		10:40
Total Vertical Depth is 60 feet. Total Linear Depth is 85 feet.			LEGEND: CLAY		
			SAND		






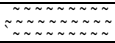

SWMU 3

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LITHOLOGIC LOG			BORING/WELL ID 003-001-ASB		Page 1 of 1
Facility: Paducah Gaseous Diffusion Plant, Paducah, KY			Site: SWMU- 003		
Project: BGOU RI			Client: USDOE/PRS		
Drilling Contractor: Chase Environmental, LLC			Driller: Jeff Brownfield		
Start Time/Date: 9:30/4-9-07			End Time/Date: 15:30/4-9-07		
Borehole Diameter: 6.25"			Drilling Method: Direct Push Through Augers		
Sampling Method: DT-21 Dual Tube / 2.25" ID Augers			Drill Rig: Geoprobe® 6620DT		
Total Depth (Vertical): 60 feet			Angle: 45°		
Logged By: Mark Gartner			Protective Level: Modified Level D		
Coordinates: E -6072.453 N -814.337			Direction (plant grid): Southeast		
Depth (ft)	SAMPLE		LITHOLOGIC DESCRIPTION	GRAPHIC LOG	COMMENTS
	INTERVAL	NUMBER			
0	0-7	N/A	Core not retrieved	N/A	N/A
5	7-11	003001SA010	CLAY, brown, 7.5YR 6/4, firm, moist, with gray and black streaks	~~~~~	9:40
10	11-14	003001SA015	CLAY, brown, 7.5YR 5/2, firm, moist, mottled, with gray and orange streaks	~~~~~	9:47
15					
20	14-28	N/A	Core not retrieved	N/A	N/A
25					
	28-30		CLAY, light gray, 7.5YR 7/1, soft, moist, with orange streaks	~~~~~	
30	30-32	003001SA030	CLAY, strong brown, 7.5YR 5/8, firm, dry, with gray streaks: 5% sand	~~~~~	13:02
35	32-42	N/A	Core not retrieved	N/A	N/A
40					
	42-46	003001SA045	CLAY, strong brown, 7.5YR 5/6, firm, moist, with gray and orange streaks	~~~~~	14:15
45					
50	46-57	N/A	Core not retrieved	N/A	N/A
55					
	57-60	003001SA060	CLAY, brown, 7.5YR 5/4, firm moist, mottled, with black streaks	~~~~~	15:30
Total Vertical Depth is 60 feet. Total Linear Depth is 85 feet.			LEGEND: CLAY		~~~~~
			SAND		~~~~~

LITHOLOGIC LOG			BORING/WELL ID 003-002-ASB		Page 1 of 1
Facility: Paducah Gaseous Diffusion Plant, Paducah, KY			Site: SWMU- 003		
Project: BGOU RI			Client: USDOE/PRS		
Drilling Contractor: Chase Environmental, LLC			Driller: Jeff Brownfield		
Start Time/Date: 10:30/4-10-07			End Time/Date: 16:22/4-10-07		
Borehole Diameter: 6.25"			Drilling Method: Direct Push Through Augers		
Sampling Method: DT-21 Dual Tube / 2.25" ID Augers			Drill Rig: Geoprobe® 6620DT		
Total Depth (Vertical): 60 feet			Angle: 45°		
Logged By: Mark Gartner			Protective Level: Modified Level D		
Coordinates: E -5904.505 N -808.997			Direction (plant grid): Southwest:		
Depth (ft)	SAMPLE		LITHOLOGIC DESCRIPTION	GRAPHIC LOG	COMMENTS
	INTERVAL	NUMBER			
0	0-7	N/A	Core not retrieved	N/A	N/A
5	7-11	003002SA010	CLAY, gray, 7.5YR 5/1, hard, dry, mottled, roots, with orange streaks	~~~~~ ~~~~~ ~~~~~	10:37
10	11-14	003002SA015	CLAY, brown, 7.5YR 5/4, soft, moist, with gray and orange streaks	~~~~~ ~~~~~ ~~~~~	10:42
15					
20	14-28	N/A	Core not retrieved	N/A	N/A
25					
30	28-32	003002SA030	CLAY, gray, 7.5YR 5/1, firm, moist: sand (10%)	~~~~~ ~~~~~ ~~~~~	14:37
35	32-42	N/A	Core not retrieved	N/A	N/A
40	42-46	003002SA045	CLAY, strong brown, 7.5YR 4/6, firm, moist: sand (5%) gray	~~~~~ ~~~~~ ~~~~~	15:47
45					
50	46-57	N/A	Core not retrieved	N/A	N/A
55	57-60	003002SA060	CLAY, white, 7.5YR 8/1, hard, moist, with orange streaks: sand (5%)	~~~~~ ~~~~~ ~~~~~	16:22
Total Vertical Depth is 60 feet. Total Linear Depth is 85 feet.			LEGEND: CLAY		~~~~~ ~~~~~ ~~~~~
			SAND		~~~~~ ~~~~~ ~~~~~

LITHOLOGIC LOG			BORING/WELL ID 003-003-ASB		Page 1 of 1
Facility: Paducah Gaseous Diffusion Plant, Paducah, KY			Site: SWMU- 003		
Project: BGOU RI			Client: USDOE/PRS		
Drilling Contractor: Chase Environmental, LLC			Driller: Jeff Brownfield		
Start Time/Date: 13:29/4-12-07			End Time/Date: 10:22/4-16-07		
Borehole Diameter: 6.25"			Drilling Method: Direct Push Through Augers		
Sampling Method: DT-21 Dual Tube / 2.25" ID Augers			Drill Rig: Geoprobe® 6620DT		
Total Depth (Vertical): 60 feet			Angle: 45°		
Logged By: Mark Gartner			Protective Level: Modified Level D		
Coordinates: E -5755.259 N -802.323			Direction (plant grid):		
Depth (ft)	SAMPLE		LITHOLOGIC DESCRIPTION	GRAPHIC LOG	COMMENTS
	INTERVAL	NUMBER			
0	0-7	N/A	Core not retrieved	N/A	N/A
5	7-11	003003SA010	CLAY, brown, 7.5YR 5/4, firm, moist, with gray and orange streaks	~~~~~	13:38
10	11-14	003003SA015	CLAY, light gray, 7.5YR 7/1, firm, moist, with orange streaks	~~~~~	13:42
15					
20	14-28	N/A	Core not retrieved	N/A	N/A
25					
30	28-32	003003SA030 003003WA060	CLAY, white, 7.5YR 8/1, soft, dry, with brown streaks	~~~~~	16:52
35					
40	32-42	N/A	Core not retrieved	N/A	N/A
45	42-46	003003SA045	CLAY, gray, 7.5YR 6/1, very soft, wet, with orange streaks: sand (20%)	~~~~~	9:05 / 4-13-07 Duplicate
			CLAY, gray, 7.5YR 6/1, very soft, wet, with orange streaks: sand (20%)	~~~~~	9:20
50	46-57	N/A	Core not retrieved	N/A	N/A
55	57-60	003003SD060 003003SA060	CLAY, brown, 7.5YR 5/4, hard, moist, with orange, gray and black streaks	~~~~~	10:12 / Duplicate
Total Vertical Depth is 60 feet. Total Linear Depth is 85 feet.			LEGEND: CLAY	~~~~~	
			SAND	~~~~~	

LITHOLOGIC LOG			BORING/WELL ID 003-004-ASB		Page 1 of 1
Facility: Paducah Gaseous Diffusion Plant, Paducah, KY			Site: SWMU- 003		
Project: BGOU RI			Client: USDOE/PRS		
Drilling Contractor: Chase Environmental, LLC			Driller: Jeff Brownfield		
Start Time/Date: 15:10/4-16-07			End Time/Date: 13:41/4-17-07		
Borehole Diameter: 6.25"			Drilling Method: Direct Push Through Augers		
Sampling Method: DT-21 Dual Tube / 2.25" ID Augers			Drill Rig: Geoprobe® 6620DT		
Total Depth (Vertical): 60 feet			Angle: 45°		
Logged By: Mark Gartner			Protective Level: Modified Level D		
Coordinates: E -5634.547 N -910.438			Direction (plant grid): West		
Depth (ft)	SAMPLE		LITHOLOGIC DESCRIPTION	GRAPHIC LOG	COMMENTS
	INTERVAL	NUMBER			
0	0-7	N/A	Core not retrieved	N/A	N/A
5	7-11	003004SA010	CLAY, gray, 7.5YR 6/1, firm, moist, with brown and orange streaks		15:25
10	11-14	003004SA015	CLAY, strong brown, 7.5YR 5/6, firm, moist, with gray and orange streaks		15:30
15					
20	14-28	N/A	Core not retrieved	N/A	N/A
25					
30	28-32	003004SA030 003004WA060	SAND, reddish brown, fine grained, well sorted, with gray streaks		16:07
35	32-42	N/A	Core not retrieved	N/A	N/A
40					
45	42-46	003004SA045	CLAY, brown, 7.5YR 5/4, firm, moist, with gray streaks		12:55 / 4-17-07
50	46-57	N/A	Core not retrieved	N/A	N/A
55	57-60	003004SA060	CLAY, brown, 7.5YR 4/4, hard, moist, with gray streaks		13:41
Total Vertical Depth is 60 feet. Total Linear Depth is 85 feet.			LEGEND: CLAY		
			SAND		

LITHOLOGIC LOG			BORING/WELL ID 003-005-VSB		Page 1 of 1
Facility: Paducah Gaseous Diffusion Plant, Paducah, KY			Site: SWMU- 003		
Project: BGOU RI			Client: USDOE/PRS		
Drilling Contractor: Chase Environmental, LLC			Driller: Jeff Brownfield		
Start Time/Date: 12:54 / 2-7-07			End Time/Date: 13:54 / 2-7-07		
Borehole Diameter: 2.25"			Drilling Method: Direct Push		
Sampling Method: DT-21 Dual Tube			Drill Rig: Geoprobe® 6620DT		
Total Depth (Vertical): 15 feet			Angle: Vertical		
Logged By: Mark Gartner			Protective Level: Modified Level D		
Coordinates: E -5372.989 N -899.566			Direction (plant grid): N/A		
Depth (ft)	SAMPLE		LITHOLOGIC DESCRIPTION	GRAPHIC LOG	COMMENTS
	INTERVAL	NUMBER			
0	0-1	003005SA001	CLAY, gray, 7.5YR 6/1, soft, moist, mottled with orange streaks	~~~~~	13:22
	1-5	003005SA005	CLAY, light gray, 7.5YR 7/1, soft, moist with orange streaks	~~~~~	13:30
5	5-10	003005SA010 003005SD010	CLAY, light brown, 7.5YR 6/3, firm, moist with orange and gray streaks	~~~~~	13:36 Duplicate sample was collected.
10	10-15	003005SA015	CLAY, light gray, 7.5YR 7/1, firm, moist with orange streaks	~~~~~	13:54
Total Depth is 15 feet.			LEGEND: CLAY		~~~~~
			SAND		~~~~~

LITHOLOGIC LOG			BORING/WELL ID 003-006-VSB		Page 1 of 1
Facility: Paducah Gaseous Diffusion Plant, Paducah, KY			Site: SWMU- 003		
Project: BGOU RI			Client: USDOE/PRS		
Drilling Contractor: Chase Environmental, LLC			Driller: Jeff Brownfield		
Start Time/Date: 10:05 / 2-7-07			End Time/Date: 11:10 / 2-7-07		
Borehole Diameter: 2.25"			Drilling Method: Direct Push		
Sampling Method: DT-21 Dual Tube			Drill Rig: Geoprobe® 6620DT		
Total Depth (Vertical): 15 feet			Angle: Vertical		
Logged By: Mark Gartner			Protective Level: Modified Level D		
Coordinates: E -5296.92 N -723.924			Direction (plant grid): N/A		
Depth (ft)	SAMPLE		LITHOLOGIC DESCRIPTION	GRAPHIC LOG	COMMENTS
	INTERVAL	NUMBER			
0	0-1	003006SA001	CLAY, brown, 7.5YR 5/3, firm, moist, mottled with gray and orange streaks: Chert (40%)	~~~~~ ~~~~~ ~~~~~	10:36
	1-5				
5	5-10	003006SA010	CLAY, brown, 7.5YR 5/4, soft, moist with few orange and gray streaks	~~~~~ ~~~~~ ~~~~~	11:04
10	10-15	003006SA015	CLAY, light brown, 7.5YR 6/4, soft, moist with orange streaks	~~~~~ ~~~~~ ~~~~~	11:10
Total Depth is 15 feet.			LEGEND: CLAY	~~~~~ ~~~~~ ~~~~~	
			SAND	~~~~~ ~~~~~ ~~~~~	

LITHOLOGIC LOG			BORING/WELL ID 003-007-VSB		Page 1 of 1
Facility: Paducah Gaseous Diffusion Plant, Paducah, KY			Site: SWMU- 003		
Project: BGOU RI			Client: USDOE/PRS		
Drilling Contractor: Chase Environmental, LLC			Driller: Jeff Brownfield		
Start Time/Date: 8:51 / 2-7-07			End Time/Date: 9:20 / 2-7-07		
Borehole Diameter: 2.25"			Drilling Method: Direct Push		
Sampling Method: DT-21 Dual Tube			Drill Rig: Geoprobe® 6620DT		
Total Depth (Vertical): 15 feet			Angle: Vertical		
Logged By: Mark Gartner			Protective Level: Modified Level D		
Coordinates: E -5210.803 N -533.498			Direction (plant grid): N/A		
Depth (ft)	SAMPLE		LITHOLOGIC DESCRIPTION	GRAPHIC LOG	COMMENTS
	INTERVAL	NUMBER			
0	0-1	003007SA001	CLAY, brown, 7.5YR 5/3, hard, moist, mottled with gray and orange streaks	~~~~~ ~~~~~ ~~~~~	8:51
	1-5				
		003007SA005	CLAY, brown, 7.5YR 5/4, firm, moist, with orange streaks	~~~~~ ~~~~~ ~~~~~	9:00
5	5-10	003007SA010	CLAY, light brown, 7.5YR 6/3, firm, moist with few black streaks	~~~~~ ~~~~~ ~~~~~	9:09
10	10-15	003007SA015	CLAY, light brown, 7.5YR 6/4, firm, moist with orange streaks	~~~~~ ~~~~~ ~~~~~	9:28
Total Depth is 15 feet.			LEGEND: CLAY	~~~~~ ~~~~~ ~~~~~	
			SAND	~~~~~ ~~~~~ ~~~~~	

LITHOLOGIC LOG			BORING/WELL ID 003-008-VSB		Page 1 of 1
Facility: Paducah Gaseous Diffusion Plant, Paducah, KY			Site: SWMU- 003		
Project: BGOU RI			Client: USDOE/PRS		
Drilling Contractor: Chase Environmental, LLC			Driller: Jeff Brownfield		
Start Time/Date: 13:24 / 2-6-07			End Time/Date: 13:47 / 2-6-07		
Borehole Diameter: 2.25"			Drilling Method: Direct Push		
Sampling Method: DT-21 Dual Tube			Drill Rig: Geoprobe® 6620DT		
Total Depth (Vertical): 15 feet			Angle: Vertical		
Logged By: Mark Gartner			Protective Level: Modified Level D		
Coordinates: E -5132.269 N -363.783			Direction (plant grid): N/A		
Depth (ft)	SAMPLE		LITHOLOGIC DESCRIPTION	GRAPHIC LOG	COMMENTS
	INTERVAL	NUMBER			
0	0-1	003008SA001	CLAY, light brown, 7.5YR 6/3, firm, moist, mottled, roots with gray and orange streaks	~~~~~	13:24
	1-5				
5	5-10	003008SA010	CLAY, reddish yellow, 7.5YR 6/8, firm, moist with gray, white, and black streaks	~~~~~	13:39
10	10-15	003008SA015	CLAY, light brown, 7.5YR 6/4, soft, moist with gray and orange streaks	~~~~~	13:47
Total Depth is 15 feet.			LEGEND: CLAY	~~~~~	
			SAND	~~~~~	




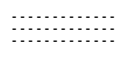



LITHOLOGIC LOG			BORING/WELL ID 003-009-VSB		Page 1 of 1
Facility: Paducah Gaseous Diffusion Plant, Paducah, KY			Site: SWMU- 003		
Project: BGOU RI			Client: USDOE/PRS		
Drilling Contractor: Chase Environmental, LLC			Driller: Jeff Brownfield		
Start Time/Date: 11:48 / 2-6-07			End Time/Date: 12:27 / 2-6-07		
Borehole Diameter: 2.25"			Drilling Method: Direct Push		
Sampling Method: DT-21 Dual Tube			Drill Rig: Geoprobe® 6620DT		
Total Depth (Vertical): 15 feet			Angle: Vertical		
Logged By: Mark Gartner			Protective Level: Modified Level D		
Coordinates: E -4938.326 N -363.171			Direction (plant grid): N/A		
Depth (ft)	SAMPLE		LITHOLOGIC DESCRIPTION	GRAPHIC LOG	COMMENTS
	INTERVAL	NUMBER			
0	0-1	003009SA001	CLAY, brown, 7.5YR 5/3, firm, moist, mottled, roots with a few gray and orange streaks	~~~~~ ~~~~~ ~~~~~	11:48
	1-5				
5	5-10	003009SA010	CLAY, light brown, 7.5YR 6/3, firm, moist, mottled with gray and orange streaks	~~~~~ ~~~~~ ~~~~~	12:17
10	10-15	003009SA015	CLAY, brown, 7.5YR 5/4, soft, moist with gray streaks	~~~~~ ~~~~~ ~~~~~	12:27
Total Depth is 15 feet.			LEGEND: CLAY	~~~~~ ~~~~~ ~~~~~	
			SAND	~~~~~ ~~~~~ ~~~~~	

LITHOLOGIC LOG			BORING/WELL ID 003-010-VSB		Page 1 of 1
Facility: Paducah Gaseous Diffusion Plant, Paducah, KY			Site: SWMU- 003		
Project: BGOU RI			Client: USDOE/PRS		
Drilling Contractor: Chase Environmental, LLC			Driller: Jeff Brownfield		
Start Time/Date: 9:26 / 2-6-07			End Time/Date: 10:05 / 2-6-07		
Borehole Diameter: 2.25"			Drilling Method: Direct Push		
Sampling Method: DT-21 Dual Tube			Drill Rig: Geoprobe® 6620DT		
Total Depth (Vertical): 15 feet			Angle: Vertical		
Logged By: Mark Gartner			Protective Level: Modified Level D		
Coordinates: E -4804.221 N -361.385			Direction (plant grid): N/D		
Depth (ft)	SAMPLE		LITHOLOGIC DESCRIPTION	GRAPHIC LOG	COMMENTS
	INTERVAL	NUMBER			
0	0-1	003010SA001	CLAY, brown, 7.5YR 5/3, hard, moist with a few gray and black streaks	~~~~~	9:26
	1-5	003010SA005	CLAY, brown, 7.5YR 5/4, soft, moist, with gray and orange streaks	~~~~~	9:45
5	5-10	003010SA010	CLAY, brown, 7.5YR 5/4, soft, moist, mottled with gray, black and orange streaks	~~~~~	9:55
10	10-15	003010SA015	CLAY, brown, 7.5YR 5/3, very soft, moist with orange streaks	~~~~~	10:05
Total Depth is 15 feet.			LEGEND: CLAY		~~~~~
			SAND		~~~~~

SWMU 5

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LITHOLOGIC LOG			BORING/WELL ID 005-101-ASB		Page 1 of 1
Facility: Paducah Gaseous Diffusion Plant, Paducah, KY			Site: SWMU- 005		
Project: BGOU RI			Client: USDOE/PRS		
Drilling Contractor: Chase Environmental, LLC			Driller: Ryan Kulik		
Start Time/Date: 11:42/5-17-07			End Time/Date: 8:45/5-18-07		
Borehole Diameter: 6.25"			Drilling Method: Direct Push Through Augers		
Sampling Method: DT-21 Dual Tube / 2.25" ID Augers			Drill Rig: Geoprobe® 6620DT		
Total Depth (Vertical): 60 feet			Angle: 40°		
Logged By: Mark Gartner			Protective Level: Modified Level D		
Coordinates: E -6615.283 N 194.077			Direction (plant grid): South		
Depth (ft)	SAMPLE		LITHOLOGIC DESCRIPTION	GRAPHIC LOG	COMMENTS
	INTERVAL	NUMBER			
0	0-7	N/A	Core not retrieved	N/A	N/A
5	7-11	005101SA010	CLAY, light brown, 7.5YR 6/4, soft, moist with gray and orange streaks	~~~~~	12:01
10	11-14	005101SA015	CLAY, light brown, 7.5YR 6/4, soft, moist, with gray and orange streaks	~~~~~	12:10
15					
20	14-28	N/A	Core not retrieved	N/A	N/A
25					
30	28-32	005101SA030	CLAY, brown, 7.5YR 5/4, firm, moist with gray, black and orange streaks	~~~~~	13:39
35	32-42	N/A	Core not retrieved	N/A	N/A
40					
45	42-46	005101SA45 005101WA060	SAND, tan, medium grained, well sorted	15:00
50	46-57	N/A	Core not retrieved	N/A	N/A
55					
	57-60	005101SA060	SAND, reddish brown, fine grained, well sorted	8:45 / 5-18-07
Total Vertical Depth is 60 feet. Total Linear Depth is 85 feet.			LEGEND: CLAY		~~~~~
			SAND	

LITHOLOGIC LOG			<i>BORING/WELL ID 005-102-ASB</i>		Page 1 of 1
Facility: Paducah Gaseous Diffusion Plant, Paducah, KY			Site: SWMU- 005		
Project: BGOU RI			Client: USDOE/PRS		
Drilling Contractor: Chase Environmental, LLC			Driller: Jeff Brownfield		
Start Time/Date: 8:50/4-21-07			End Time/Date: 14:50/4-21-07		
Borehole Diameter: 6.25"			Drilling Method: Direct Push Through Augers		
Sampling Method: DT-21 Dual Tube / 2.25" ID Augers			Drill Rig: Geoprobe® 6620DT		
Total Depth (Vertical): 60 feet			Angle: 45°		
Logged By: Mark Gartner			Protective Level: Modified Level D		
Coordinates: E -6352.979 N 86.614			Direction (plant grid): West		
Depth (ft)	SAMPLE		LITHOLOGIC DESCRIPTION	GRAPHIC LOG	COMMENTS
	INTERVAL	NUMBER			
0	0-7	N/A	Core not retrieved	N/A	N/A
5	7-11	005102SA010	CLAY, brown, 7.5YR 5/4, soft, moist, with black organic layers and gray streaks		9:05
10	11-14	005102SA015	CLAY, light gray, 7.5YR 7/1, soft, moist, with orange streaks		9:12
15					
20	14-28	N/A	Core not retrieved	N/A	N/A
25					
30	28-32	005102SA030	CLAY, strong brown, 7.5YR 5/4, firm, moist with gray streaks		10:04
35	32-40	N/A	Core not retrieved	N/A	N/A
40	40-41	005102WA060	Core not retrieved	N/A	N/A
	41-42	N/A	Core not retrieved	N/A	N/A
	42-46	005102SA045	SAND, light gray, fine grained: Clay (30%), black layers		12:06
45					
50	46-57	N/A	Core not retrieved	N/A	N/A
55	57-60	005102SA060	CLAY, brown, 7.5YR 5/4, soft, moist: sand (25%)		14:50
Total Vertical Depth is 60 feet. Total Linear Depth is 85 feet.			LEGEND: CLAY		
			SAND		

LITHOLOGIC LOG			BORING/WELL ID 005-103-ASB		Page 1 of 1
Facility: Paducah Gaseous Diffusion Plant, Paducah, KY			Site: SWMU- 005		
Project: BGOU RI			Client: USDOE/PRS		
Drilling Contractor: Chase Environmental, LLC			Driller: Jeff Brownfield		
Start Time/Date: 15:18/4-18-07			End Time/Date: 4-19-07		
Borehole Diameter: 6.25"			Drilling Method: Direct Push Through Augers		
Sampling Method: DT-21 Dual Tube / 2.25" ID Augers			Drill Rig: Geoprobe® 6620DT		
Total Depth (Vertical): 60 feet			Angle: 45°		
Logged By: Mark Gartner			Protective Level: Modified Level D		
Coordinates: E -6350.415 N 3.873			Direction (plant grid): West		
Depth (ft)	SAMPLE		LITHOLOGIC DESCRIPTION	GRAPHIC LOG	COMMENTS
	INTERVAL	NUMBER			
0	0-7	N/A	Core not retrieved	N/A	N/A
5	7-11	005103SA010	CLAY, brown, 7.5YR 5/4, soft, moist, with gray and orange streaks	~~~~~ ~~~~~ ~~~~~	15:30
10	11-14	005103SA015	CLAY, brown, 7.5YR 5/4, firm, dry, with gray streaks	~~~~~ ~~~~~ ~~~~~	15:32
15					
20	14-28	N/A	Core not retrieved	N/A	N/A
25					
30	28-32	005103SA030	Sand, fine grained with gray streaks: Clay (35%)	----- ----- -----	16:33
35	32-42	N/A	Core not retrieved	N/A	N/A
40					
45	42-46	005103SA045	CLAY, brown, 7.5YR 5/4, firm, dry, with gray streaks	~~~~~ ~~~~~ ~~~~~	4-19-07
50	46-57	N/A	Core not retrieved	N/A	N/A
55					
	57-60	005103SA060 005103SD060	CLAY, brown, 7.5YR 5/4, firm, dry, with gray streaks	~~~~~ ~~~~~ ~~~~~	Duplicate
			SAND, medium grained and wet	----- ----- -----	
Total Vertical Depth is 60 feet. Total Linear Depth is 85 feet.			LEGEND: CLAY		~~~~~ ~~~~~ ~~~~~
			SAND		----- ----- -----

LITHOLOGIC LOG			BORING/WELL ID 005-103B-ASB		Page 1 of 1
Facility: Paducah Gaseous Diffusion Plant, Paducah, KY			Site: SWMU- 005		
Project: BGOU RI			Client: USDOE/PRS		
Drilling Contractor: Chase Environmental, LLC			Driller: Jeff Brownfield		
Start Time/Date: 11:49/4-18-07			End Time/Date: 13:35 / 4-18-07		
Borehole Diameter: 6.25"			Drilling Method: Direct Push Through Augers		
Sampling Method: DT-21 Dual Tube / 2.25" ID Augers			Drill Rig: Geoprobe® 6620DT		
Total Depth (Vertical): 60 feet			Angle: 45°		
Logged By: Mark Gartner			Protective Level: Modified Level D		
Coordinates: E N			Direction (plant grid): Southwest		
Depth (ft)	SAMPLE		LITHOLOGIC DESCRIPTION	GRAPHIC LOG	COMMENTS
	INTERVAL	NUMBER			
0	0-7	N/A	Core not retrieved	N/A	N/A
5	7-11	005-103B-8-10 ft	CLAY, reddish yellow, 7.5YR 6/6, firm, moist, with gray and black streaks	~~~~~	12:00 Sample not submitted to lab*
10	11-14	005-103B-13-15 ft	CLAY, brown, 7.5YR 5/4, hard, moist, with gray and black streaks	~~~~~	12:04 Sample not submitted to lab*
15					
20	14-28	N/A	Core not retrieved	N/A	N/A
25					
	28-32	005-103B-28-30 ft	CLAY, strong brown, 7.5YR 5/6, firm, moist, with gray and orange streaks: Sand (10%)	~~~~~	13:35
Total Vertical Depth is 30 feet. Total Linear Depth is 45 feet.			LEGEND: CLAY	~~~~~	
			SAND	~~~~~	





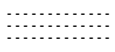


* Angled boring 005-103B was drilled in the wrong direction. Samples 005-103B-8-10 ft and 005-103B-13-15 ft were discarded because the samples were not collected from below buried waste in SWMU 5.

SWMU 6

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LITHOLOGIC LOG			BORING/WELL ID 006-101-ASB		Page 1 of 1
Facility: Paducah Gaseous Diffusion Plant, Paducah, KY			Site: SWMU- 006		
Project: BGOU RI			Client: USDOE/PRS		
Drilling Contractor: Chase Environmental, LLC			Driller: Jeff Brownfield		
Start Time/Date: 15:45/2-26-07			End Time/Date: 9:46/2-28-07		
Borehole Diameter: 6.25"			Drilling Method: Direct Push Through Augers		
Sampling Method: DT-21 Dual Tube / 2.25" ID Augers			Drill Rig: Geoprobe® 6620DT		
Total Depth (Vertical): 60 feet			Angle: 45°		
Logged By: Mark Gartner			Protective Level: Modified Level D		
Coordinates: E -6224.07 N 156.469			Direction (plant grid): South		
Depth (ft)	SAMPLE		LITHOLOGIC DESCRIPTION	GRAPHIC LOG	COMMENTS
	INTERVAL	NUMBER			
0	0-7	N/A	Core not retrieved	N/A	N/A
5	7-11	006101SA010	CLAY, brown, 7.5YR 5/3, firm, moist, with gray and orange streaks	~~~~~	16:00
10	11-14	006101SA015	CLAY, light brown, 7.5YR 6/4, firm, moist, mottled, with orange streaks	~~~~~	16:12
15					
20	14-28	N/A	Core not retrieved	N/A	N/A
25					
30	28-32	006101SA030 006101SD030	CLAY, reddish yellow, 7.5YR 6/6, firm, moist, mottled, with gray and orange streaks	~~~~~	8:30 / 2-27-07
35	32-42	N/A	Core not retrieved	N/A	N/A
40					
45	42-46	006101SA045 006101WA060	CLAY, brown, 7.5YR 4/4, hard, moist, with gray streaks	~~~~~	9:44
50	46-57	N/A	Core not retrieved	N/A	N/A
55	57-60	006101SA060	SAND, fine to medium grained, poorly sorted	9:46 / 2-28-07
Total Vertical Depth is 60 feet. Total Linear Depth is 85 feet.			LEGEND: CLAY		~~~~~
			SAND	

LITHOLOGIC LOG			BORING/WELL ID 006-102-ASB		Page 1 of 1
Facility: Paducah Gaseous Diffusion Plant, Paducah, KY			Site: SWMU- 006		
Project: BGOU RI			Client: USDOE/PRS		
Drilling Contractor: Chase Environmental, LLC			Driller: Jeff Brownfield		
Start Time/Date: 9:25/2-22-07			End Time/Date: 2-26-07		
Borehole Diameter: 6.25"			Drilling Method: Direct Push Through Augers		
Sampling Method: DT-21 Dual Tube / 2.25" ID Augers			Drill Rig: Geoprobe® 6620DT		
Total Depth (Vertical): 60 feet			Angle: 45°		
Logged By: Mark Gartner			Protective Level: Modified Level D		
Coordinates: E -6275.763 N 95.128			Direction (plant grid): East		
Depth (ft)	SAMPLE		LITHOLOGIC DESCRIPTION	GRAPHIC LOG	COMMENTS
	INTERVAL	NUMBER			
0	0-7	N/A	Core not retrieved	N/A	N/A
5	7-11	006102SA010	CLAY, brown, 7.5YR 4/4, firm, moist, mottled, with gray and orange streaks	~~~~~	9:34
10	11-14	006102SA015 006102SD015	CLAY, brown, 7.5YR 5/4, hard, moist, mottled, with gray and orange streaks	~~~~~	9:39
15	14-18	N/A	Core not retrieved	N/A	N/A
	18-19	006102WA060	Core not retrieved	N/A	N/A
20	19-28	N/A	Core not retrieved	N/A	N/A
25					
30	28-32	006102SA030	CLAY, strong brown, 7.5YR 5/6, hard, moist, mottled, with gray and orange streaks	~~~~~	11:00
35	32-42	N/A	Core not retrieved	N/A	N/A
40	42-46	006102SA045	SAND, fine to coarse grained, poorly sorted	14:15 1" of Recovery
45					
50	46-57	N/A	Core not retrieved	N/A	N/A
55	57-60	006102SA060	SAND, fine to medium grained, poorly sorted	2-26-07
Total Vertical Depth is 60 feet. Total Linear Depth is 85 feet.			LEGEND: CLAY		~~~~~
			SAND	

LITHOLOGIC LOG			BORING/WELL ID 006-103-ASB		Page 1 of 1
Facility: Paducah Gaseous Diffusion Plant, Paducah, KY			Site: SWMU- 006		
Project: BGOU RI			Client: USDOE/PRS		
Drilling Contractor: Chase Environmental, LLC			Driller: Jeff Brownfield		
Start Time/Date: 11:40/2-8-07			End Time/Date: 10:11/2-14-07		
Borehole Diameter: 6.25"			Drilling Method: Direct Push Through Augers		
Sampling Method: DT-21 Dual Tube / 2.25" ID Augers			Drill Rig: Geoprobe® 6620DT		
Total Depth (Vertical): 60 feet			Angle: 45°		
Logged By: Mark Gartner			Protective Level: Modified Level D		
Coordinates: E -6217.781 N 68.719			Direction (plant grid): North		
Depth (ft)	SAMPLE		LITHOLOGIC DESCRIPTION	GRAPHIC LOG	COMMENTS
	INTERVAL	NUMBER			
0	0-7	N/A	Core not retrieved	N/A	N/A
5	7-11	006103SA010	CLAY, light brown, 7.5YR 6/4, firm, moist, with gray, black, and orange streaks		12:02
10	11-14	006103SA015	CLAY, light brown, 7.5YR 6/4, firm, moist, with gray, black, and orange streaks		12:08
15					
20	14-28	N/A	Core not retrieved	N/A	N/A
25					
30	28-32	006103SA030	CLAY, brown, 7.5YR 5/4, firm, moist, with gray streaks: Sand (5%)		13:37
35	32-42	N/A	Core not retrieved	N/A	N/A
40					
45	42-46	006103SA045	CLAY, strong brown, 7.5YR 5/6, hard, moist, with gray and orange streaks		14:04 / 2-9-07
45	46-50	N/A	Core not retrieved	N/A	N/A
50	50-51	006103WA060	Core not retrieved	N/A	N/A
50	46-57	N/A	Core not retrieved	N/A	N/A
55					
	57-60	006103SA060	CLAY, strong brown, 7.5YR 5/6, firm, moist: sand (10%), white, fine grained		10:11 / 2-14-07
Total Vertical Depth is 60 feet. Total Linear Depth is 85 feet.			LEGEND: CLAY		
			SAND		

LITHOLOGIC LOG			BORING/WELL ID 006-104-ASB		Page 1 of 1
Facility: Paducah Gaseous Diffusion Plant, Paducah, KY			Site: SWMU- 006		
Project: BGOU RI			Client: USDOE/PRS		
Drilling Contractor: Chase Environmental, LLC			Driller: Jeff Brownfield		
Start Time/Date: 9:50/2-19-07			End Time/Date: 14:05/2-21-07		
Borehole Diameter: 6.25"			Drilling Method: Direct Push Through Augers		
Sampling Method: DT-21 Dual Tube / 2.25" ID Augers			Drill Rig: Geoprobe® 6620DT		
Total Depth (Vertical): 60 feet			Angle: 45°		
Logged By: Mark Gartner / Todd Mills			Protective Level: Modified Level D		
Coordinates: E -6178.211 N 93.911			Direction (plant grid): Northeast		
Depth (ft)	SAMPLE		LITHOLOGIC DESCRIPTION	GRAPHIC LOG	COMMENTS
	INTERVAL	NUMBER			
0	0-7	N/A	Core not retrieved	N/A	N/A
5	7-11	006104SA010	CLAY, brown, 7.5YR 5/4, firm, moist, with gray streaks	~~~~~	10:11
10	11-14	006104SA015	CLAY, brown, 7.5YR 5/3, hard, moist, with gray streaks	~~~~~	
15					
20	14-28	N/A	Core not retrieved	N/A	N/A
25					
30	28-32	006104SA030	CLAY, strong brown, 7.5YR 4/6, hard, dry, with sand streaks	~~~~~	13:41
35					
40	32-42	N/A	Core not retrieved	N/A	N/A
45	42-46	006104SA045 006104WA060	CLAY, pinkish gray, 7.5YR 7/2, very stiff, moist, from 43 to 45 feet	~~~~~	
50					
55	46-57	N/A	Core not retrieved	N/A	N/A
	57-60	N/A	Core not retrieved	N/A	13:09 Attempt failed 57-60' sample
60	60-63	N/A	Core not retrieved	N/A	13:50 Attempt failed 64-66' sample
	63-65	006104SA060	SAND, fine grained, reddish brown, well sorted, black streaks	14:05
Total Vertical Depth is 60 feet. Total Linear Depth is 85 feet.			LEGEND: CLAY	~~~~~	
			SAND	

SWMU 7

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LITHOLOGIC LOG			BORING/WELL ID 007-001-ASB		Page 1 of 1
Facility: Paducah Gaseous Diffusion Plant, Paducah, KY			Site: SWMU- 007		
Project: BGOU RI			Client: USDOE/PRS		
Drilling Contractor: Chase Environmental, LLC			Driller: Ryan Kulik		
Start Time/Date: 09:54/3-28-07			End Time/Date: 09:41/3-29-07		
Borehole Diameter: 6.25"			Drilling Method: Direct Push Through Augers		
Sampling Method: DT-21 Dual Tube / 2.25" ID Augers			Drill Rig: Geoprobe® 6620DT		
Total Depth (Vertical): 60 feet			Angle: 45°		
Logged By: Mark Gartner			Protective Level: Modified Level D		
Coordinates: E -6271.047 N 913.003			Direction (plant grid): Southeast		
Depth (ft)	SAMPLE		LITHOLOGIC DESCRIPTION	GRAPHIC LOG	COMMENTS
	INTERVAL	NUMBER			
0	0-7	N/A	Core not retrieved	N/A	N/A
5	7-11	007001SD010	CLAY, strong brown, 7.5YR 5/8, hard, moist, mottled, with gray and orange streaks	~~~~~	10:14 Duplicate
10		007001SA010	CLAY, brown, 7.5YR 5/3, firm, moist, mottled, with black and orange streaks	~~~~~	10:17
		007001SA015	CLAY, brown, 7.5YR 5/3, firm, moist, mottled, with black and orange streaks	~~~~~	10:24
15	14-28	N/A	Core not retrieved	N/A	N/A
20					
25					
30	28-32	007001SA030	CLAY, gray, 7.5YR 6/1, firm, moist, with orange streaks: Sand (5%)	~~~~~	12:45
35	32-42	N/A	Core not retrieved	N/A	N/A
40	42-46	007001SA045 007001WA060	SAND, gray, medium grained, poorly sorted: Chert (10%) angular to subangular, ranging from ¼ to ½ inch across	13:44
45					
50	46-57	N/A	Core not retrieved	N/A	N/A
55	57-60	007001SA060	SAND, gray, medium grained, poorly sorted: Chert (4 inch seam), angular, ½ to ¾ inch across	9:22 / 3-29-07 Min. sample recovered.
60	60-62.5		SAND, gray, medium grained, poorly sorted: Chert (4 inch seam), angular, ½ to ¾ inch across	9:41: Additional sample required.
	62.5-65		N/A	Core not retrieved	N/A
Total Vertical Depth is 62.5 feet. Total Linear Depth is 87.5 feet.			LEGEND: CLAY		~~~~~
			SAND	

LITHOLOGIC LOG			BORING/WELL ID 007-002-ASB		Page 1 of 1
Facility: Paducah Gaseous Diffusion Plant, Paducah, KY			Site: SWMU- 007		
Project: BGOU RI			Client: USDOE/PRS		
Drilling Contractor: Chase Environmental, LLC			Driller: Ryan Kulik		
Start Time/Date: 12:04/3-27-07			End Time/Date: 16:42/3-27-07		
Borehole Diameter: 6.25"			Drilling Method: Direct Push Through Augers		
Sampling Method: DT-21 Dual Tube / 2.25" ID Augers			Drill Rig: Geoprobe® 6620DT		
Total Depth (Vertical): 60 feet			Angle: 45°		
Logged By: Mark Gartner			Protective Level: Modified Level D		
Coordinates: E -6328.252 N 859.906			Direction (plant grid): Southeast		
Depth (ft)	SAMPLE		LITHOLOGIC DESCRIPTION	GRAPHIC LOG	COMMENTS
	INTERVAL	NUMBER			
0	0-7	N/A	Core not retrieved	N/A	N/A
5	7-11	007002SA010	CLAY, gray, 7.5YR 6/1, firm, moist, mottled, with gray streaks	~~~~~ ~~~~~ ~~~~~	12:25
10	11-14	007002SA015	CLAY, light brown, 7.5YR 6/4, firm, moist, mottled, with gray and orange streaks	~~~~~ ~~~~~ ~~~~~	12:39
15					
20	14-28	N/A	Core not retrieved	N/A	N/A
25					
30	28-32	007002SA030	CLAY, strong brown, 7.5YR 5/6, firm, moist, mottled with gray and orange streaks: Sand (5%)	~~~~~ ~~~~~ ~~~~~	14:38
35	32-42	N/A	Core not retrieved	N/A	N/A
40	42-46	007002SA045	SAND, gray, medium grained, well sorted, moist	----- ----- -----	15:38
45	46-50	N/A			
50	50-51	007002WA060			
	51-57	N/A	Core not retrieved	N/A	N/A
55	57-60	007002SA060	CLAY, light brown, 7.5YR 6/3, soft, moist with gray and orange streaks: Sand (15%) fine grained	~~~~~ ~~~~~ ~~~~~	16:42
Total Vertical Depth is 60 feet. Total Linear Depth is 85 feet.			LEGEND: CLAY		~~~~~ ~~~~~ ~~~~~
			SAND		----- ----- -----

LITHOLOGIC LOG			BORING/WELL ID 007-003-ASB		Page 1 of 1
Facility: Paducah Gaseous Diffusion Plant, Paducah, KY			Site: SWMU- 007		
Project: BGOU RI			Client: USDOE/PRS		
Drilling Contractor: Chase Environmental, LLC			Driller: Ryan Kulik		
Start Time/Date: 8:35/3-19-07			End Time/Date: 11:08/3-21-07		
Borehole Diameter: 6.25"			Drilling Method: Direct Push Through Augers		
Sampling Method: DT-21 Dual Tube / 2.25" ID Augers			Drill Rig: Geoprobe® 6620DT		
Total Depth (Vertical): 60 feet			Angle: 45°		
Logged By: Mark Gartner			Protective Level: Modified Level D		
Coordinates: E N			Direction (plant grid): Southeast		
Depth (ft)	SAMPLE		LITHOLOGIC DESCRIPTION	GRAPHIC LOG	COMMENTS
	INTERVAL	NUMBER			
0	0-7	N/A	Core not retrieved	N/A	N/A
5	7-11	007003SA010	CLAY, brown, 7.5YR 5/2, soft, moist, with few orange streaks	~~~~~	8:40
10	11-14	007003SA015	CLAY, pinkish gray, 7.5YR 6/2, firm, moist, mottled with gray and orange streaks	~~~~~	8:47
15	14-28	N/A	Core not retrieved	N/A	N/A
20	28-32	007003SA030	CLAY, strong brown, 7.5YR 5/6, soft, moist, mottled with gray and orange streaks	~~~~~	9:28
25	32-42	N/A	Core not retrieved	N/A	N/A
30	42-46	007003SA045 007003WA060	SAND, fine to medium grained, poorly sorted, water with angular chert up to ¼ across	13:48 / 3-20-07
35	46-57	N/A	Core not retrieved	N/A	N/A
40	57-60	007003SA060	CLAY, reddish yellow, 7.5YR 6/8, hard, moist with gray streaks	~~~~~	11:08 / 3-21-07
Total Vertical Depth is 60 feet. Total Linear Depth is 85 feet.			LEGEND: CLAY		~~~~~
			SAND	

LITHOLOGIC LOG			BORING/WELL ID 007-004-ASB		Page 1 of 1
Facility: Paducah Gaseous Diffusion Plant, Paducah, KY			Site: SWMU- 007		
Project: BGOU RI			Client: USDOE/PRS		
Drilling Contractor: Chase Environmental, LLC			Driller: Ryan Kulik		
Start Time/Date: 8:56/3-26-07			End Time/Date: 15:30/3-26-07		
Borehole Diameter: 6.25"			Drilling Method: Direct Push Through Augers		
Sampling Method: DT-21 Dual Tube / 2.25" ID Augers			Drill Rig: Geoprobe® 6620DT		
Total Depth (Vertical): 60 feet			Angle: 45°		
Logged By: Mark Gartner			Protective Level: Modified Level D		
Coordinates: E -6711.864 N 794.304			Direction (plant grid): East		
Depth (ft)	SAMPLE		LITHOLOGIC DESCRIPTION	GRAPHIC LOG	COMMENTS
	INTERVAL	NUMBER			
0	0-7	N/A	Core not retrieved	N/A	N/A
5	7-11	007004SA010	CLAY, gray, 7.5YR 6/1, firm, moist, with few orange streaks	~~~~~	9:12
10	11-14	007004SA015	CLAY, gray, 7.5YR 5/1, soft, moist with orange streaks	~~~~~	9:18 40% recovery
15	14-18		CLAY, gray, 7.5YR 5/1, soft, moist with orange streaks	~~~~~	Additional sample
20	18-28	N/A	Core not retrieved	N/A	N/A
25	28-32	007004SA030	SAND, medium to coarse grained, poorly sorted: Chert (25%), angular, up to ½ inch across	12:12
35	32-42	N/A	Core not retrieved	N/A	N/A
40	42-46	007004SA045	CLAY, light brown, 7.5YR6/3, soft, moist, with gray streaks	13:30
45	46-57	N/A	Core not retrieved	N/A	N/A
55	57-60	007004SA060	CLAY, brown, 7.5YR 5/3, hard, moist, mottled with black streaks	~~~~~	15:30
Total Vertical Depth is 60 feet. Total Linear Depth is 85 feet.			LEGEND: CLAY		~~~~~
			SAND	

LITHOLOGIC LOG			BORING/WELL ID 007-005-ASB		Page 1 of 1		
Facility: Paducah Gaseous Diffusion Plant, Paducah, KY			Site: SWMU- 007				
Project: BGOU RI			Client: USDOE/PRS				
Drilling Contractor: Chase Environmental, LLC			Driller: Jeff Brownfield				
Start Time/Date: 15:24/3-12-07			End Time/Date: 3-13-07				
Borehole Diameter: 6.25"			Drilling Method: Direct Push Through Augers				
Sampling Method: DT-21 Dual Tube / 2.25" ID Augers			Drill Rig: Geoprobe® 6620DT				
Total Depth (Vertical): 60 feet			Angle: 45°				
Logged By: Mark Gartner			Protective Level: Modified Level D				
Coordinates: E -6713.919 N 838.269			Direction (plant grid): Southwest				
Depth (ft)	SAMPLE		LITHOLOGIC DESCRIPTION	GRAPHIC LOG	COMMENTS		
	INTERVAL	NUMBER					
0	0-7	N/A	Core not retrieved	N/A	N/A		
5	7-11	007005SA010	CLAY, light gray, 7.5YR 7/1, firm, moist, mottled with orange streaks	~~~~~	15:37		
10	11-14	007005SA015	GRAVEL	Gravel collapsed the sample liner – no recovery		
15	14-18		CLAY, light brown, 7.5YR 6/4, firm, moist, mottled with gray streaks	~~~~~	16:20		
20	18-28	N/A	Core not retrieved	N/A	N/A		
25	28-32	007005SA030	CLAY, gray, 7.5YR 6/1, soft, moist, mottled with orange streaks	~~~~~	9:04 / 3-13-07		
30	32-42	N/A	Core not retrieved	N/A	N/A		
35	42-46	007005SA045 007005WA060	SAND, fine to coarse grained, poorly sorted, wet, iron stained: Chert (5%), subangular	9:54		
40	46-57	N/A	Core not retrieved	N/A	N/A		
45	57-60	007005SA060	CLAY, strong brown, 7.5YR 5/6, firm, moist, mottled with gray streaks with sand seams	~~~~~			
Total Vertical Depth is 60 feet. Total Linear Depth is 85 feet.			LEGEND :	GRAVEL	CLAY	~~~~~
						SAND

LITHOLOGIC LOG			BORING/WELL ID 007-006-ASB		Page 1 of 1
Facility: Paducah Gaseous Diffusion Plant, Paducah, KY			Site: SWMU- 007		
Project: BGOU RI			Client: USDOE/PRS		
Drilling Contractor: Chase Environmental, LLC			Driller: Ryan Kulik		
Start Time/Date: 15:42/3-21-07			End Time/Date: 15:15 / 3-23-07		
Borehole Diameter: 6.25"			Drilling Method: Direct Push Through Augers		
Sampling Method: DT-21 Dual Tube / 2.25" ID Augers			Drill Rig: Geoprobe® 6620DT		
Total Depth (Vertical): 60 feet			Angle: 45°		
Logged By: Mark Gartner			Protective Level: Modified Level D		
Coordinates: E -6786.053 N 876.721			Direction (plant grid): Northeast		
Depth (ft)	SAMPLE		LITHOLOGIC DESCRIPTION	GRAPHIC LOG	COMMENTS
	INTERVAL	NUMBER			
0	0-7	N/A	Core not retrieved	N/A	N/A
5	7-11	007006SA010	CLAY, brown, 7.5YR 5/4, hard, moist with gray and orange streaks	~~~~~	16:21
10	11-14	007006SA015	CLAY, light brown, 7.5YR 6/4, firm, moist with gray streaks	~~~~~	16:45
15					
20	14-28	N/A	Core not retrieved	N/A	N/A
25					
30	28-32	007006SA030	CLAY, reddish yellow, 7.5YR 6/6, hard, moist with gray and orange streaks: Sand (5%)	~~~~~	9:22 / 3-22-07
35	32-42	N/A	Core not retrieved	N/A	N/A
40					
	42-46	N/A	N/A	N/A	No recovery
45	46-51	007006SA045	Clay, brown, 7.5YR 5/3, hard, mist, mottled, with orange and gray streaks	~~~~~	13:00 / 3-23-07
50	51-57	N/A	Core not retrieved	N/A	N/A
55	57-60	007006SA060	Clay, brown, 7.5YR 5/3, hard, mist, mottled, with orange and gray streaks	~~~~~	15:15
Total Vertical Depth is 60 feet. Total Linear Depth is 85 feet.			LEGEND: CLAY		~~~~~
			SAND		~~~~~

LITHOLOGIC LOG			BORING/WELL ID 007-007-ASB		Page 1 of 1
Facility: Paducah Gaseous Diffusion Plant, Paducah, KY			Site: SWMU- 007		
Project: BGOU RI			Client: USDOE/PRS		
Drilling Contractor: Chase Environmental, LLC			Driller: Jeff Brownfield		
Start Time/Date: 13:56/3-14-07			End Time/Date: 10:10 / 3-16-07		
Borehole Diameter: 6.25"			Drilling Method: Direct Push Through Augers		
Sampling Method: DT-21 Dual Tube / 2.25" ID Augers			Drill Rig: Geoprobe® 6620DT		
Total Depth (Vertical): 60 feet			Angle: 45°		
Logged By: Mark Gartner			Protective Level: Modified Level D		
Coordinates: E -6602.836 N 849.08			Direction (plant grid): North		
Depth (ft)	SAMPLE		LITHOLOGIC DESCRIPTION	GRAPHIC LOG	COMMENTS
	INTERVAL	NUMBER			
0	0-7	N/A	Core not retrieved	N/A	N/A
5	7-11	007007SA010	CLAY, gray, 7.5YR 6/1, soft, moist	~~~~~ ~~~~~ ~~~~~	14:38
10	11-14	007007SA015	CLAY, light brown, 7.5YR 6/4, firm, moist with gray and orange streaks	~~~~~ ~~~~~ ~~~~~	14:42
15	14-28	N/A	Core not retrieved	N/A	N/A
20	28-32	007007SA030	CLAY, reddish yellow, 7.5YR 6/6, firm, moist with gray and orange streaks: Sand (5%)	~~~~~ ~~~~~ ~~~~~	16:32
25	32-42	N/A	Core not retrieved	N/A	N/A
30	42-46	007007SA045 007007SD045 007007WA060 007007WD060	SAND, fine grained up to 1/2" across, poorly sorted, water	13:26 / 3-15-07
35	46-57	N/A	Core not retrieved	N/A	N/A
40	57-60	007007SA060	SAND , fine to medium grained with 5% clay	10:10 / 3-16-07
Total Vertical Depth is 60 feet.			LEGEND: CLAY		~~~~~ ~~~~~ ~~~~~
Total Linear Depth is 83 feet.			SAND	

LITHOLOGIC LOG			BORING/WELL ID 007-008-ASB		Page 1 of 1
Facility: Paducah Gaseous Diffusion Plant, Paducah, KY			Site: SWMU- 007		
Project: BGOU RI			Client: USDOE/PRS		
Drilling Contractor: Chase Environmental, LLC			Driller: Jeff Brownfield		
Start Time/Date: 9:20/4-2-07			End Time/Date: 16:41 / 4-2-07		
Borehole Diameter: 6.25"			Drilling Method: Direct Push Through Augers		
Sampling Method: DT-21 Dual Tube / 2.25" ID Augers			Drill Rig: Geoprobe® 6620DT		
Total Depth (Vertical): 60 feet			Angle: 45°		
Logged By: Mark Gartner			Protective Level: Modified Level D		
Coordinates: E -6064.711 N 934.282			Direction (plant grid): Southeast		
Depth (ft)	SAMPLE		LITHOLOGIC DESCRIPTION	GRAPHIC LOG	COMMENTS
	INTERVAL	NUMBER			
0	0-7	N/A	Core not retrieved	N/A	N/A
5	7-11	007008SA010	CLAY, gray, 7.5YR 6/1, soft, moist, with orange streaks	~~~~~	9:30
10	11-14	007008SA015	CLAY, light gray, 7.5YR 7/1, firm, moist, mottled, with orange streaks	~~~~~	9:35
15	14-28	N/A	Core not retrieved	N/A	N/A
20	28-32	007008SA030	CLAY, strong brown, 7.5YR 5/6, firm, moist, mottled with gray streaks	~~~~~	10:34
25	32-42	N/A	Core not retrieved	N/A	N/A
30	42-46	007008SA045 007008WA060	SAND, medium grained, wet: Chert (5%) angular, 1/8" across	12:42
35	46-57	N/A	Core not retrieved	N/A	N/A
40	57-60	007008SA060	SAND, medium grained with 10% clay: Chert (few), angular up to 1/2" across	16:41
Total Vertical Depth is 60 feet. Total Linear Depth is 83 feet.			LEGEND: CLAY		~~~~~
			SAND	

LITHOLOGIC LOG			BORING/WELL ID 007-009-VSB		Page 1 of 1
Facility: Paducah Gaseous Diffusion Plant, Paducah, KY			Site: SWMU- 007		
Project: BGOU RI			Client: USDOE/PRS		
Drilling Contractor: Chase Environmental, LLC			Driller: Jeff Brownfield		
Start Time/Date: 12:05/4-30-07			End Time/Date: 5-8-07		
Borehole Diameter: 6.25"			Drilling Method: Direct Push Through Augers		
Sampling Method: DT-21 Dual Tube / 2.25" ID Augers			Drill Rig: CME High Torque 55 and Geoprobe® 6620DT		
Total Depth (Vertical): 96 feet			Angle: Vertical		
Logged By: Mark Gartner			Protective Level: Modified Level D		
Coordinates: E -6785.376 N 964.982			Direction (plant grid): N/A		
Depth (ft)	SAMPLE		LITHOLOGIC DESCRIPTION	GRAPHIC LOG	COMMENTS
	INTERVAL	NUMBER			
0	0-1	007009SA001	CLAY, light brown, 7.5YR 6/3, hard, dry with black and orange streaks	~~~~~	12:15: Pused two additional liners
	1-5				
5	5-10	007009SA010	CLAY, light brown, 7.5YR 6/3, firm, moist with gray and orange streaks	~~~~~	12:24
10	10-15	007009SA015	CLAY, light brown, 7.5YR 6/4, soft, moist with orange streaks	~~~~~	12:27
15	15-25	N/A	Core not retrieved	N/A	N/A
20					
25	25-30	007009SA030	CLAY, light brown, 7.5YR 6/3, firm, moist, mottled with gray and orange streaks	~~~~~	12:49
30	30-40	N/A	Core not retrieved	N/A	N/A
35					
40	40-45	007009SA045	CLAY, light brown, 7.5YR 4/3, very hard, moist, with gray streaks	~~~~~	13:37
45	45-55	007009WA030	Core not retrieved	N/A	N/A
50					
55	55-60	007009SA60	CLAY, brown, 7.5YR 4/4, firm, moist, with gray streaks	~~~~~	10:29 / 5-1-07
Total Vertical Depth is 96 feet. Water samples collected at the following depths: 69, 80, 90, 90D ft.			LEGEND: CLAY		~~~~~
			SAND		~~~~~

LITHOLOGIC LOG			BORING/WELL ID 007-010-VSB		Page 1 of 1
Facility: Paducah Gaseous Diffusion Plant, Paducah, KY			Site: SWMU- 007		
Project: BGOU RI			Client: USDOE/PRS		
Drilling Contractor: Chase Environmental, LLC			Driller: Jeff Brownfield		
Start Time/Date: 8:45/5-10-07			End Time/Date: 5-15-07		
Borehole Diameter: 6.25"			Drilling Method: Direct Push Through Augers		
Sampling Method: DT-21 Dual Tube / 2.25" ID Augers			Drill Rig: CME High Torque 55 and Geoprobe® 6620DT		
Total Depth (Vertical): 100 feet			Angle: Vertical		
Logged By: Mark Gartner			Protective Level: Modified Level D		
Coordinates: E -6543.575 N 957.997			Direction (plant grid): N/A		
Depth (ft)	SAMPLE		LITHOLOGIC DESCRIPTION	GRAPHIC LOG	COMMENTS
	INTERVAL	NUMBER			
0	0-1	007010SA001	Gravel – fill material	8:45: Pushed 4 liners. 780.00 dpm
	1-5	007010SA005	CLAY, white, 7.5YR 8/1, soft, moist	~~~~~	8:50
5	5-10	007010SA010	CLAY, white, 7.5YR 8/1, firm, moist, mottled with orange streaks	~~~~~	9:20
10	10-15	007010SA015	CLAY, pinkish white, 7.5YR 6/4, soft, moist with gray streaks	~~~~~	12:19
15	15-25	N/A	Core not retrieved	N/A	N/A
20					
25	25-30	007010SA030	CLAY, brown, 7.5YR 5/4, hard, moist, mottled with gray and orange streaks	~~~~~	12:35
30	30-40	N/A	Core not retrieved	N/A	N/A
35					
40	40-45	0070110SA045 007010WA030	CLAY, brown, 7.5YR 5/4, firm, moist, with orange streaks: Sand (10%)	~~~~~	9:30 / 5-11-07
45	45-55	N/A	Core not retrieved	N/A	N/A
50					
55	55-60	007010SA060 007010WA060	SAND, fine grained reddish brown with black streaks: Silt (10%)	15:45
Total Vertical Depth is 100 ft Water samples collected at the following depths: 66, 80, 90, 100 ft			LEGEND: GRAVEL	
				CLAY	~~~~~
				SAND

LITHOLOGIC LOG			BORING/WELL ID 007-011-VSB		Page 1 of 1
Facility: Paducah Gaseous Diffusion Plant, Paducah, KY			Site: SWMU- 007		
Project: BGOU RI			Client: USDOE/PRS		
Drilling Contractor: Chase Environmental, LLC			Driller: Jeff Brownfield		
Start Time/Date: 14:21 / 4-24-07			End Time/Date: 4-26-07		
Borehole Diameter: 8.25"			Drilling Method: 4.25" ID Augers		
Sampling Method: Split Spoons			Drill Rig: CME High Torque 55		
Total Depth (Vertical): 100 feet			Angle: Vertical		
Logged By: Mark Gartner			Protective Level: Modified Level D		
Coordinates: E -6282.4 N 869.1			Direction (plant grid): N/A		
Depth (ft)	SAMPLE		LITHOLOGIC DESCRIPTION	GRAPHIC LOG	COMMENTS
	INTERVAL	NUMBER			
0	0-1	N/A	Sample not Collected – Soil all rock and Gravel	15:09
	1-5	007011SA005	CLAY, gray, 7.5YR 6/1, firm, moist, mottled with orange streaks	~~~~~	15:20
5	5-10	007010SA010	CLAY, white, 7.5YR 8/1, firm, moist, mottled with orange streaks	~~~~~	15:30
10	10-15	007010SA015	CLAY, light gray, 7.5YR 7/1, soft, moist, variegated gray and orange	~~~~~	15:53
15	15-25	N/A	Core not retrieved	N/A	N/A
20					
25	25-30	007011SA030 007011SD030	CLAY, light gray, 7.5YR 7/1, soft, moist, variegated gray and orange	~~~~~	16:04
30	30-40	N/A	Core not retrieved	N/A	N/A
35					
40	40-45	007011SA045 007011WA030	SAND, fine grained, gray: Chert (15%) sub angular, 1/8" to 1/2" across	8:20 / 4-25-07
45	45-55	N/A	Core not retrieved	N/A	N/A
50					
55	55-60	007011SA060 007011WA060	SAND, medium grained, gray: Chert (5%), angular, 1/8" to 3" across	13:35
Total Vertical Depth is 100 feet. Water samples collected at the following depths: 70, 80, 90 ft.			LEGEND: GRAVEL	
				CLAY	~~~~~
				SAND

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SWMU 30

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LITHOLOGIC LOG			BORING/WELL ID 030-001-ASB		Page 1 of 1
Facility: Paducah Gaseous Diffusion Plant, Paducah, KY			Site: SWMU- 030		
Project: BGOU RI			Client: USDOE/PRS		
Drilling Contractor: Chase Environmental, LLC			Driller: Jeff Brownfield		
Start Time/Date: 8:34/3-5-07			End Time/Date: 14:40/3-5-07		
Borehole Diameter: 6.25"			Drilling Method: Direct Push Through Augers		
Sampling Method: DT-21 Dual Tube / 2.25" ID Augers			Drill Rig: Geoprobe® 6620DT		
Total Depth (Vertical): 60 feet			Angle: 45°		
Logged By: Mark Gartner			Protective Level: Modified Level D		
Coordinates: E -7187.038 N 976.622			Direction (plant grid): Southeast		
Depth (ft)	SAMPLE		LITHOLOGIC DESCRIPTION	GRAPHIC LOG	COMMENTS
	INTERVAL	NUMBER			
0	0-7	N/A	Core not retrieved	N/A	N/A
5	7-11	030001SA010	CLAY, light brown, 7.5YR 6/4, firm, moist, mottled, with gray and orange streaks	~~~~~ ~~~~~ ~~~~~	8:52
10	11-14	030001SA015	CLAY, light brown, 7.5YR 6/4, firm, moist, mottled, with gray and orange streaks	~~~~~ ~~~~~ ~~~~~	8:56
15	14-18		CLAY, light brown, 7.5YR 6/4, firm, moist, mottled, with gray and orange streaks		9:04 Additional sample was required
20	18-28	N/A	Core not retrieved	N/A	N/A
25					
30	28-32	030001SA030	SAND, iron stained, fine grained, well sorted, slight moisture	9:31
35	32-42	N/A	Core not retrieved	N/A	N/A
40					
45	42-46	030001SA045	CLAY, light brown, 7.5YR 6/3, hard, moist, with red and black streaks	~~~~~ ~~~~~ ~~~~~	10:20
50	46-57	N/A	Core not retrieved	N/A	N/A
55					
	57-60	030001SA060	CLAY, light brown, 7.5YR 6/4, soft, moist, with gray and orange streaks: Sand from 84 to 85', fine to medium grained , poorly sorted	~~~~~ ~~~~~ ~~~~~	14:40
Total Vertical Depth is 60 feet. Total Linear Depth is 85 feet.			LEGEND: CLAY		~~~~~ ~~~~~ ~~~~~
			SAND	

LITHOLOGIC LOG			BORING/WELL ID 030-002-ASB		Page 1 of 1
Facility: Paducah Gaseous Diffusion Plant, Paducah, KY			Site: SWMU- 030		
Project: BGOU RI			Client: USDOE/PRS		
Drilling Contractor: Chase Environmental, LLC			Driller: Jeff Brownfield		
Start Time/Date: 8:30/3-6-07			End Time/Date: 13:31/3-6-07		
Borehole Diameter: 2.25"			Drilling Method: Direct Push		
Sampling Method: DT-21 Dual Tube			Drill Rig: Geoprobe® 6620DT		
Total Depth (Vertical): 60 feet			Angle: 45°		
Logged By: Mark Gartner			Protective Level: Modified Level D		
Coordinates: E -7035.76 N 883.462			Direction (plant grid): Northwest		
Depth (ft)	SAMPLE		LITHOLOGIC DESCRIPTION	GRAPHIC LOG	COMMENTS
	INTERVAL	NUMBER			
0	0-7	N/A	Core not retrieved	N/A	N/A
5	7-11	030002SA010	CLAY, reddish yellow, 7.5YR 6/6, hard, moist, mottled, with gray and orange streaks	~~~~~	8:48
10	11-14	030002SA015	CLAY, reddish yellow, 7.5YR 6/6, firm, moist, mottled, with gray and orange streaks	~~~~~	8:54
15	14-18		CLAY, light brown, 7.5YR 6/3, firm, moist, with gray and orange streaks	~~~~~	8:56 Additional sample required
20	18-28	N/A	Core not retrieved	N/A	N/A
25	28-32	030002SA030	CLAY, light brown, 7.5YR 6/4, hard, moist, with gray and orange streaks: Sand from 40 to 41', medium to coarse grained, poorly sorted, iron stained	~~~~~	9:19
35	32-42	N/A	Core not retrieved	N/A	N/A
40	42-46	030002SA045	CLAY, brown, 7.5YR 5/3, hard, moist, mottled, with gray and orange streaks	~~~~~	10:41
45	46-50	030002SD045	CLAY, brown, 7.5YR 5/3, hard, moist, mottled, with gray and orange streaks	~~~~~	Duplicate
50	50-57	N/A	Core not retrieved	N/A	N/A
55	57-60	030002SA060	CLAY, light gray, 7.5YR 7/1, hard, moist, with orange streaks: Sand (5%)	~~~~~	13:31 / 3-6-07
Total Vertical Depth is 60 feet. Total Linear Depth is 85 feet.			LEGEND: CLAY		~~~~~
			SAND		~~~~~






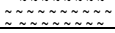
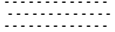
LITHOLOGIC LOG			BORING/WELL ID 030-003-ASB		Page 1 of 1
Facility: Paducah Gaseous Diffusion Plant, Paducah, KY			Site: SWMU- 030		
Project: BGOU RI			Client: USDOE/PRS		
Drilling Contractor: Chase Environmental, LLC			Driller: Jeff Brownfield		
Start Time/Date: 8:30/3-7-07			End Time/Date: 13:11/3-8-07		
Borehole Diameter: 6.25"			Drilling Method: Direct Push Through Augers		
Sampling Method: DT-21 Dual Tube / 2.25" ID Augers			Drill Rig: Geoprobe® 6620DT		
Total Depth (Vertical): 60 feet			Angle: 45°		
Logged By: Mark Gartner			Protective Level: Modified Level D		
Coordinates: E -6939.763 N 881.58			Direction (plant grid): Southeast		
Depth (ft)	SAMPLE		LITHOLOGIC DESCRIPTION	GRAPHIC LOG	COMMENTS
	INTERVAL	NUMBER			
0	0-7	N/A	Core not retrieved	N/A	N/A
5	7-11	030003SA010	CLAY, brown, 7.5YR 4/4, firm, moist, with gray streaks	~~~~~	8:35
10	11-14	030003SA015	CLAY, brown, 7.5YR 5/3, firm, moist, with orange streaks	~~~~~	8:48
15	14-28	N/A	Core not retrieved	N/A	N/A
20	28-32	030003SA030	CLAY, reddish yellow, 7.5YR 6/6, very hard, moist, with gray streak	~~~~~	14:56
25	32-42	N/A	Core not retrieved	N/A	N/A
30	42-46	030003SA045	CLAY, strong brown, 7.5YR 5/6, hard, moist, with gray and orange streaks	~~~~~	10:33 / 3-8-07
35	46-57	N/A	Core not retrieved	N/A	N/A
40	57-60	030003SA060 030003WA060	CLAY, strong brown, 7.5YR 5/6, hard, moist, mottled, with gray streaks	~~~~~	13:11
Total Vertical Depth is 60 feet. Total Linear Depth is 85 feet.			LEGEND: CLAY		~~~~~
			SAND		~~~~~

LITHOLOGIC LOG			BORING/WELL ID 030-004-ASB		Page 1 of 1
Facility: Paducah Gaseous Diffusion Plant, Paducah, KY			Site: SWMU- 030		
Project: BGOU RI			Client: USDOE/PRS		
Drilling Contractor: Chase Environmental, LLC			Driller: Jeff Brownfield		
Start Time/Date: 3-6-07			End Time/Date: 3-6-07		
Borehole Diameter: 2.75"			Drilling Method: Direct Push		
Sampling Method: DT-21 Dual Tube			Drill Rig: Geoprobe® 6620DT		
Total Depth (Vertical): 60 feet			Angle: 45°		
Logged By: Mark Gartner			Protective Level: Modified Level D		
Coordinates: E -6800.238 N 781.032			Direction (plant grid): Northwest		
Depth (ft)	SAMPLE		LITHOLOGIC DESCRIPTION	GRAPHIC LOG	COMMENTS
	INTERVAL	NUMBER			
0	0-7	N/A	Core not retrieved	N/A	N/A
5	7-11	030004SA010	CLAY, white, 7.5YR 8/1, hard, moist, with orange streaks	~~~~~ ~~~~~ ~~~~~	
10	11-14	030004SA015	CLAY, white, 7.5YR 8/1, hard, moist, with orange streaks	~~~~~ ~~~~~ ~~~~~	
15	14-28	N/A	Core not retrieved	N/A	N/A
20	28-32	030004SA030	CLAY, white, 7.5YR 8/1, hard, moist, with a few orange streaks	~~~~~ ~~~~~ ~~~~~	
25	32-42	N/A	Core not retrieved	N/A	N/A
30	42-46	030004SA045	CLAY, white, 7.5YR 8/1, hard, moist, with a few orange streaks	~~~~~ ~~~~~ ~~~~~	
35	46-57	N/A	Core not retrieved	N/A	N/A
40	57-60	030004SA060	CLAY, brown, 7.5YR 4/4, very hard, dry, with gray streaks	~~~~~ ~~~~~ ~~~~~	14:32
Total Vertical Depth is 60 feet. Total Linear Depth is 85 feet.			LEGEND: CLAY		~~~~~ ~~~~~ ~~~~~
			SAND		~~~~~ ~~~~~ ~~~~~

SWMU 145

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LITHOLOGIC LOG			BORING/WELL ID 145-101-ASB		Page 1 of 1
Facility: Paducah Gaseous Diffusion Plant, Paducah, KY			Site: SWMU- 145		
Project: BGOU RI			Client: USDOE/PRS		
Drilling Contractor: Chase Environmental, LLC			Driller: Jeff Brownfield		
Start Time/Date: 14:15/1-24-07			End Time/Date: 12:35 / 1-24-07		
Borehole Diameter: 6.25"			Drilling Method: Direct Push Through Augers		
Sampling Method: DT-21 Dual Tube / 2.25" ID Augers			Drill Rig: Geoprobe® 6620DT		
Total Depth (Vertical): 60 feet			Angle: 45°		
Logged By: Todd Mills / Mark Gartner			Protective Level: Modified Level D		
Coordinates: E -2156.26 N 4493.87			Direction (plant grid): South		
Depth (ft)	SAMPLE		LITHOLOGIC DESCRIPTION	GRAPHIC LOG	COMMENTS
	INTERVAL	NUMBER			
0	0-7	N/A	Core not retrieved	N/A	N/A
5	7-11	145101SA010	CLAY, brown, 7.5YR 4/3, medium firm, slightly moist, with gray streaks		14:22
10	11-14	145101SA015	CLAY, gray, 7.5YR 6/1, medium firm, moist with light gray streaks		14:31
15					
20	14-28	N/A	Core not retrieved	N/A	N/A
25					
30	28-32	145101SA030	CLAY, gray, 7.5YR 6/1, firm, slightly moist with gray streaks		15:15
35	32-42	N/A	Core not retrieved	N/A	N/A
40					
45	42-46	145101SA045	CLAY, brown, 7.5YR 4/4, very firm, slightly moist, with black streaks		10:15 / 1-25-07
45	46-50	N/A	Core not retrieved	N/A	N/A
50	50-53	145101SA060	CLAY, light brown, 7.5YR 6/4, hard, moist, gray and orange streaks with a 3" sand seam, white, fine grained to 1/4" across, poorly sorted		12:35
Total Vertical Depth is 53 feet. Total Linear Depth is 75 feet.			LEGEND: CLAY		
			SAND		

LITHOLOGIC LOG			BORING/WELL ID 145-102-ASB		Page 1 of 1
Facility: Paducah Gaseous Diffusion Plant, Paducah, KY			Site: SWMU- 145		
Project: BGOU RI			Client: USDOE/PRS		
Drilling Contractor: Chase Environmental, LLC			Driller: Jeff Brownfield		
Start Time/Date: 8:40/1-16-07			End Time/Date: 16:42/1-18-07		
Borehole Diameter: 2.75"			Drilling Method: Direct Push Through Augers		
Sampling Method: DT-21 Dual Tube / 2.25" ID Augers			Drill Rig: Geoprobe® 6620DT		
Total Depth (Vertical): 60 feet			Angle: 45°		
Logged By: Mark Gartner			Protective Level: Modified Level D		
Coordinates: E -1895.42 N 4042.42			Direction (plant grid): West		
Depth (ft)	SAMPLE		LITHOLOGIC DESCRIPTION	GRAPHIC LOG	COMMENTS
	INTERVAL	NUMBER			
0	0-7	N/A	Core not retrieved	N/A	N/A
5	7-11	145102SA010	CLAY, brown, 7.5YR 7/1, firm, dry, black organic material, with orange streaks		9:42
10	11-14	145102SA015	CLAY, gray, 7.5YR 7/1, firm, dry, crumbly		
15					
20	14-28	N/A	Core not retrieved	N/A	N/A
25					
30	28-32	145102SA030 145102SD030	CLAY, brown, 7.5YR 4/4, hard, moist, with orange and gray streaks		10:28
35	32-42	N/A	Core not retrieved	N/A	N/A
40					
45	42-46	145102SA045	CLAY, reddish yellow, 7.5YR 6/6, firm, moist, gray and orange streaks		13:07
50	46-57	N/A	Core not retrieved	N/A	N/A
55					
	57-60	145102SA060	CLAY (from 80 to 81.5'), strong brown, 7.5 YR 4/6, hard, moist, with black streaks SAND (from 81.5 to 83'), light gray fine grained, well sorted		16:42
Total Vertical Depth is 60 feet. Total Linear Depth is 83 feet.			LEGEND: CLAY		
			SAND		

LITHOLOGIC LOG			BORING/WELL ID 145-103-ASB		Page 1 of 1
Facility: Paducah Gaseous Diffusion Plant, Paducah, KY			Site: SWMU- 145		
Project: BGOU RI			Client: USDOE/PRS		
Drilling Contractor: Chase Environmental, LLC			Driller: Jeff Brownfield		
Start Time/Date: 11:47 / 1-9-07			End Time/Date: 15:19/1-9-07		
Borehole Diameter: 2.75"			Drilling Method: Direct Push		
Sampling Method: DT-21 Dual Tube			Drill Rig: Geoprobe® 6620DT		
Total Depth (Vertical): 60 feet			Angle: 45°		
Logged By: Mark Gartner			Protective Level: Modified Level D		
Coordinates: E -2465.84 N 3406.88			Direction (plant grid): Northwest		
Depth (ft)	SAMPLE		LITHOLOGIC DESCRIPTION	GRAPHIC LOG	COMMENTS
	INTERVAL	NUMBER			
0	0-7	N/A	Core not retrieved	N/A	N/A
5	7-11	145103SA010	CLAY, dark gray, 7.5YR 4/1, soft, moist, rock fragments	~~~~~	12:02
10	11-14	145103SA015	CLAY, gray, 7.5YR 6/1, firm, dry	~~~~~	12:12
15					
20	14-28	N/A	Core not retrieved	N/A	N/A
25					
30	28-32	145103SA030	CLAY, brown, 7.5YR 5/3, firm, moist, with gray streaks	~~~~~	12:52
35	32-42	N/A	Core not retrieved	N/A	N/A
40	42-46	145103SA045	CLAY, brown, 7.5YR 5/4, firm, moist, gray streaks: Sand (10%)	~~~~~	13:49
45					
50	46-57	N/A	Core not retrieved	N/A	N/A
55	57-60	145103SA060	CLAY, strong brown, 7.5 YR 5/6, hard, moist with black and gray streaks: Sand (10%)	~~~~~	15:19
Total Vertical Depth is 60 feet. Total Linear Depth is 85 feet.			LEGEND: CLAY		~~~~~
			SAND		~~~~~

LITHOLOGIC LOG			BORING/WELL ID 145-104-ASB		Page 1 of 1
Facility: Paducah Gaseous Diffusion Plant, Paducah, KY			Site: SWMU- 145		
Project: BGOU RI			Client: USDOE/PRS		
Drilling Contractor: Chase Environmental, LLC			Driller: Jeff Brownfield		
Start Time/Date: 14:34 / 1-10-07			End Time/Date: 12:42/1-22-07		
Borehole Diameter: 2.75"			Drilling Method: Direct Push Through Augers		
Sampling Method: DT-21 Dual Tube / 2.25" ID Augers			Drill Rig: Geoprobe® 6620DT		
Total Depth (Vertical): 60 feet			Angle: 45°		
Logged By: Mark Gartner			Protective Level: Modified Level D		
Coordinates: E -2333.16 N 3381.42			Direction (plant grid): East		
Depth (ft)	SAMPLE		LITHOLOGIC DESCRIPTION	GRAPHIC LOG	COMMENTS
	INTERVAL	NUMBER			
0	0-7	N/A	Core not retrieved	N/A	N/A
5	7-11	145104SA010	CLAY, greenish gray, GLEY1 6/10Y, soft, dry, wood fragments	~~~~~	14:57
10	11-14	145104SA015	CLAY, greenish gray, GLEY1 6/10Y, soft, dry, wood fragments	~~~~~	15:16 Stop drilling activities due to High LEL readings.
15	14-28	N/A	Core not retrieved	N/A	N/A
20	28-32	145104SA030	CLAY, brown, 7.5YR 4/4, firm, moist with orange streaks	~~~~~	9:45
25	32-42	N/A	Core not retrieved	N/A	N/A
30	42-46	145104SA045	CLAY, brown, 7.5 YR 5/4, hard, moist with gray and orange streaks	~~~~~	10:26
35	46-57	N/A	Core not retrieved	N/A	N/A
40	57-60	145104SA060	CLAY, strong brown, 7.5 YR 5/8, hard, dry, with 1" sand seam, white, fine grained, dry	~~~~~	12:42
Total Vertical Depth is 60 feet. Total Linear Depth is 82.5 feet.			LEGEND: CLAY		~~~~~
			SAND		~~~~~

LITHOLOGIC LOG			BORING/WELL ID 145-105-ASB		Page 1 of 1
Facility: Paducah Gaseous Diffusion Plant, Paducah, KY			Site: SWMU- 145		
Project: BGOU RI			Client: USDOE/PRS		
Drilling Contractor: Chase Environmental, LLC			Driller: Jeff Brownfield		
Start Time/Date: 12:38 / 1-8-07			End Time/Date: 13:41 / 1-23-07		
Borehole Diameter: 6.25"			Drilling Method: Direct Push Through Augers		
Sampling Method: DT-21 Dual Tube / 2.25" ID Augers			Drill Rig: Geoprobe® 6620DT		
Total Depth (Vertical): 60 feet			Angle: 45°		
Logged By: Mark Gartner / Todd Mills			Protective Level: Modified Level D		
Coordinates: E -2765.27 N 3183.99			Direction (plant grid): Northwest		
Depth (ft)	SAMPLE		LITHOLOGIC DESCRIPTION	GRAPHIC LOG	COMMENTS
	INTERVAL	NUMBER			
0	0-7	N/A	Core not retrieved	N/A	N/A
5	7-11	145105SA010	CLAY, brown, 7.5YR 5/3, soft, moist, mottles with gray and red streaks	~~~~~	12:52
10	11-14	145105SA015	CLAY, brown, 7.5YR 4/3, firm, moist, dark orange streaks	~~~~~	13:02
15					
20	14-28	N/A	Core not retrieved	N/A	N/A
25					
30	28-32	145105SA030	CLAY, reddish yellow, 7.5YR 6/6, firm, dry, with gray streaks: Sand seam (1'), fine grained, with a few angular chert fragments	~~~~~	14:26
35					
40	32-42	N/A	Core not retrieved	N/A	N/A
45	42-46	145105SA045	CLAY, light brown, 7.5YR 6/4, hard, dry, gray streaks:	~~~~~	15:49
50					
55	46-57	N/A	Core not retrieved	N/A	N/A
	57-60	145105SA060	SILTY CLAYEY, strong brown, 7.5YR 5/6, sand and gravel	~~~~~	14:10 / 1-23-07
Total Vertical Depth is 60 feet. Total Linear Depth is 85 feet.			LEGEND: CLAY		~~~~~
			SAND		~~~~~

LITHOLOGIC LOG			BORING/WELL ID 145-106-ASB		Page 1 of 1
Facility: Paducah Gaseous Diffusion Plant, Paducah, KY			Site: SWMU- 145		
Project: BGOU RI			Client: USDOE/PRS		
Drilling Contractor: Chase Environmental, LLC			Driller: Jeff Brownfield		
Start Time/Date: 9:20/1-26-07			End Time/Date: 14:49/1-26-07		
Borehole Diameter: 6.25"			Drilling Method: Direct Push Through Augers		
Sampling Method: DT-21 Dual Tube / 2.25" ID Augers			Drill Rig: Geoprobe® 6620DT		
Total Depth (Vertical): 43 feet			Angle: 45°		
Logged By: Todd Mills			Protective Level: Modified Level D		
Coordinates: E -3088.28 N 3545.83			Direction (plant grid): East		
Depth (ft)	SAMPLE		LITHOLOGIC DESCRIPTION	GRAPHIC LOG	COMMENTS
	INTERVAL	NUMBER			
0	0-7	N/A	Core not retrieved	N/A	N/A
5	7-11	145106SA010	CLAY, very pale brown, 10YR 7/3, firm, slightly moist, with iron staining	~~~~~	Duplicate sample
10		145106SD010	CLAY, very pale brown, 10YR 7/3, firm, slightly moist, with iron staining	~~~~~	10:13
	11-14	145106SA015	CLAY, gray, 7.5YR 7/1, firm, dry	~~~~~	10:25
15	14-28	N/A	Core not retrieved	N/A	N/A
20					
25					
	28-30	145106SA030	CLAY, brown, 7.5YR 4/3, very firm, moist, with black streaks	~~~~~	11:39
30	30-33	N/A	Core not retrieved	N/A	N/A
35	33-35	145106SA045	CLAY, brown, 7.5YR 4/4, very firm, moist	~~~~~	13:59
	38-40	145106SA060	CLAY, brown, 7.5YR 4/4, very firm, slightly moist, black streaks	~~~~~	15:00
40	40-43	N/A	Core not retrieved	N/A	N/A
Total Vertical Depth is 43 feet. Total Linear Depth is 60 feet.			LEGEND: CLAY	~~~~~	
			SAND	~~~~~	

LITHOLOGIC LOG			BORING/WELL ID 145-107-ASB		Page 1 of 1
Facility: Paducah Gaseous Diffusion Plant, Paducah, KY			Site: SWMU- 145		
Project: BGOU RI			Client: USDOE/PRS		
Drilling Contractor: Chase Environmental, LLC			Driller: Jeff Brownfield		
Start Time/Date: 13:50 / 1-29-07			End Time/Date: 11:24/1-31-07		
Borehole Diameter: 6.25"			Drilling Method: Direct Push Through Augers		
Sampling Method: DT-21 Dual Tube / 2.25" ID Augers			Drill Rig: Geoprobe® 6620DT		
Total Depth (Vertical): 60 feet			Angle: 45°		
Logged By: Mark Gartner			Protective Level: Modified Level D		
Coordinates: E -2809.44 N 4138.49			Direction (plant grid): South		
Depth (ft)	SAMPLE		LITHOLOGIC DESCRIPTION	GRAPHIC LOG	COMMENTS
	INTERVAL	NUMBER			
0	0-7	N/A	Core not retrieved	N/A	N/A
5	7-11	145107SA010	CLAY, brown, 7.5YR 4/2, firm, moist with gray and orange streaks: Quartz (5%), 1/8" to 1/4" across, rounded to subangular	~~~~~	14:02
10	11-14	145107SA015	CLAY, dark gray, 7.5YR 4/1, soft, moist	~~~~~	14:10
15	14-18		CLAY, dark gray, 7.5YR 4/1, soft, moist	~~~~~	Additional sample was required.
20	18-28	N/A	Core not retrieved	N/A	N/A
25	28-32	145107SA030	CLAY, brown, 7.5YR 4/4, firm, moist, gray streaks with 10% sand: Sand seam (2.5'), fine grained to medium grained, poorly sorted with 10% clay	~~~~~	15:11
35	32-42	N/A	Core not retrieved	N/A	N/A
40	42-46	145107SA045	CLAY, light gray, 7.5YR 7/1, very hard, dry with orange streaks	~~~~~	15:44 / 1-30-07
45	46-57	N/A	Core not retrieved	N/A	N/A
55	57-60	145107SA060	SAND, white, fine grained, dry with iron staining	11:24 / 1-31-07
Total Vertical Depth is 60 feet. Total Linear Depth is 83 feet.			LEGEND: CLAY		~~~~~
			SAND	

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APPENDIX C

ANALYTICAL DATA AND QA/QC EVALUATION RESULTS

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APPENDIX C

**ANALYTICAL DATA AND QA/QC
EVALUATION RESULTS**

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APPENDIX D

THREE DIMENSIONAL VISUALIZATION FIGURES

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APPENDIX D

THREE DIMENSIONAL VISUALIZATION FIGURES

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APPENDIX E
CONTAMINANT FATE AND TRANSPORT MODELING RESULTS
FOR THE BGOU

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ACRONYMS

AT123D	Analytical Transient 1-, 2-, 3-Dimensional Model
BGOU	Burial Grounds Operable Unit
CERCLA	Comprehensive Environmental Response, Compensation and Liability Act
COC	contaminant of concern
COPC	chemical of potential concern
DAF	dilution/attenuation factor
DCE	dichloroethene
DNAPL	dense nonaqueous-phase liquid
DQO	Data Quality Objective
ELCR	excess lifetime cancer risk
EPA	U. S. Environmental Protection Agency
f_{oc}	soil organic carbon fraction
HQ	hazard quotient
HU	hydrogeologic unit
K_d	distribution coefficient
K_{oc}	organic carbon partition coefficient
MCL	Maximum Contaminant Level
MEPAS	Multimedia Environmental Pollutant Assessment System
Paducah OREIS	Paducah Oak Ridge Environmental Information System
PCB	polychlorinated biphenyl
PGDP	Paducah Gaseous Diffusion Plant
POE	point of exposure
RBC	risk-based concentration
RCRA	Resource Conservation and Recovery Act
RESRAD	RESidual RADioactive Materials
RfD	reference dose
RG	Regional Gravel Aquifer
RGO	remedial goal option
RI	Remedial Investigation
SADA	Statistical Analysis and Decision Assistance Model
SESOIL	Seasonal Soil Compartment Model
SQL	sample quantitation limit
SSL	soil screening level
SVOC	semivolatile organic compound
SWMU	solid waste management unit
^{99}Tc	technetium-99
TCE	trichloroethene
UCL	upper confidence limit
UCRS	Upper Continental Recharge System
VOC	volatile organic compound

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E.1. INTRODUCTION

This appendix presents the methods and results of the fate and transport modeling performed for the Burial Grounds Operable Unit (BGOU) Remedial Investigation (RI), consisting of [Solid Waste Management Unit (SWMU) 2, SWMU3, SWMU 4, SWMU 5, SWMU 6, SWMU 7, SWMU 30, and SWMU 145].

The fate and transport modeling of the BGOU RI is consistent with the Tiers 1, 2 and 3 of the modeling matrix included in the Paducah Gaseous Diffusion Plant (PGDP) Risk Methods Document (DOE 2001). This modeling matrix is consistent with the Draft 2008 revision to the Risk Methods Document, since the methodologies are the same for fate and transport modeling. As indicated by this matrix (Table E.1.1), Tier 1 consists of simple screens using soil screening levels (SSLs) to identify those contaminants that may migrate from source areas to undefined downgradient points of exposure (POEs); Tier 2 consists of source delineation and transport modeling using input parameters that are unlikely to underestimate the potential for contaminant transport to undefined downgradient POEs (i.e., are conservative estimates of contaminant transport); and Tier 3 consists of source delimitation and transport modeling using input parameters that result in more accurate estimates of future contaminant concentrations at POEs beneath the SWMUs, at the PGDP plant boundary, PGDP property boundary, Little Bayou seeps, and the Ohio River.

Section 5 summarizes the modeling results documented by this appendix.

Table E.1.1. Modeling Matrix for Groundwater^a

	Values for Soil to Protect Groundwater	Model	Point of Exposure	Notes
INVESTIGATION DOCUMENTS	<p>Tier 1</p> <p>Initial analysis used to identify COPCs that might migrate from source areas and require further fate and transport analysis.</p>	<p>Concentrations in source term are the maximum detected concentrations of contaminants in the source. Contaminant concentrations compared to site screening levels and groundwater protection values in Appendix A of the PGDP Methods Document.</p>	<p>At source unit.</p>	<p>Use dilution/attenuation factor (DAF) of 1 for site screening levels unless site-specific values are available.</p> <p>Groundwater Protection value based on residential use and targets of 1E-6, 0.1, and 1 for risk, hazard, and dose, respectively.</p> <p>If site-specific DAF values are used, then the groundwater protection value should be justified.</p> <p>The depth to groundwater will be considered in the calculation.</p>
	<p>Tier 2</p> <p>Analysis is used to refine the list of COPCs that might migrate from source areas. Depending on the DQOs for the project, additional fate and transport analysis of selected COPCs might be completed.</p>	<p>Concentrations in source term for all contaminants are the lesser of the maximum and UCL95 concentration of the appropriate distribution. Fate and transport modeling completed using SESOIL and/or RESRAD.</p>	<p>At source unit.</p>	<p>Includes source delimitation.</p> <p>The analysis will recognize SESOIL limitations when modeling inorganic COPCs-refine K_ds.</p>
DECISION DOCUMENTS	<p>Tier 3</p> <p>Analysis is used for COCs identified from Tier 2 modeling. Includes consideration of COC concentrations at downgradient locations. The results of this analysis may be used to develop clean-up levels for some COCs.</p>	<p>Source term developed using SADA. Fate and transport completed using SESOIL and RESRAD with AT123D.</p>	<p>At source unit and at downgradient exposure points.</p> <p>Exposure points are at the plant boundary, the property boundary, Little Bayou seeps, and the Ohio River.</p>	<p>Uses source delimitation and refined K_ds from previous tiers.</p> <p>Contaminant migration paths will be derived using the sitewide groundwater model.</p> <p>On the Terrace (southern portion of PGDP), different points of exposure will apply and be determined using the sitewide groundwater model.</p>
	<p>Tier 4</p> <p>Analysis is used for the COCs presenting the greatest risk at downgradient exposure points. The results of this analysis may be used to develop clean-up levels for some COCs.</p>	<p>Source modeling and MODFLOW T</p>	<p>Down-gradient points</p> <p>Exposure points are at the plant boundary, the property boundary, Little Bayou Creek, and the Ohio River.</p>	<p>To be used to refine clean-up goals (if needed).</p> <p>On the Terrace (southern portion of PGDP), different points of exposure will apply and be determined using the sitewide groundwater model.</p>

^a Adapted from Table 3.2 of the PGDP Risk Methods Document (DOE/OR/07-1506&D2).

AT123D = Analytical Transient 1-, 2-, 3-Dimensional

COC = contaminant of concern

COPC = chemical of potential concern

DAF = dilution/attenuation factor

DQO = Data Quality Objective

K_d = distribution coefficient

RESRAD = Residual Radioactive Materials

SADA = Statistical Analysis and Decision Assistance

SESOIL = Seasonal Soil Compartment Model

UCL = upper confidence level

E.2. RESULTS FROM PREVIOUS MODELING EFFORTS

E.2.1 SUMMARY

Transport modeling results contained in previous investigations and risk assessments were examined to determine the types of models completed previously and the results of those modeling activities. All reports considered were from work completed between 1990 and 2004.

As part of this summary, previously completed transport models were categorized into one of the four modeling tiers described in Table 3.2 in the Risk Methods Document (DOE 2001). These tiers and their descriptions are as follows:

- Tier 1: Results are derived using simple comparisons between sampling results and soil screening levels for groundwater protection. No source-term calculations are performed. Results are used for scoping investigation activities. The POE considered is at the source unit.
- Tier 2: Results are derived using analytical models such as the Multimedia Environmental Pollutant Assessment System (MEPAS), Residual Radioactive Materials (RESRAD), SESOIL, and AT123D. Source-terms are derived by assuming that the source-term volume consists of all areas with a detected result and that the source-term concentration is equal to the maximum detected concentration over all samples. Results are used to determine if a response action should be considered for the source. The POE considered is at the source unit.
- Tier 3: Results are derived using analytical models such as MEPAS, RESRAD, SESOIL, and AT123D. Source-terms are less conservatively derived than under Tier 2 by using three-dimensional plots and/or computer programs that can perform geospatial modeling (e.g., SADA). The source concentration is assumed to be the average concentration over all detected concentrations within the source volume. Results are used in decision documents to select among possible response actions and to derive cleanup levels. The POEs considered are at the source unit and at downgradient points (e.g., the PGDP boundary, property boundary, and either Little Bayou Creek or the Ohio River).
- Tier 4: Results are derived using numerical models, such as MODFLOW T. Similar to Tier 3, source-terms are derived using three-dimensional plots and/or computer programs that can perform geospatial modeling. The source concentration is assumed to be the average concentration over all detected concentrations within the source volume. Results are used in decision documents to design a selected response action, such as in refining cleanup levels and selecting monitoring points. The POEs considered are at the source unit and at downgradient points (e.g., the PGDP boundary, property boundary, and either Little Bayou Creek or the Ohio River).

Generally, all previous modeling that has been performed for the burial grounds falls within Tier 2; however, in most cases, modeling to downgradient POEs (i.e., the PGDP boundary and/or property boundary) was included. Modeling to the downgradient points is similar to the Tier 3 requirement. No modeling to Little Bayou Creek or the Ohio River has been completed previously for the burial grounds.

Table E.2.1 and the following text summarize previous modeling performed for each burial ground. No previous modeling has been performed for SWMU 145. All risk and hazard estimates presented are for hypothetical residential use of groundwater obtained from the Regional Gravel Aquifer (RGA) at locations such as the plant boundary and property boundary.

Table E.2.1. Summary of Previous Modeling Performed for Burial Grounds at PGDP

Unit	Tier/Model Used	Report	PGDP Boundary		Property Boundary		River/Little Bayou Creek Seeps	
			Total risk/hazard	COCs	Total risk/hazard	COCs	Total risk/hazard	COCs
SWMU 2	Tier 1— None	<i>Results of the Public Health and Ecological Assessment, Phase II, at the Paducah Gaseous Diffusion Plant, Paducah, Kentucky, KY/SUB/13B-9777C P-03/1991/1, December 1991.</i>	Not calculated; qualitative determination	TCE, ⁹⁹ Tc, Beryllium, Chromium, Lead	Not calculated	NA	Not calculated	NA
			Not calculated; qualitative determination	⁹⁹ Tc, Uranium, metals	Not calculated	NA	Not calculated	NA
			VOCs Risk = 3E-05 Hazard = <1 Dose = NA Based on predicted maximum concentration at PGDP boundary at 35 years from present.	TCE	Risk = 2E-05 Hazard = <1 Dose = NA Based on predicted maximum concentration at PGDP boundary 35 years from present.	TCE	Not calculated	NA
SWMU 3	Tier 1— None	<i>Results of the Public Health and Ecological Assessment, Phase II, at the Paducah Gaseous Diffusion Plant, Paducah, Kentucky, KY/SUB/13B-9777C P-03/1991/1, December 1991.</i>	Not calculated; qualitative determination	TCE, ⁹⁹ Tc, Beryllium, Chromium, Lead	Not calculated	NA	Not calculated	NA
			Not calculated; qualitative determination	⁹⁹ Tc, Uranium, metals	Not calculated	NA	Not calculated	NA
			VOCs Risk = 3E-05 Hazard = <1 Dose = NA Based on predicted maximum concentration at PGDP boundary at 35 years from present.	TCE	Risk = 2E-05 Hazard = <1 Dose = NA Based on predicted maximum concentration at PGDP boundary 35 years from present.	TCE	Not calculated	NA
SWMU 2	Tier 2— MEPAS	<i>Data Summary and Interpretation Report for Interim Remedial Design at Solid Waste Management Unit 2 of Waste Area Grouping 22 at the Paducah Gaseous Diffusion Plant, Paducah, Kentucky, DOE/OR/07-1549&D1, February 1997b.</i>	Metals Risk = 1E-05 Hazard = <1 Dose = NA Based on predicted maximum concentration at PGDP boundary at 1,505 years from present.	Arsenic	Risk = 1E-05 Hazard = <1 Dose = NA Based on predicted maximum concentration at boundary at more than 1,000 years from present.	Arsenic	Not calculated	NA
			Not calculated	NA	Risk = 6E-03 Hazard = 1,000 Dose = <1 mrem/year	Risk: TCE, vinyl chloride, ⁹⁹ Tc Hazard: <i>cis</i> -1,2-DCE; TCE	Not calculated	NA
			Not calculated; qualitative determination	TCE, ⁹⁹ Tc, Beryllium, Chromium, Lead	Not calculated	NA	Not calculated	NA
SWMU 3	Tier 1— None	<i>Results of the Public Health and Ecological Assessment, Phase II, at the Paducah Gaseous Diffusion Plant, Paducah, Kentucky, KY/SUB/13B-9777C P-03/1991/1, December 1991.</i>	Not calculated; qualitative determination	TCE, ⁹⁹ Tc, Beryllium, Chromium, Lead	Not calculated	NA	Not calculated	NA
			Not calculated; qualitative determination	⁹⁹ Tc, Uranium, metals	Not calculated	NA	Not calculated	NA
			VOCs Risk = 3E-05 Hazard = <1 Dose = NA Based on predicted maximum concentration at PGDP boundary at 35 years from present.	TCE	Risk = 2E-05 Hazard = <1 Dose = NA Based on predicted maximum concentration at PGDP boundary 35 years from present.	TCE	Not calculated	NA

Table E.2.1.1. Summary of Previous Modeling Performed for Burial Grounds at PGDP (Continued)

Unit	Tier/Model Used	Report	PGDP Boundary		Property Boundary		River/Little Bayou Creek Seeps	
			Total risk/hazard	COCs	Total risk/hazard	COCs	Total risk/hazard	COCs
SWMU 3	Tier 2— SESOL/ AT123D	<i>Sitewide Risk Assessment Model and Environmental Baseline for the Paducah Gaseous Diffusion Plant, Paducah, Kentucky, DOE/OR/07-2104&D0, September 2003.</i>	Not calculated	NA	Risk = 7E-06 Hazard = 2 Dose = <1 mrem/year	Risk: ⁹⁹ Tc Hazard: naphthalene	Not calculated	NA
SWMU 4	Tier 2— MEPAS	<i>Remedial Investigation Report for Waste Area Grouping 3 at the Paducah Gaseous Diffusion Plant, Paducah, Kentucky, DOE/OR/07-1895&D1, September 2000.</i>	TCE and solvents Risk = 6E-02 Hazard = 2,000 (Assumed 100 years from present)	Risk: 1,1-DCE; TCE; vinyl chloride; carbon tetrachloride Hazard: 1,1-DCE; TCE	Not calculated; however, a comparison of concentrations indicates that risks and hazards would be about one order of magnitude less than those calculated for the PGDP boundary.	Assumed the same as PGDP boundary COCs	Not calculated	NA
			Metals and radionuclides Risk = 6E-03 Hazard = 400 (Assumed at >1,000 years from present)	Risk: Arsenic, ²³⁷ Np, ²³⁹ Pu, ⁹⁹ Tc, ²³⁴ U, ²³⁵ U, ²³⁸ U Hazard: Arsenic, Cobalt, Copper, Iron, Manganese				
	Tier 2— SESOL/ AT123D	<i>Sitewide Risk Assessment Model and Environmental Baseline for the Paducah Gaseous Diffusion Plant, Paducah, Kentucky, DOE/OR/07-2104&D0, September 2003.</i>	Dose = Not calculated Not calculated	NA NA	Risk = >1 Hazard = 2,000,000 Dose = 2 mrem/year	Risk: carbon tetrachloride; chloroform; 1,1-DCE; TCE; vinyl chloride; ⁹⁹ Tc Hazard: carbon tetrachloride; chloroform; <i>cis</i> -1,2-DCE; <i>trans</i> -1,2-DCE; 1,1-DCE; TCE; vinyl chloride Dose: ⁹⁹ Tc	Not calculated	NA

Table E.2.1. Summary of Previous Modeling Performed for Burial Grounds at PGDP (Continued)

Unit	Tier/Model Used	Report	PGDP Boundary		Property Boundary		River/Little Bayou Creek Seeps	
			Total risk/hazard	COCs	Total risk/hazard	COCs	Total risk/hazard	COCs
SWMU 5	Tier 2—MEPAS	Remedial Investigation Report for Waste Area Grouping 3 at the Paducah Gaseous Diffusion Plant, Paducah, Kentucky, DOE/OR/07-1895&D1, September 2000.	TCE and solvents Risk = <1E-06 Hazard = <1	Risk: none Hazard: none	Not calculated; however, a comparison of concentrations indicates that risks and hazards would be about one order of magnitude less than those calculated for the PGDP boundary.	Assumed the same as PGDP boundary COCs	Not calculated	NA
			Metals and radionuclides Risk = <1E-06 Hazard = 100 (Assumed at >1,000 years from present) Dose = Not calculated	Risk: none Hazard: Iron, Manganese				
SWMU 6	Tier 2—SES/OIL/AT123D	Site-wide Risk Assessment Model and Environmental Baseline for the Paducah Gaseous Diffusion Plant, Paducah, Kentucky, DOE/OR/07-2104&D0, September 2003.	Not calculated	NA	Risk = 5E-03 Hazard = 100 Dose = <1 mrem/year	Risk: 1,1-DCE and ⁹⁹ Tc Hazard: naphthalene Dose: None	Not calculated	NA
			TCE and solvents Risk = <1E-06 Hazard = <1 Metals and radionuclides Risk = <1E-06 Hazard = 20 (Assumed at >1,000 years from present) Dose = Not calculated	Risk: none Hazard: none Risk: none Hazard: Iron				
SWMU 7	Tier 2—SES/OIL/AT123D	Site-wide Risk Assessment Model and Environmental Baseline for the Paducah Gaseous Diffusion Plant, Paducah, Kentucky, DOE/OR/07-2104&D0, September 2003.	Not calculated	NA	Risk = 3E-05 Hazard = <1 Dose = <1 mrem/year	Risk: ⁹⁹ Tc Hazard: none Dose: none	Not calculated	NA
			Not calculated; qualitative determination	TCE; 1,2-DCE; vinyl chloride; ⁹⁹ Tc; Arsenic; Chromium; Nickel				
SWMU 7	Tier 1—None	Results of the Public Health and Ecological Assessment, Phase II, at the Paducah Gaseous Diffusion Plant, Paducah, Kentucky, KY/SUB/13B-9777C P-03/1991/1.	Not calculated; qualitative determination	TCE; 1,2-DCE; vinyl chloride; ⁹⁹ Tc; Arsenic; Chromium; Nickel	Not calculated	NA	Not calculated	NA

Table E.2.1. Summary of Previous Modeling Performed for Burial Grounds at PGDP (Continued)

Unit	Tier/Model Used	Report	PGDP Boundary		Property Boundary		River/Little Bayou Creek Seeps	
			Total risk/hazard	COCs	Total risk/hazard	COCs	Total risk/hazard	COCs
SWMU 7	Tier 2— SESOL/ AT123D	Remedial Investigation Report for Solid Waste Management Units 7 and 30 of Waste Area Grouping 22 at Paducah Gaseous Diffusion Plant, Paducah, Kentucky, DOE/OR/07-1604&D2, January 1998.	Not calculated	NA	Risk = 2E-04 Hazard = <1 Dose = Not calculated (Results are for sources at both SWMUs 7 and 30 and are for 100 years from present.)	Risk: vinyl chloride, ⁹⁹ Tc Hazard: none Dose: NA	Not calculated	NA
			Risk = Not calculated Hazard = NA Dose = Not calculated [Results are maximum contribution from the incinerator area (Area Z) in SWMU 7.]	Maximum concentration of ⁹⁹ Tc was 63 pCi/L at 20 years from present	Maximum concentration of ⁹⁹ Tc was 11 pCi/L at 25 years from present.	Not calculated	NA	
SWMU 30	Tier 2— SESOL/ AT123D	Site-wide Risk Assessment Model and Environmental Baseline for the Paducah Gaseous Diffusion Plant, Paducah, Kentucky, DOE/OR/07-2104&D0, September 2003.	Not calculated	NA	Risk = 8E-04 Hazard = 30 Dose = 11 mrem/year	Risk: benzene, chloroform, ethylbenzene, ⁹⁹ Tc Hazard: Copper, benzene, naphthalene Dose: ⁹⁹ Tc NA	Not calculated	NA
			Not calculated; qualitative determination	TCE; 1,2-DCE; vinyl chloride; ⁹⁹ Tc; Arsenic; Chromium; Nickel	Not calculated	Not calculated	Not calculated	NA
SWMU 30	Tier 1— None	Results of the Public Health and Ecological Assessment, Phase II, at the Paducah Gaseous Diffusion Plant, Paducah, Kentucky, KY/SUB/13B-9777C P-03/1991/1.	Not calculated	NA	Risk = 2E-04 Hazard = <1 Dose = Not calculated (Results are for sources at both SWMUs 7 and 30 and are for 100 years from present.)	Risk: vinyl chloride, ⁹⁹ Tc Hazard: none Dose: NA	Not calculated	NA
			Not calculated	NA	Not calculated	Not calculated	Not calculated	NA

Table E.2.1. Summary of Previous Modeling Performed for Burial Grounds at PGDP (Continued)

Unit	Tier/Model Used	Report	PGDP Boundary		Property Boundary		River/Little Bayou Creek Creeks	
			Total risk/hazard	COCs	Total risk/hazard	COCs	Total risk/hazard	COCs
SWMU 30	Tier 3— SESOL/ AT123D	<i>Technetium-99 Transport Modeling Results for Sources at SWMUs 7 and 30 at the Paducah Gaseous Diffusion Plant, Paducah, Kentucky, KY/EM-266, March 1998.</i>	Risk = Not calculated Hazard = NA Dose = Not calculated (Results are maximum contribution from Pits B/C in SWMU 30.)	Maximum concentration of ⁹⁹ Tc was 122 pCi/L at 20 years from present.	Risk = Not calculated Hazard = NA Dose = Not calculated (Results are maximum contribution from Pits B/C in SWMU 30.)	Maximum concentration of ⁹⁹ Tc was 21 pCi/L at 25 years from present	Not calculated	NA
	Tier 2— SESOL/ AT123D	<i>Site-wide Risk Assessment Model and Environmental Baseline for the Paducah Gaseous Diffusion Plant, Paducah, Kentucky, DOE/OR/07-2104&D0, September 2003.</i>	Not calculated	NA	Risk = 3E-04 Hazard = 8 Dose = 5 mrem/year	Risk: ⁹⁹ Tc Hazard: naphthalene Dose: ⁹⁹ Tc	Not calculated	NA

COC = contaminant of concern

DCE = dichloroethene

MEPAS = Multimedia Environmental Pollutant Assessment System

mrem = millirem

NA = Not applicable

²³⁷Np = neptunium-237

²³⁹Pu = Plutonium-239

SESOL = Seasonal Soil Compartment Model

⁹⁹Tc = technetium-99

TCE = trichloroethene

²³⁴U = uranium-234

²³⁵U = uranium-235

²³⁸U = uranium-238

VOC = volatile organic compound

More modeling results are available for SWMU 2 than other BGOU SWMUs; however, no modeling has extended to Tier 3. Tier 2 modeling results, which have included modeling to the PGDP boundary and property boundary POEs, have concluded that this unit may be a potential contributor of trichloroethene (TCE) and other volatile organic compounds (VOCs) to groundwater. In addition, this unit may be a contributor of technetium-99 (^{99}Tc), but the risks due to ^{99}Tc levels are two orders of magnitude less (i.e., equal to $3\text{E-}05$) than those from solvents ($5\text{E-}03$). It is unlikely that this unit is a contributor of metals to groundwater, and an extensive analysis in *Data Summary and Interpretation Report for Interim Remedial Design at Solid Waste Management 2 of Waste Area Grouping 22 at the Paducah Gaseous Diffusion Plant, Paducah, Kentucky*, DOE/OR/07-1549&D1, (DOE 1997) determined that the uranium metal present in the burial ground is unlikely to contribute to groundwater contamination.

SWMU 3 may contribute contaminants to groundwater; however, no modeling has extended to Tier 3. Tier 2 modeling results, at the property boundary POE, have concluded that this unit is a minor contributor of ^{99}Tc to groundwater (Risk = $7\text{E-}06$). Naphthalene also has been identified as a contaminant of concern (COC) (for hazard), but this result is suspect due to conservative source-term development. This unit has not been shown to be a contributor of metals to groundwater.

SWMU 4 previous modeling identified a risk over 1 and hazard over 2,000,000 at the property boundary POE using Tier 2 modeling. The maximum risks are predicted to be from chloroform (> 1), 1,1-DCE ($3.98\text{E-}01$), carbon tetrachloride ($1.22\text{E-}01$), TCE ($2.37\text{E-}02$), and vinyl chloride ($1.46\text{E-}02$). The maximum hazards are predicted to be from chloroform (1,710,000), *cis*-1,2-DCE (789,000), *trans*-1,2-DCE (16,900), and carbon tetrachloride (11,600) (DOE 2003a). COCs include VOCs [TCE; 1,1-dichloroethene (DCE); vinyl chloride; 1,2-DCE; carbon tetrachloride, and chloroform], metals (arsenic, cobalt, copper, iron, and manganese), and radionuclides [neptunium-237 (^{237}Np), plutonium-239 (^{239}Pu), ^{99}Tc , uranium-234 (^{234}U), uranium-235 (^{235}U), and uranium-238 (^{238}U)]. Although Tier 2 modeling derived elevated risk and hazard, these results are highly uncertain because of the conservative source term used in the modeling.

SWMU 5 previous modeling identified COCs for risk as 1,1-DCE and ^{99}Tc and identified COCs for hazard as naphthalene, Mn, and Fe. Although Tier 2 modeling derived an elevated risk ($5\text{E-}05$) and hazard (100), these results are highly uncertain due to the conservative Tier 2 source-term used in the modeling.

SWMU 6 Tier 2 modeling derived elevated risk from ^{99}Tc ($3\text{E-}05$) and hazard from iron (20); however, these results are highly uncertain due to the conservative source-term used in the modeling.

SWMU 7 may contribute contaminants to groundwater; however, Tiers 2 and 3 modeling results indicate that the contamination contributed is probably not significant. While early Tier 2 modeling identified SWMU 7 as a potential source of ^{99}Tc and vinyl chloride, later Tier 3 modeling determined that the level of ^{99}Tc that might reach a receptor at the PGDP boundary or property boundary (maximum of 63 and 11 pCi/L) is well below the Maximum Contaminant Level (MCL) (900 pCi/L). Later Tier 2 modeling (i.e., that from the sitewide risk model) did identify additional COCs; however, this result is highly uncertain given the conservative source-term used.

SWMU 30 may contribute contaminants to groundwater; however, Tiers 2 and 3 modeling results indicate that the contamination contributed probably is not significant. While early Tier 2 modeling identified SWMU 30 as a potential source of ^{99}Tc and vinyl chloride; later Tier 3 modeling determined that the level of ^{99}Tc that might reach a receptor at the PGDP boundary or property boundary (maximum of 122 and 21 pCi/L) is well below the MCL (900 pCi/L). Later Tier 2 modeling (i.e., that from the sitewide risk model) did identify ^{99}Tc as an important COC; however, this result is highly uncertain given the conservative source-term used.

Attachment E.1 provides more detailed summaries and excerpts of reports of previous modeling for the BGOU RI SWMUs.

E.3. MODELING COMPLETED AS PART OF BGOU RI

E.3.1 GROUNDWATER MODELING

The BGOU RI performed fate and transport modeling using the SADA, SESOIL, MODFLOW/MODPATH, and AT123D models. In general, the selected POEs where groundwater concentrations of the analytes were estimated were below the SWMU, at the plant boundary, at the property boundary, at the Little Bayou seeps, and the Ohio River; however, not all SWMUs have transport pathways to all of the POEs. For example, SWMU 145 is located outside of the plant boundary and does not contribute to the Little Bayou seeps.

Modelers used the following approach to evaluate the migration of the selected analytes from the BGOU SWMUs to groundwater and subsequently to the POEs.

1. Develop a conceptual model of each SWMU including estimated depths to the RGA and the derived flow paths and distances to the POEs using MODFLOW/MODPATH and the PGDP sitewide groundwater model.
2. Refine the source zones for each analyte in a SWMU using the SADA model.
3. Perform leachate modeling using SESOIL to estimate the rate of contaminant loading over time from each source area in a SWMU.
4. Perform saturated flow and transport modeling with AT123D using contaminant loading information from SESOIL.

Contaminant migration may have impacted three hydrogeologic units underlying the source areas at the SWMUs comprising the BGOU. These units, which control the flow of groundwater and contaminant migration at these SWMU source areas, are as follows (in descending order):

1. Upper Continental Recharge System (UCRS)—approximately 60 ft of silt and clay with horizons of sand and gravel;
2. RGA—approximately 40 ft of gravel, sand, and silt deposits that overlie the McNairy Formation; and
3. McNairy Formation—approximately 225 ft of sand and silt with some clay.

Previous work has shown that groundwater flow in the UCRS is primarily vertical and that the lateral groundwater flow in the McNairy is significantly slower than that in the RGA. The primary contaminant pathway considered in the fate and transport modeling is vertical migration through the UCRS followed by lateral migration in the RGA to the POEs.

SADA. Spatial Analysis and Decision Assistance model (UT 2005) was used to estimate the source volumes of analytes from the sample results through geospatial interpolation techniques. (See Attachment E.1 for additional information on source delineation). Surface and subsurface sampling results were taken from the Paducah Oak Ridge Environmental Information System (Paducah OREIS). The limitations of this data include the lack of sampling results for the waste, which may exhibit higher concentrations than the surrounding soil samples. In addition, a portion of the waste, which was not modeled in this report, is contained in drums at several SWMUs. The limitations of the data used in the analyses are presented in detail in Section E.3.3. Information for each result included the sample and station identifier, the date of

sample collection, the location and depth at which the sample was taken, whether the analyte was detected or not detected at the sample quantitation limit (SQL), and the result. The results initially were screened against the risk-based SSLs for protection of RGA groundwater for significant analytes at PGDP contained in the Risk Methods Document (DOE 2001) (see Appendix F).

Modelers divided each analyte source area into rows and columns with a uniform spacing of 20 ft for SWMUs 2, 3, 4, 5, 6, 7, and 30 and a uniform spacing of 100 ft for SWMU 145. Multiple domains with varying depths were used to characterize the analyte source areas vertically in relation to the existing aquifers; therefore, the domain was further discretized into horizontal layers. Analyte results for each domain were compiled, and analyte concentrations in each cell of the domain were predicted using geospatial interpolation (see Appendix E Attachment 2 for details).

The techniques in SADA that can be used for source term development in a three-dimensional system are nearest neighbor, inverse distance, and kriging. The nearest neighbor technique was selected for source zone refinement because it yielded results that were most compatible with the conceptual site model of contaminant release, as described in Attachment 2 to Appendix E.

As shown in the source term tables of following sections, the size of sources varied between the layers. The SESOIL input parameter for contaminant sources allows only one value for the source area; therefore, for each contaminant, the area of the SADA layer with the highest contaminant mass was used as the SESOIL input for source zone area. The analyte concentrations in the other layers were normalized to the area of the layer with the maximum mass.

SESOIL. Seasonal Soil Compartment Model (Bonazountas and Wagner 1984) was used for leachate modeling. SESOIL estimates contaminant concentrations in the soil profile following introduction via direct application and/or interaction with other media. The model defines the soil compartment as a soil column extending from the ground surface through the unsaturated zone to the top of the saturated soil zone/water table. Processes simulated in SESOIL are categorized in three cycles—the hydrologic cycle, sediment cycle, and pollutant cycle. Each cycle is a separate submodule in the SESOIL code. The hydrologic cycle includes rainfall, surface runoff, infiltration, soil-water content, evapotranspiration, and groundwater recharge. The sediment cycle includes sediment washload as a result of rainstorms (i.e., soil erosion that results from surface runoff). The pollutant cycle includes convective transport, volatilization, adsorption/desorption, and degradation/decay. A contaminant in SESOIL can partition in up to four phases (liquid, adsorbed, air, and pure). Output of the SESOIL model includes contaminant concentrations at various soil depths and contaminant loss from the unsaturated soil zone in terms of surface runoff, percolation to groundwater, volatilization, and degradation. SESOIL predicts the monthly contaminant load to the water table from the area of concern that can be directly input into the AT123D model for contaminant migration in the saturated zone to selected downgradient POEs.

The hydrologic modeling parameter values used in the SESOIL modeling were based on representative conditions at the PGDP and site specific values for the individual SWMU (Table E.3.1). The modeling parameters were selected so that they could account for expected variability in the hydraulic system and would be unlikely to underestimate contaminant release and transport. However, SESOIL does have limitations in regard to modeling waste that may potentially be located in the water table in the UCRS and contaminant transformations such as that resulting from radionuclide decay. These issues and their contribution to the uncertainty in the analyses are addressed in Section E.3.3.

Table E.3.1. Soil Parameters Used in SESOIL Modeling for the BGOU RI

Input Parameter	Value	Source
Soil type	Silty clay	PGDP site-specific
Bulk density (g/cm ³)	1.46	Laboratory analysis
Percolation rate (cm/year)	11	PGDP calibrated model
Intrinsic permeability (cm ²)	1.6E-10	Calibrated
Disconnectedness index	10	Calibrated
Porosity	0.45	Laboratory analysis
Depth to water table (m)		Site specific (to RGA) based on field observation
SWMU 2	19.5	
SWMU 3	19.8	
SWMU 4	19.2	
SWMU 5	18.3	
SWMU 6	19.2	
SWMU 7	18.3	
SWMU 30	18.6	
SWMU 145	17.7	
Fraction of organic carbon (%)	0.08	Laboratory analysis
Frendlich equation exponent	1	SESOIL default value

The chemical-specific parameters used in the SESOIL modeling included each analyte's solubility in water, organic carbon partition coefficient (K_{oc}), Henry's Law constant, distribution coefficient (K_d), diffusion coefficients in air and water, and, for TCE and radionuclides, degradation rate constant. The chemical-specific parameters are presented for each SWMU analyte in Sections E.3.1.1 through E.3.1.8. The K_d values for organic compounds were derived using the following relationship.

$$K_d = K_{oc} \times f_{oc}$$

where: K_d is the distribution coefficient,

K_{oc} is the organic carbon partition coefficient, and

f_{oc} is the fraction of organic carbon for source area soils.

The f_{oc} used for the unsaturated zone at PGDP was 0.08 (DOE 1998a).

AT123D. AT123D Simulation of Waste Transport in the Aquifer System (Yeh 1981) was used for saturated flow and contaminant transport modeling. AT123D computes the spatial-temporal concentration distribution of chemicals in the aquifer system and predicts the transient spread of a chemical plume through a groundwater aquifer. The fate and transport processes accounted for in AT123D are advection, dispersion, adsorption/retardation, and decay. This model can be used as a tool for estimating the dissolved concentration of a chemical in three dimensions in the groundwater resulting from a mass release (either continuous or instant or depleting source) from a source. In the present modeling, the time varying mass loading was transferred from the SESOIL output file, and the concentrations of analytes were estimated at the selected POEs. The chemical-specific parameters match those used in SESOIL modeling, except no degradation of TCE was assumed in the RGA. The chemical-specific parameters are presented for each SWMU analyte in Sections E.3.1.1 through E.3.1.8. Excluding the distance to the POEs, Table E.3.2 presents the hydrogeologic parameters used for saturated flow and contaminant transport modeling for the BGOU RI. The AT123D flow model was abstracted from the calibrated flow model of the site developed in MODFLOW by using the flow parameters for hydraulic gradients and hydraulic conductivities of the RGA from the numerical model as described below.

Table E.3.2. Hydrogeologic Parameters Used in AT123D Modeling for the BGOU RI

Input Parameter	Value	Source
Bulk density (kg/m ³)	1,670	Laboratory analysis
Effective porosity	0.3	PGDP sitewide model calibrated value
Hydraulic conductivity (m/hour)		PGDP sitewide model calibrated value
SWMUs 2, 3, 4, 5, 6, 7, and 30	19.05	
SWMU 145	6.35	
Hydraulic gradient (m/m)		PGDP sitewide model calibrated value
SWMUs 2 and 3	0.0002	
SWMU 4	0.0002	
SWMU 5	0.0002	
SWMUs 6 and 145	0.0008	
SWMU 7	0.0003	
SWMU 30	0.00036	
Aquifer thickness	9.14 m 30 ft	Site average
Longitudinal dispersivity (m)	15	Approximate values used in the past
Density of water (kg/m ³)	1,000	Default
Fraction of organic carbon (%)	0.02 ^a	Laboratory analysis
Source Area	Variable	These dimensions were derived from the SADA Analysis for each analyte.

^a UCRS soils were assigned an f_{oc} value of 0.08% while the RGA was assigned an f_{oc} value of 0.02%

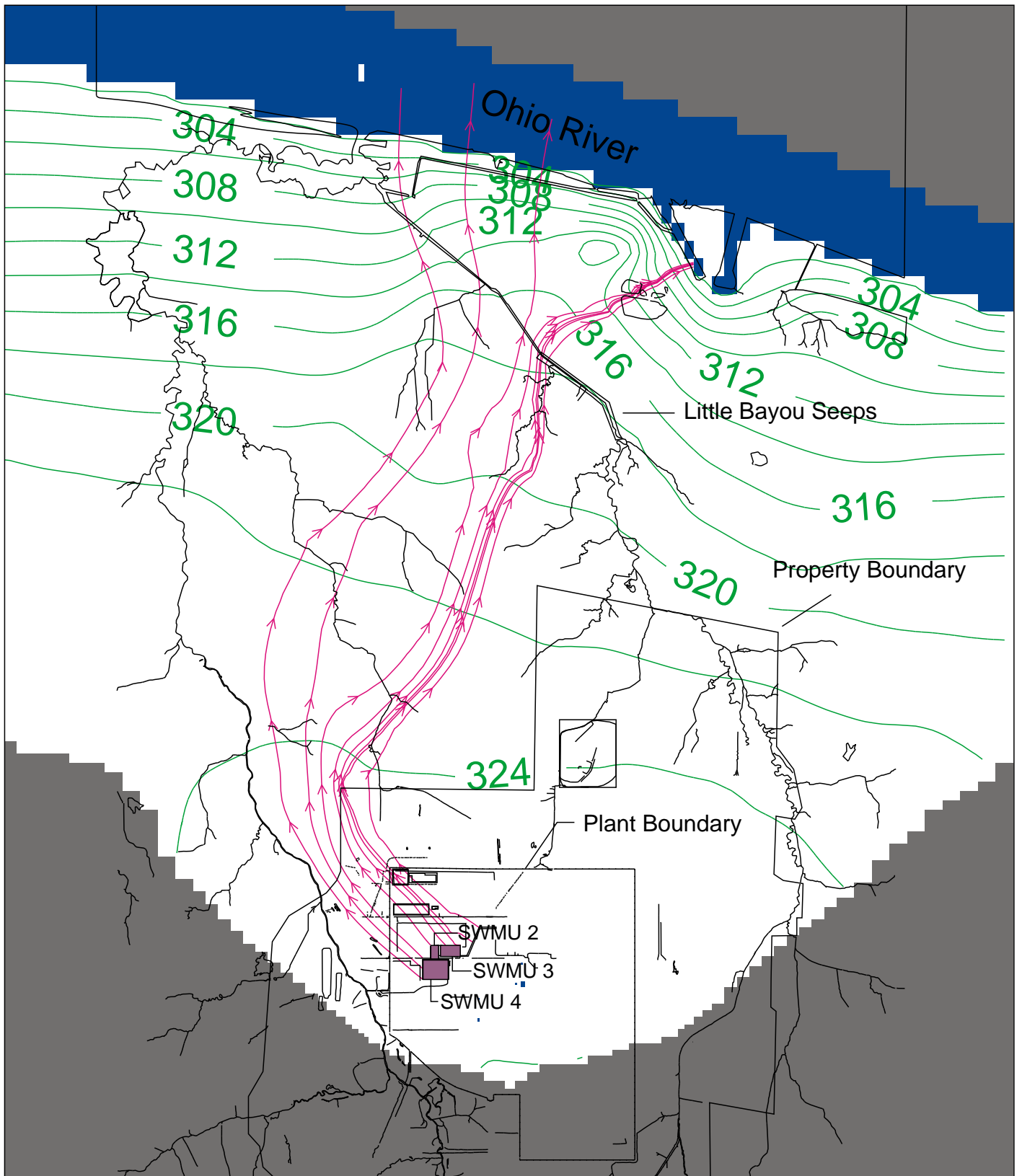
MODFLOW/MODPATH. The U.S. Geological Survey’s Modular Three-Dimensional Finite-Difference Ground-Water Model/A Particle-Tracking Postprocessor Model for MODFLOW (USGS 2005) computer codes were used to evaluate the particle tracks from selected BGOU SWMUs and to determine the distances to the POEs, hydraulic gradients, and hydraulic conductivities of the RGA for input into the AT123D model. Figure E.3.1 shows MODPATH particle tracks for all of the BGOU SWMUs.

E.3.1.1 SWMU 2

The C-749 Uranium Burial Ground (SWMU 2) is located within the west-central section of the plant. SWMU 2 was used from 1951 to 1977 for the disposal of uranium and uranium-contaminated wastes. Disposal records for SWMU 2 indicate that 270 tons of uranium, 59,000 gal of oils, and 450 gal of TCE were disposed of in the unit (DOE 1999a).

E.3.1.1.1 Conceptual model for source areas at SWMU 2

SWMU 2 occupies an area of approximately 32,000 ft² (0.73 acres); with approximate dimensions of 160 ft by 200 ft. The thickness of the UCRS is estimated to be 64 ft (depth to the top of the RGA). The primary waste at SWMU 2 consists of uranium and uranium alloys, placed in pits that were excavated to depths of 7 to 17 ft. Other wastes at the unit consist of uranyl fluoride and TCE.



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U.S. DEPARTMENT OF ENERGY
DOE PORTSMOUTH/PADUCAH PROJECT OFFICE
PADUCAH GASEOUS DIFFUSION PLANT

PADUCAH
Remediation Services
A Portage Shaw Joint Venture Company

Paducah Gaseous Diffusion Plant

- █ River
- █ Particle Tracks
- █ Hydraulic Head (ft)

0 100 500 1000
METERS

TRUE NORTH
PLANT NORTH
20 deg

Figure E.3.1. Particle Tracking Results for SWMUs 2, 3, and 4
E-25

The uranium buried at PGDP most likely is in the metallic state or is coated with uranium (IV) oxide. Neither of these forms of uranium is very susceptible to leaching. The kinetics of dissolution of the buried metal and uranium (IV) oxide is controlled by the amount of oxygen and carbon dioxide that leaches through the waste. Site records show that much of the metal was coated with oil. Petroleum-based oils are resistant to chemical and biological degradation and from leaching by percolating waters. In addition, oils consume oxygen as they slowly degrade, which lowers the oxidation-reduction potential. Under such conditions, uranium dissolution is negligible (ORNL 1998).

The conceptual model for SWMU 2 is that contaminants in the disposal site directly impacted soils below and adjacent to the areas where the material was buried and, through vertical infiltration in soil, potentially may impact the groundwater underlying these sources. The infiltrating groundwater migrates vertically through the UCRS and laterally in the RGA, which could transport the contaminants to the POEs.

E.3.1.1.2 Contaminant transport modeling for SWMU 2 using SESOIL and AT123D

SESOIL allows for the input of 4 soil layers with up to 10 sublayers within each soil layer for contaminant source input. For this modeling effort, the soil zones were arranged in four layers. The first soil layer represented the SADA surface soil data from 0 to 1 ft deep. The second soil layer was 9 ft thick representing SADA layer 2 from 1 to 9 ft in depth, the third layer was subdivided into 4 sublayers of equal thickness (10 ft each) representing SADA layers 3 through 6. SESOIL soil layer 4 was subdivided into 4 sublayers of equal thickness (3.5 ft each) to represent the total thickness of SADA layer 7 from 50 ft to 64 ft in depth. Figure E.3.2 provides an illustration of the SADA and SESOIL contaminant loading layers. Table E.3.3 presents the analytes remaining after the screening process and the source terms for each analyte source zone at SWMU 2. Table E.3.4 lists the chemical-specific parameters applicable to the SESOIL model of SWMU 2. The distances to the POEs used in the AT123D model for SWMU 2 are 1,528 ft to the plant boundary, 3,753 ft to the property boundary, and 21,126 ft to the Ohio River. SWMU 2 particle tracks do not travel to the Little Bayou seeps.

E.3.1.1.3 Groundwater modeling results for SWMU 2

Table E.3.5 summarizes the predicted maximum groundwater concentrations at the POEs for the analytes modeled at SWMU 2. These contaminants were predicted by SESOIL to reach the water table within the 1,000 year period in concentrations that were greater than the groundwater background or greater than the groundwater child no action levels. Several contaminants that originally passed the screening for groundwater did not reach the water table in 1,000 years [i.e., antimony, benzo(a)pyrene, mercury, nickel, ²³⁵U, vanadium, PCB-1254, and Pu-239.] or exhibited groundwater concentrations that were less than the groundwater background or the groundwater child no action levels (i.e., zinc) (see Section 5.4 of the main text).

As shown in Table E.3.5, the predicted maximum groundwater concentrations of *cis*-1,2-DCE and TCE at the plant boundary, property boundary, and Ohio River are predicted to exceed the MCL in the future. None of the other analytes are expected to attain concentrations that exceed their respective MCLs at any of the POEs.

SESOIL LAYER	SESOIL SUB LAYER	SADA LAYER
1 (0-1 ft)	1 (1 ft)	L1
2 (1-10 ft)	1 (9 ft)	L2
3 (10-50 ft)	1 (10 ft)	L3
	2 (10 ft)	L4
	3 (10 ft)	L5
	4 (10 ft)	L6
4 (50-64 ft)	1 (3.5 ft)	L7
	2 (3.5 ft)	
	3 (3.5 ft)	
	4 (3.5 ft)	

Figure E.3.2. Conceptualization of the SADA and SESOIL Layers for Contaminant Loading

Table E.3.3. Summary of Source Term Characteristics Developed by SADA for SWMU 2

SADA Layer	Depth (ft)	Average (mg/kg) ^a	Area (ft ²)	Volume (ft ³)	Mass (gm) ^a	Concentration Factor	Adjusted Average (mg/kg) ^a
Antimony							
L1	0-1	11.60	1.77E+05	1.77E+05	8.48E+04	1.15	13.28
L2	01-10	10.05	1.54E+05	1.54E+06	6.41E+05	1.00	10.05
L3	10-20	0.00	0.00	0.00	0.00	0.00	0.00
L4	20-30	0.00	0.00	0.00	0.00	0.00	0.00
L5	30-40	0.00	0.00	0.00	0.00	0.00	0.00
L6	40-50	0.00	0.00	0.00	0.00	0.00	0.00
L7	50-64	0.00	0.00	0.00	0.00	0.00	0.00
Total Mass					7.26E+05		
Arsenic							
L1	0-1	22.10	4.80E+04	4.80E+04	4.39E+04	1.21	26.79
L2	01-10	8.36	3.96E+04	3.96E+05	1.37E+05	1.00	8.36
L3	10-20	6.85	3.96E+04	4.36E+05	1.23E+05	1.00	6.85
L4	20-30	5.77	3.96E+04	4.36E+05	1.04E+05	1.00	5.77
L5	30-40	6.47	3.96E+04	4.36E+05	1.16E+05	1.00	6.47
L6	40-50	5.87	3.96E+04	4.36E+05	1.06E+05	1.00	5.87
L7	50-64	4.92	3.96E+04	3.56E+05	7.26E+04	1.00	4.92
Total Mass					7.03E+05		
Benzo(a)pyrene							
L1	0-1	0.14	1.88E+04	1.88E+04	1.09E+02	1.00	0.14
L2	01-10	0.00	0.00	0.00	0.00	0.00	0.00
L3	10-20	0.00	0.00	0.00	0.00	0.00	0.00
L4	20-30	0.00	0.00	0.00	0.00	0.00	0.00
L5	30-40	0.00	0.00	0.00	0.00	0.00	0.00
L6	40-50	0.00	0.00	0.00	0.00	0.00	0.00
L7	50-64	0.00	0.00	0.00	0.00	0.00	0.00
Total Mass					1.09E+02		
cis-1,2-DCE							
L1	0-1	0.00	0.00E+00	0.00E+00	0.00E+00	0.00	0.00
L2	01-10	1.06	1.64E+04	1.64E+05	7.17E+03	2.41	2.55
L3	10-20	114.72	6.80E+03	7.48E+04	3.55E+05	1.00	114.72
L4	20-30	70.45	9.60E+03	1.06E+05	3.08E+05	1.41	99.46
L5	30-40	80.50	8.40E+03	9.24E+04	3.08E+05	1.24	99.44
L6	40-50	44.87	1.16E+04	1.28E+05	2.37E+05	1.71	76.54
L7	50-64	37.17	1.12E+04	1.01E+05	1.55E+05	1.65	61.22
Total Mass					1.37E+06		
Manganese							
L1	0-1	372.83	4.80E+04	4.80E+04	7.40E+05	1.21	451.92
L2	01-10	369.68	3.96E+04	3.96E+05	6.05E+06	1.00	369.68
L3	10-20	386.99	3.96E+04	4.36E+05	6.97E+06	1.00	386.99
L4	20-30	378.32	3.96E+04	4.36E+05	6.81E+06	1.00	378.32
L5	30-40	377.03	3.96E+04	4.36E+05	6.79E+06	1.00	377.03
L6	40-50	316.18	3.96E+04	4.36E+05	5.69E+06	1.00	316.18
L7	50-64	277.65	3.96E+04	3.56E+05	4.09E+06	1.00	277.65
Total Mass					3.71E+07		

Table E.3.3. Summary of Source Term Characteristics Developed by SADA for SWMU 2 (Continued)

SADA Layer	Depth (ft)	Average (mg/kg) ^a	Area (ft ²)	Volume (ft ³)	Mass (gm) ^a	Concentration Factor	Adjusted Average (mg/kg) ^a
Mercury							
L1	0-1	0.29	3.72E+04	3.72E+04	4.41E+02	1.00	0.29
L2	01-10	0.02	1.60E+03	1.60E+04	1.19E+01	0.04	0.0008
L3	10-20	0.02	8.00E+02	8.80E+03	6.18E+00	0.02	0.0004
L4	20-30	0.02	8.00E+02	8.80E+03	6.18E+00	0.02	0.0004
L5	30-40	0.00	0.00	0.00	0.00	0.00	0.00
L6	40-50	0.00	0.00	0.00	0.00	0.00	0.00
L7	50-64	0.00	0.00	0.00	0.00	0.00	0.00
Total Mass					4.65E+02		
Naphthalene							
L1	0-1	0.27	3.72E+04	3.72E+04	4.17E+02	1.00	0.27
L2	01-10	0.00	0.00	0.00	0.00	0.00	0.00
L3	10-20	0.00	0.00	0.00	0.00	0.00	0.00
L4	20-30	0.00	0.00	0.00	0.00	0.00	0.00
L5	30-40	0.00	0.00	0.00	0.00	0.00	0.00
L6	40-50	0.00	0.00	0.00	0.00	0.00	0.00
L7	50-64	0.00	0.00	0.00	0.00	0.00	0.00
Total Mass					4.17E+02		
Nickel							
L1	0-1	29.10	4.80E+04	4.80E+04	5.77E+04	1.32	38.37
L2	01-10	11.31	3.56E+04	3.56E+05	1.66E+05	0.98	11.06
L3	10-20	11.97	3.64E+04	4.00E+05	1.98E+05	1.00	11.97
L4	20-30	10.76	3.20E+04	3.52E+05	1.57E+05	0.88	9.46
L5	30-40	10.88	3.28E+04	3.61E+05	1.62E+05	0.90	9.81
L6	40-50	10.95	2.92E+04	3.21E+05	1.45E+05	0.80	8.78
L7	50-64	10.04	2.88E+04	2.59E+05	1.08E+05	0.79	7.94
Total Mass					9.94E+05		
⁹⁹Tc							
L1	0-1	1.82	5.20E+04	5.20E+04	3.92E+09	1.67	3.04
L2	01-10	0.58	3.12E+04	3.12E+05	7.44E+09	1.00	0.58
L3	10-20	0.32	1.96E+04	2.16E+05	2.86E+09	0.63	0.20
L4	20-30	0.10	1.68E+04	1.85E+05	7.80E+08	0.54	0.06
L5	30-40	0.16	1.20E+04	1.32E+05	8.59E+08	0.38	0.06
L6	40-50	0.07	1.72E+04	1.89E+05	5.42E+08	0.55	0.04
L7	50-64	0.07	1.32E+04	1.19E+05	3.27E+08	0.42	0.03
Total Mass					1.67E+10		
TCE							
L1	0-1	0.00	0.00E+00	0.00E+00	0.00E+00		
L2	01-10	0.13	9.60E+03	9.60E+04	5.11E+02	1.04	0.13
L3	10-20	42.65	9.20E+03	1.01E+05	1.78E+05	1.00	42.65
L4	20-30	24.28	1.16E+04	1.28E+05	1.28E+05	1.26	30.61
L5	30-40	14.58	1.16E+04	1.28E+05	7.69E+04	1.26	18.39
L6	40-50	8.94	1.28E+04	1.41E+05	5.20E+04	1.39	12.44
L7	50-64	0.20	1.08E+04	9.72E+04	8.06E+02	1.17	0.24
Total Mass					4.37E+05		

Table E.3.3. Summary of Source Term Characteristics Developed by SADA for SWMU 2 (Continued)

SADA Layer	Depth (ft)	Average (mg/kg) ^a	Area (ft ²)	Volume (ft ³)	Mass (gm) ^a	Concentration Factor	Adjusted Average (mg/kg) ^a
²³⁹Pu							
L1	0-1	1.36	8.70E+05	8.70E+05	4.89E+04	1.01	1.37
L2	01-10	1.36	8.70E+05	8.70E+06	4.89E+05	1.01	1.37
L3	10-20	1.34	8.64E+05	9.50E+06	5.28E+05	1.00	1.34
L4	20-30	1.52	7.44E+05	8.18E+06	5.16E+05	0.86	1.31
L5	30-40	1.47	7.53E+05	8.28E+06	5.04E+05	0.87	1.28
L6	40-50	1.44	7.55E+05	8.30E+06	4.93E+05	0.87	1.26
L7	50-64	1.36	7.70E+05	6.93E+06	3.91E+05	0.89	1.22
			Total Mass		2.97E+06		
PCB-1248							
L1	0-1	0.06	2.56E+04	2.56E+04	6.14E+01	0.82	0.05
L2	01-10	0.06	2.48E+04	2.48E+05	5.95E+02	0.79	0.05
L3	10-20	0.88	3.12E+04	3.43E+05	1.25E+04	1.00	0.88
L4	20-30	0.81	3.04E+04	3.34E+05	1.12E+04	0.97	0.79
L5	30-40	0.69	3.08E+04	3.39E+05	9.65E+03	0.99	0.68
L6	40-50	0.59	3.32E+04	3.65E+05	8.94E+03	1.06	0.63
L7	50-64	0.00	0.00E+00	0.00E+00	0.00E+00	0.00	0.00
			Total Mass		7.76E+05		
PCB-1260							
L1	0-1	0.11	1.98E+05	1.98E+05	8.91E+02	1.04	0.11
L2	01-10	0.11	1.97E+05	1.97E+06	8.84E+03	1.03	0.11
L3	10-20	0.11	1.92E+05	2.11E+06	9.31E+03	1.00	0.11
L4	20-30	0.11	1.88E+05	2.06E+06	9.05E+03	0.98	0.10
L5	30-40	0.10	1.78E+05	1.95E+06	8.38E+03	0.93	0.10
L6	40-50	0.10	1.70E+05	1.87E+06	7.92E+03	0.89	0.10
L7	50-64	0.10	1.60E+05	1.44E+06	5.98E+03	0.83	0.08
			Total Mass		5.04E+04		
²³⁴U							
L1	0-1	16.01	5.20E+04	5.20E+04	3.44E+10	1.25	20.01
L2	01-10	14.33	4.16E+04	4.16E+05	2.46E+11	1.00	14.33
L3	10-20	0.81	4.28E+04	4.71E+05	1.57E+10	1.03	0.83
L4	20-30	0.76	4.04E+04	4.44E+05	1.39E+10	0.97	0.73
L5	30-40	0.83	4.12E+04	4.53E+05	1.56E+10	0.99	0.82
L6	40-50	0.72	4.08E+04	4.49E+05	1.34E+10	0.98	0.71
L7	50-64	0.64	4.20E+04	3.78E+05	9.92E+09	1.01	0.64
			Total Mass		3.49E+11		
²³⁵U							
L1	0-1	2.73	5.20E+04	5.20E+04	5.86E+09	1.30	3.55
L2	01-10	3.43	4.00E+04	4.00E+05	5.67E+10	1.00	3.43
L3	10-20	0.09	4.00E+04	4.40E+05	1.72E+09	1.00	0.09
L4	20-30	0.08	4.00E+04	4.40E+05	1.48E+09	1.00	0.08
L5	30-40	0.07	4.00E+04	4.40E+05	1.34E+09	1.00	0.07
L6	40-50	0.07	4.00E+04	4.40E+05	1.27E+09	1.00	0.07
L7	50-64	0.00	0.00	0.00	0.00E+00	0.00	0.00
			Total Mass		6.83E+10		
²³⁸U							
L1	0-1	88.34	5.20E+04	5.20E+04	1.90E+11	1.25	110.43
L2	01-10	83.85	4.16E+04	4.16E+05	1.44E+12	1.00	83.85
L3	10-20	1.49	4.28E+04	4.71E+05	2.90E+10	1.03	1.53
L4	20-30	1.10	4.04E+04	4.44E+05	2.02E+10	0.97	1.07

Table E.3.3. Summary of Source Term Characteristics Developed by SADA for SWMU 2 (Continued)

SADA Layer	Depth (ft)	Average (mg/kg) ^a	Area (ft ²)	Volume (ft ³) ²³⁸ U	Mass (gm) ^a	Concentration Factor	Adjusted Average (mg/kg) ^a
L5	30-40	1.02	4.12E+04	4.53E+05	1.92E+10	0.99	1.01
L6	40-50	0.88	4.08E+04	4.49E+05	1.64E+10	0.98	0.87
L7	50-64	0.71	4.20E+04	3.78E+05	1.11E+10	1.01	0.71
			Total Mass		1.73E+12		
Uranium							
L1	0-1	167.67	4.80E+04	4.80E+04	3.33E+05	7.06	1,183.53
L2	01-10	798.80	6.80E+03	6.80E+04	2.25E+06	1.00	798.80
L3	10-20	14.51	8.80E+03	9.68E+04	5.81E+04	1.29	18.78
L4	20-30	18.13	1.44E+04	1.58E+05	1.19E+05	2.12	38.39
L5	30-40	13.60	1.12E+04	1.23E+05	6.93E+04	1.65	22.40
L6	40-50	12.45	8.00E+03	8.80E+04	4.53E+04	1.18	14.65
L7	50-64	5.79	9.20E+03	8.28E+04	1.98E+04	1.35	7.84
			Total Mass		2.89E+06		
Vanadium							
L1	0-1	28.55	4.80E+04	4.80E+04	5.67E+04	1.26	36.06
L2	01-10	19.14	3.80E+04	3.80E+05	3.01E+05	1.00	19.14
L3	10-20	20.37	3.80E+04	4.18E+05	3.52E+05	1.00	20.37
L4	20-30	17.44	3.80E+04	4.18E+05	3.01E+05	1.00	17.44
L5	30-40	17.44	3.96E+04	4.36E+05	3.14E+05	1.04	18.17
L6	40-50	16.62	3.96E+04	4.36E+05	2.99E+05	1.04	17.32
L7	50-64	15.05	3.96E+04	3.56E+05	2.22E+05	1.04	15.68
			Total Mass		1.85E+06		
Zinc							
L1	0-1	104.88	4.80E+04	4.80E+04	2.08E+05	1.00	104.88
L2	01-10	32.00	3.20E+03	3.20E+04	4.23E+04	0.07	2.13
L3	10-20	35.78	4.00E+03	4.40E+04	6.51E+04	0.08	2.98
L4	20-30	34.70	3.20E+03	3.52E+04	5.05E+04	0.07	2.31
L5	30-40	35.30	4.80E+03	5.28E+04	7.71E+04	0.10	3.53
L6	40-50	35.72	4.80E+03	5.28E+04	7.80E+04	0.10	3.57
L7	50-64	34.10	4.80E+03	4.32E+04	6.09E+04	0.10	3.41
			Total Mass		5.82E+05		

^a Radionuclides are in units of pCi/g for concentrations and pCi for mass.

Table E.3.4. Chemical-Specific Parameters of the Analytes Used in SESOIL Modeling of SWMU 2

Analyte	Mol. Wt. (MW) (g/mol)	Solubility in water (mg/L)	Diffusion in air (cm ² /s)	Diffusion in water (m ² /hr)	Henry's Constant (atm.m3/mol)	Koc (L/kg)	Kd ^a (L/kg)	Half Life (years)
Antimony	121.75	1.00E+07	NA	3.60E-07	NA	NA	45	infinite
Arsenic	74.92	1.00E+07	NA	3.60E-07	NA	NA	29	infinite
Benzo(a)pyrene	252.32	1.62E-03	4.3E-02	3.24E-06	1.13E-06	9.69E+05	772	infinite
<i>cis</i> -1,2-DCE	96.94	3.50E+03	0.07	4.07E-06	4.08E-03	35.5	0.028	infinite
Manganese	54.94	1.00E+07	NA	1.29E-07	NA	NA	65	infinite
Mercury	200.59	6.00E-02	3.07E-02	2.27E-06	2.44E-02	NA	52	infinite
Naphthalene	128.16	31.0	0.059	2.70E-06	4.83E-04	1.19E+03	0.95	infinite
Nickel	58.69	1.00E+07	NA	3.60E-07	NA	NA	300	infinite
PCB-1248	288	1.70E-02	1.75E-02	2.38E-06	1.60E-04	2.51E+04	20	Infinite
PCB-1260	375.7	2.70E-02	1.38E-02	1.56E-06	7.40E-05	2.07E+05	165.6	Infinite
²³⁹ Pu	239	1.00E+07	NA	3.60E-07	NA	NA	550	2.41E+04
⁹⁹ Tc	99	1.00E+07	NA	3.60E-07	NA	NA	0.2	2.13E+05
TCE	131	1,100	0.08	3.28E-06	0.0103	94	0.0752	26.6
²³⁴ U	234	1.00E+07	NA	3.60E-07	NA	NA	66.8	2.44E+05
²³⁵ U	235	1.00E+07	NA	3.60E-07	NA	NA	66.8	7.04E+08
²³⁸ U	238	1.00E+07	NA	3.60E-07	NA	NA	66.8	4.47E+09
Uranium	238	1.00E+07	NA	3.60E-07	NA	NA	66.8	4.47E+09
Vanadium	50.94	1.00E+07	NA	3.60E-07	NA	NA	1,000	infinite
Zinc	67.41	1.00E+07	NA	3.60E-07	NA	NA	62	infinite

^a The soil/water distribution coefficient (Kd) of an organic compound depends on the soil's organic content (f_{oc}) and compound's organic partition coefficient (Koc). Kd values presented for organic compounds are for UCRS soils (with f_{oc} value of 0.08%) only. Kds used in AT123D are different due to the f_{oc} of 0.02% in the RGA.

Table E.3.5. Concentrations of the Analytes in Groundwater Predicted in SESOIL and AT123D Modeling of SWMU 2

Analyte	Predicted Maximum Groundwater Concentration ^{a,b}				MCL (mg/L or pCi/L)
	SWMU (mg/L)	Plant Boundary (mg/L)	Property Boundary (mg/L)	Ohio River (mg/L)	
Arsenic	<i>3.54E-02</i>	2.91E-03	8.35E-09	0	0.01
<i>cis</i> -1,2-DCE	<i>1.15E+01</i>	<i>1.74E+00</i>	<i>8.58E-01</i>	<i>3.38E-01</i>	0.07
Manganese	<i>7.16E-01</i>	1.86E-05	0	0	^d
Naphthalene	9.38E-04	1.57E-04	8.27E-05	3.42E-05	^d
PCB-1248	1.54E-03	1.28E-09	0	0	^d
PCB-1260	8.73E-05	0	0	0	^d
⁹⁹ Tc	1.02E+02	1.59E+01	8.06E+00	3.11E+00	900 ^c
TCE	<i>1.48E+00</i>	<i>2.17E-01</i>	<i>1.10E-01</i>	<i>4.12E-02</i>	0.005
²³⁴ U	1.58E+00	1.75E-05	0	0	20
²³⁸ U	1.81E+00	2.03E-05	0	0	20
Uranium	9.86E-03	8.33E-08	0	0	0.03

^a Values in bold, italic font exceed the analyte's MCL

^b Radionuclide concentrations are in pCi/L

^c ⁹⁹Tc MCL based on a critical organ dose at 4 mrem/yr from drinking water consumption

^d MCLs not available for these contaminants

The hazard quotients (HQs) and excess lifetime cancer risks (ELCRs) calculated in accordance with the Risk Methods Document (DOE 2001) are presented in Table E.3.6. (Appendix F provides a full

description of the risk assessment methodology and calculations.) The predicted TCE concentrations result in the greatest HQs and cancer risks; therefore, TCE is the most important analyte for contaminant migration at SWMU 2, while *cis*-1,2-DCE also provides (HQs) greater than 1. Arsenic also provides a cancer risk greater than 10^{-5} .

Table E.3.6. Hazard and Cancer Risk Predicted from Maximum Groundwater Concentrations Derived in Modeling of SWMU 2 Using SESOIL and AT123D^a

Analyte	SWMU		Plant Boundary		Property Boundary		Ohio River	
	Hazard Quotient	Cancer Risk	Hazard Quotient	Cancer Risk	Hazard Quotient	Cancer Risk	Hazard Quotient	Cancer Risk
Arsenic	11.3	9.38E-04	0.9	7.7E-05	<0.1	<1.0E-06	^b	^b
<i>cis</i> -1,2-DCE	607	^b	91.9	^b	45.3	^b	17.9	^b
Manganese	1.52	^b	<0.1	^b	^b	^b	^b	^b
Naphthalene	0.47	^b	0.1	^b	<0.1	^b	<0.1	^b
PCB-1248	^b	1.70E-04	^b	<1.00E-06	^b	^b	^b	^b
PCB-1260	^b	6.95E-05	^b	^b	^b	^b	^b	^b
⁹⁹ Tc	^b	5.60E-06	^b	<1.0E-06	^b	<1.0E-06	^b	<1.0E-06
TCE	676	3.09E-02	99.1	6.7E-03	50.3	3.4E-03	4.6	1.3E-03
²³⁴ U	^b	2.23E-06	^b	<1.0E-06	^b	^b	^b	^b
²³⁸ U	^b	2.68E-06	^b	<1.0E-06	^b	^b	^b	^b
Uranium	1.58	^b	0.1	^b	^b	^b	^b	^b

^a Contaminants with a HQ greater than 0.1 or a cancer risk greater than 1.00E-06 are considered analytes.

^b Value not calculated since the groundwater concentrations was reported as zero at this POE by AT123D, or the contaminant did not have a reported cancer slope factor or chemical toxicity RfD.

Figures 5.2 through 5.4 in Section 5 of the main text, show the predicted concentrations over time at each POE for analytes with a HQ greater than 0.1 and/or a risk greater than 1.0E-06. As shown in these figures, arsenic is predicted to continue rising in concentration at 1,000 years at the plant boundary, but has not reached the property boundary or Ohio River in the 1,000 year period. Both *cis*-1,2-DCE and TCE are predicted to exceed the their MCLs at all POEs within approximately 100 years and then decline in concentration below the MCLs.

E.3.1.2 SWMU 3

The C-404 Low-Level Radioactive Waste Burial Ground (SWMU 3) is located in the west-central section of PGDP. PGDP operated SWMU 3 as a surface impoundment from approximately 1952 until early 1957. During this time, all influents to the impoundment originated from C-400. In 1957, the C-404 surface impoundment was converted to a solid waste disposal facility for solid uranium-contaminated wastes. Approximately 6,615,000 lb of uranium-contaminated wastes were disposed of at SWMU 3. The total volume is approximately 260,000 ft³. Some uranium-contaminated waste also is contaminated with TCE, radionuclides, and metals. In 1986, the disposal of all waste at C-404 Landfill was halted, and a portion of the disposed waste was found to be Resource Conservation and Recovery Act (RCRA)-hazardous. The landfill was covered with a RCRA multilayered cap and certified closed in 1987 (DOE 1987; DOE 1989).

E.3.1.2.1 Conceptual model for source areas at SWMU 3

SWMU 3 occupies an area of approximately 53,200 ft² (1.2 acres), with approximate dimensions of 140 by 380 ft. The thickness of the UCRS is estimated to be 65 ft (depth to the top of the RGA). The primary wastes at SWMU 3 consist of uranium precipitated from aqueous solutions, uranium tetrafluoride, uranium metal, uranium oxides, and radioactively contaminated trash. There are no records documenting the cleanout of sludges and sediments from the pond when it was converted to a landfill. A partial clay cap was installed on the eastern end of the landfill in 1982 (DOE 1987).

The conceptual model for SWMU 3 is that contaminants in the disposal site directly impacted soils below and adjacent to the areas where the material was landfilled and, through vertical infiltration in soil, may potentially impact the groundwater underlying these sources. The infiltrating groundwater migrates vertically through the UCRS and laterally in the RGA, which could transport the contaminants to the POEs.

E.3.1.2.2 Contaminant transport modeling for SWMU 3 using SESOIL and AT123D

For this modeling effort, the soil zones were arranged in four layers. Although SWMU 3 waste is contained in a mounded area, the mounding was not modeled in SESOIL. Instead the waste was assumed to be located at grade. The first soil layer represented the SADA surface soil data from 0 to 1 ft deep. The second soil layer was 9 ft thick representing SADA layer 2 from 1 to 9 ft in depth, the third layer was subdivided into 4 sublayers of equal thickness (10 ft each) representing SADA layers 3 through 6. SESOIL soil layer 4 was subdivided into 4 sublayers of equal thickness (3.75 ft each) to represent the total thickness of SADA layer 7 from 50 ft to 65 ft in depth. Table E.3.7 presents the analytes remaining after the screening process and the source terms for each analyte. Table E.3.8 lists the chemical-specific parameters applicable to the SESOIL model of SWMU 3. The distances to the POEs used in the AT123D model for SWMU 3 are 2,049 ft to the plant boundary, 4,455 ft to the property boundary, and 16,598 ft to the Little Bayou seeps.

Table E.3.7. Summary of Source Term Characteristics Developed by SADA for SWMU 3

SADA Layer	Depth (ft)	Average (mg/kg) ^a	Area (ft ²)	Volume (ft ³)	Mass (gm) ^a	Concentration Factor	Adjusted Average (mg/kg) ^a
Arsenic							
L1	0-1	4.97	4.20E+05	4.20E+05	8.62E+04	0.51	2.54
L2	01-10	3.89	8.20E+05	8.20E+06	1.32E+06	1.00	3.89
L3	10-20	2.42	7.70E+05	8.47E+06	8.49E+05	0.94	2.28
L4	20-30	2.50	7.70E+05	8.47E+06	8.75E+05	0.94	2.35
L5	30-40	2.44	7.70E+05	8.47E+06	8.53E+05	0.94	2.29
L6	40-50	2.44	7.70E+05	8.47E+06	8.53E+05	0.94	2.29
L7	50-65	2.28	7.60E+05	6.84E+06	6.44E+05	0.93	2.11
Total Mass					5.48E+06		
Manganese							
L1	0-1	359.57	4.20E+05	4.20E+05	6.24E+06	0.50	179.79
L2	01-10	276.44	8.40E+05	8.40E+06	9.60E+07	1.00	276.44
L3	10-20	177.46	8.40E+05	9.24E+06	6.78E+07	1.00	177.46
L4	20-30	184.17	8.40E+05	9.24E+06	7.04E+07	1.00	184.17
L5	30-40	185.80	8.40E+05	9.24E+06	7.10E+07	1.00	185.80
L6	40-50	184.36	8.40E+05	9.24E+06	7.04E+07	1.00	184.36
L7	50-65	178.20	8.40E+05	7.56E+06	5.57E+07	1.00	178.20
Total Mass					4.37E+08		
Mercury							
L1	0-1	0.02	1.40E+05	1.40E+05	1.34E+02	1.08	0.02
L2	01-10	0.02	1.20E+05	1.20E+06	9.92E+02	0.92	0.02
L3	10-20	0.02	1.20E+05	1.32E+06	1.09E+03	0.92	0.02
L4	20-30	0.02	1.30E+05	1.43E+06	1.18E+03	1.00	0.02
L5	30-40	0.02	1.30E+05	1.43E+06	1.18E+03	1.00	0.02
L6	40-50	0.02	1.30E+05	1.43E+06	1.16E+03	1.00	0.02
L7	50-65	0.02	1.40E+05	1.26E+06	1.04E+03	1.08	0.02
Total Mass					6.77E+03		
Molybdenum							
L1	0-1	4.35	2.80E+05	2.80E+05	5.03E+04	1.87	8.12
L2	01-10	3.78	1.50E+05	1.50E+06	2.34E+05	1.00	3.78
L3	10-20	0.00	0.00	0.00	0.00	0.00	0.00
L4	20-30	0.00	0.00	0.00	0.00	0.00	0.00
L5	30-40	0.00	0.00	0.00	0.00	0.00	0.00
L6	40-50	0.00	0.00	0.00	0.00	0.00	0.00
L7	50-65	0.00	0.00	0.00	0.00	0.00	0.00
Total Mass					2.85E+05		
Nickel							
L1	0-1	7.90	4.20E+05	4.20E+05	1.37E+05	0.51	4.05
L2	01-10	9.89	7.20E+05	7.20E+06	2.94E+06	0.88	8.69
L3	10-20	8.20	8.20E+05	9.02E+06	3.06E+06	1.00	8.20
L4	20-30	8.17	8.10E+05	8.91E+06	3.01E+06	0.99	8.07
L5	30-40	8.11	8.10E+05	8.91E+06	2.99E+06	0.99	8.01
L6	40-50	8.04	8.30E+05	9.13E+06	3.03E+06	1.01	8.14
L7	50-65	8.04	8.20E+05	7.38E+06	2.45E+06	1.00	8.04
Total Mass					1.76E+07		

Table E.3.7. Summary of Source Term Characteristics Developed by SADA for SWMU 3 (Continued)

SADA Layer	Depth (ft)	Average (mg/kg) ^a	Area (ft ²)	Volume (ft ³)	Mass (gm) ^a	Concentration Factor	Adjusted Average (mg/kg) ^a
⁹⁹Tc							
L1	0-1	12.58	2.60E+05	2.60E+05	1.35E+11	0.74	9.34
L2	01-10	26.86	3.50E+05	3.50E+06	3.89E+12	1.00	26.86
L3	10-20	0.00	0.00E+00	0.00E+00	0.00E+00	0.00	0.00
L4	20-30	2.40	1.00E+04	1.10E+05	1.09E+10	0.03	0.07
L5	30-40	2.40	1.00E+04	1.10E+05	1.09E+10	0.03	0.07
L6	40-50	2.40	1.00E+04	1.10E+05	1.09E+10	0.03	0.07
L7	50-65	0.00	0.00E+00	0.00E+00	0.00E+00	0.00	0.00
Total Mass					4.05E+12		
TCE							
L1	0-1	0.0063	3.00E+04	3.00E+04	7.85E+00	3.00	0.02
L2	01-10	0.0152	1.00E+04	1.00E+05	6.28E+01	1.00	0.02
L3	10-20	0.0152	1.00E+04	1.10E+05	6.91E+01	1.00	0.02
L4	20-30	0.0000	0.00E+00	0.00E+00	0.00E+00	0.00	0.00
L5	30-40	0.0000	0.00E+00	0.00E+00	0.00E+00	0.00	0.00
L6	40-50	0.0000	0.00E+00	0.00E+00	0.00E+00	0.00	0.00
L7	50-65	0.0000	0.00E+00	0.00E+00	0.00E+00	0.00	0.00
Total Mass					1.40E+02		
²³⁸U							
L1	0-1	1.29	4.20E+05	4.20E+05	2.24E+10	0.70	0.90
L2	01-10	6.67	6.00E+05	6.00E+06	1.65E+12	1.00	6.67
L3	10-20	12.63	1.90E+05	2.09E+06	1.09E+12	0.32	4.00
L4	20-30	12.63	1.90E+05	2.09E+06	1.09E+12	0.32	4.00
L5	30-40	12.26	1.90E+05	2.09E+06	1.06E+12	0.32	3.88
L6	40-50	12.26	1.90E+05	2.09E+06	1.06E+12	0.32	3.88
L7	50-65	10.53	2.00E+05	1.80E+06	7.84E+11	0.33	3.51
Total Mass					6.76E+12		
Uranium							
L1	0-1	15.97	2.90E+05	2.90E+05	1.92E+05	0.52	8.27
L2	01-10	20.56	5.60E+05	5.60E+06	4.76E+06	1.00	20.56
L3	10-20	40.45	1.80E+05	1.98E+06	3.31E+06	0.32	13.00
L4	20-30	38.59	1.90E+05	2.09E+06	3.33E+06	0.34	13.09
L5	30-40	36.09	1.80E+05	1.98E+06	2.95E+06	0.32	11.60
L6	40-50	36.09	1.80E+05	1.98E+06	2.95E+06	0.32	11.60
L7	50-65	40.62	1.50E+05	1.35E+06	2.27E+06	0.27	10.88
Total Mass					1.98E+07		
Vanadium							
L1	0-1	25.30	4.20E+05	4.20E+05	4.39E+05	0.50	12.65
L2	01-10	19.01	8.40E+05	8.40E+06	6.60E+06	1.00	19.01
L3	10-20	17.04	8.40E+05	9.24E+06	6.51E+06	1.00	17.04
L4	20-30	17.24	8.40E+05	9.24E+06	6.58E+06	1.00	17.24
L5	30-40	17.17	8.40E+05	9.24E+06	6.56E+06	1.00	17.17
L6	40-50	17.23	8.40E+05	9.24E+06	6.58E+06	1.00	17.23
L7	50-65	16.99	8.40E+05	7.56E+06	5.31E+06	1.00	16.99
Total Mass					3.86E+07		

Table E.3.7. Summary of Source Term Characteristics Developed by SADA for SWMU 3 (Continued)

SADA Layer	Depth (ft)	Average (mg/kg) ^a	Area (ft ²)	Volume (ft ³)	Mass (gm) ^a	Concentration Factor	Adjusted Average (mg/kg) ^a
Zinc							
L1	0-1	25.73	3.90E+05	3.90E+05	4.15E+05	0.60	15.44
L2	01-10	31.35	6.50E+05	6.50E+06	8.42E+06	1.00	31.35
L3	10-20	30.78	3.80E+05	4.18E+06	5.32E+06	0.58	17.99
L4	20-30	32.05	3.60E+05	3.96E+06	5.25E+06	0.55	17.75
L5	30-40	31.78	3.50E+05	3.85E+06	5.06E+06	0.54	17.11
L6	40-50	31.78	3.50E+05	3.85E+06	5.06E+06	0.54	17.11
L7	50-65	32.27	3.50E+05	3.15E+06	4.20E+06	0.54	17.38
Total Mass					3.37E+07		

^a Radionuclides are in units of pCi/g for concentrations and pCi for mass.

Table E.3.8. Chemical-Specific Parameters of the Analytes Used in SESOIL Modeling of SWMU 3

Analyte	Mol. Wt. (MW) (g/mol)	Solubility in water (mg/L)	Diffusion in air (cm ² /s)	Diffusion in water (m ² /hr)	Henry's Constant (atm.m ³ /mol)	Koc (L/kg)	Kd ^a (L/kg)	Half Life (years)
Arsenic	74.92	1.00E+07	NA	3.60E-07	NA	NA	29	infinite
Manganese	54.94	1.00E+07	NA	1.29E-07	NA	NA	65	infinite
Mercury	200.59	6.00E-02	3.07E-02	2.27E-06	2.44E-02	NA	52	infinite
Molybdenum	95.9	1.00E+07	NA	3.60E-07	NA	NA	10	infinite
Nickel	58.69	1.00E+07	NA	3.60E-07	NA	NA	300	infinite
⁹⁹ Tc	99	1.00E+07	NA	3.60E-07	NA	NA	0.2	2.13E+05
TCE	131	1,100	0.08	3.28E-06	0.0103	94	0.0752	26.6
²³⁸ U	238	1.00E+07	NA	3.60E-07	NA	NA	66.8	4.47E+09
Uranium	238	1.00E+07	NA	3.60E-07	NA	NA	66.8	4.47E+09
Vanadium	50.94	1.00E+07	NA	3.60E-07	NA	NA	1000	infinite
Zinc	67.41	1.00E+07	NA	3.60E-07	NA	NA	62	infinite

^a The soil/water distribution coefficient (Kd) of an organic compound depends on the soil's organic content (f_{oc}) and compound's organic partition coefficient (Koc). Kd values presented for organic compounds are for UCRS soils (with f_{oc} value of 0.08%) only. Kds used in AT123D are different due to the f_{oc} of 0.02% in the RGA.

E.3.1.2.3 Groundwater modeling results for SWMU 3

Table E.3.9 summarizes the predicted maximum groundwater concentrations at the POEs for the analytes modeled at SWMU 3. These contaminants were predicted by SESOIL to reach the water table within the 1,000 year period in concentrations that were greater than the groundwater background or greater than the groundwater child no action levels. Several contaminants that originally passed the screening for groundwater did not reach the water table in 1,000 years (i.e., molybdenum, nickel, and vanadium) or exhibited groundwater concentrations that were less than the groundwater background or the groundwater child no action levels (i.e., mercury, TCE, and zinc) (see Section 5.4 of the main text).

Table E.3.9. Concentrations of the Analytes in Groundwater Predicted in SESOIL and AT123D Modeling of SWMU 3

Analyte	Predicted Maximum Groundwater Concentration ^{a,b}				
	SWMU (mg/L)	Plant Boundary (mg/L)	Property Boundary (mg/L)	Little Bayou seeps (mg/L)	MCL (mg/L or pCi/L)
Arsenic	<i>3.29E-02</i>	1.22E-03	0	0	0.01
Manganese	<i>8.95E-01</i>	4.08E-10	0	0	
⁹⁹ Tc	<i>5.560E+03</i>	1.81E+03	1.36E+03	8.04E+02	900 ^c
²³⁸ U	1.59E+01	7.32E-11	0	0	20
Uranium	<i>4.89E-02</i>	2.27E-13	0	0	0.03

^a Values in bold, italic font exceed the analyte's MCL

^b Radionuclide concentrations are in pCi/L

^c ⁹⁹Tc MCL based on a critical organ dose at 4 mrem/yr from drinking water consumption

^d MCLs not available for these contaminants

As shown in Table E.3.9, the predicted maximum groundwater concentrations for all analytes except ⁹⁹Tc are less than the MCLs for the contaminants at the POEs. The HQs and cancer risks calculated in accordance with the Risk Methods Document (DOE 2001) are presented in Table E.3.10. The predicted ⁹⁹Tc concentrations result in the greatest cancer risks; therefore, ⁹⁹Tc is the most important analyte for contaminant migration at SWMU 3, while arsenic also provides an elevated cancer risk.

Table E.3.10. Hazard and Cancer Risk Predicted from Maximum Groundwater Concentrations Derived in Modeling of SWMU 3 Using SESOIL and AT123D^a

Analyte	SWMU		Plant Boundary		Property Boundary		Little Bayou seeps	
	Hazard Quotient	Cancer Risk	Hazard Quotient	Cancer Risk	Hazard Quotient	Cancer Risk	Hazard Quotient	Cancer Risk
Arsenic	10.5	8.72E-04	0.4	3.2E-05	b	b	b	b
Manganese	3.65	b	<0.1	b	b	b	b	b
⁹⁹ Tc	b	3.05E-04	b	9.9E-05	b	7.5E-05	b	4.4E-05
²³⁸ U	b	<1.0E-06	b	b	b	b	b	b
Uranium	7.82	b	<0.1	b	b	b	b	b

^a Contaminants with a HQ greater than 0.1 or a cancer risk greater than 1.00E-06 are considered analytes.

^b Value not calculated since the groundwater concentrations was reported as zero at this POE by AT123D, or the contaminant did not have a reported cancer slope factor or chemical toxicity RfD.

Figures 5.5 and 5.6 in Section 5 of the main text, show the predicted concentrations over time at each POE for analytes with a HQ greater than 0.1 and/or a risk greater than 1.0E-06 (Table E.3.10 and Table 5.4). As shown in these figures, arsenic is predicted to continue rising in concentration at 1,000 years at the plant boundary, but has not reached the property boundary or Little Bayou seeps in the 1,000 year period. ⁹⁹Tc is predicted to exceed the MCL within 200 years and then decline in concentration at the plant and property POEs.

E.3.1.3 SWMU 4

The C-747 Contaminated Burial Yard and the C-748-B Burial Area (SWMU 4) is located in the western section of the plant area. PGDP used the C-747 Burial Yard from 1951 to 1958 for the disposal of radiologically contaminated and uncontaminated debris originating from the C-410 uranium hexafluoride feed plant. The area consists of two pits covering an area of approximately 8,300 ft² (50 ft by 15 ft and 50 ft by 150 ft) (Union Carbide 1978) of the total 92,000 ft². The C-748-B Burial Area is listed in the 1973 Union Carbide document on waste disposal as a Proposed Chemical Landfill Site and is located on the west side of C-747. SWMU 4 also may have received sludges designated for disposal at the C-404 Burial Grounds. These sludges potentially included uranium-contaminated solid waste and ⁹⁹Tc contaminated

magnesium fluoride. Potential contaminants associated with this SWMU include uranium, ⁹⁹Tc, metals, and TCE (DOE 1998b).

E.3.1.3.1 Conceptual model for source areas at SWMU 4

SWMU 4 occupies an area of approximately 286,700 ft² (6.6 acres). The thickness of the UCRS is estimated to be 63 ft (depth to the top of the RGA). The conceptual model for SWMU 4 is that potentially contaminated trash and scrap was buried in waste pits at SWMU 4. Subsequently, contaminants in the disposed material directly impacted soils below or adjacent to the areas where material was buried and, through vertical infiltration in soil, contaminated the groundwater underlying these sources. The infiltrating groundwater migrates vertically through the UCRS and laterally in the RGA, which could transport the contaminants to the POEs. Previous work on the Southwest Plume discusses a secondary TCE DNAPL source in the RGA below SWMU 4. This source was not modeled, however, further discussion is provided in Section E.3.3.7.

E.3.1.3.2 Contaminant transport modeling for SWMU 4 using SESOIL and AT123D

For this modeling effort, the soil zones were arranged in four layers. The first soil layer represented the SADA surface soil data from 0 to 1 ft deep. The second soil layer was 9 ft thick representing SADA layer 2 from 1 to 9 ft in depth, the third layer was subdivided into 4 sublayers of equal thickness (10 ft each) representing SADA layers 3 through 6. SESOIL soil layer 4 was subdivided into 4 sublayers of equal thickness (3.25 ft each) to represent the total thickness of SADA layer 7 from 50 ft to 63 ft in depth, the four sublayer division of this layer allowed for better numerical solution of the final flux to the RGA. Table E.3.11 presents the analytes remaining after the screening process and the source terms for each analyte. Table E.3.12 lists the chemical-specific parameters applicable to the SESOIL model of SWMU 4. The distances to the POEs used in the AT123D model for SWMU 4 are 984 ft to the plant boundary, 3,000 ft to the property boundary, and 22,967 ft to the Ohio River. SWMU 4 particle tracks do not travel to the Little Bayou seeps.

Table E.3.11. Summary of Source Term Characteristics Developed by SADA for SWMU 4

SADA Layer	Depth (ft)	Average (mg/kg) ^a	Area (ft ²)	Volume (ft ³)	Mass (gm) ^a	Concentration Factor	Adjusted Average (mg/kg) ^a
Arsenic							
L1	0-1	8.87	1.14E+05	1.14E+05	4.16E+04	2.12	18.79
L2	01-10	8.48	5.36E+04	5.36E+05	1.88E+05	1.00	8.48
L3	10-20	6.76	2.88E+04	3.17E+05	8.85E+04	0.54	3.63
L4	20-30	6.70	3.24E+04	3.56E+05	9.87E+04	0.60	4.05
L5	30-40	6.43	2.96E+04	3.26E+05	8.65E+04	0.55	3.55
L6	40-50	6.37	2.48E+04	2.73E+05	7.18E+04	0.46	2.95
L7	50-63	5.81	1.76E+04	1.06E+05	2.53E+04	0.33	1.91
Total Mass					6.00E+05		
cis-1,2-DCE							
L1	0-1	0.00	0.00E+00	0.00E+00	0.00E+00		
L2	01-10	3.84	1.32E+04	1.32E+05	2.10E+04	0.31	1.20
L3	10-20	1.64	3.48E+04	3.83E+05	2.60E+04	0.82	1.35
L4	20-30	1.54	4.24E+04	4.66E+05	2.97E+04	1.00	1.54
L5	30-40	1.11	4.92E+04	5.41E+05	2.49E+04	1.16	1.29
L6	40-50	0.88	4.76E+04	5.24E+05	1.90E+04	1.12	0.98
L7	50-63	0.98	4.96E+04	2.98E+05	1.21E+04	1.17	1.15
Total Mass					1.33E+05		

Table E.3.11. Summary of Source Term Characteristics Developed by SADA for SWMU 4 (Continued)

SADA Layer	Depth (ft)	Average (mg/kg) ^a	Area (ft ²)	Volume (ft ³)	Mass (gm) ^a	Concentration Factor	Adjusted Average (mg/kg) ^a
Manganese							
L1	0-1	475.31	3.47E+05	3.47E+05	6.82E+06	1.50	711.33
L2	01-10	327.05	2.32E+05	2.32E+06	3.14E+07	1.00	327.05
L3	10-20	320.36	2.32E+05	2.55E+06	3.38E+07	1.00	320.36
L4	20-30	193.55	2.32E+05	2.55E+06	2.04E+07	1.00	193.55
L5	30-40	178.70	2.32E+05	2.55E+06	1.89E+07	1.00	178.70
L6	40-50	234.60	2.32E+05	2.55E+06	2.48E+07	1.00	234.60
L7	50-63	133.92	2.32E+05	1.39E+06	7.71E+06	1.00	133.92
Total Mass					1.44E+08		
Nickel							
L1	0-1	21.47	2.72E+05	2.72E+05	2.41E+05	1.36	29.10
L2	01-10	16.79	2.00E+05	2.00E+06	1.39E+06	1.00	16.79
L3	10-20	12.83	1.93E+05	2.12E+06	1.12E+06	0.96	12.34
L4	20-30	14.05	1.22E+05	1.35E+06	7.82E+05	0.61	8.58
L5	30-40	15.75	1.06E+05	1.17E+06	7.59E+05	0.53	8.33
L6	40-50	13.23	1.21E+05	1.33E+06	7.29E+05	0.60	8.00
L7	50-63	10.99	1.14E+05	6.84E+05	3.11E+05	0.57	6.25
Total Mass					5.34E+06		
⁹⁹Tc							
L1	0-1	39.00	1.36E+04	1.36E+04	2.19E+10	0.17	6.53
L2	01-10	49.65	8.12E+04	8.12E+05	1.67E+12	1.00	49.65
L3	10-20	0.00	0.00E+00	0.00E+00	0.00E+00	0.00	0.00
L4	20-30	0.00	0.00E+00	0.00E+00	0.00E+00	0.00	0.00
L5	30-40	0.00	0.00E+00	0.00E+00	0.00E+00	0.00	0.00
L6	40-50	0.00	0.00E+00	0.00E+00	0.00E+00	0.00	0.00
L7	50-63	0.00	0.00E+00	0.00E+00	0.00E+00	0.00	0.00
Total Mass					1.69E+12		
TCE							
L1	0-1	0.00	0.00E+00	0.00E+00	0.00E+00		
L2	01-10	2.39	8.24E+04	8.24E+05	8.16E+04	0.86	2.06
L3	10-20	2.85	6.60E+04	7.26E+05	8.55E+04	0.69	1.96
L4	20-30	3.02	7.24E+04	7.96E+05	9.93E+04	0.75	2.28
L5	30-40	2.56	9.60E+04	1.06E+06	1.12E+05	1.00	2.56
L6	40-50	2.45	9.84E+04	1.08E+06	1.10E+05	1.03	2.51
L7	50-63	3.15	9.96E+04	5.98E+05	7.77E+04	1.04	3.26
Total Mass					5.65E+05		
²³⁹Pu							
L1	0-1	0.64	1.59E+05	1.59E+05	4.24E+09	1.02	0.66
L2	01-10	0.65	1.58E+05	1.58E+06	4.24E+10	1.01	0.66
L3	10-20	0.64	1.56E+05	1.72E+06	4.51E+10	1.00	0.64
L4	20-30	0.65	1.53E+05	1.68E+06	4.49E+10	0.98	0.63
L5	30-40	0.62	1.51E+05	1.66E+06	4.27E+10	0.97	0.60
L6	40-50	0.59	1.47E+05	1.61E+06	3.95E+10	0.94	0.56
L7	50-63	0.58	1.42E+05	1.14E+06	2.75E+10	0.91	0.53
Total Mass					2.46E+11		

Table E.3.11. Summary of Source Term Characteristics Developed by SADA for SWMU 4 (Continued)

SADA Layer	Depth (ft)	Average (mg/kg)^a	Area (ft²)	Volume (ft³)	Mass (gm)^a	Concentration Factor	Adjusted Average (mg/kg)^a
²³⁴U							
L1	0-1	15.57	3.47E+05	3.47E+05	2.24E+11	2.03	31.66
L2	01-10	27.69	1.75E+05	1.75E+06	2.01E+12	1.03	28.40
L3	10-20	27.59	1.71E+05	1.88E+06	2.14E+12	1.00	27.59
L4	20-30	28.03	1.67E+05	1.83E+06	2.13E+12	0.98	27.38
L5	30-40	28.26	1.62E+05	1.78E+06	2.08E+12	0.95	26.74
L6	40-50	28.76	1.58E+05	1.73E+06	2.06E+12	0.92	26.53
L7	50-63	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00	0.00
Total Mass					1.06E+13		
²³⁸U							
L1	0-1	30.55	3.47E+05	3.47E+05	4.39E+11	2.03	62.10
L2	01-10	52.13	1.75E+05	1.75E+06	3.78E+12	1.03	53.48
L3	10-20	51.72	1.71E+05	1.88E+06	4.02E+12	1.00	51.72
L4	20-30	52.47	1.67E+05	1.83E+06	3.98E+12	0.98	51.25
L5	30-40	52.77	1.62E+05	1.78E+06	3.88E+12	0.95	49.93
L6	40-50	53.71	1.58E+05	1.73E+06	3.85E+12	0.92	49.56
L7	50-63	0.00	0.00E+00	0.00E+00	0.00E+00	0.00	0.00
Total Mass					1.99E+13		
Uranium							
L1	0-1	118.94	3.47E+05	3.47E+05	1.71E+06	2.03	241.79
L2	01-10	884.57	1.75E+05	1.75E+06	6.41E+07	1.03	907.36
L3	10-20	827.57	1.71E+05	1.88E+06	6.43E+07	1.00	827.57
L4	20-30	807.78	1.67E+05	1.83E+06	6.13E+07	0.98	788.86
L5	30-40	789.77	1.62E+05	1.78E+06	5.80E+07	0.95	747.23
L6	40-50	768.17	1.58E+05	1.73E+06	5.51E+07	0.92	708.80
L7	50-63	0.00	0.00E+00	0.00E+00	0.00E+00	0.00	0.00
Total Mass					3.04E+08		
Vanadium							
L1	0-1	25.21	3.47E+05	3.47E+05	3.62E+05	1.50	37.73
L2	01-10	21.69	2.32E+05	2.32E+06	2.08E+06	1.00	21.69
L3	10-20	22.83	2.32E+05	2.55E+06	2.41E+06	1.00	22.83
L4	20-30	26.28	2.32E+05	2.55E+06	2.77E+06	1.00	26.28
L5	30-40	25.06	2.32E+05	2.55E+06	2.64E+06	1.00	25.06
L6	40-50	22.81	2.32E+05	2.55E+06	2.41E+06	1.00	22.81
L7	50-63	20.40	2.32E+05	1.39E+06	1.17E+06	1.00	20.40
Total Mass					1.38E+07		
Vinyl Chloride							
L1	0-1	0.00	0.00E+00	0.00E+00	0.00E+00	0.00	0.00
L2	01-10	0.16	6.40E+03	6.40E+04	4.15E+02	0.37	0.06
L3	10-20	0.08	1.64E+04	1.80E+05	6.05E+02	0.95	0.08
L4	20-30	0.09	9.20E+03	1.01E+05	3.95E+02	0.53	0.05
L5	30-40	0.19	1.72E+04	1.89E+05	1.49E+03	1.00	0.19
L6	40-50	0.18	1.76E+04	1.94E+05	1.43E+03	1.02	0.18
L7	50-63	0.20	2.08E+04	1.25E+05	1.04E+03	1.21	0.24
Total Mass					5.37E+03		

Table E.3.11. Summary of Source Term Characteristics Developed by SADA for SWMU 4 (Continued)

SADA Layer	Depth (ft)	Average (mg/kg) ^a	Area (ft ²)	Volume (ft ³)	Mass (gm) ^a	Concentration Factor	Adjusted Average (mg/kg) ^a
Zinc							
L1	0-1	40.73	3.26E+05	3.26E+05	5.50E+05	1.00	40.73
L2	01-10	8.48	5.36E+04	5.36E+05	1.88E+05	0.16	1.39
L3	10-20	6.76	2.88E+04	3.17E+05	8.85E+04	0.09	0.60
L4	20-30	6.70	3.24E+04	3.56E+05	9.87E+04	0.10	0.67
L5	30-40	6.43	2.96E+04	3.26E+05	8.65E+04	0.09	0.58
L6	40-50	6.37	2.48E+04	2.73E+05	7.18E+04	0.08	0.48
L7	50-63	5.81	1.76E+04	1.06E+05	2.53E+04	0.05	0.31
Total Mass					1.11E+06		

^a Radionuclides are in units of pCi/g for concentrations and pCi for mass.

Table E.3.12. Chemical-Specific Parameters of the Analytes Used in SESOIL Modeling of SWMU 4

Analyte	Mol. Wt. (MW) (g/mol)	Solubility in water (mg/L)	Diffusion in air (cm ² /s)	Diffusion in water (m ² /hr)	Henry's Constant (atm.m3/mol)	Koc (L/kg)	Kd ^a (L/kg)	Half Life (years)
Arsenic	74.92	1.00E+07	NA	3.60E-07	NA	NA	29	infinite
cis-1,2-DCE	96.94	3.50E+03	0.07	4.07E-06	4.08E-03	35.5	0.028	infinite
Manganese	54.94	1.00E+07	NA	1.29E-07	NA	NA	65	infinite
Nickel	58.69	1.00E+07	NA	3.60E-07	NA	NA	300	infinite
²³⁹ Pu	239	1.00E+07	NA	3.60E-07	NA	NA	550	2.41E+04
⁹⁹ Tc	99	1.00E+07	NA	3.60E-07	NA	NA	0.2	2.13E+05
TCE	131	1,100	0.08	3.28E-06	0.0103	94	0.0752	26.6
²³⁴ U	234	1.00E+07	NA	3.60E-07	NA	NA	66.8	2.44E+05
²³⁸ U	238	1.00E+07	NA	3.60E-07	NA	NA	66.8	4.47E+09
Uranium	238	1.00E+07	NA	3.60E-07	NA	NA	66.8	4.47E+09
Vanadium	50.94	1.00E+07	NA	3.60E-07	NA	NA	1000	infinite
Zinc	67.41	1.00E+07	NA	3.60E-07	NA	NA	62	infinite
Vinyl Chloride	63	2,760	0.11	4.43E-07	0.0270	18.8	0.0152	26.6

^a The soil/water distribution coefficient (Kd) of an organic compound depends on the soil's organic content (f_{oc}) and compound's organic partition coefficient (Koc). Kd values presented for organic compounds are for UCRS soils (with f_{oc} value of 0.08%) only. Kds used in AT123D are different due to the f_{oc} of 0.02% in the RGA.

E.3.1.3.3 Groundwater modeling results for SWMU 4

Table E.3.13 summarizes the predicted maximum groundwater concentrations at the POEs for the analytes modeled at SWMU 4. These contaminants were predicted by SESOIL to reach the water table within the 1,000 year period in concentrations that were greater than the groundwater background or greater than the groundwater child no action levels. Several contaminants that originally passed the screening for groundwater did not reach the water table in 1,000 years (i.e., antimony, nickel, ²³⁹Pu, ²³⁴U, ²³⁸U, uranium, and vanadium) or exhibited groundwater concentrations that were less than the groundwater background or the groundwater child no action levels (i.e., zinc) (see Section 5.4 of the main text).

Table E.3.13. Concentrations of the Analytes in Groundwater Predicted in SESOIL and AT123D Modeling of SWMU 4

Analyte	Predicted Maximum Groundwater Concentration ^{a,b}					MCL (mg/L or pCi/L)
	SWMU (mg/L)	Plant Boundary (mg/L)	Property Boundary (mg/L)	Ohio River (mg/L)		
Arsenic	<i>1.77E-02</i>	2.70E-03	4.89E-06	0	0.01	
<i>cis</i> -1,2-DCE	<i>6.68E-01</i>	<i>1.96E-01</i>	<i>8.94E-02</i>	3.16E-02	0.07	
Manganese	<i>5.76E-01</i>	5.01E-03	0	0	^d	
⁹⁹ Tc	<i>9.008E+03</i>	<i>2.50E+03</i>	<i>1.20E+03</i>	3.79E+02	900 ^c	
TCE	<i>1.18E+00</i>	<i>4.22E-01</i>	<i>2.14E-01</i>	<i>7.67E-02</i>	0.005	
Vinyl Chloride	<i>2.61E-02</i>	<i>5.95E-03</i>	<i>2.53E-03</i>	7.82E-04	0.002	

^a Values in bold, italic font exceed the analyte's MCL

^b Radionuclide concentrations are in pCi/L

^c ⁹⁹Tc MCL based on a critical organ dose at 4 mrem/yr from drinking water consumption

^d MCLs not available for these contaminants

As shown in Table E.3.13, the predicted maximum groundwater concentrations for *cis*-1,2-DCE, ⁹⁹Tc, TCE and vinyl chloride exceed the MCLs at the plant and property boundary. TCE also is predicted to exceed the MCL at the Ohio River. The HQs and cancer risks calculated in accordance with the Risk Methods Document (DOE 2001) are presented in Table E.3.14. The predicted TCE, vinyl chloride, and ⁹⁹Tc concentrations result in the greatest cancer risks; with TCE being the most important analyte for contaminant migration at SWMU 4. TCE also exhibits the highest HQ for SWMU 4.

Table E.3.14. Hazard and Cancer Risk Predicted from Maximum Groundwater Concentrations Derived in Modeling of SWMU 4 Using SESOIL and AT123D^a

Analyte	SWMU		Plant Boundary		Property Boundary		Ohio River	
	Hazard Quotient	Cancer Risk	Hazard Quotient	Cancer Risk	Hazard Quotient	Cancer Risk	Hazard Quotient	Cancer Risk
Arsenic	5.67	4.69E-04	0.9	7.2E-05	<0.1	<1.0E-06	^b	^b
<i>cis</i> -1,2-DCE	35.3	^b	10.4	^b	4.7	^b	0.6	^b
Manganese	1.23	^b	<0.1	^b	^b	^b	^b	^b
⁹⁹ Tc	^b	4.94E-04	^b	1.4E-04	^b	6.6E-05	^b	2.1E-05
TCE	539	3.67E-02	193	2.0E-02	97.7	6.6E-03	32.7	2.4E-03
Vinyl Chloride	1.21	1.65E-02	0.3	1.9E-04	0.1	7.4E-05	<0.1	2.3E-05

^a Contaminants with a HQ greater than 0.1 or a cancer risk greater than 1.00E-06 are considered analytes.

^b Value not calculated since the groundwater concentrations was reported as zero at this POE by AT123D, or the contaminant did not have a reported cancer slope factor or chemical toxicity RfD.

Figures 5.7 through 5.11 in Section 5 of the main text, show the predicted concentrations over time at each POE for analytes with a HQ greater than 0.1 and/or a risk greater than 1.0E-06 for contaminants migrating from SWMU 4. As shown in these figures, manganese is predicted to continue rising in concentration at 1,000 years at the plant boundary, but has not reached the property boundary or Ohio River in the 1,000 year period. *Cis*-1,2-DCE; ⁹⁹Tc; TCE; and vinyl chloride are predicted to exceed the MCL at the plant and property boundaries within 100 years. TCE also is predicted to exceed the MCL at the Ohio River within 100 years.

E.3.1.4 SWMU 5

The C-746-F Burial Yard is located in the northwestern section of the PGDP secured area, adjacent to SWMU 6 to the east. Disposal pits were located on a grid system. Documentation of the size of these grids ranges from 10 ft by 10 ft cells to 20 ft by 20 ft cells excavated to a depth of 6 to 15 ft bgs. SWMU 5 was in operation from 1965 to 1987. The burial pits were used for the burial of components from the

“Work for Others” activities, some radionuclide-contaminated scrap metal, and slag from the nickel and aluminum smelters. Metals and radioisotopes are the primary potential contaminants of interest at this SWMU.

E.3.1.4.1 Conceptual model for source areas at SWMU 5

SWMU 5 occupies an area of approximately 197,400 ft² (4.5 acres). The thickness of the UCRS is estimated to be 60 ft (depth to the top of the RGA). Metals and radionuclides were buried in pits at SWMU 5. The conceptual model for SWMU 5 is that contaminants in disposed material directly impacted soils below or adjacent to the areas where material was buried and, through vertical infiltration in soil, contaminated the groundwater underlying these sources. The infiltrating groundwater migrates vertically through the UCRS and laterally in the RGA, which could transport the contaminants to the POEs.

E.3.1.4.2 Contaminant transport modeling for SWMU 5 using SESOIL and AT123D

For this modeling effort, the soil zones were arranged in four layers. The first soil layer represented the SADA surface soil data from 0 to 1 ft deep. The second soil layer was 9 ft thick representing SADA layer 2 from 1 to 9 ft in depth, the third layer was subdivided into 4 sublayers of equal thickness (10 ft each) representing SADA layers 3 through 6. SESOIL soil layer 4 was subdivided into 4 sublayers of equal thickness (2.5 ft each) to represent the total thickness of SADA layer 7 from 50 ft to 60 ft in depth. Table E.3.15 presents the analytes remaining after the screening process and the source terms for each analyte.

Table E.3.15. Summary of Source Term Characteristics Developed by SADA for SWMU 5

SADA Layer	Depth (ft)	Average (mg/kg) ^a	Area (ft ²)	Volume (ft ³)	Mass (gm) ^a	Concentration Factor	Adjusted Average (mg/kg) ^a
Acenaphthene							
L1	0-1	2.71	1.54E+05	1.54E+05	1.73E+04	1.00E+00	2.71
L2	01-10	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L3	10-20	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L4	20-30	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L5	30-40	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L6	40-50	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L7	50-60	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
Total Mass					1.73E+04		
Anthracene							
L1	0-1	3.51	1.80E+05	1.80E+05	2.61E+04	1.00E+00	3.51
L2	01-10	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L3	10-20	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L4	20-30	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L5	30-40	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L6	40-50	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L7	50-60	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
Total Mass					2.61E+04		
Arsenic							
L1	0-1	8.78	1.04E+05	1.04E+05	3.79E+04	1.76E+00	15.49
L2	01-10	2.51	4.52E+04	4.07E+05	4.22E+04	7.64E-01	1.91
L3	10-20	2.31	4.12E+04	4.12E+05	3.94E+04	6.96E-01	1.61
L4	20-30	3.47	5.92E+04	5.92E+05	8.49E+04	1.00E+00	3.47
L5	30-40	1.65	3.48E+04	3.48E+05	2.38E+04	5.88E-01	0.97
L6	40-50	1.79	3.72E+04	3.72E+05	2.75E+04	6.28E-01	1.12
L7	50-60	1.50	3.48E+04	3.48E+05	2.16E+04	5.88E-01	0.88
Total Mass					2.77E+05		
Benzo(a)pyrene							
L1	0-1	6.14	2.52E+05	2.52E+05	6.40E+04	1.00E+00	6.14
L2	01-10	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L3	10-20	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L4	20-30	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L5	30-40	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L6	40-50	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L7	50-60	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
Total Mass					6.40E+04		
Dibenz(a,h)anthracene							
L1	0-1	0.44	2.96E+04	2.96E+04	5.40E+02	1.00E+00	0.44
L2	01-10	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L3	10-20	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L4	20-30	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L5	30-40	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L6	40-50	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L7	50-60	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
Total Mass					5.40E+02		

Table E.3.15. Summary of Source Term Characteristics Developed by SADA for SWMU 5 (Continued)

SADA Layer	Depth (ft)	Average (mg/kg) ^a	Area (ft ²)	Volume (ft ³)	Mass (gm) ^a	Concentration Factor	Adjusted Average (mg/kg) ^a
Fluoranthene							
L1	0-1	14.25	1.79E+05	1.79E+05	1.06E+05	1.00E+00	14.25
L2	01-10	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L3	10-20	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L4	20-30	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L5	30-40	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L6	40-50	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L7	50-60	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
Total Mass					1.06E+05		
Fluorene							
L1	0-1	3.29	1.32E+05	1.32E+05	1.80E+04	1.00E+00	3.29
L2	01-10	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L3	10-20	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L4	20-30	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L5	30-40	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L6	40-50	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L7	50-60	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
Total Mass					1.80E+04		
Manganese							
L1	0-1	3.82E+02	4.03E+05	4.03E+05	6.36E+06	2.21E+00	845.91
L2	01-10	1.79E+02	1.82E+05	1.64E+06	1.21E+07	1.00E+00	179.07
L3	10-20	1.85E+02	1.82E+05	1.82E+06	1.39E+07	1.00E+00	185.24
L4	20-30	1.56E+02	1.82E+05	1.82E+06	1.17E+07	1.00E+00	155.86
L5	30-40	1.54E+02	1.82E+05	1.82E+06	1.16E+07	1.00E+00	154.05
L6	40-50	2.00E+02	1.82E+05	1.82E+06	1.50E+07	1.00E+00	199.52
L7	50-60	2.58E+02	1.82E+05	1.82E+06	1.94E+07	1.00E+00	258.26
Total Mass					9.02E+07		
Naphthalene							
L1	0-1	3.80	8.00E+03	8.00E+03	1.26E+03	1.00E+00	3.80
L2	01-10	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L3	10-20	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L4	20-30	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L5	30-40	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L6	40-50	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L7	50-60	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
Total Mass					1.26E+03		
Nickel							
L1	0-1	2.79E+01	3.88E+05	3.88E+05	4.47E+05	2.44E+00	68.15
L2	01-10	1.03E+01	1.43E+05	1.29E+06	5.50E+05	9.02E-01	9.32
L3	10-20	1.04E+01	1.42E+05	1.42E+06	6.14E+05	8.97E-01	9.36
L4	20-30	1.12E+01	1.59E+05	1.59E+06	7.33E+05	1.00E+00	11.16
L5	30-40	9.20E+00	1.39E+05	1.39E+06	5.28E+05	8.74E-01	8.04
L6	40-50	1.01E+01	1.50E+05	1.50E+06	6.29E+05	9.47E-01	9.58
L7	50-60	9.87E+00	1.54E+05	1.54E+06	6.29E+05	9.70E-01	9.58
Total Mass					4.13E+06		

Table E.3.15. Summary of Source Term Characteristics Developed by SADA for SWMU 5 (Continued)

SADA Layer	Depth (ft)	Average (mg/kg) ^a	Area (ft ²)	Volume (ft ³)	Mass (gm) ^a	Concentration Factor	Adjusted Average (mg/kg) ^a
PCB-1260							
L1	0-1	0.15	1.34E+05	1.34E+05	8.23E+02	1.00E+00	0.15
L2	01-10	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L3	10-20	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L4	20-30	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L5	30-40	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L6	40-50	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L7	50-60	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
Total Mass					8.23E+02		
Pyrene							
L1	0-1	8.29	2.82E+05	2.82E+05	9.68E+04	1.00E+00	8.29
L2	01-10	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L3	10-20	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L4	20-30	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L5	30-40	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L6	40-50	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L7	50-60	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
Total Mass					9.68E+04		
Selenium							
L1	0-1	1.17	2.56E+04	2.56E+04	1.23E+03	1.00E+00	1.17
L2	01-10	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L3	10-20	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L4	20-30	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L5	30-40	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L6	40-50	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L7	50-60	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
Total Mass					1.23E+03		
TCE							
L1	0-1	0.0000	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L2	01-10	0.0028	5.92E+04	5.33E+05	6.26E+01	1.00E+00	0.0028
L3	10-20	0.0029	6.04E+04	6.04E+05	7.34E+01	1.02E+00	0.0030
L4	20-30	0.0030	5.96E+04	5.96E+05	7.35E+01	1.01E+00	0.0030
L5	30-40	0.0030	5.92E+04	5.92E+05	7.43E+01	1.00E+00	0.0030
L6	40-50	0.0031	5.64E+04	5.64E+05	7.33E+01	9.53E-01	0.0030
L7	50-60	0.0032	5.44E+04	5.44E+05	7.28E+01	9.19E-01	0.0030
Total Mass					4.30E+02		
Vanadium							
L1	0-1	20.65	4.03E+05	4.03E+05	3.44E+05	2.21	45.71
L2	01-10	23.18	1.82E+05	1.64E+06	1.57E+06	1.00	23.18
L3	10-20	22.14	1.82E+05	1.82E+06	1.67E+06	1.00	22.14
L4	20-30	24.70	1.82E+05	1.82E+06	1.86E+06	1.00	24.70
L5	30-40	18.45	1.82E+05	1.82E+06	1.39E+06	1.00	18.45
L6	40-50	19.13	1.82E+05	1.82E+06	1.44E+06	1.00	19.13
L7	50-60	18.00	1.82E+05	1.82E+06	1.35E+06	1.00	18.00
Total Mass					9.62E+06		

Table E.3.15. Summary of Source Term Characteristics Developed by SADA for SWMU 5 (Continued)

SADA Layer	Depth (ft)	Average (mg/kg) ^a	Area (ft ²)	Volume (ft ³)	Mass (gm) ^a	Concentration Factor	Adjusted Average (mg/kg) ^a
Zinc							
L1	0-1	70.69	3.21E+05	3.21E+05	9.38E+05	2.37	167.74
L2	01-10	32.20	1.30E+05	1.17E+06	1.56E+06	0.96	30.96
L3	10-20	32.87	1.33E+05	1.33E+06	1.81E+06	0.99	32.38
L4	20-30	34.66	1.27E+05	1.27E+06	1.82E+06	0.94	32.51
L5	30-40	37.01	1.15E+05	1.15E+06	1.76E+06	0.85	31.53
L6	40-50	42.44	1.28E+05	1.28E+06	2.25E+06	0.95	40.30
L7	50-60	40.94	1.35E+05	1.35E+06	2.29E+06	1.00	40.94
Total Mass					1.24E+07		
¹³⁷Cs							
L1	0-1	0.05	1.04E+05	1.04E+05	2.04E+08	1.04	0.05
L2	01-10	0.05	1.04E+05	9.32E+05	1.83E+09	1.03	0.05
L3	10-20	0.05	1.00E+05	1.00E+06	1.98E+09	1.00	0.05
L4	20-30	0.05	9.64E+04	9.64E+05	1.89E+09	0.96	0.05
L5	30-40	0.05	9.24E+04	9.24E+05	1.80E+09	0.92	0.04
L6	40-50	0.05	8.64E+04	8.64E+05	1.69E+09	0.86	0.04
L7	50-60	0.04	3.16E+04	2.84E+05	4.17E+08	0.31	0.01
Total Mass					9.81E+09		
⁹⁹Tc							
L1	0-1	7.16	1.42E+05	1.42E+05	4.22E+10	1.00	7.16
L2	01-10	0.00	0.00E+00	0.00E+00	0.00E+00	0.00	0.00
L3	10-20	0.00	0.00E+00	0.00E+00	0.00E+00	0.00	0.00
L4	20-30	0.00	0.00E+00	0.00E+00	0.00E+00	0.00	0.00
L5	30-40	0.00	0.00E+00	0.00E+00	0.00E+00	0.00	0.00
L6	40-50	0.00	0.00E+00	0.00E+00	0.00E+00	0.00	0.00
L7	50-60	3.89	9.60E+03	9.60E+04	1.54E+10	0.07	0.26
Total Mass					5.76E+10		
Uranium							
L1	0-1	233.55	7.48E+04	7.48E+04	7.22E+11	0.99	231.08
L2	01-10	233.54	7.48E+04	6.73E+05	6.50E+12	0.99	231.07
L3	10-20	225.20	7.56E+04	7.56E+05	7.04E+12	1.00	225.20
L4	20-30	230.77	7.28E+04	7.28E+05	6.95E+12	0.96	222.22
L5	30-40	221.96	7.52E+04	7.52E+05	6.90E+12	0.99	220.78
L6	40-50	228.41	7.16E+04	7.16E+05	6.76E+12	0.95	216.32
L7	50-60	236.98	4.76E+04	4.28E+05	4.20E+12	0.63	149.21
Total Mass					3.91E+13		

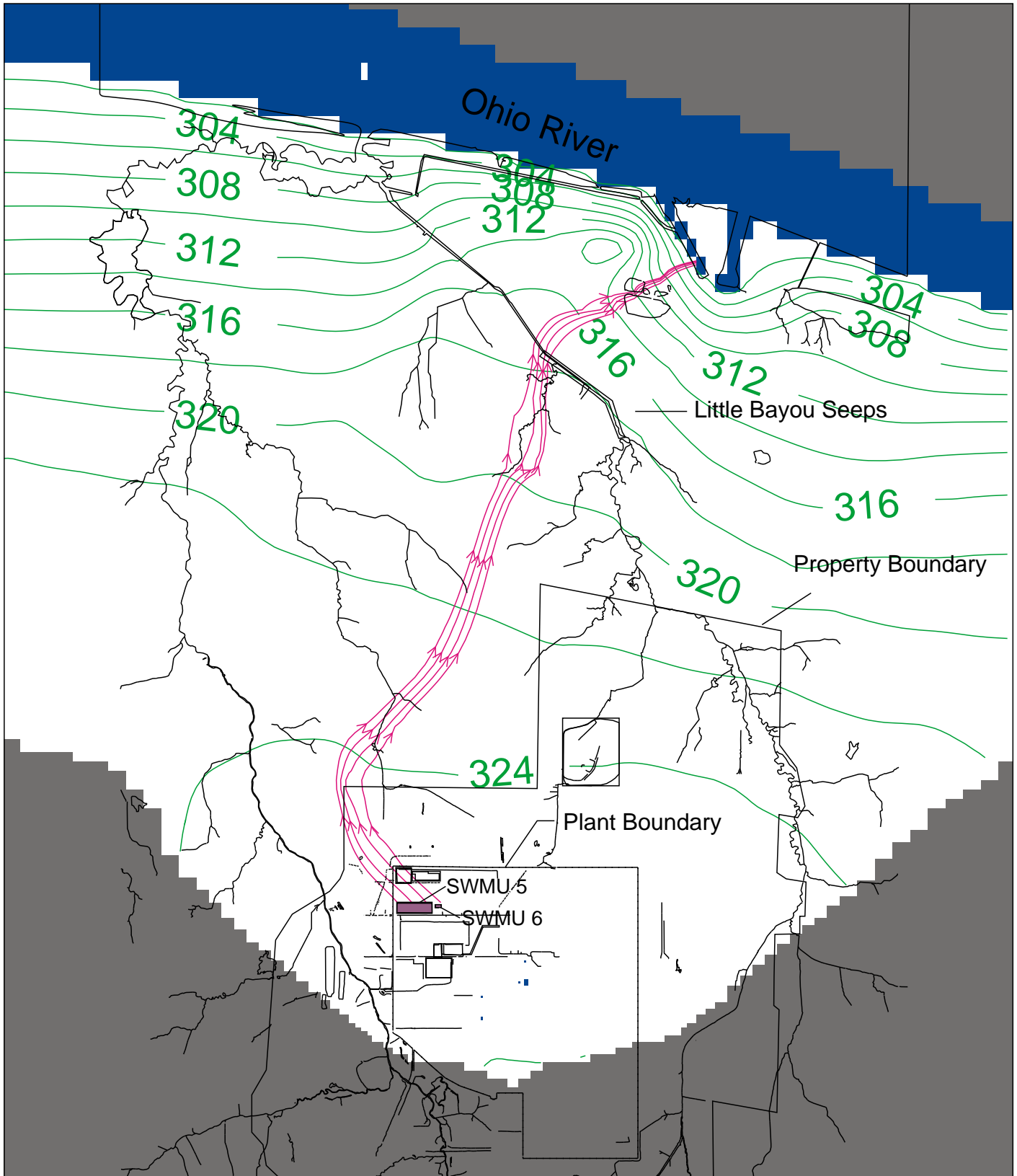
^a Radionuclides are in units of pCi/g for concentrations and pCi for mass.

Table E.3.16 lists the chemical-specific parameters applicable to the SESOIL model of SWMU 5. Figure E.3.3 shows the particle tracks that were modeled for SWMUs 5 and 6. The distances to the POEs used in the AT123D model for SWMU 5 are 778 ft to the plant boundary, 2,293 ft to the property boundary, and 19,844 ft to the Ohio River. SWMU 5 particle tracks do not travel to the Little Bayou seeps.

Table E.3.16. Chemical-Specific Parameters of the Analytes Used in SESOIL Modeling of SWMU 5

Analyte	Mol. Wt. (MW) (g/mol)	Solubility in water (mg/L)	Diffusion in air (cm ² /s)	Diffusion in water (m ² /hr)	Henry's Constant (atm.m3/mol)	Koc (L/kg)	Kd ^a (L/kg)	Half Life (years)
Acenaphthene	154.0	4.20	0.04	2.77E-6	1.60E-04	4.90E+03	3.9	infinite
Anthracene	178.24	0.043	0.032	2.79E-06	5.55E-05	2.04E+04	16.3	infinite
Arsenic	74.92	1.00E+07	NA	3.60E-07	NA	NA	29	infinite
Benzo(a)pyrene	252.32	1.62E-03	4.3E-02	3.24E-06	1.13E-06	9.69E+05	772	infinite
¹³⁷ Cs	137	1.00E+07	NA	3.60E-07	NA	NA	280	30.17
Dibenzo(a,h)anthracene	278.33	0.0025	0.020	1.86E-06	1.47E-08	1.78E+06	1,424	infinite
Fluoranthene	202.26	0.206	0.030	2.29E-06	1.61E-05	4.91E+04	39.3	infinite
Fluorene	166.0	1.90	0.061	2.84E-06	7.7E-05	7.9E+03	6.3	infinite
Manganese	54.94	1.00E+07	NA	1.29E-07	NA	NA	65	infinite
Naphthalene	128.16	31.0	0.059	2.70E-06	4.83E-04	1.19E+03	0.95	infinite
Nickel	58.69	1.00E+07	NA	3.60E-07	NA	NA	300	infinite
PCB-1260	375.70	0.027	0.014	4.32E-06	7.40E-05	2.07E+05	165.6	infinite
Pyrene	202.3	0.135	0.0272	2.61E-06	1.1E-05	6.8E+04	54.4	infinite
Selenium	80.98	1.00E+07	NA	3.60E-07	NA	NA	5	infinite
⁹⁹ Tc	99	1.00E+07	NA	3.60E-07	NA	NA	0.2	2.13E+05
TCE	131	1,100	0.08	3.28E-06	0.0103	94	0.0752	26.6
Uranium	238	1.00E+07	NA	3.60E-07	NA	NA	66.8	4.47E+09
Vanadium	50.94	1.00E+07	NA	3.60E-07	NA	NA	1,000	infinite
Zinc	67.41	1.00E+07	NA	3.60E-07	NA	NA	62	infinite

^a The soil/water distribution coefficient (Kd) of an organic compound depends on the soil's organic content (f_{oc}) and compound's organic partition coefficient (Koc). Kd values presented for organic compounds are for UCRS soils (with f_{oc} value of 0.08%) only. Kds used in AT123D are different due to the f_{oc} of 0.02% in the RGA.



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U.S. DEPARTMENT OF ENERGY
DOE PORTSMOUTH/PADUCAH PROJECT OFFICE
PADUCAH GASEOUS DIFFUSION PLANT

Paducah Gaseous Diffusion Plant

- River
- Particle Tracks
- Hydraulic Head (ft)

TRUE NORTH
PLANT NORTH
20°

0 100 500 1000
METERS

Figure E.3.3. Particle Tracks for SWMUs 5 and 6
E-50

E.3.1.4.3 Groundwater modeling results for SWMU 5

Table E.3.17 summarizes the predicted maximum groundwater concentrations at the POEs for the analytes modeled at SWMU 5. These contaminants were predicted by SESOIL to reach the water table within the 1,000 year period in concentrations that were greater than the groundwater background or greater than the groundwater child no action levels. Several contaminants that originally passed the screening for groundwater did not reach the water table in 1,000 years [i.e., benzo(a)pyrene, dibenzo(a,h)anthracene, fluoranthene, PCB-1260, pyrene, Cs-137, and vanadium] or exhibited groundwater concentrations that were less than the groundwater background or the groundwater child no action levels (i.e., acenaphthene, anthracene, fluorine, nickel, selenium, TCE, and zinc) (see Section 5.4 of the main text).

Table E.3.17. Concentrations of the Analytes in Groundwater Predicted in SESOIL and AT123D Modeling of SWMU 5

Analyte	Predicted Maximum Groundwater Concentration ^{a,b}				
	SWMU (mg/L)	Plant Boundary (mg/L)	Property Boundary (mg/L)	Ohio River (mg/L)	MCL (mg/L)
Arsenic	9.25E-03	1.78E-03	1.27E-04	0	0.01
Manganese	<i>1.01E+00</i>	<i>8.69E-02</i>	2.30E-11	0	^d
Naphthalene	5.55E-03	9.82E-04	3.72E-04	1.08E-04	NA
⁹⁹ Tc	1.27E+02	4.99E+01	2.64E+01	8.72E+00	900 ^c
Uranium	4.60E-01	3.32E-02	4.65E-11	0	0.03

^a Values in bold, italic font exceed the analyte's MCL.

^b Radionuclide concentrations are in pCi/L.

^c ⁹⁹Tc MCL based on a critical organ dose at 4 mrem/yr from drinking water consumption.

^d MCLs not available for these contaminants.

All analytes are less than their MCL (note acenaphthene, manganese, and naphthalene do not have MCLs). The HQs and cancer risks calculated in accordance with the Risk Methods Document (DOE 2001) are presented in Table E.3.18. The predicted uranium concentration at the plant boundary results in the greatest HQ, with arsenic, naphthalene and manganese also exhibiting a HQ above 0.1. Arsenic presents the highest cancer risk followed by ⁹⁹Tc.

Table E.3.18. Hazard and Cancer Risk Predicted from Maximum Groundwater Concentrations Derived in Modeling of SWMU 5 Using SESOIL and AT123D^a

Analyte	SWMU		Plant Boundary		Property Boundary		Ohio River	
	Hazard Quotient	Cancer Risk	Hazard Quotient	Cancer Risk	Hazard Quotient	Cancer Risk	Hazard Quotient	Cancer Risk
Arsenic	2.96	2.45E-04	0.6	4.7E-05	<0.1	3.4E-06	^b	^b
Manganese	2.15	^b	0.2	^b	<0.1	^b	^b	^b
Naphthalene	2.80	^b	0.5	^b	0.2	^b	<0.1	^b
⁹⁹ Tc	^b	6.97E-06	^b	2.7E-06	^b	1.4E-06	^b	<1.0E-06
Uranium	73.6	^b	5.31	^b	<0.1	^b	^b	^b

^a Contaminants with a HQ greater than 1 or a cancer risk greater than 1.00E-06 are considered analytes.

^b Value not calculated since the groundwater concentrations was reported as zero at this POE by AT123D, or the contaminant did not have a reported cancer slope factor or chemical toxicity RfD.

Figures 5.12 through 5.15 in Section 5 of the main text, show the predicted concentrations over time at each POE for analytes with a HQ greater than 0.1 and/or a risk greater than 1.0E-06 (see Table E.3.18) for contaminants migrating from SWMU 5. As shown in these figures, manganese is predicted to continue rising in concentration at 1,000 years at the plant boundary, but has not reached the property boundary or Ohio River in the 1,000 year period. Arsenic also is increasing in concentration at the plant boundary at 1,000 years; however, the concentrations are less than the MCL. ⁹⁹Tc is not predicted to exceed the MCL at the POEs.

E.3.1.5 SWMU 6

The C-747-B Burial Ground is located in the northwest section of the plant area east of SWMU5. PGDP buried waste at SWMU 6 in five separate burial cells between 1960 and 1976. The contents of each cell are as follows (DOE 2000):

- Area H—Magnesium Scrap Burial Area. The scrap buried at this location is magnesium, in various shapes generated in the machine shop.
- Area I—Exhaust Fan Burial Area. Eight exhaust hood blowers removed from C-710 were discarded in this pit. These blowers, which were about 15 inches in diameter and weighed about 100 lb each, were discarded in 1966 because of contamination with perchloric acid. Each blower was spaced about 4 ft apart in the hole. Area I-2 is a 6 ft by 6 ft pit just north of the main Area I pit that was used in 1976 for the disposal of additional exhaust fans.
- Area J—Contaminated Aluminum Burial Area. The contaminated scrap buried in this hole involved aluminum scrap in the form of nuts, bolts, plates, trimmings, etc., that were generated in the converter and compressor shop. This scrap was buried about 1960 or 1962.
- Area K—Magnesium Scrap Burial Area. The scrap buried at this location is magnesium in various shapes generated in the machine shop.
- Area L—Modine Trap Burial Area. A single contaminated modine trap was buried in this area. The cold trap was about 4 ft in diameter, approximately 15 ft long, and weighed about 5,000 lb.

Approximately 50% of the surface area of SWMU 6 formerly has been used to store radioactively-contaminated equipment and materials. These items include industrial forklifts and transport carts, flatbed trailers, generators, concrete pipes, and other miscellaneous items (DOE 2000).

E.3.1.5.1 Conceptual model for source areas at SWMU 6

SWMU 6 occupies an area of approximately 13,500 ft² (0.31 acres). The thickness of the UCRS is estimated to be 63 ft (depth to the top of the RGA). The conceptual model for SWMU 6 is that the waste cells contain potentially contaminated materials. Subsequently, contaminants in the disposed material directly impacted soils below or adjacent to the areas where material was buried and, through vertical infiltration in soil, contaminated the groundwater underlying these sources. The infiltrating groundwater migrates vertically through the UCRS and laterally in the RGA, which could transport the contaminants to the POEs.

E.3.1.5.2 Contaminant transport modeling for SWMU 6 using SESOIL and AT123D

For this modeling effort, the soil zones were arranged in four layers. The first soil layer represented the SADA surface soil data from 0 to 1 ft deep. The second soil layer was 9 ft thick representing SADA layer 2 from 1 to 9 ft in depth, the third layer was subdivided into 4 sublayers of equal thickness (10 ft each) representing SADA layers 3 through 6. SESOIL soil layer 4 was subdivided into 4 sublayers of equal thickness (3.25 ft each) to represent the total thickness of SADA layer 7 from 50 ft to 63 ft in depth. Table E.3.19 presents the analytes remaining after the screening process and the source terms for each analyte. Table E.3.20 lists the chemical-specific parameters applicable to the SESOIL model of SWMU 6. Figure E.3.3 presents the head distribution and flowpaths of several particles released from the SWMU. The distances to the POEs used in the AT123D model for SWMU 6 are 1,328 ft to the plant boundary, 3,561 ft to the property boundary, 15,138 ft to the Little Bayou seeps, and 19,424 ft to the Ohio River.

Table E.3.19. Summary of Source Term Characteristics Developed by SADA for SWMU 6

SADA Layer	Depth (ft)	Average (mg/kg)^a	Area (ft²)	Volume (ft³)	Mass (gm)^a	Concentration Factor	Adjusted Average (mg/kg)^a
Arsenic							
L1	0-1	5.63	2.50E+03	2.50E+03	5.82E+02	0.22	1.22
L2	01-10	3.16	9.90E+03	9.90E+04	1.29E+04	0.86	2.72
L3	10-20	2.83	9.70E+03	1.07E+05	1.25E+04	0.84	2.39
L4	20-30	3.99	1.15E+04	1.27E+05	2.08E+04	1.00	3.99
L5	30-40	2.68	3.60E+03	3.96E+04	4.38E+03	0.31	0.84
L6	40-50	3.58	5.80E+03	6.38E+04	9.44E+03	0.50	1.80
L7	50-63	3.50	6.10E+03	4.88E+04	7.06E+03	0.53	1.86
Total Mass					6.77E+04		
Beryllium							
L1	0-1	0.58	1.88E+04	1.88E+04	4.54E+02	0.56	0.33
L2	01-10	0.93	8.90E+03	8.90E+04	3.43E+03	0.27	0.25
L3	10-20	0.87	9.30E+03	1.02E+05	3.67E+03	0.28	0.24
L4	20-30	1.06	1.37E+04	1.51E+05	6.60E+03	0.41	0.43
L5	30-40	1.32	1.65E+04	1.82E+05	9.87E+03	0.49	0.65
L6	40-50	1.62	1.79E+04	1.97E+05	1.32E+04	0.53	0.87
L7	50-63	1.69	3.35E+04	2.68E+05	1.87E+04	1.00	1.69
Total Mass					5.60E+04		
Manganese							
L1	0-1	319.80	3.52E+04	3.52E+04	4.65E+05	1.00	319.80
L2	01-10	299.12	3.52E+04	3.52E+05	4.35E+06	1.00	299.12
L3	10-20	228.05	3.52E+04	3.87E+05	3.65E+06	1.00	228.05
L4	20-30	155.04	3.52E+04	3.87E+05	2.48E+06	1.00	155.04
L5	30-40	144.52	3.52E+04	3.87E+05	2.31E+06	1.00	144.52
L6	40-50	133.14	3.52E+04	3.87E+05	2.13E+06	1.00	133.14
L7	50-63	139.65	3.52E+04	2.81E+05	1.62E+06	1.00	139.65
Total Mass					1.70E+07		
Nickel							
L1	0-1	13.03	2.27E+04	2.27E+04	1.22E+04	0.64	8.28
L2	01-10	10.26	1.76E+04	1.76E+05	7.46E+04	0.49	5.06
L3	10-20	10.09	1.51E+04	1.66E+05	6.93E+04	0.42	4.27
L4	20-30	13.02	1.40E+04	1.54E+05	8.29E+04	0.39	5.11
L5	30-40	20.85	1.08E+04	1.19E+05	1.02E+05	0.30	6.31
L6	40-50	16.36	1.36E+04	1.50E+05	1.01E+05	0.38	6.23
L7	50-63	15.26	3.57E+04	2.86E+05	1.80E+05	1.00	15.26
Total Mass					6.23E+05		
TCE							
L1	0-1	0.0000	0.00E+00	0.00E+00	0.00E+00	0.00	0.0000
L2	01-10	0.0033	4.30E+03	4.30E+04	5.80E+00	1.00	0.0033
L3	10-20	0.0045	4.30E+03	4.73E+04	8.75E+00	1.00	0.0045
L4	20-30	0.0052	4.30E+03	4.73E+04	1.01E+01	1.00	0.0052
L5	30-40	0.0056	3.90E+03	4.29E+04	9.97E+00	0.91	0.0051
L6	40-50	0.0062	3.10E+03	3.41E+04	8.80E+00	0.72	0.0045
L7	50-63	0.0066	2.10E+03	1.68E+04	4.61E+00	0.49	0.0032
Total Mass					4.81E+01		
Uranium							
L1	0-1	114.00	5.90E+03	5.90E+03	2.78E+04	1.00	114.00
L2	01-10	1.06	3.00E+03	3.00E+04	1.31E+03	0.51	0.54
L3	10-20	1.06	2.80E+03	3.08E+04	1.35E+03	0.47	0.50
L4	20-30	1.05	2.90E+03	3.19E+04	1.39E+03	0.49	0.52

Table E.3.19. Summary of Source Term Characteristics Developed by SADA for SWMU 6 (Continued)

SADA Layer	Depth (ft)	Average (mg/kg) ^a	Area (ft ²)	Volume (ft ³)	Mass (gm) ^a	Concentration Factor	Adjusted Average (mg/kg) ^a
Uranium							
L5	30-40	1.06	4.30E+03	4.73E+04	2.06E+03	0.73	0.77
L6	40-50	1.06	5.30E+03	5.83E+04	2.56E+03	0.90	0.95
L7	50-63	1.09	3.70E+03	2.96E+04	1.33E+03	0.63	0.68
Total Mass					3.78E+04		
Vanadium							
L1	0-1	17.71	2.88E+04	2.88E+04	2.11E+04	0.54	9.61
L2	01-10	23.24	2.72E+04	2.72E+05	2.61E+05	0.51	11.91
L3	10-20	21.39	2.72E+04	2.99E+05	2.65E+05	0.51	10.96
L4	20-30	27.18	2.72E+04	2.99E+05	3.36E+05	0.51	13.92
L5	30-40	30.16	2.72E+04	2.99E+05	3.73E+05	0.51	15.45
L6	40-50	23.99	2.72E+04	2.99E+05	2.97E+05	0.51	12.29
L7	50-63	22.81	5.31E+04	4.25E+05	4.01E+05	1.00	22.81
Total Mass					1.95E+06		
Zinc							
L1	0-1	56.69	2.17E+04	2.17E+04	5.09E+04	0.57	32.46
L2	01-10	31.37	1.90E+04	1.90E+05	2.46E+05	0.50	15.73
L3	10-20	34.49	1.42E+04	1.56E+05	2.23E+05	0.37	12.92
L4	20-30	34.84	1.47E+04	1.62E+05	2.33E+05	0.39	13.51
L5	30-40	44.67	1.25E+04	1.38E+05	2.54E+05	0.33	14.73
L6	40-50	55.70	1.60E+04	1.76E+05	4.05E+05	0.42	23.52
L7	50-63	54.62	3.79E+04	3.03E+05	6.85E+05	1.00	54.62
Total Mass					2.10E+06		
⁹⁹Tc							
L1	0-1	1.45E+01	6.40E+03	6.40E+03	3.84E+09	1.00	14.5
L2	01-10	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00	0.0
L3	10-20	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00	0.0
L4	20-30	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00	0.0
L5	30-40	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00	0.0
L6	40-50	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00	0.0
L7	50-63	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00	0.0
Total Mass					3.84E+09		

^a Radionuclides are in units of pCi/g for concentrations and pCi for mass.

Table E.3.20. Chemical-Specific Parameters of the Analytes Used in SESOIL Modeling of SWMU 6

Analyte	Mol. Wt. (MW) (g/mol)	Solubility in water (mg/L)	Diffusion in air (cm ² /s)	Diffusion in water (m ² /hr)	Henry's Constant (atm.m ³ /mol)	Koc (L/kg)	Kd ^a (L/kg)	Half Life (years)
Arsenic	74.92	1.00E+07	NA	3.60E-07	NA	NA	29	infinite
Beryllium	9.01	1.00E+07	NA	3.60E-07	NA	NA	250	infinite
Manganese	54.94	1.00E+07	NA	3.60E-07	NA	NA	65	Infinite
Nickel	58.69	1.00E+07	NA	3.60E-07	NA	NA	300	infinite
TCE	131	1,100	0.08	3.28E-06	0.0103	94	0.0752	26.6
⁹⁹ Tc	99	1.00E+07	NA	3.60E-07	NA	NA	0.2	2.13E+05
Uranium	238	1.00E+07	NA	3.60E-07	NA	NA	66.8	4.47E+09
Vanadium	50.94	1.00E+07	NA	3.60E-07	NA	NA	1,000	infinite
Zinc	67.41	1.00E+07	NA	3.60E-07	NA	NA	62	infinite

^a The soil/water distribution coefficient (Kd) of an organic compound depends on the soil's organic content (f_{oc}) and compound's organic partition coefficient (Koc). Kd values presented for organic compounds are for UCRS soils (with f_{oc} value of 0.08%) only. Kds used in AT123D are different due to the f_{oc} of 0.02% in the RGA.

E.3.1.5.3 Groundwater modeling results for SWMU 6

All of the analytes modeled at SWMU 6 that originally passed the screening steps for groundwater did not reach the water table in 1,000 years (i.e., beryllium, nickel, vanadium, and ⁹⁹Tc) or exhibited groundwater concentrations that were less than the groundwater background or the groundwater child no action levels (i.e., arsenic, TCE, uranium, and zinc) (see Section 5.4 of the main text). The only exception was for manganese at the SWMU boundary with an HQ of 0.18.

E.3.1.6 SWMU 7

The C-747-A area is located in the northwest corner of the PGDP secured area and comprises the eastern two-thirds of C-747-A. SWMU 7 includes five discrete burial pit areas (DOE 1998c) used for the disposal of wastes from 1957 to 1979. The following summarizes what is known about the size and disposed waste in the burial pits.

- Pit B—This pit measures approximately 60 ft by 172 ft in area. Buried material includes noncombustible trash and contaminated and noncombustible material and equipment. According to the Phase II PGDP Site Investigation geophysical survey (CH2M HILL 1992), the actual excavation extends beyond the designated boundaries and may connect with the adjacent burial pit (Pit C). A geophysical survey conducted for this RI interprets B and C as separate pits.
- Pit C—This pit is approximately the same size as Pit B. Historic records indicate that both Pit B and C received the same material.
- Pit D—This pit underlies an area of approximately 15 ft by 99 ft. Documented buried material consists of uranium-contaminated concrete pieces of reactor tray bases from C-410, used during the fluorination process of uranium tetrafluoride to uranium hexafluoride.
- Pit E (outside the eastern boundary of SWMU 7, within an adjacent scrap yard)—This pit measures approximately 15 ft by 143 ft. Documented buried material consists of uranium-contaminated concrete pieces of reactor tray bases.
- Pits F1-F5—These five pits are all small (average size of each pit is approximately 20 ft by 80 ft). Engineering drawings indicate a sixth “F” pit that was not labeled. Documented buried material consists of uranium-contaminated scrap metal and equipment and empty uranium and magnesium powder.
- Pit G—This pit extends approximately 27 ft by 122 ft in area. Documented buried material consists of noncombustible trash and contaminated and noncombustible material and equipment.

E.3.1.6.1 Conceptual model for source areas at SWMU 7

SWMU 7 occupies an area of approximately 240,900 ft² (5.5 acres). The thickness of the UCRS was estimated to be 60 ft (depth to the top of the RGA). The conceptual model for SWMU 7 is that potentially contaminated materials were buried in waste pits at SWMU 7. Subsequently, contaminants in the disposed material directly impacted soils below or adjacent to the areas where material was buried and, through vertical infiltration in soil, contaminated the groundwater underlying these sources. The infiltrating groundwater migrates vertically through the UCRS and laterally in the RGA, which could transport the contaminants to the POEs.

E.3.1.6.2 Contaminant transport modeling for SWMU 7 using SESOIL and AT123D

For this modeling effort, the soil zones were arranged in four layers. The first soil layer represented the SADA surface soil data from 0 to 1 ft deep. The second soil layer was 9 ft thick representing SADA layer 2 from 1 to 9 ft in depth, the third layer was subdivided into 4 sublayers of equal thickness (10 ft each) representing SADA layers 3 through 6. SESOIL soil layer 4 was subdivided into 4 sublayers of equal thickness (2.5 ft each) to represent the total thickness of SADA layer 7 from 50 ft to 60 ft in depth, the four sublayer division of this layer allowed for better numerical solution of the final flux to the RGA. Table E.3.21 presents the analytes remaining after the screening process and the source terms for each analyte. Table E.3.22 lists the chemical-specific parameters applicable to the SESOIL model of SWMU 7. Figure E.3.4 presents the head distribution and flowpaths of several particles released from the SWMU. The distances to the POEs used in the AT123D model for SWMU 7 are 97 ft to the plant boundary, 2,367 ft to the property boundary, and 14,283 ft to the Little Bayou seeps.

Table E.3.21. Summary of Source Term Characteristics Developed by SADA for SWMU 7

SADA Layer	Depth (ft)	Average (mg/kg) ^a	Area (ft ²)	Volume (ft ³)	Mass (gm) ^a	Concentration Factor	Adjusted Average (mg/kg) ^a
1,1-DCE							
L1	0-1	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L2	01-10	0.87	2.80E+04	2.52E+05	9.01E+03	1.00E+00	0.87
L3	10-20	0.77	2.76E+04	2.76E+05	8.74E+03	9.86E-01	0.75
L4	20-30	0.80	2.68E+04	2.68E+05	8.83E+03	9.57E-01	0.76
L5	30-40	0.67	2.60E+04	2.60E+05	7.18E+03	9.29E-01	0.62
L6	40-50	0.68	2.48E+04	2.48E+05	7.00E+03	8.86E-01	0.60
L7	50-60	0.53	2.32E+04	2.32E+05	5.08E+03	8.29E-01	0.44
Total Mass					4.58E+04		
Arsenic							
L1	0-1	6.17	2.38E+05	2.38E+05	6.08E+04	1.59E+00	9.80
L2	01-10	3.60	1.50E+05	1.35E+06	2.01E+05	1.00E+00	3.60
L3	10-20	3.35	1.42E+05	1.42E+06	1.97E+05	9.49E-01	3.18
L4	20-30	3.43	1.41E+05	1.41E+06	2.00E+05	9.39E-01	3.22
L5	30-40	3.20	1.34E+05	1.34E+06	1.78E+05	8.96E-01	2.86
L6	40-50	3.43	1.26E+05	1.26E+06	1.78E+05	8.40E-01	2.88
L7	50-60	3.14	1.29E+05	1.29E+06	1.67E+05	8.59E-01	2.69
Total Mass					1.18E+06		
Benzo(a)pyrene							
L1	0-1	1.13	1.42E+05	1.42E+05	6.64E+03	1.00E+00	1.13
L2	01-10	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L3	10-20	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L4	20-30	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L5	30-40	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L6	40-50	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L7	50-60	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
Total Mass					6.64E+03		
Cadmium							
L1	0-1	0.56	1.22E+05	1.22E+05	2.80E+03	1.00E+00	0.558
L2	01-10	0.15	1.76E+04	1.58E+05	9.81E+02	1.45E-01	0.022
L3	10-20	0.15	1.68E+04	1.68E+05	1.03E+03	1.38E-01	0.020
L4	20-30	0.15	1.60E+04	1.60E+05	9.82E+02	1.32E-01	0.020
L5	30-40	0.15	1.48E+04	1.48E+05	9.28E+02	1.22E-01	0.018
L6	40-50	0.16	9.60E+03	9.60E+04	6.22E+02	7.89E-02	0.012
L7	50-60	0.16	7.60E+03	7.60E+04	4.96E+02	6.25E-02	0.010
Total Mass					7.84E+03		

Table E.3.21. Summary of Source Term Characteristics Developed by SADA for SWMU 7 (Continued)

SADA Layer	Depth (ft)	Average (mg/kg) ^a	Area (ft ²)	Volume (ft ³)	Mass (gm) ^a	Concentration Factor	Adjusted Average (mg/kg) ^a
cis-1,2-DCE							
L1	0-1	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.000
L2	01-10	0.01	3.56E+04	3.20E+05	1.37E+02	1.02E+00	0.011
L3	10-20	0.05	3.36E+04	3.36E+05	6.98E+02	9.66E-01	0.049
L4	20-30	0.06	3.64E+04	3.64E+05	8.72E+02	1.05E+00	0.061
L5	30-40	0.10	3.48E+04	3.48E+05	1.45E+03	1.00E+00	0.101
L6	40-50	0.07	3.88E+04	3.88E+05	1.09E+03	1.11E+00	0.076
L7	50-60	0.11	2.48E+04	2.48E+05	1.09E+03	7.13E-01	0.075
			Total Mass			5.33E+03	
Fluoranthene							
L1	0-1	1.79	1.68E+05	1.68E+05	1.25E+04	1.00E+00	1.79
L2	01-10	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L3	10-20	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L4	20-30	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L5	30-40	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L6	40-50	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L7	50-60	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
			Total Mass			1.25E+04	
Manganese							
L1	0-1	286.91	2.38E+05	2.38E+05	2.82E+06	1.35E+00	387.98
L2	01-10	210.86	1.76E+05	1.58E+06	1.38E+07	1.00E+00	210.86
L3	10-20	180.78	1.76E+05	1.76E+06	1.32E+07	1.00E+00	180.78
L4	20-30	168.55	1.76E+05	1.76E+06	1.23E+07	1.00E+00	168.55
L5	30-40	150.16	1.76E+05	1.76E+06	1.09E+07	1.00E+00	150.16
L6	40-50	123.56	1.76E+05	1.76E+06	8.99E+06	1.00E+00	123.56
L7	50-60	127.42	1.76E+05	1.76E+06	9.27E+06	1.00E+00	127.42
			Total Mass			7.12E+07	
Mercury							
L1	0-1	0.058	1.54E+05	1.54E+05	3.72E+02	1.00E+00	0.058
L2	01-10	0.023	3.68E+04	3.31E+05	3.19E+02	2.38E-01	0.006
L3	10-20	0.022	3.64E+04	3.64E+05	3.38E+02	2.36E-01	0.005
L4	20-30	0.021	2.92E+04	2.92E+05	2.53E+02	1.89E-01	0.004
L5	30-40	0.021	3.00E+04	3.00E+05	2.58E+02	1.94E-01	0.004
L6	40-50	0.023	2.08E+04	2.08E+05	2.01E+02	1.35E-01	0.003
L7	50-60	0.024	2.00E+04	2.00E+05	1.95E+02	1.30E-01	0.003
			Total Mass			1.94E+03	
Nickel							
L1	0-1	25.94	2.38E+05	2.38E+05	2.55E+05	1.99E+00	51.62
L2	01-10	12.64	1.27E+05	1.14E+06	5.96E+05	1.06E+00	13.40
L3	10-20	12.52	1.20E+05	1.20E+06	6.19E+05	1.00E+00	12.52
L4	20-30	13.51	9.64E+04	9.64E+05	5.38E+05	8.06E-01	10.89
L5	30-40	13.51	9.00E+04	9.00E+05	5.03E+05	7.53E-01	10.17
L6	40-50	12.56	8.84E+04	8.84E+05	4.59E+05	7.39E-01	9.29
L7	50-60	12.84	8.72E+04	8.72E+05	4.63E+05	7.29E-01	9.36
			Total Mass			3.43E+06	

Table E.3.21. Summary of Source Term Characteristics Developed by SADA for SWMU 7 (Continued)

SADA Layer	Depth (ft)	Average (mg/kg) ^a	Area (ft ²)	Volume (ft ³)	Mass (gm) ^a	Concentration Factor	Adjusted Average (mg/kg) ^a
PCB-1254							
L1	0-1	0.130	1.88E+04	1.88E+04	1.01E+02	8.87E-01	0.115
L2	01-10	0.034	2.20E+04	1.98E+05	2.79E+02	1.04E+00	0.035
L3	10-20	0.033	2.12E+04	2.12E+05	2.92E+02	1.00E+00	0.033
L4	20-30	0.033	1.92E+04	1.92E+05	2.66E+02	9.06E-01	0.030
L5	30-40	0.035	1.68E+04	1.68E+05	2.45E+02	7.92E-01	0.028
L6	40-50	0.039	9.60E+03	9.60E+04	1.55E+02	4.53E-01	0.018
L7	50-60	0.040	7.60E+03	7.60E+04	1.25E+02	3.58E-01	0.014
			Total Mass		1.46E+03		
PCB-1260							
L1	0-1	0.63	1.80E+05	1.80E+05	4.71E+03	1.00E+00	0.63
L2	01-10	2.45	8.00E+02	7.20E+03	7.29E+02	4.43E-03	0.01
L3	10-20	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L4	20-30	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L5	30-40	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L6	40-50	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L7	50-60	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
			Total Mass		5.44E+03		
Pyrene							
L1	0-1	2.07	1.68E+05	1.68E+05	1.44E+04	1.00E+00	2.07
L2	01-10	0.03	1.00E+04	9.00E+04	1.23E+02	5.95E-02	0.002
L3	10-20	0.03	9.20E+03	9.20E+04	1.26E+02	5.48E-02	0.002
L4	20-30	0.03	8.80E+03	8.80E+04	1.20E+02	5.24E-02	0.002
L5	30-40	0.03	8.80E+03	8.80E+04	1.20E+02	5.24E-02	0.002
L6	40-50	0.03	6.40E+03	6.40E+04	8.73E+01	3.81E-02	0.001
L7	50-60	0.03	5.20E+03	5.20E+04	7.09E+01	3.10E-02	0.001
			Total Mass		1.50E+04		
Selenium							
L1	0-1	0.66	1.00E+05	1.00E+05	2.73E+03	3.85E+00	2.54
L2	01-10	0.55	2.60E+04	2.34E+05	5.29E+03	1.00E+00	0.55
L3	10-20	0.54	2.12E+04	2.12E+05	4.69E+03	8.15E-01	0.44
L4	20-30	0.53	1.84E+04	1.84E+05	4.06E+03	7.08E-01	0.38
L5	30-40	0.52	1.64E+04	1.64E+05	3.51E+03	6.31E-01	0.33
L6	40-50	0.52	1.12E+04	1.12E+05	2.41E+03	4.31E-01	0.22
L7	50-60	0.49	8.00E+03	8.00E+04	1.62E+03	3.08E-01	0.15
			Total Mass		2.43E+04		
⁹⁹Tc							
L1	0-1	54.05	2.65E+05	2.65E+05	5.93E+11	1.00	54.05
L2	01-10	2.17	4.84E+04	4.36E+05	3.91E+10	0.18	0.40
L3	10-20	2.22	4.88E+04	4.88E+05	4.48E+10	0.18	0.41
L4	20-30	2.21	4.12E+04	4.12E+05	3.77E+10	0.16	0.34
L5	30-40	2.22	3.80E+04	3.80E+05	3.49E+10	0.14	0.32
L6	40-50	1.94	2.64E+04	2.64E+05	2.12E+10	0.10	0.19
L7	50-60	2.05	2.60E+04	2.60E+05	2.21E+10	0.10	0.20
			Total Mass		7.92E+11		

Table E.3.21. Summary of Source Term Characteristics Developed by SADA for SWMU 7 (Continued)

SADA Layer	Depth (ft)	Average (mg/kg) ^a	Area (ft ²)	Volume (ft ³)	Mass (gm) ^a	Concentration Factor	Adjusted Average (mg/kg) ^a
Tetrachloroethene							
L1	0-1	0.0000	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.0000
L2	01-10	0.0062	3.60E+03	3.24E+04	8.30E+00	9.00E-01	0.0056
L3	10-20	0.0062	2.00E+03	2.00E+04	5.13E+00	5.00E-01	0.0031
L4	20-30	0.0062	4.00E+03	4.00E+04	1.03E+01	1.00E+00	0.0062
L5	30-40	0.0062	1.60E+03	1.60E+04	4.10E+00	4.00E-01	0.0025
L6	40-50	0.0062	3.20E+03	3.20E+04	8.20E+00	8.00E-01	0.0050
L7	50-60	0.0062	1.20E+03	1.20E+04	3.08E+00	3.00E-01	0.0019
Total Mass					3.91E+01		
TCE							
L1	0-1	0.0000	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.0000
L2	01-10	0.0071	2.00E+04	1.80E+05	5.27E+01	5.56E-01	0.0039
L3	10-20	0.0318	2.04E+04	2.04E+05	2.68E+02	5.67E-01	0.0180
L4	20-30	0.0357	2.96E+04	2.96E+05	4.37E+02	8.22E-01	0.0294
L5	30-40	0.0550	3.20E+04	3.20E+05	7.28E+02	8.89E-01	0.0489
L6	40-50	0.0881	3.60E+04	3.60E+05	1.31E+03	1.00E+00	0.0881
L7	50-60	0.0793	2.48E+04	2.48E+05	8.13E+02	6.89E-01	0.0546
Total Mass					3.61E+03		
²³⁴U							
L1	0-1	61.35	2.38E+05	2.38E+05	6.04E+11	1.00	61.35
L2	01-10	3.12	1.06E+05	9.58E+05	1.23E+11	0.45	1.39
L3	10-20	12.13	9.12E+04	9.12E+05	4.58E+11	0.38	4.65
L4	20-30	13.64	8.64E+04	8.64E+05	4.87E+11	0.36	4.95
L5	30-40	13.21	8.44E+04	8.44E+05	4.61E+11	0.35	4.69
L6	40-50	11.24	8.48E+04	8.48E+05	3.94E+11	0.36	4.01
L7	50-60	8.23	7.60E+04	7.60E+05	2.58E+11	0.32	2.63
Total Mass					2.79E+12		
²³⁵U							
L1	0-1	7.75	2.38E+05	2.38E+05	7.62E+10	1.00	7.75
L2	01-10	0.40	5.72E+04	5.15E+05	8.43E+09	0.24	0.10
L3	10-20	0.50	4.00E+04	4.00E+05	8.29E+09	0.17	0.08
L4	20-30	0.50	3.36E+04	3.36E+05	6.93E+09	0.14	0.07
L5	30-40	0.52	2.72E+04	2.72E+05	5.80E+09	0.11	0.06
L6	40-50	0.51	1.64E+04	1.64E+05	3.46E+09	0.07	0.04
L7	50-60	0.58	1.12E+04	1.12E+05	2.67E+09	0.05	0.03
Total Mass					1.12E+11		
²³⁸U							
L1	0-1	387.67	2.38E+05	2.38E+05	3.81E+12	1.00	387.67
L2	01-10	8.67	6.88E+04	6.19E+05	2.22E+11	0.29	2.51
L3	10-20	23.85	5.52E+04	5.52E+05	5.44E+11	0.23	5.53
L4	20-30	26.37	5.48E+04	5.48E+05	5.97E+11	0.23	6.07
L5	30-40	25.20	5.04E+04	5.04E+05	5.25E+11	0.21	5.34
L6	40-50	24.50	4.08E+04	4.08E+05	4.13E+11	0.17	4.20
L7	50-60	22.16	2.92E+04	2.92E+05	2.68E+11	0.12	2.72
Total Mass					6.38E+12		

Table E.3.21. Summary of Source Term Characteristics Developed by SADA for SWMU 7 (Continued)

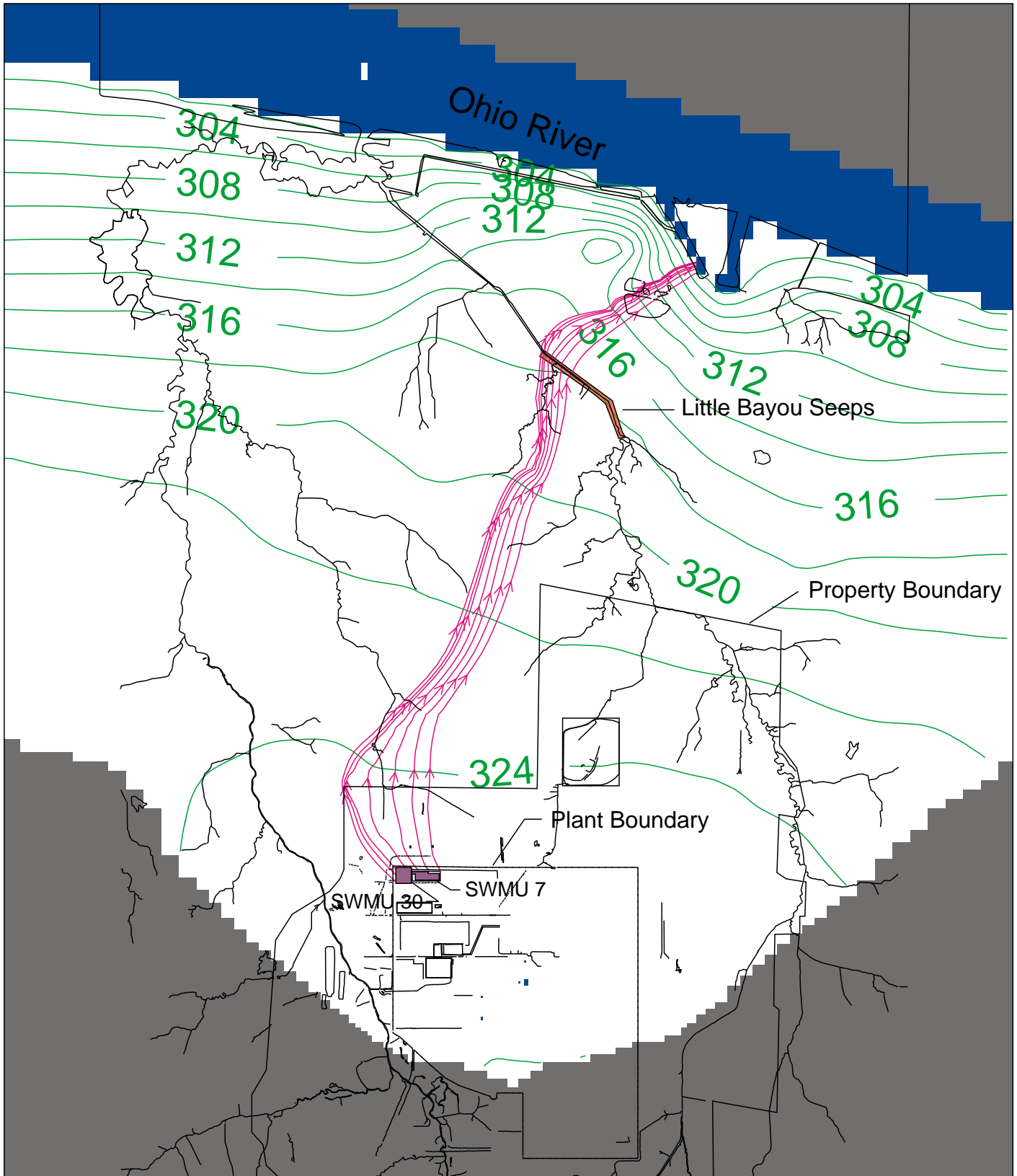
SADA Layer	Depth (ft)	Average (mg/kg) ^a	Area (ft ²)	Volume (ft ³)	Mass (gm) ^a	Concentration Factor	Adjusted Average (mg/kg) ^a
Uranium							
L1	0-1	374.74	2.22E+05	2.22E+05	3.45E+06	1.00E+00	374.74
L2	01-10	16.16	4.72E+04	4.25E+05	2.84E+05	2.12E-01	3.43
L3	10-20	21.38	2.84E+04	2.84E+05	2.51E+05	1.28E-01	2.73
L4	20-30	16.18	3.40E+04	3.40E+05	2.27E+05	1.53E-01	2.47
L5	30-40	17.66	2.88E+04	2.88E+05	2.10E+05	1.29E-01	2.29
L6	40-50	12.34	2.88E+04	2.88E+05	1.47E+05	1.29E-01	1.60
L7	50-60	14.84	1.84E+04	1.84E+05	1.13E+05	8.27E-02	1.23
Total Mass					4.68E+06		
Vanadium							
L1	0-1	26.44	2.38E+05	2.38E+05	2.60E+05	1.59	41.95
L2	01-10	12.92	1.56E+05	1.40E+06	7.48E+05	1.04	13.40
L3	10-20	12.37	1.50E+05	1.50E+06	7.67E+05	1.00	12.37
L4	20-30	11.82	1.53E+05	1.53E+06	7.47E+05	1.02	12.04
L5	30-40	11.00	1.49E+05	1.49E+06	6.77E+05	0.99	10.91
L6	40-50	10.02	1.52E+05	1.52E+06	6.28E+05	1.01	10.13
L7	50-60	10.39	1.48E+05	1.48E+06	6.34E+05	0.98	10.23
Total Mass					4.46E+06		
Vinyl Chloride							
L1	0-1	0.000	0.00E+00	0.00E+00	0.00E+00	0.00	0.0000
L2	01-10	0.005	7.20E+03	6.48E+04	1.46E+01	0.64	0.0035
L3	10-20	0.13	9.20E+03	9.20E+04	5.05E+02	0.82	0.11
L4	20-30	0.15	9.20E+03	9.20E+04	5.88E+02	0.82	0.13
L5	30-40	0.23	1.12E+04	1.12E+05	1.07E+03	1.00	0.23
L6	40-50	0.59	2.40E+03	2.40E+04	5.80E+02	0.21	0.13
L7	50-60	0.59	3.20E+03	3.20E+04	7.74E+02	0.29	0.17
Total Mass					3.53E+03		
Zinc							
L1	0-1	82.26	2.38E+05	2.38E+05	8.09E+05	2.27	186.82
L2	01-10	32.01	1.05E+05	9.43E+05	1.25E+06	1.00	32.01
L3	10-20	31.23	1.05E+05	1.05E+06	1.35E+06	1.00	31.23
L4	20-30	33.48	8.52E+04	8.52E+05	1.18E+06	0.81	27.22
L5	30-40	34.59	7.80E+04	7.80E+05	1.12E+06	0.74	25.75
L6	40-50	35.59	7.08E+04	7.08E+05	1.04E+06	0.68	24.04
L7	50-60	37.33	7.96E+04	7.96E+05	1.23E+06	0.76	28.35
Total Mass					7.98E+06		

^a Radionuclides are in units of pCi/g for concentrations and pCi for mass.

Table E.3.22. Chemical-Specific Parameters of the Analytes Used in SESOIL Modeling of SWMU 7

Analyte	Mol. Wt. (MW) (g/mol)	Solubility in water (mg/L)	Diffusion in air (cm ² /s)	Diffusion in water (m ² /hr)	Henry's Constant (atm.m ³ /mol)	Koc (L/kg)	Kd ^a (L/kg)	Half Life (years)
1,1-DCE	97	2.25E+03	0.09	3.74E-06	0.0261	65	0.013	infinite
Arsenic	74.92	1.00E+07	NA	3.60E-07	NA	NA	29	infinite
Benzo(a)pyrene	252.32	1.62E-03	4.3E-02	3.24E-06	1.13E-06	9.69E+05	772	infinite
Cadmium	112.41	1.00E+07	NA	3.60E-07	NA	NA	75	infinite
<i>cis</i> -1,2-DCE	96.94	3.50E+03	0.07	4.07E-06	4.08E-03	35.5	0.028	infinite
Fluoranthene	202.26	0.206	0.030	2.29E-06	1.61E-05	4.91E+04	39.3	infinite
Manganese	54.94	1.00E+07	NA	1.29E-07	NA	NA	65	infinite
Mercury	200.59	6.00E-02	3.07E-02	2.27E-06	2.44E-02	NA	52	infinite
Nickel	58.69	1.00E+07	NA	3.60E-07	NA	NA	300	infinite
PCB-1254	327	7.00E-02	1.56E-02	1.80E-06	3.40E-04	4.25E+04	34	infinite
PCB-1260	375.7	2.70E02	1.38E-02	1.56E-06	7.40E-05	2.07E+05	165.6	infinite
Pyrene	202.3	0.135	0.0272	2.61E-06	1.1E-05	6.8E+04	54.4	infinite
Selenium	80.98	1.00E+07	NA	3.60E-07	NA	NA	5	infinite
⁹⁹ Tc	99	1.00E+07	NA	3.60E-07	NA	NA	0.2	2.13E+05
Tetrachloroethene	165.8	200	0.072	2.95E-06	0.0184	265	0.053	infinite
TCE	131	1,100	0.08	3.28E-06	0.0103	94	0.0752	26.6
²³⁴ U	234	1.00E+07	NA	3.60E-07	NA	NA	66.8	2.44E+05
²³⁵ U	235	1.00E+07	NA	3.60E-07	NA	NA	66.8	7.04E+08
²³⁸ U	238	1.00E+07	NA	3.60E-07	NA	NA	66.8	4.47E+09
Uranium	238	1.00E+07	NA	3.60E-07	NA	NA	66.8	4.47E+09
Vanadium	50.94	1.00E+07	NA	3.60E-07	NA	NA	1,000	infinite
Vinyl Chloride	63	2,760	0.11	4.43E-07	0.0270	18.8	0.0152	infinite
Zinc	67.41	1.00E+07	NA	3.60E-07	NA	NA	62	infinite

^a The soil/water distribution coefficient (Kd) of an organic compound depends on the soil's organic content (f_{oc}) and compound's organic partition coefficient (Koc). Kd values presented for organic compounds are for UCRS soils (with f_{oc} value of 0.08%) only. Kds used in AT123D are different due to the f_{oc} of 0.02% in the RGA.



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U.S. DEPARTMENT OF ENERGY
DOE PORTSMOUTH/PADUCAH PROJECT OFFICE
PADUCAH GASEOUS DIFFUSION PLANT

Paducah Gaseous Diffusion Plant

- █ River
- █ Particle Tracks
- █ Hydraulic Head (ft)

0 100 500 1000
METERS

Figure E.3.4. Particle Tracks for SWMUs 7 and 30

E.3.1.6.3 Groundwater modeling results for SWMU 7

Table E.3.23 summarizes the predicted maximum groundwater concentrations at the POEs for the analytes modeled at SWMU 7. These contaminants were predicted by SESOIL to reach the water table within the 1,000 year period in concentrations that were greater than the groundwater background or greater than the groundwater child no action levels. Several contaminants that originally passed the screening for groundwater did not reach the water table in 1,000 years [i.e., benzo(a)pyrene, fluoranthene, nickel, PCB-1260, Np-237, and vanadium] or exhibited groundwater concentrations that were less than the groundwater background or the groundwater child no action levels (i.e., cadmium, mercury, pyrene, selenium, tetrachloroethene, zinc and ²³⁵U) (see Section 5.4 of the main text).

As shown in Table E.3.23, the predicted maximum groundwater concentrations for 1,1-DCE; arsenic; manganese; TCE; and vinyl chloride exceed the MCLs at the plant boundary. All analyte groundwater concentrations are less than the MCLs at the property boundary, and Little Bayou seeps. The HQs and cancer risks calculated in accordance with the Risk Methods Document (DOE 2001) are presented in Table E.3.24. The predicted TCE and arsenic concentrations result in the greatest HQs. 1,1-DCE, arsenic, TCE and vinyl chloride provides the highest cancer risk for SWMU 7. ⁹⁹Tc also was predicted to present cancer risks in the 10⁻⁵ range.

Figures 5.16 through 5.25 in Section 5 of the main text, show the predicted concentrations over time at each POE for analytes with a HQ greater than 0.1 and/or a risk greater than 1.0E-06 for contaminants migrating from SWMU 7. As shown in these figures, arsenic, TCE and vinyl chloride are predicted to exceed their respective MCLs at the plant boundary. No analytes were predicted to exceed their respective MCLs at the other POEs.

Table E.3.23. Concentrations of the Analytes in Groundwater Predicted in SESOIL and AT123D Modeling of SWMU 7

Analyte	Predicted Maximum Groundwater Concentration ^{a,b}				
	SWMU (mg/L)	Plant Boundary (mg/L)	Property Boundary (mg/L)	Little Bayou seeps (mg/L)	MCL (mg/L or pCi/L)
1,1-DCE	<i>8.98E-02</i>	<i>8.24E-02</i>	1.10E-02	4.02E-03	0.07
Arsenic	<i>1.78E-02</i>	<i>1.26E-02</i>	2.35E-03	0	0.01
<i>cis</i> ,-1,2-DCE	2.35E-02	2.15E-02	3.13E-03	1.17E-03	0.07
Manganese	<i>3.32E-01</i>	<i>2.41E-01</i>	1.05E-06	0	^d
PCB-1254	5.23E-05	3.09E-05	3.05E-06	1.32E-12	^d
⁹⁹ Tc	<i>9.09E+02</i>	8.25E+02	2.70E+02	1.32E+02	900 ^c
TCE	<i>1.09E-02</i>	<i>9.87E-03</i>	1.42E-03	5.06E-04	0.005
²³⁴ U	7.94E+00	5.79E+00	5.84E-06	0	20
²³⁸ U	7.59E+00	5.58E+00	5.85E-06	0	20
Uranium	3.46E-03	2.53E-03	2.68E-09	0	0.03
Vinyl Chloride	<i>1.35E-02</i>	<i>1.24E-02</i>	1.21E-03	4.13E-04	0.002

^a Values in bold, italic font exceed the analyte's MCL.

^b Radionuclide concentrations are in pCi/L.

^c ⁹⁹Tc MCL based on a critical organ dose at 4 mrem/yr from drinking water consumption.

^d MCLs not available for these contaminants.

Table E.3.24. Hazard and Cancer Risk Predicted from Maximum Groundwater Concentrations Derived in Modeling of SWMU 7 Using SESOIL and AT123D^a

Analyte	SWMU		Plant Boundary		Property Boundary		Little Bayou seeps	
	Hazard Quotient	Cancer Risk	Hazard Quotient	Cancer Risk	Hazard Quotient	Cancer Risk	Hazard Quotient	Cancer Risk
1,1-DCE	0.85	2.08E-03	0.8	1.9E-03	0.1	2.5E-04	<0.1	9.3E-05
Arsenic	5.70	4.72E-04	4.0	3.3E-04	0.8	6.2E-05	^b	^b
<i>cis</i> ,-1,2-DCE	1.24	^b	1.1	^b	0.2	^b	<0.1	^b
Manganese	0.71	^b	0.5	^b	<0.1	^b	^b	^b
PCB-1254	4.20	7.09E-06	2.5	4.8E-06	0.2	<1.0E-06	<0.1	<1.0E-06
⁹⁹ Tc	^b	4.99E-05	^b	4.5E-05	^b	1.5E-05	^b	7.3E-06
TCE	4.98	1.27E-04	4.5	3.1E-04	0.6	4.4E-05	0.2	1.6E-05
²³⁴ U	^b	1.11E-05	^b	8.2E-06	^b	<1.0E-06	^b	^b
²³⁸ U	^b	1.32E-05	^b	9.7E-06	^b	<1.0E-06	^b	^b
Uranium	0.55	^b	0.40	^b	<0.1	^b	^b	^b
Vinyl Chloride	0.64	3.72E-04	0.6	3.6E-04	<0.1	3.6E-05	<0.1	1.2E-05

^a Contaminants with a HQ greater than 0.1 or a cancer risk greater than 1.00E-06 are considered analytes.

^b Value not calculated since the groundwater concentrations was reported as zero at this POE by AT123D, or the contaminant did not have a reported cancer slope factor or chemical toxicity RfD.

E.3.1.7 SWMU 30

SWMU 30 consists of an historical burn-and-burial pit and the location of a former incinerator, used from 1951 to 1970 to burn combustible trash which may have contained uranium contamination. The pit is reported to have been excavated to a depth of 12 ft. SWMU 30 is bounded on the east side by C-747-A Burial Ground (SWMU 7).

E.3.1.7.1 Conceptual model for source areas at SWMU 30

SWMU 30 occupies an area of approximately 128,000 ft² (2.9 acres). The thickness of the UCRS was estimated to be 61 ft (depth to the top of the RGA). The conceptual model for SWMU 30 is that potentially contaminated materials were buried in waste pits at SWMU 30. Subsequently, contaminants in disposed material directly impacted soils below or adjacent to the areas where material was buried and, through vertical infiltration in soil, contaminated the groundwater underlying these sources. The infiltrating groundwater migrates vertically through the UCRS and laterally in the RGA, which could transport the contaminants to the POEs.

E.3.1.7.2 Contaminant transport modeling for SWMU 30 using SESOIL and AT123D

For this modeling effort, the soil zones were arranged in four layers. The first soil layer represented the SADA surface soil data from 0 to 1 ft deep. The second soil layer was 9 ft thick representing SADA layer 2 from 1 to 9 ft in depth, the third layer was subdivided into 4 sublayers of equal thickness (10 ft each) representing SADA layers 3 through 6. SESOIL soil layer 4 was subdivided into 4 sublayers of equal thickness (2.75 ft each) to represent the total thickness of SADA layer 7 from 50 ft to 61 ft in depth, the four sublayer division of this layer allowed for better numerical solution of the final flux to the RGA Table E.3.25 presents the analytes remaining after the screening process and the source terms for each analyte. Table E.3.26 lists the chemical-specific parameters applicable to the SESOIL model of SWMU 30. Figure E.3.4 presents the head distribution and flowpaths of several particles released from the SWMU. The distances to the POEs used in the AT123D model for SWMU 30 are 107 ft to the plant boundary, 2,127 ft to the property boundary, and 13,013 ft to the Little Bayou seeps.

Table E.3.25. Summary of Source Term Characteristics Developed by SADA for SWMU 30

SADA Layer	Depth (ft)	Average (mg/kg)^a	Area (ft²)	Volume (ft³)	Mass (gm)^a	Concentration Factor	Adjusted Average (mg/kg)^a
1,1-DCE							
L1	0-1	0.00	0.00E+00	0.00E+00	0.00E+00	0.00	0.00
L2	01-10	0.00	0.00E+00	0.00E+00	0.00E+00	0.00	0.00
L3	10-20	0.00	4.00E+02	4.00E+03	0.00E+00	0.13	0.00
L4	20-30	0.00	4.00E+02	4.00E+03	0.00E+00	0.13	0.00
L5	30-40	0.01	1.60E+03	1.60E+04	3.31E+00	0.50	0.003
L6	40-50	0.01	2.00E+03	2.00E+04	4.13E+00	0.63	0.003
L7	50-61	0.01	3.20E+03	3.52E+04	7.28E+00	1.00	0.01
Total Mass					1.47E+01		
Acenaphthene							
L1	0-1	0.28	5.16E+04	5.16E+04	6.06E+02	1.00E+00	0.28
L2	01-10	0.017	3.20E+03	2.88E+04	2.02E+01	6.20E-02	0.0011
L3	10-20	0.017	3.20E+03	3.20E+04	2.25E+01	6.20E-02	0.0011
L4	20-30	0.017	3.20E+03	3.20E+04	2.25E+01	6.20E-02	0.0011
L5	30-40	0.017	2.40E+03	2.40E+04	1.69E+01	4.65E-02	0.0008
L6	40-50	0.017	1.20E+03	1.20E+04	8.43E+00	2.33E-02	0.0004
L7	50-61	0.017	1.20E+03	1.32E+04	9.28E+00	2.33E-02	0.0004
Total Mass					7.06E+02		

Table E.3.25. Summary of Source Term Characteristics Developed by SADA for SWMU 30 (Continued)

SADA Layer	Depth (ft)	Average (mg/kg)^a	Area (ft²)	Volume (ft³)	Mass (gm)^a	Concentration Factor	Adjusted Average (mg/kg)^a
Arsenic							
L1	0-1	4.55	7.15E+04	7.15E+04	1.35E+04	0.86	3.92
L2	01-10	4.49	7.25E+04	6.52E+05	1.21E+05	0.87	3.93
L3	10-20	4.36	7.44E+04	7.44E+05	1.34E+05	0.90	3.91
L4	20-30	4.23	7.72E+04	7.72E+05	1.35E+05	0.93	3.94
L5	30-40	4.10	8.04E+04	8.04E+05	1.36E+05	0.97	3.98
L6	40-50	4.04	8.30E+04	8.30E+05	1.39E+05	1.00	4.04
L7	50-61	3.95	8.59E+04	9.45E+05	1.54E+05	1.04	4.09
Total Mass					8.33E+05		
Benzo(a)pyrene							
L1	0-1	1.00	9.72E+04	9.72E+04	4.01E+03	1.00E+00	1.00
L2	01-10	0.05	3.20E+03	2.88E+04	6.19E+01	3.29E-02	0.0017
L3	10-20	0.05	3.20E+03	3.20E+04	6.88E+01	3.29E-02	0.0017
L4	20-30	0.05	3.20E+03	3.20E+04	6.88E+01	3.29E-02	0.0017
L5	30-40	0.05	2.40E+03	2.40E+04	5.16E+01	2.47E-02	0.0013
L6	40-50	0.05	1.20E+03	1.20E+04	2.58E+01	1.23E-02	0.0006
L7	50-61	0.05	1.20E+03	1.32E+04	2.84E+01	1.23E-02	0.0006
Total Mass					4.31E+03		
Cadmium							
L1	0-1	1.92	4.68E+04	4.68E+04	3.72E+03	1.00E+00	1.925
L2	01-10	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.000
L3	10-20	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.000
L4	20-30	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.000
L5	30-40	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.000
L6	40-50	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.000
L7	50-61	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.000
Total Mass					3.72E+03		
Dibenzoanthracene							
L1	0-1	0.33	5.64E+04	5.64E+04	7.61E+02	1.00E+00	0.327
L2	01-10	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.000
L3	10-20	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.000
L4	20-30	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.000
L5	30-40	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.000
L6	40-50	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.000
L7	50-61	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.000
Total Mass					7.61E+02		
Fluorene							
L1	0-1	0.25	4.56E+04	4.56E+04	4.72E+02	1.00E+00	0.25
L2	01-10	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L3	10-20	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L4	20-30	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L5	30-40	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L6	40-50	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L7	50-61	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
Total Mass					4.72E+02		

Table E.3.25. Summary of Source Term Characteristics Developed by SADA for SWMU 30 (Continued)

SADA Layer	Depth (ft)	Average (mg/kg)^a	Area (ft²)	Volume (ft³)	Mass (gm)^a	Concentration Factor	Adjusted Average (mg/kg)^a
Manganese							
L1	0-1	338.66	1.22E+05	1.22E+05	1.71E+06	1.46E+00	495.84
L2	01-10	250.12	8.36E+04	7.52E+05	7.78E+06	1.00E+00	250.12
L3	10-20	242.52	8.36E+04	8.36E+05	8.38E+06	1.00E+00	242.52
L4	20-30	238.27	8.36E+04	8.36E+05	8.24E+06	1.00E+00	238.27
L5	30-40	221.98	8.36E+04	8.36E+05	7.67E+06	1.00E+00	221.98
L6	40-50	206.94	8.36E+04	8.36E+05	7.15E+06	1.00E+00	206.94
L7	50-61	212.37	8.36E+04	9.20E+05	8.07E+06	1.00E+00	212.37
Total Mass					4.90E+07		
Mercury							
L1	0-1	0.111	8.48E+04	8.48E+04	3.88E+02	1.00E+00	0.111
L2	01-10	0.062	1.40E+04	1.26E+05	3.24E+02	1.65E-01	0.0103
L3	10-20	0.055	1.24E+04	1.24E+05	2.81E+02	1.46E-01	0.0080
L4	20-30	0.053	1.04E+04	1.04E+05	2.28E+02	1.23E-01	0.0065
L5	30-40	0.040	9.20E+03	9.20E+04	1.52E+02	1.08E-01	0.0043
L6	40-50	0.022	6.80E+03	6.80E+04	6.14E+01	8.02E-02	0.0018
L7	50-61	0.022	6.80E+03	7.48E+04	6.75E+01	8.02E-02	0.0018
Total Mass					1.50E+03		
Naphthalene							
L1	0-1	0.31	8.00E+03	8.00E+03	1.03E+02	1.00E+00	0.31
L2	01-10	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L3	10-20	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L4	20-30	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L5	30-40	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L6	40-50	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
L7	50-61	0.00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00
Total Mass					1.03E+02		
Nickel							
L1	0-1	60.18	1.22E+05	1.22E+05	3.05E+05	1.65E+00	99.00
L2	01-10	17.40	7.44E+04	6.70E+05	4.82E+05	1.00E+00	17.40
L3	10-20	16.86	7.32E+04	7.32E+05	5.10E+05	9.84E-01	16.58
L4	20-30	16.42	7.32E+04	7.32E+05	4.97E+05	9.84E-01	16.16
L5	30-40	15.69	7.28E+04	7.28E+05	4.72E+05	9.78E-01	15.36
L6	40-50	15.44	7.28E+04	7.28E+05	4.65E+05	9.78E-01	15.11
L7	50-61	15.11	7.44E+04	8.18E+05	5.11E+05	1.00E+00	15.11
Total Mass					3.24E+06		
PCB-1254							
L1	0-1	0.200	3.00E+04	3.00E+04	2.48E+02	1.00E+00	0.200
L2	01-10	0.028	5.60E+03	5.04E+04	5.83E+01	1.87E-01	0.0052
L3	10-20	0.028	5.60E+03	5.60E+04	6.48E+01	1.87E-01	0.0052
L4	20-30	0.028	5.60E+03	5.60E+04	6.48E+01	1.87E-01	0.0052
L5	30-40	0.028	5.60E+03	5.60E+04	6.48E+01	1.87E-01	0.0052
L6	40-50	0.028	5.60E+03	5.60E+04	6.48E+01	1.87E-01	0.0052
L7	50-61	0.028	5.20E+03	5.72E+04	6.62E+01	1.73E-01	0.0049
Total Mass					6.32E+02		

E.3.25. Summary of Source Term Characteristics Developed by SADA for SWMU 30 (Continued)

SADA Layer	Depth (ft)	Average (mg/kg)^a	Area (ft²)	Volume (ft³)	Mass (gm)^a	Concentration Factor	Adjusted Average (mg/kg)^a
PCB-1260							
L1	0-1	1.54	1.22E+05	1.22E+05	7.81E+03	1.00E+00	1.54
L2	01-10	0.08	3.00E+04	2.70E+05	8.40E+02	2.45E-01	0.018
L3	10-20	0.07	2.76E+04	2.76E+05	7.92E+02	2.25E-01	0.016
L4	20-30	0.07	2.52E+04	2.52E+05	6.93E+02	2.06E-01	0.014
L5	30-40	0.06	2.12E+04	2.12E+05	5.27E+02	1.73E-01	0.010
L6	40-50	0.05	1.84E+04	1.84E+05	4.17E+02	1.50E-01	0.008
L7	50-61	0.05	1.68E+04	1.85E+05	4.14E+02	1.37E-01	0.007
Total Mass					1.15E+04		
Pyrene							
L1	0-1	2.16	1.08E+05	1.08E+05	9.66E+03	1.00E+00	2.16
L2	01-10	0.043	3.04E+04	2.74E+05	4.82E+02	2.81E-01	0.012
L3	10-20	0.042	2.96E+04	2.96E+05	5.16E+02	2.74E-01	0.012
L4	20-30	0.042	2.80E+04	2.80E+05	4.81E+02	2.59E-01	0.011
L5	30-40	0.041	2.56E+04	2.56E+05	4.32E+02	2.37E-01	0.010
L6	40-50	0.040	2.40E+04	2.40E+05	4.02E+02	2.22E-01	0.009
L7	50-61	0.040	2.20E+04	2.42E+05	4.00E+02	2.04E-01	0.008
Total Mass					1.24E+04		
Selenium							
L1	0-1	0.66	2.12E+04	2.12E+04	5.78E+02	1.56E+00	1.03
L2	01-10	0.91	1.40E+04	1.26E+05	4.73E+03	1.03E+00	0.94
L3	10-20	0.91	1.36E+04	1.36E+05	5.09E+03	1.00E+00	0.91
L4	20-30	0.90	1.32E+04	1.32E+05	4.93E+03	9.71E-01	0.88
L5	30-40	0.92	1.20E+04	1.20E+05	4.56E+03	8.82E-01	0.81
L6	40-50	0.96	1.08E+04	1.08E+05	4.27E+03	7.94E-01	0.76
L7	50-61	0.95	1.04E+04	1.14E+05	4.51E+03	7.65E-01	0.73
Total Mass					2.87E+04		
⁹⁹Tc							
L1	0-1	20.79	1.22E+05	1.22E+05	1.05E+11	1.00	20.79
L2	01-10	1.78	3.56E+04	3.20E+05	2.36E+10	0.29	0.52
L3	10-20	1.54	3.32E+04	3.32E+05	2.12E+10	0.27	0.42
L4	20-30	1.42	3.08E+04	3.08E+05	1.80E+10	0.25	0.36
L5	30-40	1.14	2.68E+04	2.68E+05	1.26E+10	0.22	0.25
L6	40-50	0.89	2.40E+04	2.40E+05	8.80E+09	0.20	0.17
L7	50-61	0.86	2.20E+04	2.42E+05	8.56E+09	0.18	0.15
Total Mass					1.98E+11		
TCE							
L1	0-1	0.0000	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.000
L2	01-10	0.037	4.40E+03	3.96E+04	6.12E+01	1.00E+00	0.037
L3	10-20	0.037	4.40E+03	4.40E+04	6.80E+01	1.00E+00	0.037
L4	20-30	0.037	3.60E+03	3.60E+04	5.57E+01	8.20E-01	0.031
L5	30-40	0.037	3.20E+03	3.20E+04	4.95E+01	7.30E-01	0.027
L6	40-50	0.037	2.40E+03	2.40E+04	3.71E+01	5.40E-01	0.020
L7	50-61	0.037	2.40E+03	2.64E+04	4.08E+01	5.40E-01	0.020
Total Mass					3.12E+02		

E.3.25. Summary of Source Term Characteristics Developed by SADA for SWMU 30 (Continued)

SADA Layer	Depth (ft)	Average (mg/kg)^a	Area (ft²)	Volume (ft³)	Mass (gm)^a	Concentration Factor	Adjusted Average (mg/kg)^a
²³⁴U							
L1	0-1	42.53	1.22E+05	1.22E+05	2.15E+11	1.00	42.53
L2	01-10	4.39	6.56E+04	5.90E+05	1.07E+11	0.54	2.36
L3	10-20	4.64	6.76E+04	6.76E+05	1.30E+11	0.55	2.56
L4	20-30	4.54	6.64E+04	6.64E+05	1.25E+11	0.54	2.46
L5	30-40	4.04	6.80E+04	6.80E+05	1.14E+11	0.56	2.25
L6	40-50	3.99	6.60E+04	6.60E+05	1.09E+11	0.54	2.15
L7	50-61	3.50	7.00E+04	7.70E+05	1.11E+11	0.57	2.00
Total Mass					9.11E+11		
²³⁵U							
L1	0-1	4.44	1.22E+05	1.22E+05	2.25E+10	1.00	4.44
L2	01-10	0.31	3.80E+04	3.42E+05	4.34E+09	0.31	0.10
L3	10-20	0.33	3.44E+04	3.44E+05	4.64E+09	0.28	0.09
L4	20-30	0.31	3.40E+04	3.40E+05	4.41E+09	0.28	0.09
L5	30-40	0.34	2.84E+04	2.84E+05	4.01E+09	0.23	0.08
L6	40-50	0.35	2.72E+04	2.72E+05	3.92E+09	0.22	0.08
L7	50-61	0.36	2.36E+04	2.60E+05	3.90E+09	0.19	0.07
Total Mass					4.77E+10		
²³⁸U							
L1	0-1	103.92	1.22E+05	1.22E+05	5.26E+11	1.00	103.92
L2	01-10	7.61	5.96E+04	5.36E+05	1.69E+11	0.49	3.71
L3	10-20	9.37	5.80E+04	5.80E+05	2.25E+11	0.47	4.44
L4	20-30	9.55	5.32E+04	5.32E+05	2.10E+11	0.43	4.15
L5	30-40	8.75	5.16E+04	5.16E+05	1.87E+11	0.42	3.69
L6	40-50	8.99	4.60E+04	4.60E+05	1.71E+11	0.38	3.38
L7	50-61	8.64	4.48E+04	4.93E+05	1.76E+11	0.37	3.16
Total Mass					1.66E+12		
Uranium							
L1	0-1	797.22	2.88E+04	2.88E+04	9.49E+05	1.00E+00	797.22
L2	01-10	4.39	5.28E+04	4.75E+05	8.62E+04	1.83E+00	8.05
L3	10-20	4.20	5.40E+04	5.40E+05	9.38E+04	1.88E+00	7.88
L4	20-30	4.19	5.20E+04	5.20E+05	9.01E+04	1.81E+00	7.57
L5	30-40	3.97	5.40E+04	5.40E+05	8.87E+04	1.88E+00	7.45
L6	40-50	4.06	5.24E+04	5.24E+05	8.79E+04	1.82E+00	7.38
L7	50-61	3.87	5.40E+04	5.94E+05	9.50E+04	1.88E+00	7.25
Total Mass					1.49E+06		
Vanadium							
L1	0-1	25.32	1.22E+05	1.22E+05	1.28E+05	1.47	37.25
L2	01-10	14.34	8.32E+04	7.49E+05	4.44E+05	1.00	14.34
L3	10-20	13.51	8.32E+04	8.32E+05	4.65E+05	1.00	13.51
L4	20-30	12.98	8.36E+04	8.36E+05	4.49E+05	1.00	13.05
L5	30-40	12.05	8.36E+04	8.36E+05	4.16E+05	1.00	12.11
L6	40-50	11.28	8.36E+04	8.36E+05	3.90E+05	1.00	11.33
L7	50-61	11.04	8.36E+04	9.20E+05	4.20E+05	1.00	11.09
Total Mass					2.71E+06		

E.3.25. Summary of Source Term Characteristics Developed by SADA for SWMU 30 (Continued)

SADA Layer	Depth (ft)	Average (mg/kg) ^a	Area (ft ²)	Volume (ft ³)	Mass (gm) ^a	Concentration Factor	Adjusted Average (mg/kg) ^a
Zinc							
L1	0-1	105.49	1.22E+05	1.22E+05	5.34E+05	1.84	194.45
L2	01-10	42.25	6.72E+04	6.05E+05	1.06E+06	1.01	42.76
L3	10-20	41.98	6.60E+04	6.60E+05	1.15E+06	0.99	41.73
L4	20-30	42.44	6.52E+04	6.52E+05	1.14E+06	0.98	41.67
L5	30-40	42.67	6.44E+04	6.44E+05	1.14E+06	0.97	41.39
L6	40-50	43.94	6.36E+04	6.36E+05	1.16E+06	0.96	42.08
L7	50-61	44.35	6.64E+04	7.30E+05	1.34E+06	1.00	44.35
Total Mass					7.51E+06		

^a Radionuclides are in units of pCi/g for concentrations and pCi for mass.

Table E.3.26. Chemical-Specific Parameters of the Analytes Used in SESOIL Modeling of SWMU 30

Analyte	Mol. Wt. (MW) (g/mol)	Solubility in water (mg/L)	Diffusion in air (cm ² /s)	Diffusion in water (m ² /hr)	Henry's Constant (atm.m3/mol)	Koc (L/kg)	Kd ^a (L/kg)	Half Life (years)
1,1-DCE	97	2.25E+03	0.09	3.74E-06	0.0261	65	0.013	infinite
Acenaphthene	154.0	4.20	0.04	2.77E-6	1.60E-04	4.90E+03	3.9	infinite
Arsenic	74.92	1.00E+07	NA	3.60E-07	NA	NA	29	infinite
Benzo(a)pyrene	252.32	1.62E-03	4.3E-02	3.24E-06	1.13E-06	9.69E+05	772	infinite
Cadmium	112.41	1.00E+07	NA	3.60E-07	NA	NA	75	infinite
Dibenzo(a,h)anthracene	278.33	0.0025	0.020	1.86E-06	1.47E-08	1.78E+06	1424	infinite
Fluorene	166.0	1.90	0.061	2.84E-06	7.7E-05	7.9E+03	6.3	infinite
Manganese	54.94	1.00E+07	NA	1.29E-07	NA	NA	65	infinite
Mercury	200.59	6.00E-02	3.07E-02	2.27E-06	2.44E-02	NA	52	infinite
Naphthalene	128.16	31.0	0.059	2.70E-06	4.83E-04	1.19E+03	0.95	infinite
Nickel	58.69	1.00E+07	NA	3.60E-07	NA	NA	300	infinite
PCB-1254	327	7.00E-02	1.56E-02	1.80E-06	3.40E-04	4.25E+04	34	infinite
PCB-1260	375.7	2.70E02	1.38E-02	1.56E-06	7.40E-05	2.07E+05	165.6	infinite
Pyrene	202.3	0.135	0.0272	2.61E-06	1.1E-05	6.8E+04	54.4	infinite
Selenium	80.98	1.00E+07	NA	3.60E-07	NA	NA	5	infinite
⁹⁹ Tc	99	1.00E+07	NA	3.60E-07	NA	NA	0.2	2.13E+05
TCE	131	1,100	0.08	3.28E-06	0.0103	94	0.0752	26.6
²³⁴ U	234	1.00E+07	NA	3.60E-07	NA	NA	66.8	2.44E+05
²³⁵ U	235	1.00E+07	NA	3.60E-07	NA	NA	66.8	7.04E+08
²³⁸ U	238	1.00E+07	NA	3.60E-07	NA	NA	66.8	4.47E+09
Uranium	238	1.00E+07	NA	3.60E-07	NA	NA	66.8	4.47E+09
Vanadium	50.94	1.00E+07	NA	3.60E-07	NA	NA	1,000	infinite
Zinc	67.41	1.00E+07	NA	3.60E-07	NA	NA	62	infinite

^a The soil/water distribution coefficient (Kd) of an organic compound depends on the soil's organic content (f_{oc}) and compound's organic partition coefficient (Koc). Kd values presented for organic compounds are for UCRS soils (with f_{oc} value of 0.08%) only. Kds used in AT123D are different due to the f_{oc} of 0.02% in the RGA.

E.3.1.7.3 Groundwater modeling results for SWMU 30

Table E.3.27 summarizes the predicted maximum groundwater concentrations at the POEs for the analytes modeled at SWMU 30. These contaminants were predicted by SESOIL to reach the water table within the 1,000 year period in concentrations that were greater than the groundwater background or greater than the groundwater child no action levels. Several contaminants that originally passed the screening for groundwater did not reach the water table in 1,000 years [i.e., benzo(a)pyrene, cadmium, dibenzo(a,h)anthracene, nickel, and vanadium] or exhibited groundwater concentrations that were less than the groundwater background or the groundwater child no action levels (i.e., acenaphthene, fluorine, mercury, naphthalene, PCB-1254, PCB-1260, pyrene, zinc and ²³⁵U) (see Section 5.4 of the main text) including acenaphthene, benzo(a)pyrene, cadmium, dibenzo(a,h)anthracene, fluorine, mercury, naphthalene, nickel, PCB-1254, PCB-1260, pyrene, ²³⁵U, vanadium, and zinc.

Table E.3.27. Concentrations of the Analytes in Groundwater Predicted in SESOIL and AT123D Modeling of SWMU 30

Analyte	Predicted Maximum Groundwater Concentration ^{a,b}				MCL (mg/L or pCi/L)
	SWMU (mg/L)	Plant Boundary (mg/L)	Property Boundary (mg/L)	Little Bayou seeps (mg/L)	
1,1-DCE	8.18E-05	7.65E-05	6.14E-06	1.86E-06	0.07
Arsenic	<i>1.82E-02</i>	<i>1.21E-02</i>	2.50E-03	0	0.01
Manganese	<i>3.78E-01</i>	<i>2.51E-01</i>	2.85E-04	0	^d
Selenium	1.51E-02	8.30E-03	9.21E-04	3.15E-04	0.05
⁹⁹ Tc	2.87E+02	2.64E+02	7.08E+01	2.92E+01	900 ^c
TCE	9.11E-04	8.60E-04	7.70E-05	2.60E-05	0.005
²³⁴ U	3.99E+00	2.75E+00	1.44E-03	0	20
²³⁸ U	5.91E+00	4.07E+00	1.98E-03	0	20
Uranium	8.40E-03	4.81E-03	2.41E-06	0	0.03

^a Values in bold, italic font exceed the analyte's MCL.

^b Radionuclide concentrations are in pCi/L.

^c ⁹⁹Tc MCL based on a critical organ dose at 4 mrem/yr from drinking water consumption.

^d MCLs not available for these contaminants.

As shown in Table E.3.27, the predicted maximum groundwater concentrations for arsenic and manganese exceed their respective MCLs at the plant boundary. The HQs and cancer risks calculated in accordance with the Risk Methods Document (DOE 2001) are presented in Table E.3.28. The predicted arsenic concentrations result in the greatest HQ and cancer risk for SWMU 30. TCE and ⁹⁹Tc also exhibit elevated cancer risks.

Table E.3.28. Hazard and Cancer Risk Predicted from Maximum Groundwater Concentrations Derived in Modeling of SWMU 30 Using SESOIL and AT123D^a

Analyte	SWMU		Plant Boundary		Property Boundary		Little Bayou seeps	
	Hazard Quotient	Cancer Risk	Hazard Quotient	Cancer Risk	Hazard Quotient	Cancer Risk	Hazard Quotient	Cancer Risk
1,1-DCE	<0.1	1.90E-06	<0.1	1.8E-06	<0.1	<1.0E-06	<0.1	<1.0E-06
Arsenic	5.83	4.82E-04	3.8	3.2E-04	0.8	6.2E-05	^b	^b
Manganese	0.80	^b	0.5	^b	<0.1	^b	^b	^b
Selenium	0.29	^b	0.2	^b	<0.1	^b	<0.1	^b
⁹⁹ Tc	^b	1.57E-05	^b	1.4E-05	^b	3.9E-06	^b	1.6E-06
TCE	0.42	2.83E-05	0.4	2.7E-05	<0.1	2.4E-06	<0.1	<1.0E-06
²³⁴ U	^b	5.63E-06	^b	3.9E-06	^b	<1.0E-06	^b	^b
²³⁸ U	^b	1.03E-05	^b	7.1E-06	^b	<1.0E-06	^b	^b
Uranium	1.34	^b	0.8	^b	<0.1	^b	^b	^b

^a Contaminants with a HQ greater than 0.1 or a cancer risk greater than 1.00E-06 are considered analytes.

^b Value not calculated since the groundwater concentrations was reported as zero at this POE by AT123D, or the contaminant did not have a reported cancer slope factor or chemical toxicity RfD.

Figures 5.26 through 5.33 in Section 5 of the main text, show the predicted concentrations over time at each POE for analytes with a HQ greater than 0.1 and/or a risk greater than 1.0E-06 for contaminants migrating from SWMU 30. As shown in these figures, arsenic, manganese, and TCE are predicted to exceed their respective MCLs at the plant boundary. TCE is also predicted to exceed the MCL at the property boundary, Little Bayou seeps, and Ohio River.

E.3.1.8 SWMU 145

SWMU 145, located north of PGDP, began operation in the early 1950s. A 1973 document *The Discard of Scrap Materials by Burial at the Paducah Plant* (Union Carbide 1973), states this area was used by the contractor during the construction of PGDP to discard all types of scrap and waste materials. Use of the area for discard of scrap and waste by subcontractors continued until the early 1980s. Construction debris, such as concrete, roofing materials, wire, wood, shingles with asbestos, and welding rods are expected to have been disposed of in the area. Approximately once a year, the accumulated scrap piles were moved by plant personnel into piles or earth depressions and, whenever practicable, covered with dirt. The area was later permitted for the construction and operation of the C-746-S & T Landfills (BJC 2001). Currently, the C-746-S&T Landfills are located on top of SWMU 145 (DOE 1999b). Area P (SWMU 145) is defined by the encompassing C-746-S&T Landfills (SWMUs 9 and 10, respectively).

E.3.1.8.1 Conceptual model for source areas at SWMU 145

SWMU 145 occupies an area of approximately 1,916,640 ft² (44 acre). The thickness of the UCRS was estimated to be 60 ft (depth to the top of the RGA). The conceptual model for SWMU 145 is that potentially contaminated materials were buried and landfilled at SWMU 145. Subsequently, contaminants in the disposed material directly impacted soils below or adjacent to the areas where material was buried and, through vertical infiltration in soil, contaminated the groundwater underlying these sources. The infiltrating groundwater migrates vertically through the UCRS and laterally in the RGA, which could transport the contaminants to the POEs.

E.3.1.8.2 Contaminant transport modeling for SWMU 145 using SESOIL and AT123D

For this modeling effort, the soil zones were arranged in four layers. The first soil layer represented the SADA surface soil data from 0 to 1 ft deep. The second soil layer was 9 ft thick representing SADA layer 2 from 1 to 9 ft in depth, the third layer was subdivided into 4 sublayers of equal thickness (10 ft each)

representing SADA layers 3 through 6. SESOIL soil layer 4 was subdivided into 4 sublayers of equal thickness (2.0 ft each) to represent the total thickness of SADA layer 7 from 50 ft to 58 ft in depth, the four sublayer division of this layer allowed for better numerical solution of the final flux to the RGA. Table E.3.29 presents the analytes remaining after the screening process and the source terms for each analyte. Table E.3.30 lists the chemical-specific parameters applicable to the SESOIL model of SWMU 30. Figure E.3.5 presents the head distribution and flowpaths of several particles released from the SWMU. The distances to the POEs used in the AT123D model for SWMU 145 are 2,951 ft to the property boundary and 11,489 ft to the Ohio River.

Table E.3.29. Summary of Source Term Characteristics Developed by SADA for SWMU 145

SADA Layer	Depth (ft)	Average (mg/kg)^a	Area (ft²)	Volume (ft³)	Mass (gm)^a	Concentration Factor	Adjusted Average (mg/kg)^a
Antimony							
L1	0-1	0.00	0.00	0.00	0.00	0.00	0.00
L2	01-10	11.47	1.04E+06	9.36E+06	4.44E+06	1.04	11.93
L3	10-20	11.47	9.70E+05	9.70E+06	4.60E+06	0.97	11.13
L4	20-30	11.46	1.00E+06	1.00E+07	4.74E+06	1.00	11.46
L5	30-40	11.39	9.70E+05	9.70E+06	4.57E+06	0.97	11.05
L6	40-50	11.37	9.80E+05	9.80E+06	4.61E+06	0.98	11.14
L7	50-58	11.40	9.60E+05	7.68E+06	3.62E+06	0.96	10.94
Total Mass					2.66E+07		
Arsenic							
L1	0-1	0.00	0.00	0.00	0.00	0.00	0.00
L2	01-10	4.21	1.91E+06	1.72E+07	2.99E+06	0.97	4.10
L3	10-20	4.33	1.96E+06	1.96E+07	3.51E+06	1.00	4.33
L4	20-30	4.38	1.92E+06	1.92E+07	3.48E+06	0.98	4.29
L5	30-40	4.28	1.93E+06	1.93E+07	3.41E+06	0.98	4.21
L6	40-50	4.28	1.94E+06	1.94E+07	3.43E+06	0.99	4.24
L7	50-58	4.18	1.93E+06	1.54E+07	2.67E+06	0.98	4.12
Total Mass					1.99E+07		
PCB-1254							
L1	0-1	0.00	0.00	0.00	0.00	0.00	0.00
L2	01-10	0.59	2.40E+05	2.16E+06	5.28E+04	1.00	0.59
L3	10-20	1.90	2.00E+04	2.00E+05	1.57E+04	0.08	0.16
L4	20-30	1.90	1.00E+04	1.00E+05	7.86E+03	0.04	0.08
L5	30-40	1.90	1.00E+04	1.00E+05	7.86E+03	0.04	0.08
L6	40-50	0.00	0.00	0.00	0.00	0.00	0.00
L7	50-58	0.00	0.00	0.00	0.00	0.00	0.00
Total Mass					8.57E+04		
PCB-1260							
L1	0-1	0.00	0.00	0.00	0.00	0.00	0.00
L2	01-10	12.50	2.00E+04	1.80E+05	9.30E+04	1.00	12.50
L3	10-20	12.50	2.00E+04	2.00E+05	1.03E+05	1.00	12.50
L4	20-30	12.50	2.00E+04	2.00E+05	1.03E+05	1.00	12.50
L5	30-40	12.50	2.00E+04	2.00E+05	1.03E+05	1.00	12.50
L6	40-50	12.50	1.00E+04	1.00E+05	5.17E+04	0.50	6.25
L7	50-58	12.50	1.00E+04	8.00E+04	4.13E+04	0.50	6.25
Total Mass					4.97E+05		

Table E.3.29. Summary of Source Term Characteristics Developed by SADA for SWMU 145 (Continued)

SADA Layer	Depth (ft)	Average (mg/kg)^a	Area (ft²)	Volume (ft³)	Mass (gm)^a	Concentration Factor	Adjusted Average (mg/kg)^a
Cadmium							
L1	0-1	0.00	0.00	0.00	0.00	0.00	0.00
L2	01-10	2.47	1.00E+04	9.00E+04	9.19E+03	0.20	0.49
L3	10-20	2.39	4.00E+04	4.00E+05	3.95E+04	0.80	1.91
L4	20-30	2.39	4.00E+04	4.00E+05	3.95E+04	0.80	1.91
L5	30-40	2.40	5.00E+04	5.00E+05	4.97E+04	1.00	2.40
L6	40-50	2.40	5.00E+04	5.00E+05	4.97E+04	1.00	2.40
L7	50-58	2.40	5.00E+04	4.00E+05	3.98E+04	1.00	2.40
Total Mass					2.39E+05		
Manganese							
L1	0-1	0.00	0.00	0.00	0.00	0.00	0.00
L2	01-10	146.59	2.55E+06	2.30E+07	1.39E+08	1.00	146.59
L3	10-20	147.65	2.55E+06	2.55E+07	1.56E+08	1.00	147.65
L4	20-30	143.48	2.55E+06	2.55E+07	1.51E+08	1.00	143.48
L5	30-40	140.68	2.55E+06	2.55E+07	1.48E+08	1.00	140.68
L6	40-50	170.30	2.55E+06	2.55E+07	1.80E+08	1.00	170.30
L7	50-58	170.13	2.55E+06	2.04E+07	1.43E+08	1.00	170.13
Total Mass					9.51E+08		
Mercury							
L1	0-1	0.00	0.00	0.00	0.00	0.00	0.00
L2	01-10	0.02	7.70E+05	6.93E+06	5.51E+03	0.95	0.02
L3	10-20	0.02	7.70E+05	7.70E+06	6.12E+03	0.95	0.02
L4	20-30	0.02	7.40E+05	7.40E+06	5.87E+03	0.91	0.02
L5	30-40	0.05	8.10E+05	8.10E+06	1.75E+04	1.00	0.05
L6	40-50	0.05	8.00E+05	8.00E+06	1.74E+04	0.99	0.05
L7	50-58	0.06	7.30E+05	5.84E+06	1.34E+04	0.90	0.05
Total Mass					6.96E+04		
Nickel							
L1	0-1	0.00	0.00	0.00	0.00	0.00	0.00
L2	01-10	10.21	2.00E+06	1.80E+07	7.60E+06	0.95	9.68
L3	10-20	10.22	2.00E+06	2.00E+07	8.45E+06	0.95	9.69
L4	20-30	9.87	1.95E+06	1.95E+07	7.96E+06	0.92	9.12
L5	30-40	10.50	2.11E+06	2.11E+07	9.16E+06	1.00	10.50
L6	40-50	10.08	2.11E+06	2.11E+07	8.80E+06	1.00	10.08
L7	50-58	10.20	2.06E+06	1.65E+07	6.95E+06	0.98	9.96
Total Mass					4.98E+07		
Vanadium							
L1	0-1	0.00	0.00	0.00	0.00	0.00	0.00
L2	01-10	18.43	1.21E+06	1.09E+07	8.30E+06	1.03	18.90
L3	10-20	18.47	1.20E+06	1.20E+07	9.16E+06	1.02	18.78
L4	20-30	19.44	1.18E+06	1.18E+07	9.48E+06	1.00	19.44
L5	30-40	19.94	1.18E+06	1.18E+07	9.73E+06	1.00	19.94
L6	40-50	19.55	1.18E+06	1.18E+07	9.54E+06	1.00	19.55
L7	50-58	18.71	1.19E+06	9.52E+06	7.36E+06	1.01	18.86
Total Mass					5.54E+07		

Table E.3.29. Summary of Source Term Characteristics Developed by SADA for SWMU 145 (Continued)

SADA Layer	Depth (ft)	Average (mg/kg)^a	Area (ft²)	Volume (ft³)	Mass (gm)^a	Concentration Factor	Adjusted Average (mg/kg)^a
Benzo(a)Pyrene							
L1	0-1	0.00	0.00	0.00	0.00	0.00	0.00
L2	01-10	0.00	0.00	0.00	0.00E+00	0.00	0.00
L3	10-20	0.00	0.00	0.00	0.00E+00	0.00	0.00
L4	20-30	0.00	0.00	0.00	0.00E+00	0.00	0.00
L5	30-40	0.00	0.00	0.00	0.00E+00	0.00	0.00
L6	40-50	0.00	0.00	0.00	0.00E+00	0.00	0.00
L7	50-58	0.00	0.00	0.00	0.00E+00	0.00	0.00
Total Mass					1.82E+02		
¹³⁷Cs							
L1	0-1	0.00	0.00E+00	0.00E+00	0.00E+00	0.00	0.00
L2	01-10	0.14	1.28E+05	1.28E+06	7.46E+09	1.00	0.14
L3	10-20	0.11	1.23E+05	1.23E+06	5.67E+09	0.96	0.11
L4	20-30	0.10	1.05E+05	1.05E+06	4.55E+09	0.82	0.09
L5	30-40	0.14	1.05E+05	1.05E+06	6.20E+09	0.82	0.12
L6	40-50	0.14	1.03E+05	1.03E+06	6.12E+09	0.80	0.12
L7	50-58	0.14	1.03E+05	8.20E+05	4.89E+09	0.80	0.12
Total Mass					3.69E+10		
²³⁹Pu							
L1	0-1	0.00	0.00	0.00	0.00E+00	0.00	0.00
L2	01-10	0.28	1.00E+05	9.00E+05	1.15E+10	1.00	0.28
L3	10-20	0.26	1.00E+05	1.00E+06	1.40E+10	1.00	0.26
L4	20-30	0.26	1.00E+05	1.00E+06	1.40E+10	1.00	0.26
L5	30-40	0.26	1.00E+05	1.00E+06	1.40E+10	1.00	0.26
L6	40-50	0.26	1.00E+05	1.00E+06	1.40E+10	1.00	0.26
L7	50-58	0.26	1.00E+05	8.00E+05	1.02E+10	1.00	0.26
Total Mass					7.77E+10		
⁹⁹Tc							
L1	0-1	0.00	0.00	0.00	0.00E+00	0.00	0.00
L2	01-10	26.67	1.10E+05	9.90E+05	1.09E+12	1.00	26.67
L3	10-20	15.98	1.30E+05	1.30E+06	8.59E+11	1.18	18.89
L4	20-30	15.98	1.30E+05	1.30E+06	8.59E+11	1.18	18.89
L5	30-40	15.98	1.30E+05	1.30E+06	8.59E+11	1.18	18.89
L6	40-50	11.91	1.30E+05	1.30E+06	6.40E+11	1.18	14.07
L7	50-58	11.91	1.30E+05	1.04E+06	5.12E+11	1.18	14.07
Total Mass					4.82E+12		
²³⁴U							
L1	0-1	0.00	0.00	0.00	0.00E+00	0.00	0.00
L2	01-10	0.74	2.00E+04	1.80E+05	3.28E+11	1.00	0.74
L3	10-20	0.00	0.00	0.00	2.49E+12	0.00	0.00
L4	20-30	0.00	0.00	0.00	2.50E+12	0.00	0.00
L5	30-40	0.00	0.00	0.00	2.50E+12	0.00	0.00
L6	40-50	0.00	0.00	0.00	2.46E+12	0.00	0.00
L7	50-58	0.00	0.00	0.00	1.79E+12	0.00	0.00
Total Mass					1.21E+13		

Table E.3.29. Summary of Source Term Characteristics Developed by SADA for SWMU 145 (Continued)

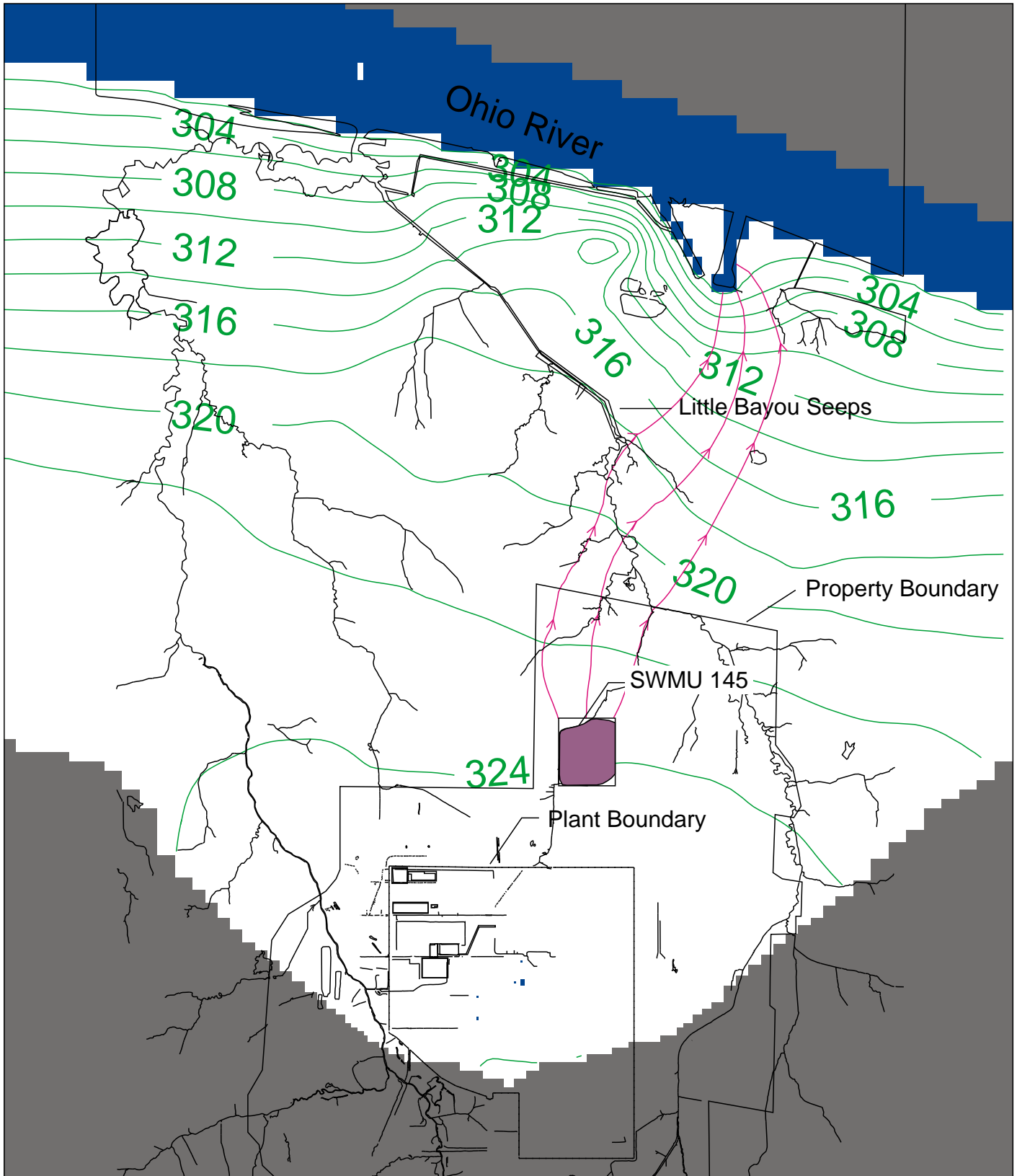
SADA Layer	Depth (ft)	Average (mg/kg) ^a	Area (ft ²)	Volume (ft ³)	Mass (gm) ^a	Concentration Factor	Adjusted Average (mg/kg) ^a
²³⁵ U							
L1	0-1	0.00	0.00	0.00	0.00E+00	0.00	0.00
L2	01-10	0.30	5.00E+04	4.50E+05	6.16E+09	1.00	0.30
L3	10-20	0.19	5.00E+04	5.00E+05	7.16E+09	1.00	0.19
L4	20-30	0.19	5.00E+04	5.00E+05	4.22E+09	1.00	0.19
L5	30-40	0.19	5.00E+04	5.00E+05	4.22E+09	1.00	0.19
L6	40-50	0.00	0.00	0.00	0.00E+00	0.00	0.00
L7	50-58	0.00	0.00	0.00	0.00E+00	0.00	0.00
Total Mass					2.18E+10		
²³⁸ U							
L1	0-1	0.00	0.00	0.00	0.00E+00	0.00	0.00
L2	01-10	2.62	1.21E+06	1.09E+07	2.02E+11	0.96	2.52
L3	10-20	6.77	1.14E+06	1.14E+07	1.36E+12	0.90	6.12
L4	20-30	6.60	1.17E+06	1.17E+07	3.51E+12	0.93	6.13
L5	30-40	6.14	1.26E+06	1.26E+07	3.51E+12	1.00	6.14
L6	40-50	5.62	1.26E+06	1.26E+07	3.52E+12	1.00	5.62
L7	50-58	5.67	1.25E+06	1.00E+07	3.22E+12	0.99	5.62
Total Mass					2.34E+12		

^a Radionuclides are in units of pCi/g for concentrations and pCi for mass.

Table E.3.30. Chemical-Specific Parameters of the Analytes Used in SESOIL Modeling of SWMU 145

Analyte	Mol. Wt. (MW) (g/mol)	Solubility in water (mg/L)	Diffusion in air (cm ² /s)	Diffusion in water (m ² /hr)	Henry's Constant (atm.m3/mol)	Koc (L/kg)	Kd ^a (L/kg)	Half Life (years)
Antimony	121.75	1.00E+07	NA	3.60E-07	NA	NA	45	infinite
PCB-1254	327	7.00E-02	1.56E-02	1.80E-06	3.40E-04	4.25E+04	34	infinite
PCB-1260	375.7	2.70E02	1.38E-02	1.56E-06	7.40E-05	2.07E+05	165.6	infinite
Arsenic	74.92	1.00E+07	NA	3.60E-07	NA	NA	29	infinite
Benzo(a)pyrene	252.32	1.62E-03	4.3E-02	3.24E-06	1.13E-06	9.69E+05	772	infinite
Cadmium	112.41	1.00E+07	NA	3.60E-07	NA	NA	75	infinite
Manganese	54.94	1.00E+07	NA	3.60E-07	NA	NA	65	infinite
Mercury	200.59	6.00E-02	3.07E-02	2.27E-06	2.44E-02	NA	52	infinite
Nickel	58.69	1.00E+07	NA	3.60E-07	NA	NA	300	infinite
Vanadium	50.94	1.00E+07	NA	3.60E-07	NA	NA	1000	infinite
¹³⁷ Cs	137	1.00E+07	NA	3.60E-07	NA	NA	280	30.17
²³⁸ U	238	1.00E+07	NA	3.60E-07	NA	NA	66.8	4.47E+09
²³⁵ U	235	1.00E+07	NA	3.60E-07	NA	NA	66.8	7.04E+08
²³⁴ U	234	1.00E+07	NA	3.60E-07	NA	NA	66.8	2.44E+05
⁹⁹ Tc	99	1.00E+07	NA	3.60E-07	NA	NA	0.2	2.13E+05
²³⁹ Pu	239	1.00E+07	NA	3.60E-07	NA	NA	550	2.41E+04

^a The soil/water distribution coefficient (Kd) of an organic compound depends on the soil's organic content (f_{oc}) and compound's organic partition coefficient (Koc). Kd values presented for organic compounds are for UCRS soils (with f_{oc} value of 0.08%) only. Kds used in AT123D are different due to the f_{oc} of 0.02% in the RGA.



d:\autocad\portage\paducah\swmu45.dwg

U.S. DEPARTMENT OF ENERGY
DOE PORTSMOUTH/PADUCAH PROJECT OFFICE
PADUCAH GASEOUS DIFFUSION PLANT

PADUCAH
Remediation Services
A Portage Shaw Joint Venture Company

Paducah Gaseous Diffusion Plant

- █ River
- █ Particle Tracks
- █ Hydraulic Head (ft)

TRUE NORTH
PLANT NORTH

0 100 500 1000
METERS

Figure E.3.5. Particle Tracks for SWMU 145

E.3.1.8.3 Groundwater modeling results for SWMU 145

Table E.3.31 summarizes the predicted maximum groundwater concentrations at the POEs for the analytes modeled at SWMU 145. These contaminants were predicted by SESOIL to reach the water table within the 1,000 year period in concentrations that were greater than the groundwater background or greater than the groundwater child no action levels. Several contaminants that originally passed the screening for groundwater did not reach the water table in 1,000 years [i.e., PCB-1254, benzo(a)pyrene, vanadium, ²³⁴U, ²³⁵U, and ²³⁹Pu] or exhibited groundwater concentrations that were less than the groundwater background or the groundwater child no action levels (i.e., cadmium, mercury, and nickel) (see Section 5.4 of the main text).

Table E.3.31. Concentrations of the Analytes in Groundwater Predicted in SESOIL and AT123D Modeling of SWMU 145

Analyte	SWMU (mg/L)	Property Boundary (mg/L)	Ohio River (mg/L)	MCL (mg/L or pCi/L)
Antimony	<i>7.99E-02</i>	1.51E-06	0	0.006
Arsenic	<i>6.21E-02</i>	1.61E-03	0	0.01
PCB-1260	1.92E-03	0	0	d
Manganese	8.44E-01	0	0	d
⁹⁹ Tc	<i>1.01E+04</i>	<i>1.84E+03</i>	<i>9.65E+02</i>	900 ^c
²³⁸ U	7.67E-02	0	0	20

^a Values in bold, italic font exceed the analyte's MCL.

^b Radionuclide concentrations are in pCi/L.

^c ⁹⁹Tc MCL based on a critical organ dose at 4 mrem/yr from drinking water consumption.

^d MCLs not available for these contaminants.

As shown in Table E.3.31, the predicted maximum groundwater concentration for ⁹⁹Tc exceeds the MCLs at the property boundary and at the Ohio River. All remaining analytes are less than their MCLs at the POEs. The HQs and cancer risks calculated in accordance with the Risk Methods Document (DOE 2001) are presented in Table E.3.32. The predicted arsenic concentrations at the property boundary results in the greatest HQ, with both arsenic and ⁹⁹Tc providing elevated cancer risks.

Table E.3.32. Hazard and Cancer Risk Predicted from Maximum Groundwater Concentrations Derived in Modeling of SWMU 145 Using SESOIL and AT123D^a

Analyte	SWMU		Property Boundary		Near Ohio River	
	Hazard Quotient	Cancer Risk	Hazard Quotient	Cancer Risk	Hazard Quotient	Cancer Risk
Antimony	20	^b	<0.1	^b	^b	^b
Arsenic	19.9	1.65E-03	0.5	4.3E-05	^b	^b
PCB-1260	^b	3.05E-02				
Manganese	1.80	^b				
⁹⁹ Tc	^b	5.54E-04	^b	1.0E-04	^b	5.3E-05
²³⁸ U	^b	1.10E-07				

^a Contaminants with a HQ greater than 0.1 or a cancer risk greater than 1.00E-06 are considered analytes.

^b Value not calculated since the groundwater concentrations was reported as zero at this POE by AT123D, or the contaminant did not have a reported cancer slope factor or chemical toxicity RfD.

Figures 5.34 through 5.36 in Section 5 of the main text, show the predicted concentrations over time at each POE for analytes with a HQ greater than 0.1 and/or a risk greater than 1.0E-06 for contaminants migrating from SWMU 145. As shown in these figures, manganese is predicted to continue rising in concentration at 1,000 years at the plant boundary exceeding the MCL, but has not reached the property

boundary or Ohio River in the 1,000 year period. Arsenic is also increasing in concentration at the plant boundary at 1,000 years, however the concentrations are less than the MCL. ⁹⁹Tc is not predicted to exceed the MCL at the POEs.

E.3.2 VAPOR TRANSPORT MODELING

The BGOU RI includes vapor transport modeling to evaluate the potential air concentrations in a residential basement for soil and groundwater contamination at the BGOU SWMUs and POEs. Modelers used the Johnson and Ettinger model (1991), coded into spreadsheets by EPA (2004), to assess the potential migration of VOCs into a residential basement.

Johnson and Ettinger (1991) introduced a screening-level model which incorporates both convective and diffusive mechanisms for estimating the transport of contaminant vapors emanating from either subsurface soils or groundwater into indoor spaces located directly above or in close proximity to the source of contamination. The Johnson and Ettinger model is a one-dimensional analytical solution to convective and diffusive vapor transport into indoor spaces and provides an estimated attenuation coefficient that relates the vapor concentration in the indoor space to the vapor concentration at the source of contamination.

Since the Johnson and Ettinger model is a screening level model, the number of parameter inputs is minimized. Table E.3.33 provides the input parameter values used in the vapor transport analysis. All analyses for the BGOU RI used the default chemical property library. The contaminant source inventories for the soil layers beneath the SWMUs were obtained from the SADA analyses presented in Section E.3.1.

Table E.3.34 presents the resulting basement air concentrations, predicted by the model. The HQs and cancer risks calculated in accordance with the Risk Methods Document (DOE 2001) are presented in Table E.3.35. The vapor transport modeling for mercury was conservatively based on the metallic form, which has a Henry's Law Constant of 1.07E-02 atm-m³/mol. Metallic mercury, with its uniquely high vapor pressure relative to other metals, can enter the atmosphere from the groundwater environment as several different gaseous compounds. The rate of vaporization of mercury and certain of its inorganic compounds decrease in the sequence Hg > Hg₂Cl₂ > HgCl₂ > HgS > HgO. The Henry's Law Constant decreases dramatically down the sequence (for example, HgCl₂ has a value of 7.09E-10 atm-m³/mol).

Table E.3.33. Vapor Transport Model Input Parameter Values

Parameter	Value	Reference
Average Soil Temperature (T _s)	15 °C	Default value
Depth below grade to bottom of enclosed space floor (L _F)	200 cm	Default value
SCS soil type	Silty Clay	Table E.3.2
Soil dry bulk density (ρ _b)	1.46 g/cm ³	Table E.3.2
Soil total porosity (n)	0.45	Table E.3.2
Soil water-filled porosity (θ _w)	0.167	Default value
Soil organic carbon fraction (f _{oc})	0.08	Table E.3.2
Enclosed space floor thickness (L _{crack})	10 cm	Default value
Soil-building pressure differential (Δ _p)	40 g/cm-s ²	Default value
Enclosed space floor length (L _B)	1,000 cm	Default value
Enclosed space floor width (W _B)	1,000 cm	Default value
Enclosed space height (H _B)	366 cm	Default value
Floor-wall seam crack width (W)	0.1 cm	Default value
Indoor air exchange rate (ER)	0.5 hr ⁻¹	Default value

Table E.3.34. Basement Air Concentrations Based on Vapor Transport Modeling Results for each BGOU SWMU

Source Area	Contaminant	Air concentration (mg/m ³)		
		On-Site	Plant Boundary	Property Boundary
SWMU 2	TCE	2.81E-02	1.09E-04	5.55E-05
	<i>cis</i> -1,2-DCE	1.95E-01	7.82E-04	3.89E-04
	Naphthalene	2.70E-07	1.56E-08	8.43E-09
SWMU 3	TCE	1.62E-05	8.52E-10	4.47E-10
	Mercury	7.22E-06	1.12E-14	0.00E+00
SWMU 4	TCE	4.90E-03	2.12E-04	1.08E-04
	<i>cis</i> -1,2-DCE	5.76E-03	8.80E-05	4.05E-05
	Vinyl Chloride	6.7E-03	1.98E-04	2.55E-06
SWMU 5	TCE	5.41E-06	1.98E-07	9.13E-08
	Acenaphthene	2.04E-07	7.47E-08	4.30E-08
	Fluorene	5.16E-08	2.37E-08	1.27E-08
	Naphthalene	3.80E-06	9.75E-08	3.79E-08
	Pyrene	2.28E-09	NA	NA
SWMU 6	TCE	9.34E-06	3.88E-09	1.92E-09
SWMU 7	TCE	8.63E-05	4.96E-06	7.16E-07
	<i>cis</i> -1,2-DCE	2.13E-04	9.66E-06	1.42E-06
	Vinyl Chloride	1.23E-02	1.25E-05	1.22E-06
	1,1-DCE	1.03E-02	6.70E-05	9.03E-06
	Mercury	9.99E-06	2.22E-09	2.41E-12
	Pyrene	7.68E-09	4.93E-12	1.31E-12
	Tetrachloroethene	2.00E-05	6.40E-08	4.70E-09
SWMU 30	TCE	6.75E-05	3.42E-07	2.96E-08
	1,1-DCE	3.36E-05	4.85E-08	3.62E-09
	Acenaphthene	2.77E-08	4.96E-09	9.22E-10
	Fluorene	3.92E-09	NA	NA
	Mercury	1.66E-05	8.91E-1	2.23E-11
	Pyrene	6.56E-10	2.47E-11	6.54E-12
	Naphthalene	3.10E-07	1.90E-08	1.85E-09
SWMU 145	Mercury	1.42E-05	7.95E-08	2.60E-14

Table E.3.35. Vapor Hazard Quotients and Risk-Based on Vapor Transport Modeling Results for Each BGOU SWMU

Source Area	Contaminant	On-Site			Plant Boundary			Property Boundary		
		HQ	ELCR	HQ	ELCR	HQ	ELCR	HQ	ELCR	
SWMU 2	TCE	3.15E+00	1.84E-03	1.22E-02	7.14E-06	6.22E-03	3.64E-06	6.22E-03	3.64E-06	
	<i>cis</i> -1,2-DCE	2.50E+01	NA	1.00E-01	NA	4.99E-02	NA	4.99E-02	NA	
	Naphthalene	4.03E-04	NA	4.99E-06	NA	1.26E-05	NA	1.26E-05	NA	
SWMU 3	TCE	1.82E-03	1.06E-06	9.63E-08	5.68E-11	5.01E-08	2.93E-11	5.01E-08	2.93E-11	
	Mercury	1.08E-01	NA	1.67E-10	NA	NA	NA	NA	NA	
SWMU 4	TCE	5.54E-01	3.23E-04	2.38E-02	1.39E-05	1.21E-02	7.07E-06	1.21E-02	7.07E-06	
	<i>cis</i> -1,2-DCE	7.38E-01	NA	1.13E-02	NA	5.19E-03	NA	5.19E-03	NA	
	Vinyl Chloride	2.99E-01	4.19E-05	8.85E-03	1.24E-06	1.14E-04	1.60E-08	1.14E-04	1.60E-08	
SWMU 5	TCE	6.06E-04	3.54E-07	2.22E-05	1.30E-08	1.02E-05	5.98E-09	1.02E-05	5.98E-09	
	Acenaphthene	4.37E-06	NA	1.60E-06	NA	9.21E-07	NA	9.21E-07	NA	
	Fluorene	1.65E-06	NA	7.57E-07	NA	4.06E-07	NA	4.06E-07	NA	
	Naphthalene	5.67E-03	NA	1.45E-03	NA	5.65E-04	NA	5.65E-04	NA	
	Pyrene	9.71E-08	NA	NA	NA	NA	NA	NA	NA	
	TCE	1.05E-03	6.12E-07	4.35E-07	2.54E-10	2.15E-07	1.26E-10	2.15E-07	1.26E-10	
	TCE	9.68E-03	5.65E-06	5.56E-04	3.25E-07	8.03E-05	4.69E-08	8.03E-05	4.69E-08	
	<i>cis</i> -1,2-DCE	2.73E-02	NA	1.24E-03	NA	1.82E-04	NA	1.82E-04	NA	
SWMU 7	Vinyl Chloride	5.48E-01	7.68E-05	5.59E-04	7.83E-08	5.45E-05	7.64E-09	5.45E-05	7.64E-09	
	1,1-DCE	2.30E-01	3.66E-04	1.50E-03	2.39E-06	2.02E-04	3.21E-07	2.02E-04	3.21E-07	
	Mercury	1.49E-01	NA	3.31E-05	NA	3.59E-08	NA	3.59E-08	NA	
SWMU 30	Pyrene	3.27E-07	NA	2.10E-10	NA	5.58E-11	NA	5.58E-11	NA	
	Tetrachloroethene	1.49E-04	8.13E-09	4.78E-07	2.60E-11	3.51E-08	1.91E-12	3.51E-08	1.91E-12	
	TCE	7.57E-03	4.42E-06	3.83E-05	2.24E-08	3.32E-06	1.94E-09	3.32E-06	1.94E-09	
	1,1-DCE	7.52E-04	1.20E-06	1.09E-06	1.73E-09	8.10E-08	1.29E-10	8.10E-08	1.29E-10	
	Acenaphthene	5.93E-07	NA	1.06E-07	NA	1.97E-08	NA	1.97E-08	NA	
SWMU 145	Fluorene	1.25E-07	NA	NA	NA	NA	NA	NA	NA	
	Mercury	2.47E-01	NA	1.33E-05	NA	3.33E-07	NA	3.33E-07	NA	
	Naphthalene	4.62E-04	NA	2.83E-05	NA	2.76E-06	NA	2.76E-06	NA	
	Pyrene	2.80E-08	NA	1.05E-09	NA	2.79E-10	NA	2.79E-10	NA	
	Mercury	2.12E-01	NA	1.19E-03	NA	3.88E-10	NA	3.88E-10	NA	

NA = not applicable

E.3.3 UNCERTAINTY ANALYSIS FOR THE TRANSPORT MODELING

The SADA SESOIL and AT123D models were used for the investigation, resulting in the use of some simplifying assumptions. These assumptions resulted in modeling uncertainties.

E.3.3.1 Source Term Development

The analyte screening analysis for the groundwater pathway and modeling is provided in Appendix E, Attachment 3. Very few contaminants detected during the sampling of the BGOU were deleted because of frequency of detection. Many more contaminants were deleted because they were not detected or because the maximum concentration did not exceed the screening criteria. Many organic compounds did not exceed the screening criteria and also could have been screened based on the frequency of detection. Conversely, many contaminants were included in COCs at various SWMUs even though the detection frequency was approximately 1 to 3 percent. These contaminants included vinyl chloride, TCE, 1,1-DCE, naphthalene, and *cis*-1,2-DCE, which were retained at COCs and contributed significantly to overall risk.

Contaminants that were screened based solely on frequency of detection included 1,1,1-trichloroethane, 1,1,2-trichloroethane, 1,2-dichloroethane, dibenzofuran, acenaphthylene, mercury, and silver. The elements or chemicals listed above were screened for frequency at only one or two SWMUs.

The source term was developed using sampling results, geospatial analyses in SADA, and considering SESOIL limitations. While the sampling results are appropriate for source identification, SESOIL requires input of the soil concentrations for each layer of interest in the UCRS. Due to SESOIL's requirement to use the same constant area for each layer, the analyte concentrations of all layers needed to be normalized against the area of the layer with the maximum estimated analyte mass; therefore, geospatial interpolation was used based on the SADA nearest neighbor algorithm to estimate the total mass in each UCRS layer based upon the sampling results.

The techniques in SADA that can be used for source term development are nearest neighbor, inverse distance, and ordinary kriging. The nearest neighbor technique was selected for source zone refinement because it yielded results that were most compatible with the conceptual site model of contaminant release, as described in Attachment 2 to Appendix E.

Each potential analyte source area was discretized using rows and columns with a uniform spacing. Multiple domains with varying depths were used to characterize the analyte source areas vertically in relation to the existing aquifers; therefore, the domain was further discretized into horizontal layers. Analyte results for each domain were compiled, and analyte concentrations in each cell of the domain were predicted using geospatial interpolation (see Appendix E Attachment 2 for details).

The source term is based on a three-dimensional, geospatial analysis of the data using nearest neighbor interpolation in SADA. Therefore, sample data was assessed both horizontally within each layer and vertically between layers. This resulted in a conservative analysis of the subsurface data, such that sample detections in a layer with no corresponding sample locations in the adjacent vertical layers, resulted in predictions of contamination in these adjacent layers. For the BGOU RI, soil samples were typically collected from angled soil borings; thus, deeper samples did not underlay shallower samples. The lack of vertical control throughout the layers tended to result in contamination being estimated throughout the depth of the vertical layers to the RGA. This is illustrated in Table E.3.36 for TCE data at SWMU 4 (presenting the highest risk for all BGOU SWMUs) in which the maximum sample detection for the layers is generally much less than the maximum concentration predicted for a layer by SADA. In this case, the maximum concentration (i.e., 41 mg/kg) in layer 5 has been interpolated into layers 2, 3, 4, and 6

from layer 5. In general, SADA provides average TCE concentration in the layers that are greater in comparison to the average of the sample detections. Due to the lack of sample data points, the nearest neighbor interpolation tends to estimate large areas of contamination for which there are no data. For example, layer 2 contains two samples with detections; however, SADA predicts that 206 cells (20 ft by 20 ft grid cells) are contaminated. Therefore, SADA provides a maximum estimate of the total contamination and mass using the nearest neighbor interpolation method.

The SADA estimated uranium mass in relation to other metals (i.e., vanadium and manganese) appears to be underestimated. The mass of metals, such as vanadium and manganese also appear to be overestimated using SADA. The SADA interpolation estimates the mass between sample points. This results in an estimated mass of vanadium and manganese in the waste volume based on sample points located outside the waste zone, since these metals tend to be ubiquitous throughout the soils. Likewise, the sample points for uranium outside the waste zone are used to interpolate the mass in the waste zone. The transport of uranium from the waste into the surrounding soils is limited due to the sorption of this metal. Since the waste was not sampled, the uranium mass estimates for the waste areas in the SADA model likely are underestimated due to the limited migration of uranium. The uncertainty in the mass of uranium present in waste will be further addressed during remedial alternatives screening in the feasibility study.

Table E.3.36. Comparison of Sample Data with SADA Predicted Concentrations for TCE at SWMU 4

SADA Layer	Depth (ft)	Number of Detects and Total Samples	Detect Average (mg/kg)	Detect Range (mg/kg)	SADA Predicted Number of Contaminated Cells ^a	SADA Predicted Average (mg/kg)	SADA Predicted Cell Contaminant Range (mg/kg)
L1	0-1	0/21	0.00	0	0	0.00	0
L2	01-10	2/67	0.006	0.004 – 0.008	206	2.39	0.0036 – 41.0
L3	10-20	6/77	0.095	0.016 – 0.4	165	2.85	0.0036 – 41.0
L4	20-30	6/73	0.29	0.0064 – 0.82	181	3.02	0.0036 – 41.0
L5	30-40	10/29	4.83	0.011 – 41.0	240	2.56	0.0064 – 41.0
L6	40-50	11/29	2.14	0.012 – 9.2	246	2.45	0.0064 – 41.0
L7	50-63	13/39	4.44	0.02 – 25.0	249	3.15	0.0064 – 25.0

^a Cells for SWMU 4 were 20 ft by 20 ft.

E.3.3.2 SESOIL and AT123D Transport Uncertainties

As noted previously, due to SESOIL’s requirement to use the same constant area for each layer, the analyte concentrations of all layers needed to be normalized against the area of the layer with the maximum estimated analyte mass. The use of this methodology and uncertainty in the predictions were evaluated using TCE at SWMU 4 as an example and running each layer separately at its initial SADA concentration and area (i.e., no area and concentration normalization) and comparing these results to the original runs (i.e., normalized area and concentration). It should be noted that the results using this methodology will not match the total results presented previously for TCE at SWMU 4. By separating the layers, diffusion and volatilization gradients are different by individual layer compared to when all layers are modeled simultaneously in SESOIL; therefore, this is a comparison by layer and not by total mass in the system. The SADA data presented in Table E.3.37 were used in the analysis.

Table E.3.37. Summary of SADA Source Term Data for TCE at SWMU 4

SADA Layer	Depth (ft)	Total Mass (g)	Non-Normalized Average (mg/kg)	Non- Normalized Area (ft ²)	Normalized Average (mg/kg)	Normalized Area (ft ²)
L1	0-1	0.00E+00	0.00	0.00E+00	0.00	9.60E+04
L2	01-10	8.16E+04	2.39	8.24E+04	2.06	9.60E+04
L3	10-20	8.55E+04	2.85	6.60E+04	1.96	9.60E+04
L4	20-30	9.93E+04	3.02	7.24E+04	2.28	9.60E+04
L5	30-40	1.12E+05	2.56	9.60E+04	2.56	9.60E+04
L6	40-50	1.10E+05	2.45	9.84E+04	2.51	9.60E+04
L7	50-63	7.77E+04	3.15	9.96E+04	3.26	9.60E+04

The results of the analysis are presented in Figures E.3.6 through E.3.11. The results indicate that the normalization of the area and concentrations for input into SESOIL has an effect on the results but the differences are not considered significant when the total uncertainty of the sources from SADA are considered.

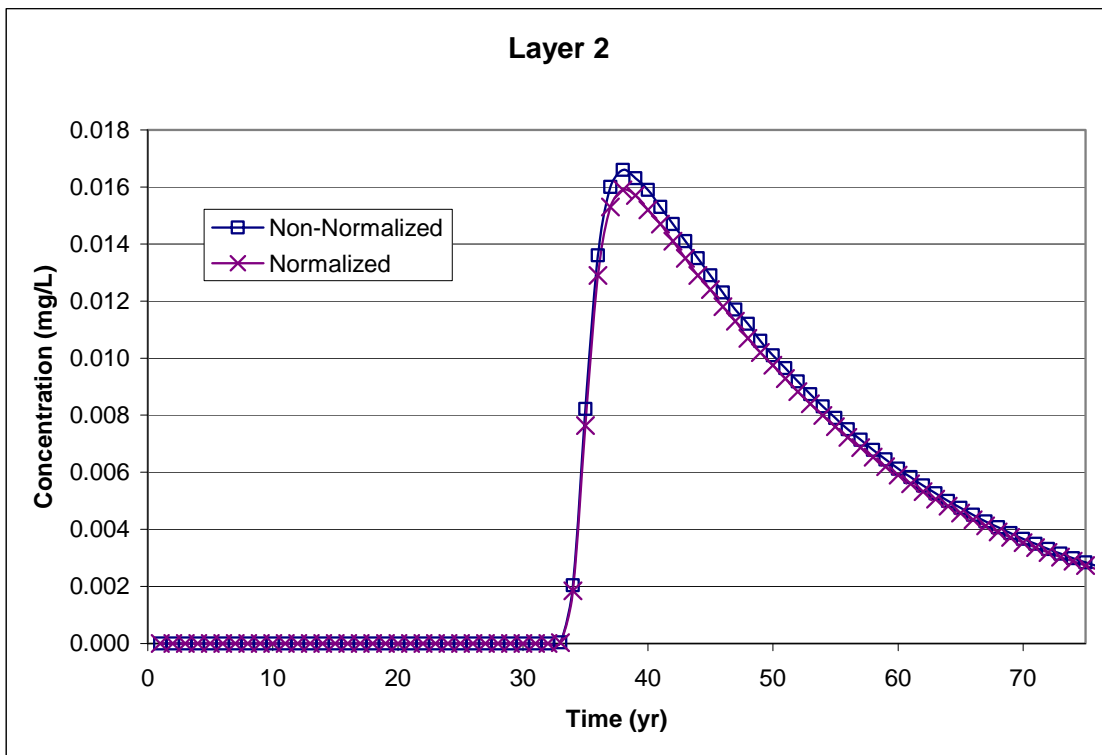


Figure E.3.6. Comparison of Predicted Groundwater Concentrations from a Layer 2 Source for TCE at SWMU 4 for the Normalized and Non-Normalized Area and Source Concentrations

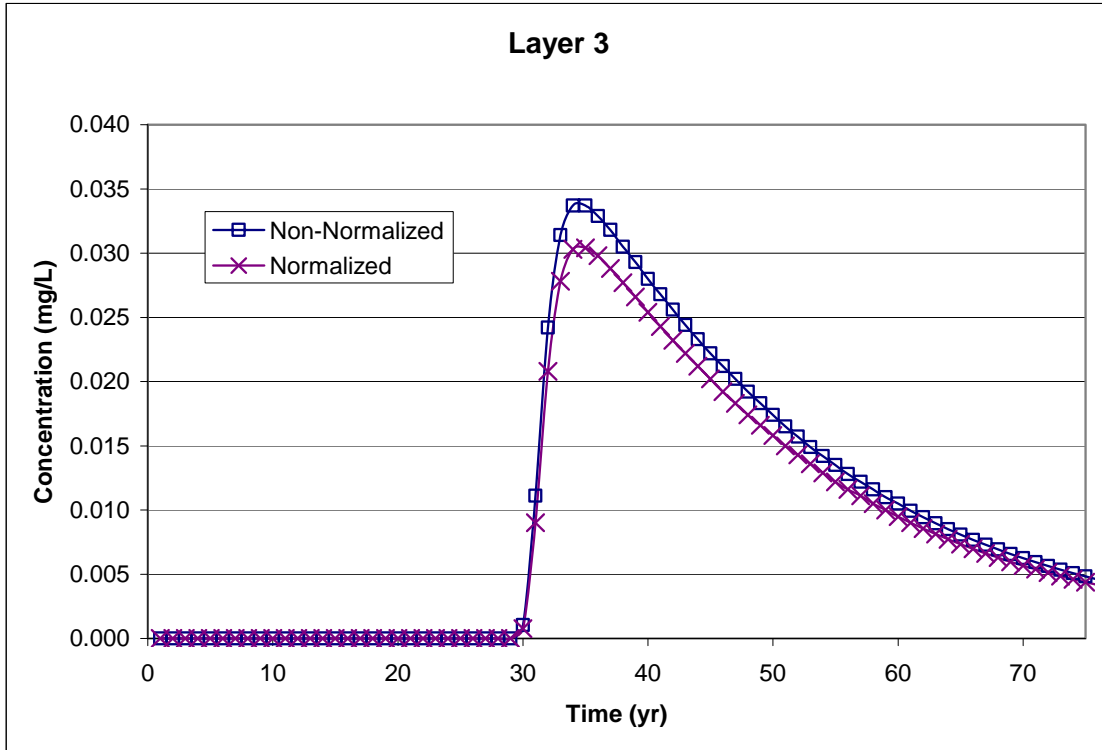


Figure E.3.7. Comparison of Predicted Groundwater Concentrations from a Layer 3 Source for TCE at SWMU 4 for the Normalized and Non-Normalized Area and Source Concentrations

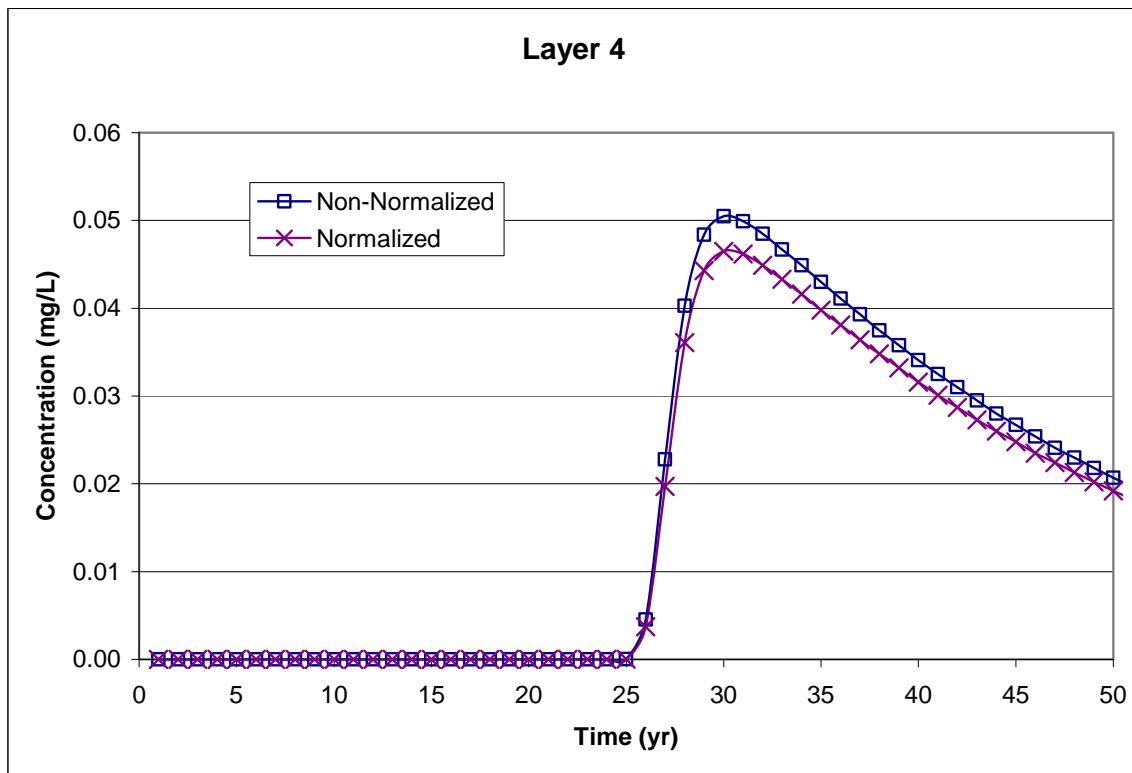


Figure E.3.8. Comparison of Predicted Groundwater Concentrations from a Layer 4 Source for TCE at SWMU 4 for the Normalized and Non-Normalized Area and Source Concentrations

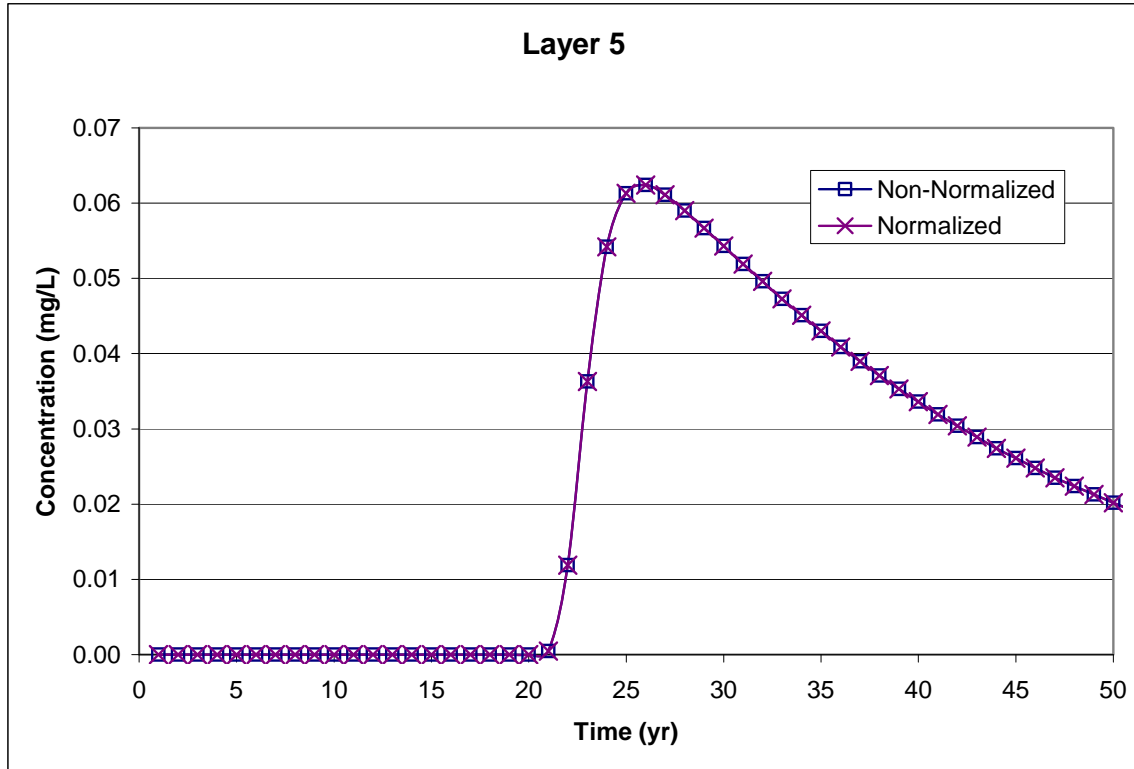


Figure E.3.9. Comparison of Predicted Groundwater Concentrations from a Layer 5 Source for TCE at SWMU 4 for the Normalized and Non-Normalized Area and Source Concentrations

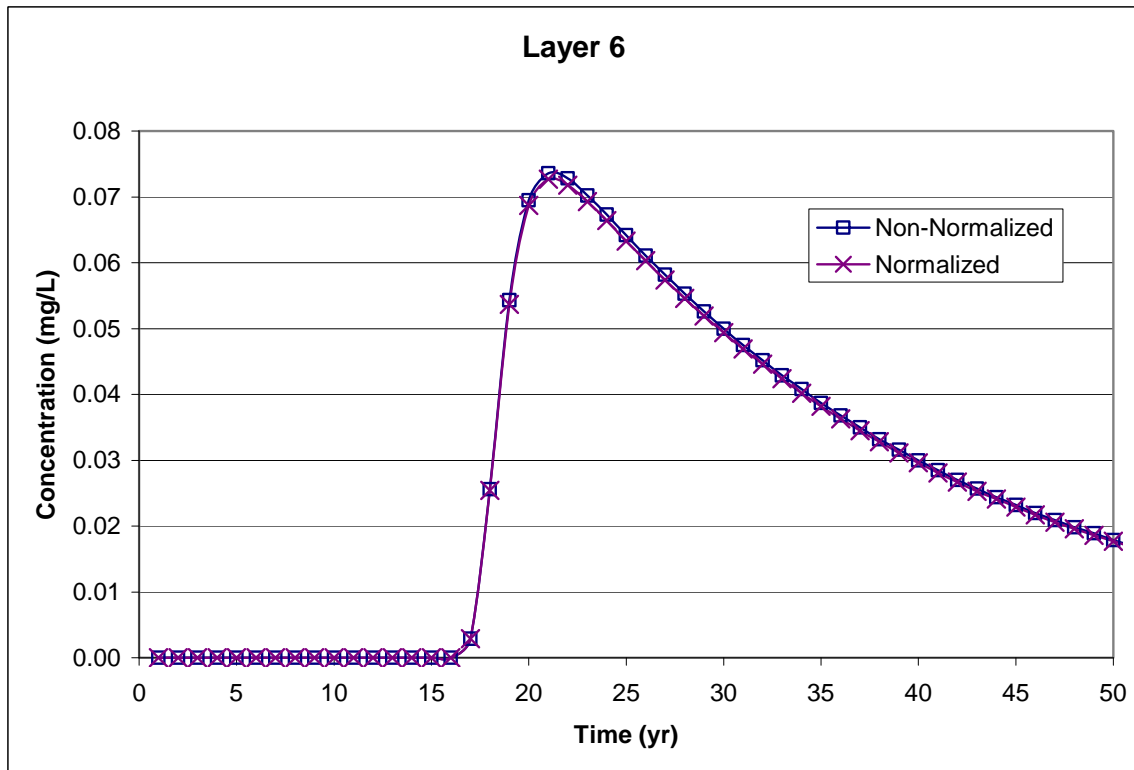


Figure E.3.10. Comparison of Predicted Groundwater Concentrations from a Layer 6 Source for TCE at SWMU 4 for the Normalized and Non-Normalized Area and Source Concentrations

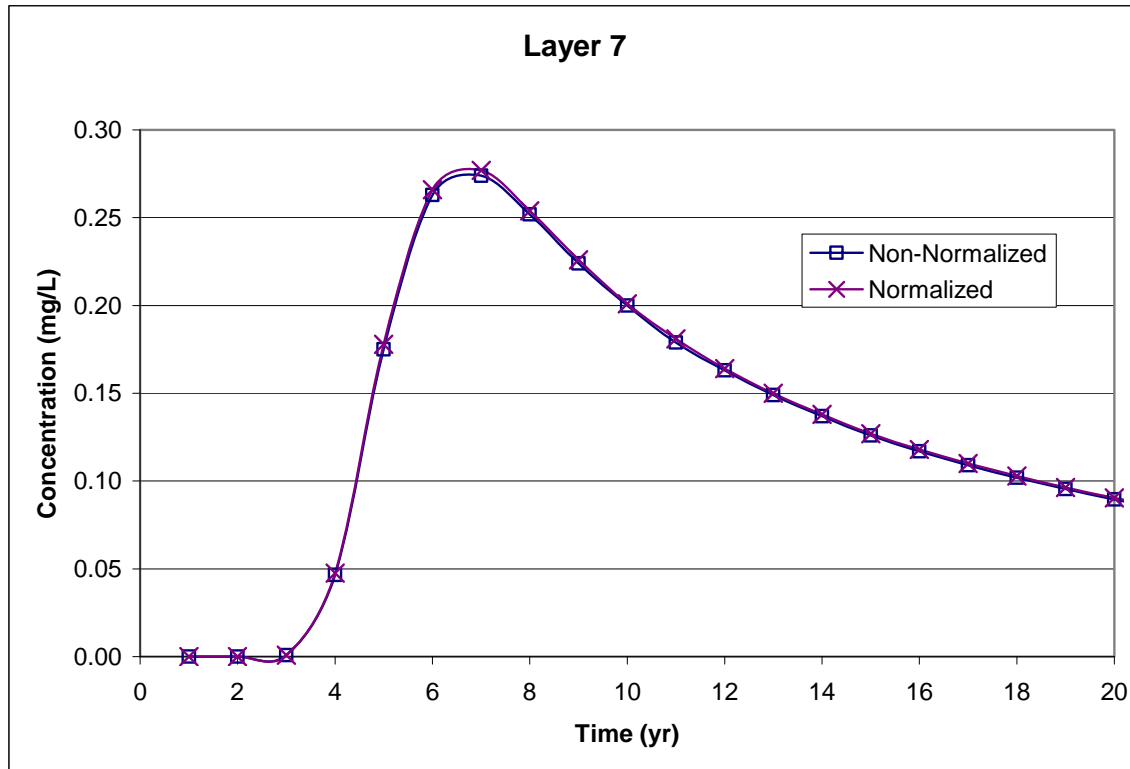


Figure E.3.11. Comparison of Predicted Groundwater Concentrations from a Layer 7 Source for TCE at SWMU 4 for the Normalized and Non-Normalized Area and Source Concentrations

An additional source of uncertainty in the AT123D modeling runs involves the use of a single hydraulic conductivity and hydraulic gradient. The hydraulic conductivity and gradient are variable from the SWMU locations to the various POEs. The MODPATH model was run to establish the steady-state head distribution in the RGA. MODPATH was used to track flowpaths of particles released from the SWMU location by using the steady-state, head distribution generated by MODFLOW. The distances from the SWMU to the POEs were taken along the flowpaths to determine the distance from the SWMU to the POEs. The hydraulic gradient from the SWMU to the property boundary was estimated using the head difference divided by the distance from the release point to the property boundary POE. The conductivity along the flowpath was also estimated for use in the AT123D model.

The selection of the sorption coefficient (K_d) for uranium in the UCRS was also evaluated for uncertainty. The sorption coefficient for sand, 66.8 L/kg was used in the analyses to provide conservatism and to account for uncertainty in the material properties directly below the SWMUs. However, site-specific measurements at PGDP have indicated that the uranium K_d in the UCRS may be much higher than used in the analyses. Therefore, the uncertainty in the uranium modeling results were evaluated using varying K_d values for uranium in the UCRS (i.e., 66.8, 133.6, 200.4, and 267.2 L/kg). The analysis was focused on SWMU 7 at the SWMU boundary and at the plant boundary. The results of the analysis for ^{238}U and ^{234}U are provided in Figures E.3.12 through E.3.15.

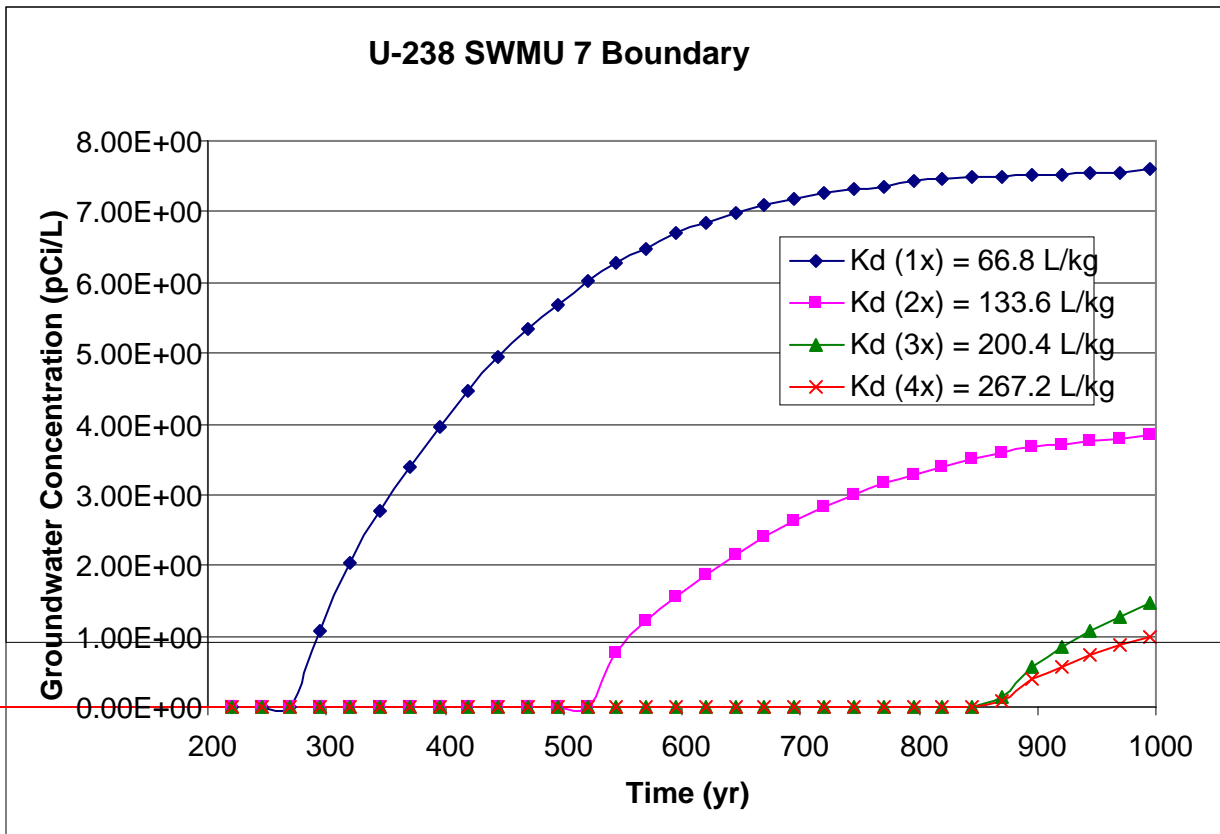


Figure E.3.12. SWMU 7 Groundwater Concentrations for ^{238}U at the SWMU Boundary for Varying UCRS K_d Values

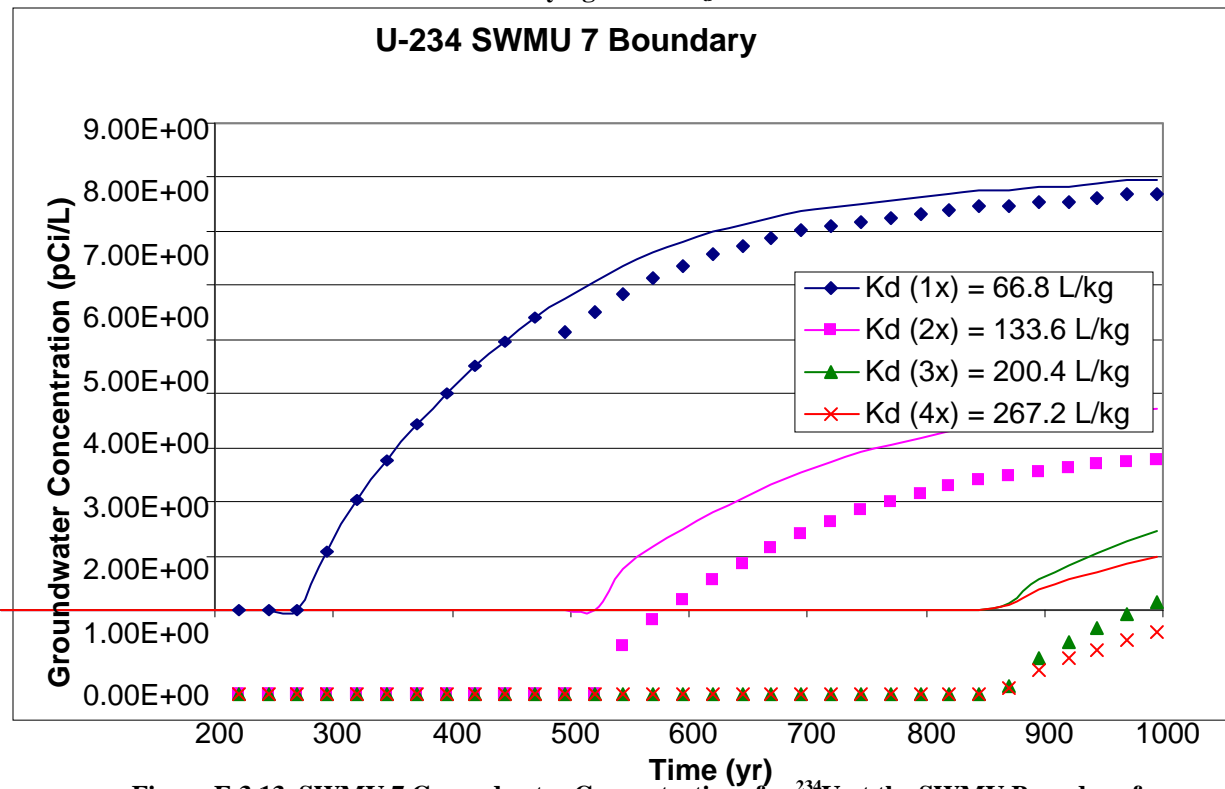


Figure E.3.13. SWMU 7 Groundwater Concentrations for ^{234}U at the SWMU Boundary for Varying UCRS K_d Values

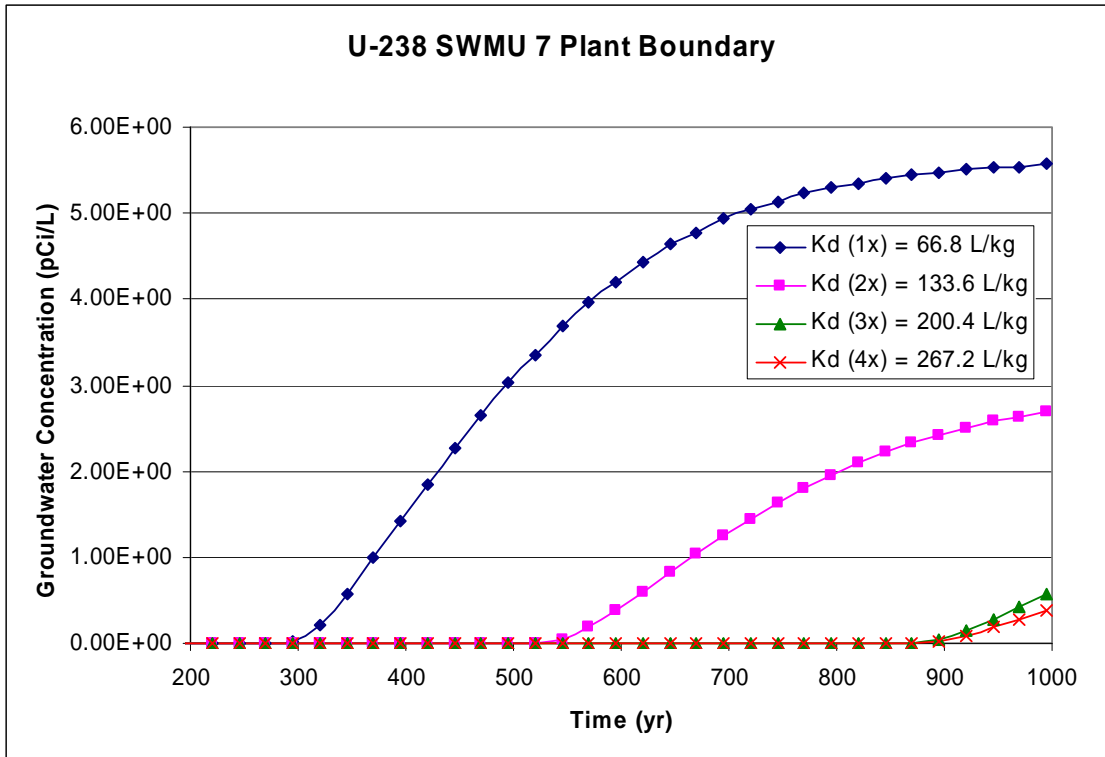


Figure E.3.14. SWMU 7 Groundwater Concentrations for ^{238}U at the Plant Boundary for Varying UCRS K_d Values

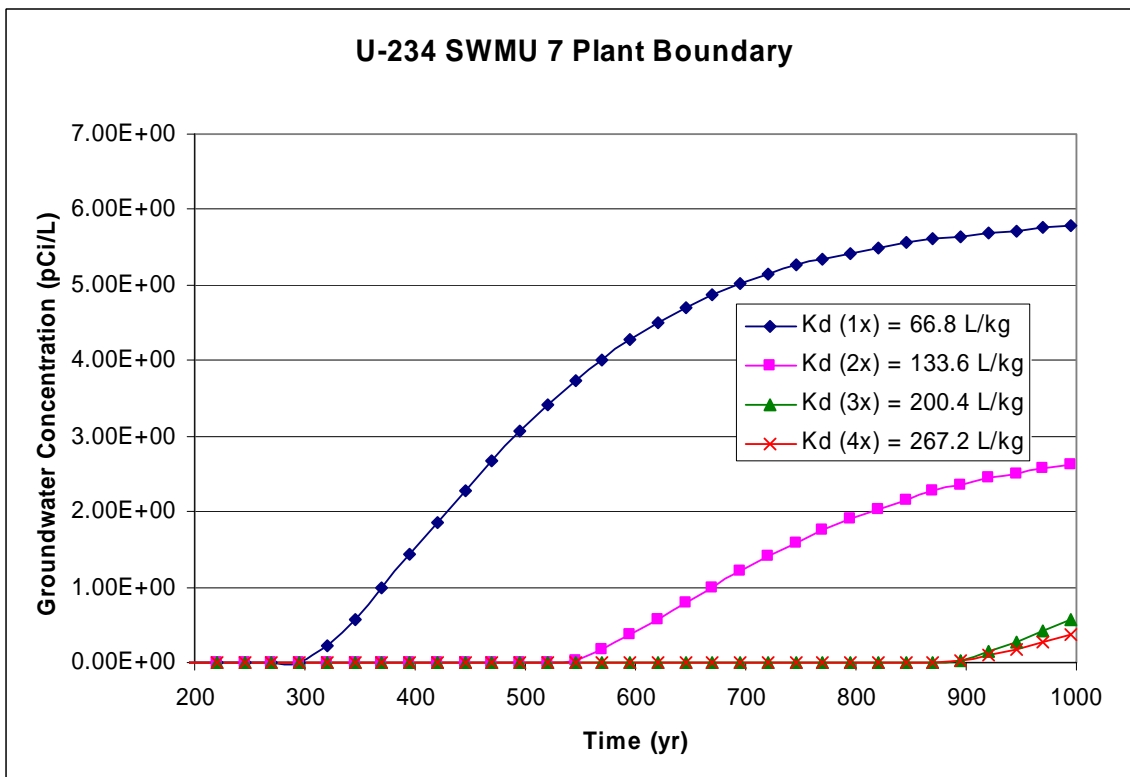


Figure E.3.15. SWMU 7 Groundwater Concentrations for ^{234}U at the Plant Boundary for Varying UCRS K_d Values

The analyses show that as the uranium K_d is increased, the arrival time of the contaminant at the POEs is shifted to later times. In addition, the maximum groundwater concentrations within the 1,000 year time period decrease. As the uranium K_d is increased beyond 4 times the original value of 66.8 L/kg, the contaminant no longer reaches the RGA water table within the 1,000 year analysis period. This analysis shows that the sand K_d of 66.8 L/kg used for the UCRS likely is low for PGDP, however, due to the lack of sample data in the waste areas, a conservative uranium K_d was chosen to estimate the risks and hazards from uranium at the SWMUs.

Another source of uncertainty in the fate and transport modeling involves earlier analyses of the SWMUs on the western side of the PGDP which indicated the presence of a water table in the UCRS that results in some waste being below the locally high water table. Site data indicate that at least some of the burial pits of the BGOU SWMUs are saturated, with the primary flow direction being down into the RGA. Insignificant horizontal flow is assumed to occur above the RGA. The modeling assumed that the soil zones above the RGA are unsaturated, with contaminants being transported vertically downward into the RGA. The assumption that these zones are unsaturated in the model may have resulted in overestimation of contaminant migration from the various sources to the RGA for SWMUs below the UCRS water table. Overestimation is the result of the interaction between layers with low vertical hydraulic conductivity (i.e., HU2 Confining and HU3) and the shallow water table. Generally, this interaction results in contaminant concentrations in pore water within each layer approaching equilibrium with soil prior to migration because the rate of migration is very slow. This phenomenon ultimately would result in rates of contaminant migration (i.e., flux) that are less than that which would result from the introduction (i.e., infiltration) of “clean” water from precipitation through an unsaturated layer.

An additional uncertainty involves the fact that SESOIL and AT123D do not consider contaminant transformation such as that for radioactive decay chain ingrowth of progeny. An analysis was conducted to evaluate the potential impact of progeny ingrowth from ^{238}U and ^{234}U at SWMU 7. To evaluate the movement of progeny, a simplified assumption was made that radioactive progeny travel at the same rate of the parent. This assumption has been shown to be conservative (Codell *et al.*, 1982) and greatly simplifies the calculations. The assumption was also made that no progeny exist at the time of waste emplacement. The concentration of the i^{th} progeny in a decay chain at the receptor location is then calculated by:

$$C_i = C_{\text{parent}} \frac{DIF_i \times R_{d \text{ parent}}}{DIF_{\text{parent}} \times R_{d i}}$$

Where

DIF_i = decay ingrowth factor of the i^{th} progeny

DIF_{parent} = decay-ingrowth factor of the parent

$R_{d i}$ = retardation factor of the parent

$R_{d \text{ parent}}$ = retardation factor of the parent

C_{parent} = groundwater concentration of the parent (pCi/L)

The sorption coefficients for sand were used in the analysis for uranium (66.8 L/kg), thorium (3200 L/kg) and radium (500 L/kg) (Sheppard and Thibault 1990).

The decay-ingrowth factor for an n member decay chain is given by Scrable *et al.*, (1974):

$$DIF_i(t) = \frac{\lambda_i}{\lambda_1} \left[\left(\prod_{i=1}^{n-1} \lambda_i \right) \sum_{i=1}^n \frac{e^{-\lambda t}}{\prod_{j=1, j \neq i}^n (\lambda_j - \lambda_i)} \right]$$

Where

λ_1 = decay constant for the parent (yr^{-1})

λ_i = decay constant for the i th progeny (yr^{-1})

t = time (years)

The results of the analysis for ^{238}U and ^{234}U at SWMU 7 are provided in Figure E.3.16 and E.3.17.

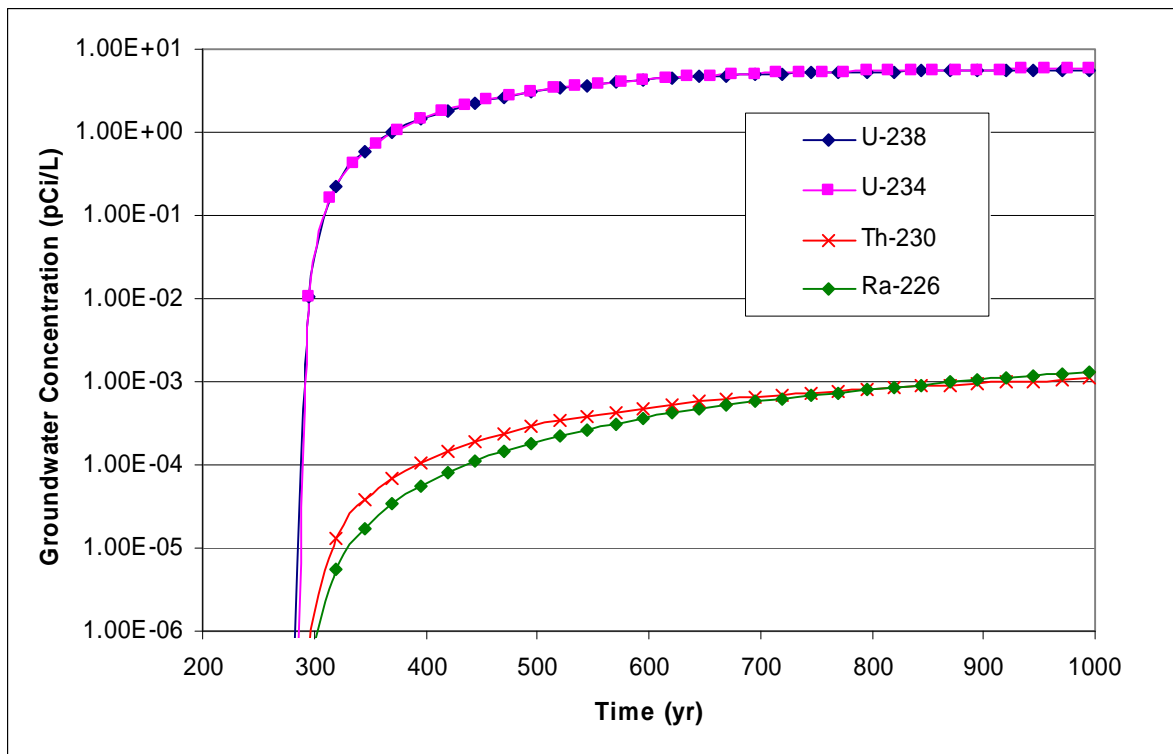


Figure E.3.16. SWMU 7 Groundwater Concentrations from Progeny Ingrowth from ^{238}U and ^{234}U at the Plant Boundary

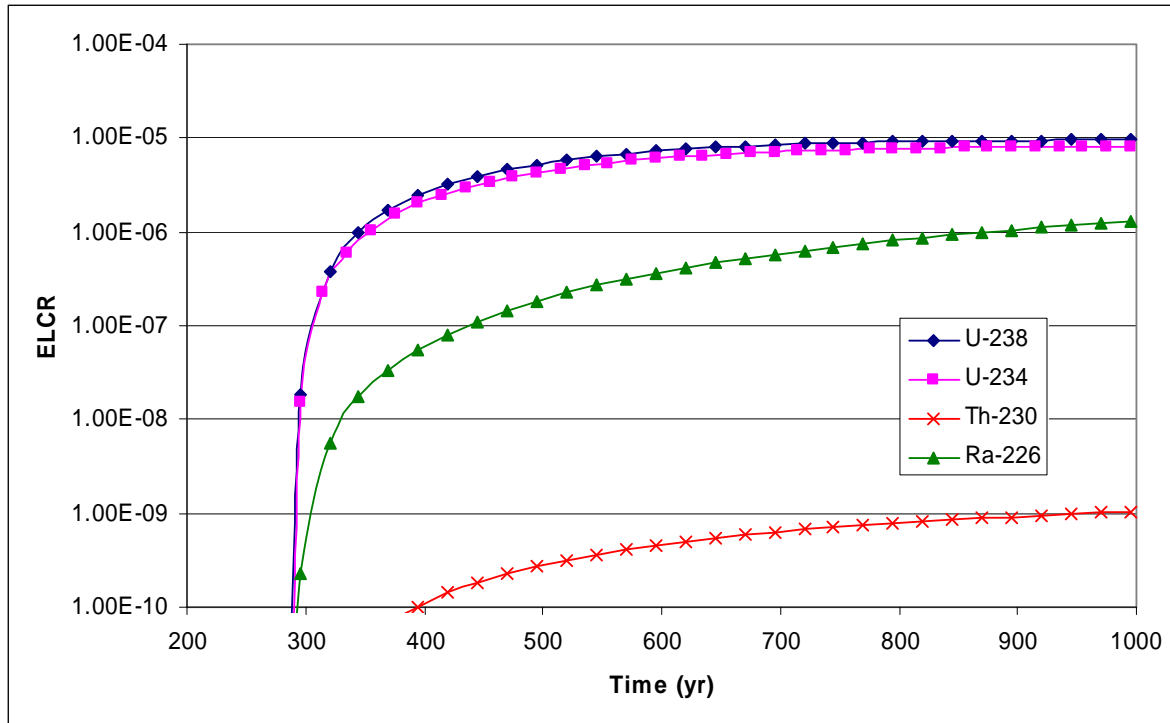


Figure E.3.17. SWMU 7 Groundwater ELCR from Progeny Ingrowth from ²³⁸U and ²³⁴U at the Plant Boundary

This analysis shows that the ingrowth of ²²⁶Ra would provide an additional 7 % to the ELCR to that estimated for ²³⁸U and ²³⁴U at SWMU 7. However, this contribution is considered a conservative estimate since the progeny were assumed to transport with the uranium parents. In reality, the higher sorption coefficients for ²³⁰Th and ²²⁶Ra in comparison to uranium would result in differential transport such that the predicted concentrations of these progeny would be less than that provided by this simplified analysis.

It is also understood that the ingrowth of progeny from uranium isotopes will, in many cases, continue to increase beyond the 1,000 year modeling period and that risk will continue to increase. The uncertainty related to this potential increase in risk will be managed in the FS and in future decision and design documents.

E.3.3.3 Potential Interaction of Sources

The simulations presented in this report for the BGOU SWMUs are based on individual simulations of each SWMU. There is a potential that source plumes from the SWMUs could interact at the POEs. According to the flow paths presented in Figure E.3.18, the contaminant plumes from a few of the BGOU SWMUs would interact. The contaminant flow paths from SWMU 6 and SWMU 30 will interact, however, as noted previously, SWMU 6 did not have any groundwater analytes. The contaminant plumes from SWMU 3 and SWMU 5 will interact, and SWMU 2 will interact with a portion of the SWMU 5 contaminant plume. The interaction of the plumes could not be assessed using the SESOIL/AT123D model, since only one SWMU can be discretized in the model for each run.

An evaluation was conducted to ensure that analytes were not eliminated from the groundwater analyses from combined source contributions in the groundwater. The screening evaluation was conducted for the

potential interaction of SWMU 2, 3, and 5. The analysis was based on the conservative summation of the maximum groundwater concentrations below each SWMU detailed in Section 5, Tables 5.5, 5.6, and 5.8. These combined contaminant concentrations are provided in Table E.3.38. These combined groundwater concentrations then were compared to the child resident NALs from Table A.18 of the 2001 Risk Methods Document and the provisional groundwater backgrounds shown in the 2001 Risk Method Document in Table A.13. The results of the conservative analysis indicate that the selection of analytes evaluated for risk and hazard would not change based on the combination of the source contributions to groundwater from SWMU 2, 3, and 5.

E.3.3.4 Location of the POEs

The POEs used in the modeling were placed at locations on the SWMU boundary, plant boundary, property boundary, Little Bayou seeps, and Ohio River where the greatest contaminant concentrations are expected in the future. By picking locations on the centerline of predicted contaminant plumes as the POEs, the modeling assumed that the hypothetical future resident would pick, by chance, the worst possible location to install a water supply well.

Based on particle tracks taken from the calibrated sitewide numerical flow model developed in MODFLOW for PGDP, SWMUs 2, 4, 5, and 145 were shown not to impact the Little Bayou seeps. If the SWMUs were to impact the seeps, it has been shown that SWMUs 2, 4, and 145 have modeled groundwater concentrations at the Ohio River that exceed MCLs for several analytes; therefore, the modeled groundwater concentrations at the Little Bayou seeps also would exceed the MCLs for these analytes. Since there is uncertainty with the flow paths and the seeps' radius of influence, the contaminants in SWMU 4 that exceed the MCL at the property boundary POE were evaluated to see if they would exceed the MCL at the Little Bayou seeps. The modeling results indicated that only one contaminant, TCE, exceeded the MCL at the Little Bayou seeps (i.e., 0.083 mg/L). TCE also exceeds the MCL at the Ohio River (i.e., 0.077 mg/L) for SWMU 4. Modeling results for SWMU 5 show that the groundwater concentrations at the property boundary do not exceed the MCLs for any analytes modeled; therefore, the groundwater concentrations at the Little Bayou seeps also would be less than the MCLs for each analyte.

Table E.3.38. Evaluation of Combined Plume Interactions on Selection of Analytes

Analyte	Background Groundwater Concentration (mg/L) ^a	Groundwater Child Resident No Action Level (mg/L) ^a	Peak Conc Below SWMU 2 (mg/L) ^a	Peak Conc Below SWMU 3 (mg/L) ^a	Peak Conc Below SWMU 5 (mg/L) ^a	SUM of Peak Conc Below SWMUs (mg/L) ^a	Retain as Analyte			
							SWMU 2	SWMU 3	SWMU 5	SUM SWMUs
Acenaphthene	NA	1.36E-02			6.10E-03	6.10E-03			N	N
Anthracene	NA	7.66E-02			8.06E-03	8.06E-03			N	N
Arsenic	5.00E-03	3.50E-05	3.54E-02	3.29E-02	9.25E-03	7.76E-02	Y	Y	Y	Y
²³⁴ U	7.00E-01	5.46E-01	1.58E+00			1.58E+00	Y			Y
²³⁸ U	7.00E-01	4.43E-01	1.81E+00	1.59E+01		1.77E+01	Y	Y		Y
⁹⁹ Tc	2.23E+01	1.40E+01	1.02E+02	5.56E+03	1.27E+02	5.79E+03	Y	Y	Y	Y
<i>cis</i> -1,2-DCE	NA	2.73E-03	1.15E+01			1.15E+01	Y			Y
Manganese	1.19E-01	3.50E-02	7.16E-01	8.95E-01	1.01E+00	2.62E+00	Y	Y	Y	Y
Mercury	2.00E-04	4.44E-04		9.29E-05		9.29E-05			N	N
Fluorene	NA	9.72E-03			3.63E-03	3.63E-03			N	N
Naphthalene	NA	2.85E-04	9.38E-04		5.55E-03	6.49E-03	Y		Y	Y
Nickel	3.05E-01	3.01E-02			2.01E-03	2.01E-03			N	N
Selenium	5.00E-03	7.54E-03			1.27E-03	1.27E-03			N	N
TCE	NA	1.60E-03	1.48E+00	3.45E-04	9.91E-04	1.48E+00	Y	N	N	Y
Uranium	2.00E-03	9.06E-04	9.86E-03	4.89E-02		5.88E-02	Y	Y		Y
Zinc	4.90E-02	4.50E-01	9.83E-03	9.30E-02	1.58E-01	2.61E-01	N	N	N	N

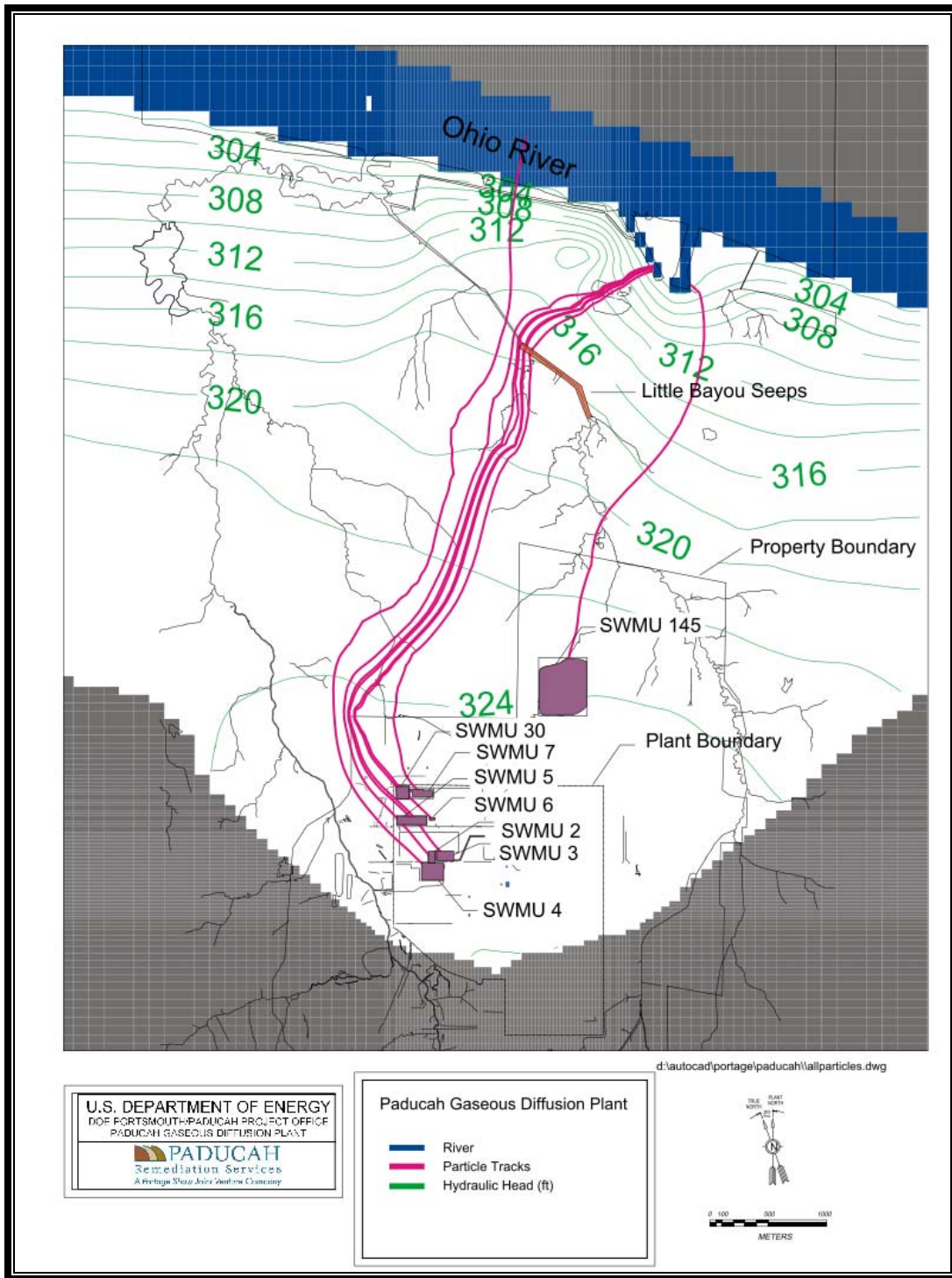


Figure E.3.18. Contaminant Plume Flow-Paths for All BGOU SWMUs

E.3.3.5 Future Environmental Changes

Several future environmental changes at the PGDP could impact the accuracy of the modeling predictions. These changes include plant shutdown and dam operation on the Ohio River. In a previous modeling effort for a landfill at PGDP, several sensitivity analyses were performed (DOE 2003b) to examine the impacts those changes may have on groundwater flow and contaminant transport. It was assumed in that sensitivity analysis that it can be expected that plant shutdown will lead to a changed recharge rate to the RGA through removal of ground cover (leading to increased recharge) and through reduced cooling water use (leading to decreased recharge); therefore, the sensitivity analysis of the groundwater travel time due to plant shutdown was studied by varying the recharge over a range of values. The results of the analysis indicated that a decrease in the recharge rate resulted in a monotonic increase in the travel time to the receptor. Thus, chemicals that have short degradation half-lives would show a decrease in concentration due to plant shutdown.

The Olmstead Dam operation is expected to increase the stage (water level) of the Ohio River; therefore, a sensitivity analysis was conducted (DOE 2003b) to assess changes in groundwater travel time in relation to dam operation, by increasing the river stage between 304.44 ft amsl and 310.04 ft amsl (the baseline river stage is 300.04 ft amsl). The results of the analysis indicated that the travel times in the aquifer changed very little in relation to the Ohio River stage; therefore, the dam operation would have little impact on the results shown in this report.

E.3.3.6 Burial Cell Waste

Sample data around and beneath the BGOU SWMUs were used to develop a source inventory of contaminants. The premise of this source inventory development is based on the inherent assumption that the contaminants around and beneath the BGOU SWMUs represent the release mass from the Burial Ground disposal cells. The groundwater transport analyses do not model potential future releases directly from the SWMU burial cells.

Waste at several SWMUs was containerized in drums before disposal. Previous inspections of buried drums at PGDP have indicated that the drums were highly corroded. It is considered unlikely that a significant portion of the drummed waste still is containerized at the BGOU SWMUs due to the length of time the drums have been buried and, thus, susceptible to a corrosive environment. The drums were not modeled in this RI report due to the overall objectives of the RI analyses and uncertainty in the degradation process. Due to the uncertainty in the degradation of the drummed waste, real measured sample data surrounding the SWMUs were used to evaluate the potential risk from the SWMU waste. This methodology resulted in the SWMUs with drummed waste exhibiting risk and hazard values that exceeded acceptable levels; therefore, the overall objectives of the RI analysis were met without requiring a detailed analysis of the degradation of drums.

E.3.3.7 SWMU 4 RGA TCE Source

The TCE source in SWMU 4 was assessed in this RI based on soil sample results. As discussed in Section 4.5.2, a discrete DNAPL zone, less than 200 ft wide, also may be present at the base of the RGA as evidenced by a discrete area with TCE concentrations greater than 10,000 µg/L in the lower RGA immediately downgradient of the SWMU. The volume of soil contaminated with TCE DNAPL at this SWMU is estimated to be approximately 31,480 yd³. This estimate assumes a source area that is 100 ft by 100 ft with a thickness of 85 ft (depth to base of RGA, which is 100 ft minus the estimated depth to base of the waste cell of 15 ft.) The DNAPL source term for TCE in the RGA at SWMU 4 was not evaluated in the modeling analyses since the RGA concentrations in the lower RGA currently exceed the MCL. The UCRS TCE source concentrations were sufficient to indicate that actions should be taken for SWMU 4.

The uncertainties related to source term size and location will be better defined in a remedial design investigation prior to the design and implementation of a remedy.

E.3.3.8 SWMU 3 UCRS Groundwater Contamination

The groundwater analyses conducted for this RI are based on soil samples obtained from soils surrounding the SWMUs and their subsequent release to the RGA and transport through the RGA. In some instances, water samples from wells in the UCRS indicated additional contaminant concentrations that were not accounted for in the analyses. For example, UCRS wells MW85, MW88, MW91, and MW94 at SWMU 3 indicated elevated levels of TCE. Figures E.3.19 through E.3.22 show the TCE concentrations trends for these wells in the UCRS.

The water data was added to the SWMU 3 TCE soil concentrations and a SADA nearest neighbor interpolation was assessed. The results of the SADA analysis are presented in Table E.3.39.

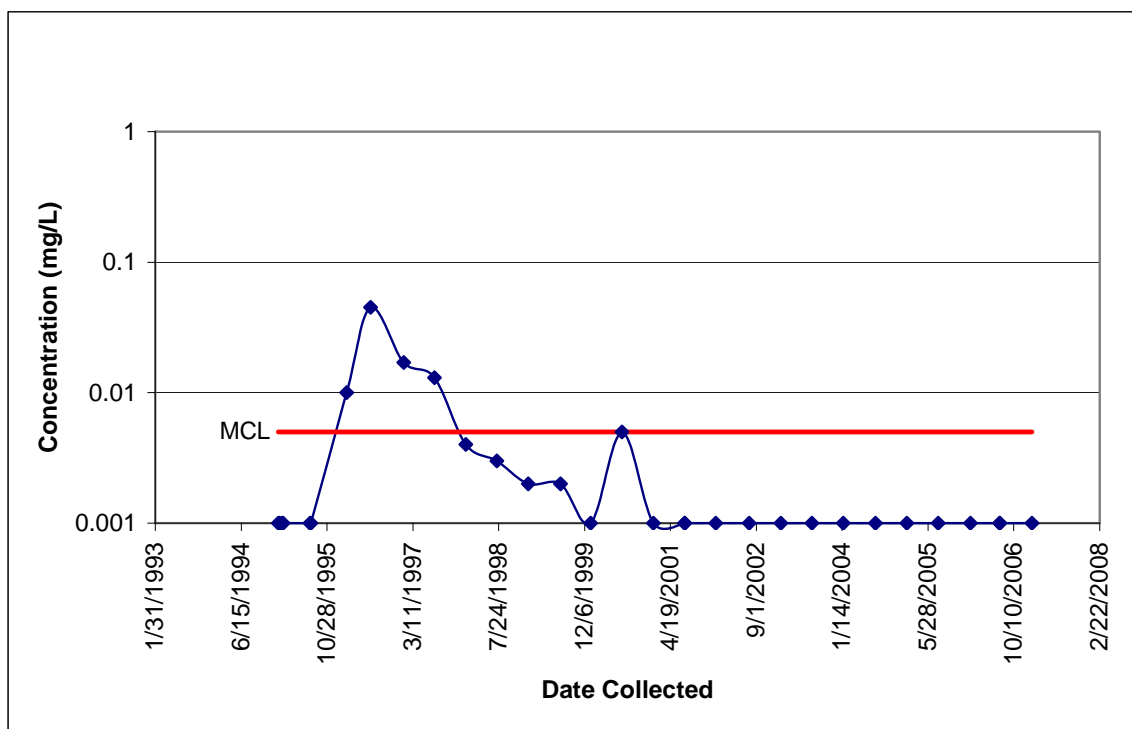


Figure E.3.19. UCRS TCE Contaminant Trend for SWMU 3 Well MW85

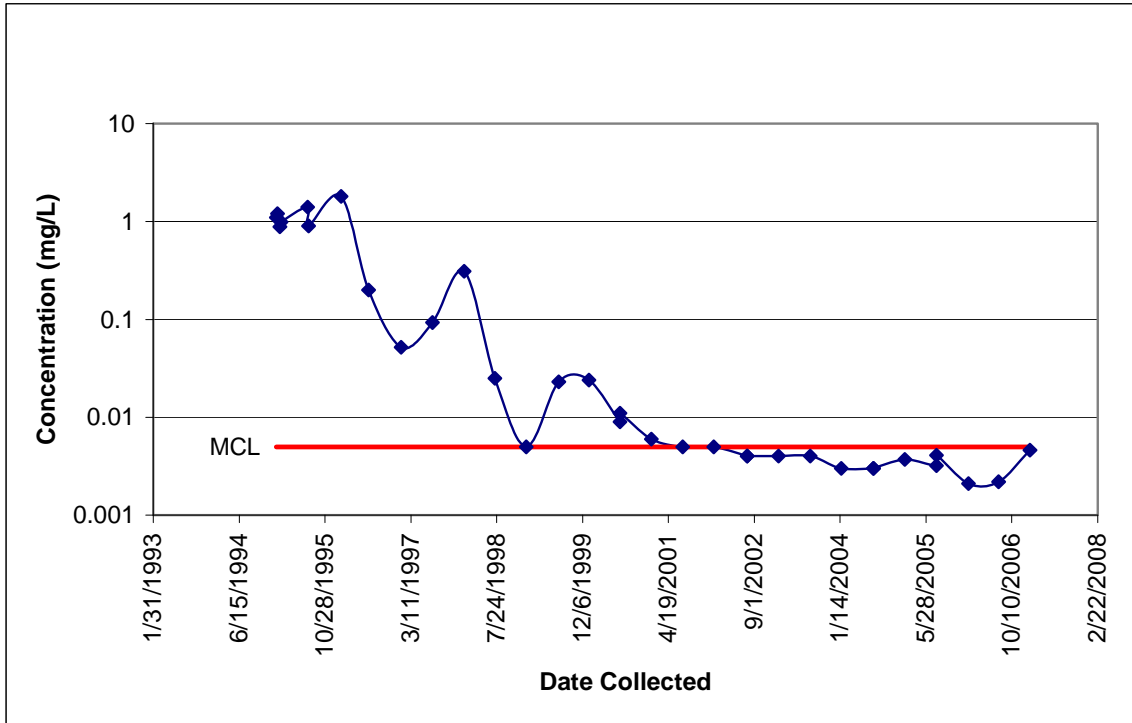


Figure E.3.20. UCRS TCE Contaminant Trend for SWMU 3 Well MW88

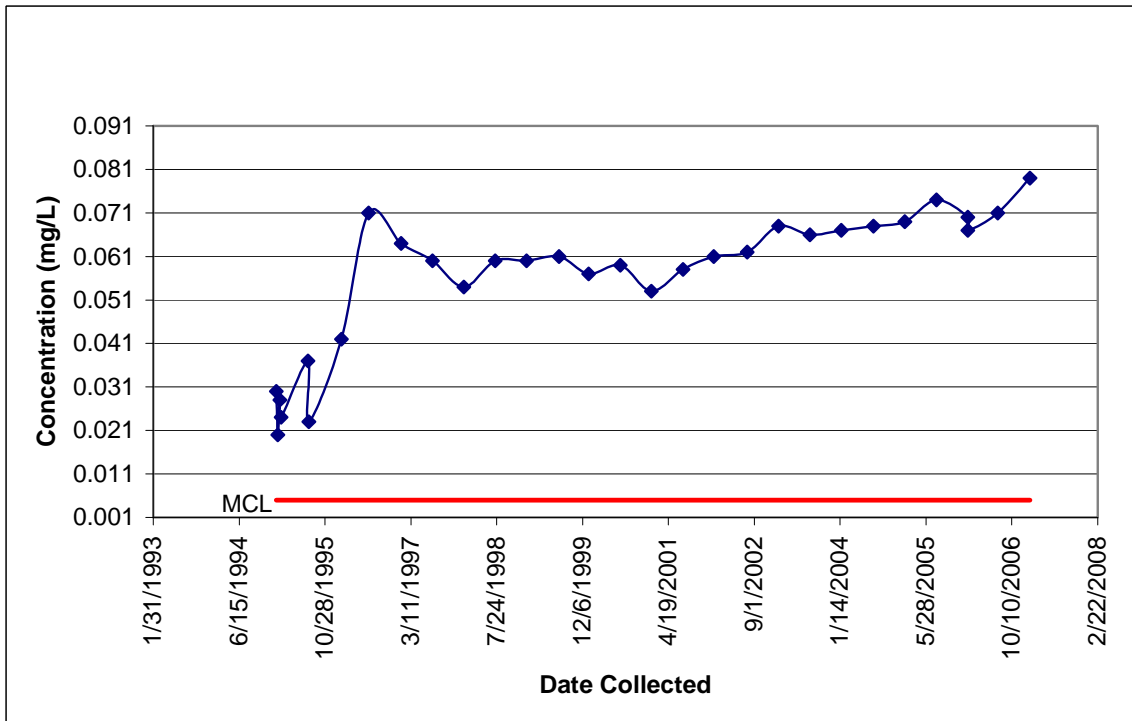


Figure E.3.21. UCRS TCE Contaminant Trend for SWMU 3 Well MW91

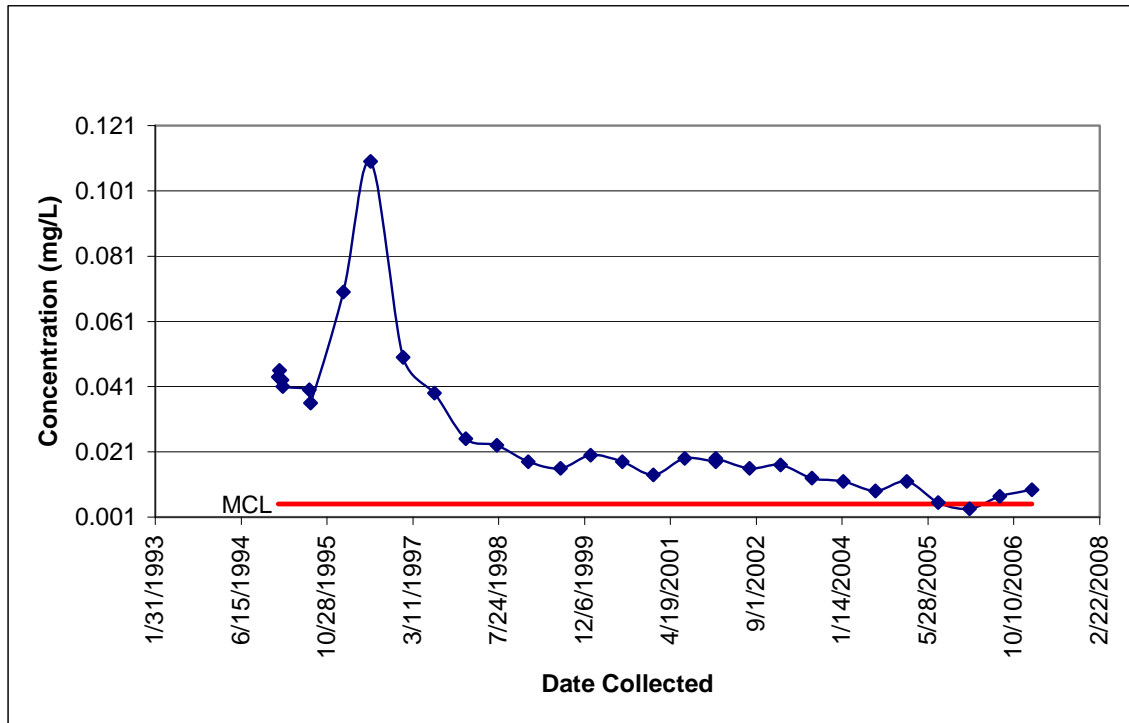


Figure E.3.22. UCRS TCE Contaminant Trend for SWMU 3 Well MW94

Table E.3.39. SWMU 3 TCE SADA Results With Well Data

Layer #	Depth	Min mg/kg	Max mg/kg	Sum mg/kg	Count #	Avg mg/kg	Area ft ²	Volume ft ³	Mass gm
	Interval ft-ft bgs								
L0	0-1	0.01	0.01	0.02	3	0.0063	3.00E+04	3.00E+04	7.85E+00
L1	01-10	0.00	0.03	0.32	21	0.0155	2.10E+05	2.10E+06	1.34E+03
L2	10-20	0.00	0.03	0.33	22	0.0148	2.20E+05	2.42E+06	1.49E+03
L3	20-30	0.00	0.03	0.33	22	0.0148	2.20E+05	2.42E+06	1.49E+03
L4	30-40	0.00	0.03	0.33	23	0.0143	2.30E+05	2.53E+06	1.49E+03
L5	40-50	0.00	0.03	0.33	23	0.0143	2.30E+05	2.53E+06	1.49E+03
L6	50-58	0.00	0.03	0.33	23	0.0143	2.30E+05	2.30E+06	1.36E+03
Total								8.67E+03	

The groundwater analysis was conducted using the SADA results for both the UCRS TCE water data and the SWMU TCE soil data. The results of the analysis are provided in Figure E.3.23.

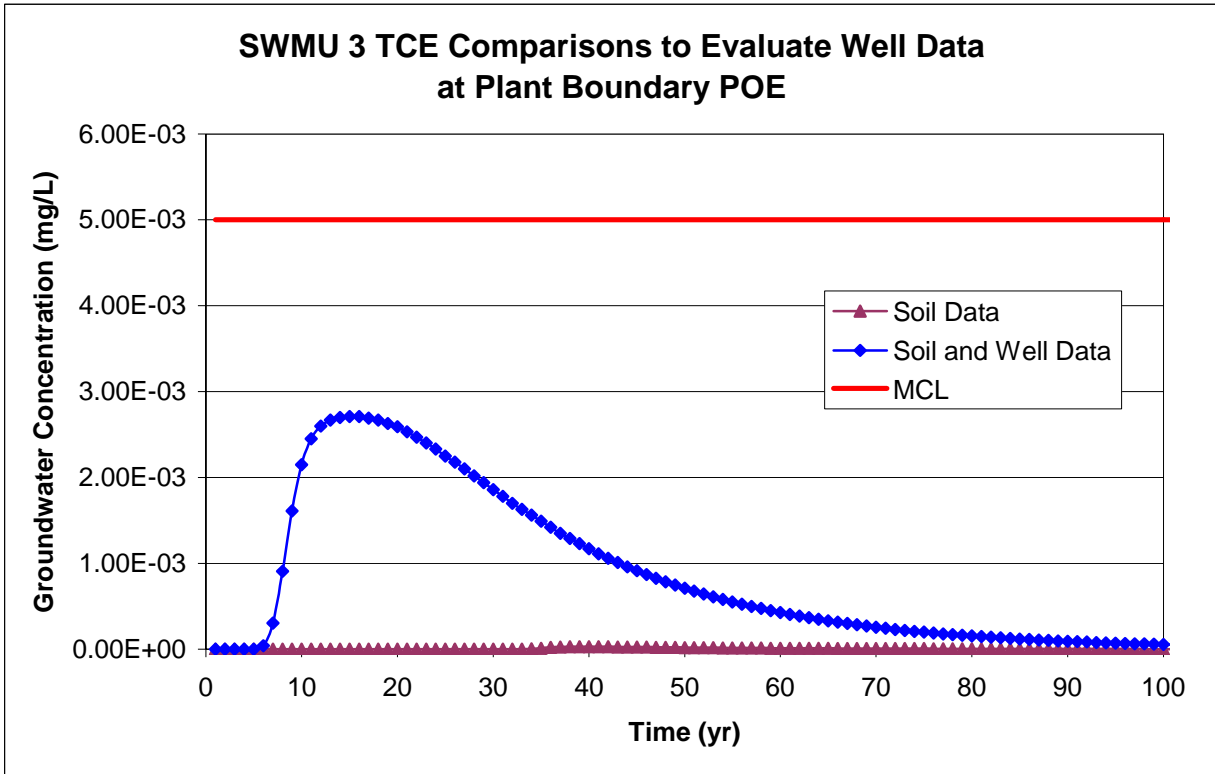


Figure E.3.23. Comparison of TCE at SWMU 3 with UCRS Well Sample Data and Soil Data

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**APPENDIX E
ATTACHMENT 1**

**PREVIOUS FATE AND TRANSPORT MODELING
FOR THE BGOU**

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E1.1. INTRODUCTION

This attachment presents more detailed summaries and excerpts of previous fate and transport modeling for the Burial Ground Operable Unit SWMUs 2, 3, 4, 5, 6, 7, and 30. No previous modeling exists for SWMU 145.

E1.2. MODELING APPEARING IN THE WASTE AREA GROUP (WAG) 22, SWMUS 2 AND 3 FEASIBILITY STUDY (FS) AND ADDENDUMS, AND THE SWMU 2 DATA SUMMARY AND INTERPRETATION REPORT

This section describes previous groundwater modeling discussed in the WAG 22 SWMU 2 and 3 FS (DOE 1995a) and in the data summary and interpretation report for the SWMU 2 interim remedial design (DOE 1997).

E1.2.1 WAG 22 SWMU 2 AND 3 FS MODELING

The following is taken from the *Feasibility Study for Solid Waste Units 2 and 3 of Waste Area Group 22 at the Paducah Gaseous Diffusion Plant, Paducah, Kentucky*, February 1995 (DOE 1995a).

GeoTrans (1992) conducted a modeling study of the hydrogeologic flow system beneath the PGDP to evaluate the feasibility and effectiveness of a proposed “pump-and-contain” groundwater extraction system. As part of this study, GeoTrans updated an existing model (GeoTrans 1990) by incorporating significant improvements in the hydrogeological characterization of the site. GeoTrans implemented the groundwater flow model using MODFLOW, the U. S. Geological Survey groundwater simulation code (McDonald 1988). The finite-difference grid for the GeoTrans model covered an area of approximately 60 km² (23 mi²) and consisted of 91 columns, 117 rows and three layers. Layer 1 of the model represented the (UCRS); Model Layer 2 represented the RGA; and Model Layer 3 represented the McNairy Formation.

GeoTrans (1992) model used water levels from August 15, 1991, which were determined during data review to be representative of steady-state conditions, to calibrate the model. The model extended from the Ohio River in the north (simulated as a constant-head boundary) to the Porters Creek Clay terrace transition area in the south (simulated as a no-flow boundary). Constant heads simulated the observed potentiometric surface in the McNairy Formation; thus, the McNairy Formation was an infinite sink or source in the model depending on the hydraulic heads simulated in the RGA.

GeoTrans (1992) calibrated the model by matching water levels observed in the UCRS and RGA. Overall, the GeoTrans model simulates hydraulic heads more closely in the RGA. In the UCRS, the model generally mismatched hydraulic heads on the order of ± 1.5 to 3 m (± 5 to 10 ft). The apparent reason for the poor fit in the UCRS most likely was due to a lack of vertical discretization and the oversimplification of the hydraulic conductivity distribution in the UCRS.

McConnell (1992a; 1992b; 1993) has developed a number of groundwater flow models to study the shallow groundwater system at the PGDP site. McConnell (1993) developed a groundwater flow model to estimate the average vertical hydraulic conductivity of Hydrogeologic Unit (HU) 3, the hydraulic

properties of the McNairy Formation, and the connection between the McNairy Formation and the Ohio River.

McConnell's model results suggested that the RGA is in direct communication with the Ohio River, and during high river stage conditions, groundwater is driven down into the McNairy Formation.

Additional models developed by McConnell simulated groundwater flow conditions at the C-747-A Burial Ground Area (SWMUs 7 and 30) (McConnell 1992a) and the C-404 Landfill Area (SWMU 4) (McConnell 1992b). The local groundwater flow model developed for the C-747-A site evaluated the effectiveness of using an impermeable cap with and without a slurry trench partially surrounding the facility to dewater the UCRS immediately beneath the landfill. The model simulated only the UCRS and RGA HUs; a no-flow boundary represented the McNairy Formation in the model. Constant heads simulated a prescribed hydraulic gradient in the RGA and no-flow boundaries around the edges of the model in the overlying units precluded the possibility of horizontal groundwater flow in the UCRS. Model results suggested that the slurry trench in combination with the cap would be required to dewater the UCRS beneath the site.

The model of the C-404 Landfill Area (SWMU 4) (McConnell 1992b) was developed to improve the conceptual model of the groundwater flow system in the C-404 area, estimate hydraulic parameters, and assess the hydraulic effect of installing an impermeable cap over the C-404 area. This model consisted of eight model layers covering an area of 7,432 m² (80,000 ft²) with minimum grid cell sizes of 15 by 15 m (50 by 50 ft). This model simulated the upper 6.1 m (20 ft) of the McNairy Formation; otherwise, boundary conditions were similar to the C-747-A model (McConnell 1992a). Due to the small size of the model and specified constant-head boundaries, the hydraulic conductivity of the RGA unit was insensitive. Model simulation results suggested that pumping in the RGA may induce leakage from the McNairy Formation, an effect inferred by Terran (1990) from the analysis of an aquifer test. Model simulations also showed that a cap installed over the landfill area reduces hydraulic heads beneath the facility, but would not be effective at dewatering it.

The objective of the McConnell study was to develop a regional model of groundwater flow at the PGDP for the purposes of testing and refining conceptual models of the groundwater flow system beneath the site and evaluating remedial alternatives for the C-749 Uranium Burial Ground, SWMU 2. Use of the model at this landfill site provided a quantitative basis for evaluating engineering alternatives and lead to informed technical and economic decisions concerning the remediation of groundwater beneath the facility. In addition, the development of this model served an important, longer range objective by providing a groundwater management tool for decision makers who needed to evaluate groundwater flow conditions at SWMUs 7 and 30, also a part of WAG 22, but which were deferred until a later date.

Previous modeling studies provided valuable information regarding hydraulic property estimates, aquifer system response, and hydrogeologic framework; however, because the previous models addressed different sets of objectives, they contained deficiencies that reduce their usefulness for this modeling study. Some of the deficiencies included the following:

Regional Model (GeoTrans 1992)

- The GeoTrans model was not satisfactorily calibrated in the UCRS layer, suggesting potential errors in the model recharge or hydraulic parameters estimates;
- The GeoTrans model did not contain sufficient vertical discretization in the UCRS and lacked sufficient horizontal discretization in the area of interest to represent heterogeneous geologic conditions beneath the PGDP;

- The GeoTrans model used a no-flow boundary condition to simulate the southern model boundary along the Porters Creek Clay terrace. Later work suggested groundwater flow may occur at this interface; and
- The hydraulic conductivity assigned to the RGA in the GeoTrans model was much higher than values that had been measured from pumping tests conducted at the site.

Site-Specific Models

- Models developed by McConnell (1992a; 1992b; 1993) did not incorporate regional groundwater flow components; and boundary conditions set close to area(s) of interest by McConnell's models overly constrained model predictions.

Geraghty & Miller's approach for their modeling study addressed the aforementioned deficiencies. The groundwater flow model resulting from this study provided a more reliable decision-making tool that incorporated regional-scale groundwater flow components in predictive simulations made at the scale appropriate for the site simulation.

To meet the objectives of the study, Geraghty & Miller developed a three-dimensional numerical model that simulates groundwater flow in the vicinity of the PGDP. Geraghty & Miller developed the model in two phases. In the first phase, Geraghty & Miller constructed and calibrated a groundwater flow model covering nearly 100 km² for the purpose of simulating groundwater flow on a regional scale in the principal water-bearing units beneath the site. The regional model simulated groundwater flow in a multi-aquifer system, consisting of the UCRS Hydrogeologic Unit (HU 2), RGA (HU 4/HU 5), and the McNairy Formation (HU 6), and incorporated detailed spatial information describing the distribution of heterogeneous sediments comprising the Upper Continental Deposits (HU 2). In the second phase, Geraghty & Miller used the regional modeling results to develop a site-scale groundwater flow model with the aim of evaluating the hydraulic effects of remedial alternatives on groundwater flow and contaminant migration pathways in the vicinity of SWMU 2.

For the simulation of groundwater flow at the PGDP, Geraghty & Miller selected the code MODFLOW, a publicly available groundwater flow simulation program developed by the U.S. Geological Survey (McDonald 1988). Using the 75 water-level targets selected for the calibration of the PGDP regional groundwater flow model, Geraghty & Miller evaluated the calibration of the model through the analysis of (1) simulated hydraulic head distribution in the HU 2A (Model Layer 1), HU 2B (Model Layer 2) and RGA units (Model Layer 3); (2) estimated hydraulic properties; and (3) residual statistics. The calibration objective for the PGDP regional groundwater flow model was to minimize the residual sum of squares computed for the 75 water-level calibration targets. The largest computed residual for the entire set of targets was -2.03 m (-9.95 ft); however, only six residuals out of the 75 targets exceeded +/- 1.5m (+/- 5 ft). Greater than 70% of the targets had residuals of +/- 0.6 m (+/- 2 ft) or less. Overall, the model showed a very good match to the measured water levels given the complex geologic conditions at the site. Residual statistics for the calibrated groundwater flow model also indicated good agreement between simulated and measured groundwater elevations. The mean was close to zero, and the residual standard deviation was less than 2% of the range of simulated water-level elevations for the entire model domain.

Following the calibration of the regional groundwater flow model for the PGDP, the second phase of model development was used to simulate the hydraulic effect(s) of postulated remedial alternatives on the groundwater flow system beneath SWMU 2. To perform detailed simulations of the remedial alternatives postulated for SWMU 2, Geraghty & Miller used a procedure known as telescopic mesh refinement or grid refinement to develop a site-scale model for the WAG 22 site. The finite-difference grid for the WAG 22 site model covered an area of 2.19 km² (0.85 mi²) and consisted of 125 columns, 113 rows, and

four layers. In the area of interest near SWMU 2, the grid cells measure 3.05 m (10 ft) on a side. Vertical discretization (layer elevations) of the WAG 22 site model was identical to the regional groundwater flow model. To preserve the characteristics of the regional groundwater flow system, the site model placed constant head boundaries along its external boundaries based on hydraulic heads simulated by the regional model. Hydraulic properties (horizontal and vertical hydraulic conductivities) and internal boundary conditions (precipitation recharge, streams and rivers, and McNair Formation constant head boundary) simulated by the site model remained unchanged from the regional model. The site model also used MODFLOW to perform the steady-state simulations of groundwater flow at the SWMU 2 site. This WAG 22 site model was used to simulate six remedial alternatives for SWMU 2.

E1.2.2 SWMU 2 DATA SUMMARY AND INTERPRETATION REPORT MODELING

The following is taken from *Data Summary and Interpretation Report for Interim Remedial Design at Solid Waste Management Unit 2 of Waste Area Grouping 22 at the Paducah Gaseous Diffusion Plant, Paducah, Kentucky*, February 1997 (DOE 1997).

E1.2.2.1 MEPAS Modeling

To quantify the potential migration of contaminants in source materials (i.e., soil, waste, and groundwater) at SWMU 2 to exposure points, MEPAS was used. For groundwater, exposure points modeled were the PGDP property boundary and the PGDP security fence. The MEPAS modeling was used to determine contaminant concentrations at exposure points over time. Tables E1.1 and E1.2 show parameters defining the environmental setting and source term used.

Transportation of contamination into the RGA and to the integrator point was modeled over a 10,000-year period. At SWMU 2, the latest sampling data showed no TCE dense nonaqueous phase liquid (DNAPL) in the RGA. However, it is known that 450 gal of TCE were disposed of at SWMU 2. To account of this material, the entire 450-gal source was modeled as being within the waste volume. It was assumed that little or no biodegradation of TCE occurs (e.g., Wilson *et al.* 1983, Kleopfer *et al.* 1985, Fetter 1993). This assumption is conservative because over long periods of time (such as the 10,000-year duration modeled in this study) TCE will probably undergo degradation into something other than vinyl chloride (Fetter 1993).

The exposure points for the groundwater flow are located at the fence and DOE boundary. From SWMU 2, the distances are 1,875 ft to the fence and 2,475 ft to the DOE boundary. The site width is perpendicular to the groundwater flow direction (north), giving a site width and length of 160 and 200 ft, respectively. For modeling, the lesser of the maximum detected concentration and the upper 95% confidence limit on the mean concentration was used for detected analytes that were on the final SAP analyte list. For analytes that were on the final SAP analyte list but were not detected, the maximum nondetected value was used. (Note, the maximum nondetected values were used because previous information indicated that each of the analytes in the final SAP analyte list should be present at SWMU 2.)

Table E1.1. MEPAS Modeling Parameters: Soil Characteristics, Hydrology, and Hydrogeology

Parameter	Waste Soil	Top Soil	HU1	HU2A	HU2 Conf	HU2B	HU3	RGA
Soil Texture ^a	9	9	9	1	11	1	11	1
% Sand ^b	5	5	5	97	10	97	8	90
% Silt ^b	75	75	75	3	40	3	38	10
% Clay ^b	20	20	20	0	50	0	54	0
% Organic Matter ^c	0.1	0.1	0.1	0.1	0.07	0.1	0.08	0.03
% Iron & Aluminum ^d	2.0	2.4	2.0	2.0	2.1	2.6	4.2	2.0
pH ^e	6.0	5.5	6.0	6.0	6.0	5.8	6.1	6.6
% Vegetative Cover ^f	---	25	---	---	---	---	---	---
Top Soil Water Capacity ^f	---	0.33	---	---	---	---	---	---
SCS Curve# ^f	---	74	---	---	---	---	---	---
Thickness ^g , ft	4.6	2.0	16.5	7.5	8	6	9	40
Bulk Density ^h , g/cm ³	1.96	---	1.96	2.24	1.96	2.24	1.96	2.16
Total Porosity ⁱ , %	33	---	33	25	40	25	38	30
Field Capacity ^f , %	25	---	25	13	35	13	30	---
Effective Porosity ^f , %	---	---	---	---	---	---	---	25
Longitudinal Dispersivity ^j (Thickness x 0.1), ft	1.65	---	1.65	0.75	0.8	0.6	0.9	---
Saturated Hydraulic Conductivity ^k , cm/s	1E-7	---	1E-7	5E-6	8E-7	1E-6	5E-7	---
Darcy Velocity ^f , ft/day	---	---	---	---	---	---	---	1.17
Travel Distance ^l	---	---	---	---	---	---	---	1,875/ 2,475
Longitudinal Dispersivity ^j (=Travel Dist*0.1), ft	---	---	---	---	---	---	---	187.5/ 247.5
Transverse Dispersivity ^j (=Long Disp*0.33), ft	---	---	---	---	---	---	---	61.9/ 81.7
Vertical Dispersivity ^j (=Long Disp*2.5E-3), ft	---	---	---	---	---	---	---	0.469/ 0.619
% of Flux into Aquifer ^m	---	---	---	---	---	---	---	100
Perpendicular Distance to Plume Centerline, ft	---	---	---	---	---	---	---	0
Vertical Distance below GW, ft	---	---	---	---	---	---	---	0

Notes:

Table taken from DOE 1997.

“---” indicates that these data are not required for that hydrogeologic layer when setting up the MEPAS model.

^a Selected from MEPAS based on descriptions of soil characteristics contained in SAIC (1994), Claussen *et al.* (1996), and SAP field investigation where 9 is silty clay loam, 1 is sand, and 11 is silty clay.

^b Values for top soil and HU1 taken from SAIC (1994). Value for HU2A taken from Claussen *et al.* (1996). Values for HU2 Confining, HU2B, HU2, and RGA are estimated based on results from SAP field investigation. Value for waste set equal to HU1.

^c Estimated from results of SAP field investigation.

^d Estimated from results of the Phases I and II Site Investigations (CH2M Hill 1991 and 1992).

^e Values for top soil, HU1, HU2B, and HU2 Confining estimated from results of SAP field investigation. Values for HU2A and HU3 from analyses for monitoring wells MW48 and MW53 as found in the PGDP Environmental Information Management System data base. Value for waste soil set equal to HU1.

^f Value is estimated from MEPAS default (Droppo *et al.* 1989) and is based on professional judgment.

^g All values, except for waste, were estimated from results of SAP field investigation. Value for waste soil was calculated as shown in Appendix B and Table 2.19 [of DOE 1997].

^h Value from McConnell (1993).

ⁱ Calculated value.

^k Calculated from values in Table 4.1 [of DOE 1997].

^l Determined from PGDP site map. First value is for migration to plant security fence; second value is for migration to DOE property boundary.

^m Value set to maximum.

Table E1.2. MEPAS Modeling Parameters: Adsorption Coefficients

Parameter	Top Soil	HU1 & Waste	HU2A	HU2 Conf	HU2B	HU3	RGA
Arsenic	19.4	19.4	5.86	19.4	5.86	19.4	5.86
Barium	2,800	2,800	530	16,000	530	16,000	530
Beryllium	1,400	1,400	70	8,000	70	8,000	70
Cadmium	423	423	14.9	56.7	14.9	56.7	14.9
Chromium	56.5	56.5	16.8	360	16.8	360	16.8
Manganese	25.3	25.3	16.5	36.9	16.5	36.9	16.5
Nickel	58.6	58.6	12.2	650	12.2	650	12.2
Silver	4	4	0.4	40	0.4	40	0.4
Thallium	0.2	0.2	0	0.8	0	0.8	0
Uranium	253	253	1,170	3640	1,170	3,640	66.8
Vanadium	100	100	50	100	50	100	50
<i>cis</i> -1,2-dichloroethene	Analyte not found in MEPAS database						
<i>trans</i> -1,2-dichloroethene	0.0278	0.0278	0.007	0.115	0.007	0.012	0.0059
Aroclor 1016	1360	1,360	134	2160	134	2,300	111
Aroclor 1221	44	44	4.33	69.7	4.33	74.2	3.59
Aroclor 1232	5.84	5.84	0.575	9.26	0.575	9.86	0.477
Aroclor 1242	47.8	47.8	4.7	75.7	4.7	80.6	3.89
Aroclor 1248	2,100	2,100	207	3,330	207	3,540	171
Aroclor 1254	4,020	4,020	395	6,360	395	6,780	328
Aroclor 1260	50,800	50,800	5,000	80,500	5,000	85,700	4,140
Trichloroethene	0.955	0.955	0.094	1.51	0.094	1.61	0.0779
Vinyl chloride	0.432	0.432	0.0425	0.685	0.0425	0.729	0.0352
²⁴¹ Am	200	200	82	1000	820	1,000	82
²³⁷ Np	3	3	3	3	3	3	3
²³⁹ Pu	100	100	10	250	10	250	10
²³⁴ Pa	50	50	0	500	0	500	0
⁹⁹ Tc	20	20	3	20	3	20	3
²³⁰ Th	500	500	100	2,700	100	2,700	100
²³⁴ Th	500	500	100	2,700	100	2,700	100
²³⁴ U	243	243	906	1,580	906	1,580	62.98
²³⁵ U	243	243	906	1,580	906	1,580	62.98
²³⁸ U	243	243	906	1,580	906	1,580	62.98

Notes:

Table taken from DOE 1997.

All adsorption coefficients except those for uranium were generated by MEPAS. Values for uranium were taken from information gathered during recently completed SWMU 7 and 30 field investigation.

The results of the MEPAS modeling are shown in Tables E1.3. The results of the MEPAS model are not the total concentrations of SAP analytes that are present in the exposure medium (e.g., groundwater) at the exposure point but only the additional contamination that may be contributed by SWMU 2 sources.

The contributed concentration of all radionuclides, and most other analytes, is much less than the analyte's respective PRGs. This result indicates that for these SAP analytes, migration from soil and waste cells through groundwater to the exposure point at the security fence is not of concern over the 10,000 years modeled by MEPAS. However, the maximum contributed concentration of arsenic, Aroclor 1221, Aroclor 1232, Aroclor 1242, TCE, 1,1-dichloroethene, and vinyl chloride exceeds these analytes' respective human health risk-based PRG. In addition, the contributed concentration of TCE exceeds its regulatory value (i.e., MCL). Note, because similar results were obtained when MEPAS was used to model the maximum contributed concentration of SAP Analytes in RGA groundwater at the plant boundary, these results are not shown.

Table E1.3. MEPAS Results—Comparison of Estimated Maximum Concentrations of Contaminants in RGA Water at the PGDP Fence Line Originating from Soil and Waste Cells to Residential Preliminary Remediation Goals

Analyte	Maximum Concentration ^a	Time of Maximum Concentration ^b	ELCR ^c	HI ^d	Regulatory Value ^e	Background/ ^f	Criteria Exceeded ^g	Units
Arsenic	4.93E-04	1,505	3.50E-06	4.52E-04	5.00E-02	1.10E-02	k	mg/L
Barium	0.00E+00	35		1.04E-01	2.00E+00	2.90E-01	No	mg/L
Beryllium	6.45E-33	9,975	1.05E-06	6.61E-03	4.00E-03	9.30E-03	No	mg/L
Cadmium	2.75E-07	9,975		6.61E-04	5.00E-03	2.10E-02	No	mg/L
Chromium	8.20E-06	9,975		7.05E-03	1.00E-01	1.30E-01	No	mg/L
Manganese	1.74E-02	2,765		6.81E-02	1.59E-01*	1.60E-01	No	mg/L
Nickel	3.48E-06	9,975		3.01E-02	6.19E-02	6.20E-02	No	mg/L
Silver	1.97E-04	1,715		7.50E-03	1.00E-01*	1.10E-01	No	mg/L
Thallium	1.07E-03	35			2.00E-03	1.10E-01	No	mg/L
Uranium	4.86E-03	665		4.53E-03	2.00E-02*		No	mg/L
Vanadium	3.08E+04	8,015		9.25E-03		1.40E-01	No	mg/L
Aroclor 1016	3.22E-31	9,975		4.69E-05	5.00E-04		No	mg/L
Aroclor 1221	1.37E-06	4,305	5.83E-07		5.00E-04		k, h	mg/L
Aroclor 1232	9.95E-06	595	6.67E-07		5.00E-04		k, h	mg/L
Aroclor 1242	1.26E-06	4,725	6.40E-07		5.00E-04		k, h	mg/L
Aroclor 1248	8.13E-40	9,975	4.03E-07		5.00E-04		No	mg/L
Aroclor 1254	3.43E-43	9,975	4.13E-07	4.30E-05	5.00E-04		No	mg/L
Aroclor 1260	0.00E+00	35	2.27E-07		5.00E-04		No	mg/L
1,1-dichloroethene	4.78E-06	35	1.62E-06	1.34E-02	7.00E-03		g, h	mg/L
1,2-dichloroethene	5.35E-05	35	1.49E-02 ⁱ		7.00E-02 ⁱ		No	mg/L
Trichloroethene	5.64E-02	105	2.01E-04	7.86E-03	5.00E-03		k, L	mg/L
Vinyl chloride	7.74E-05	35	2.04E-06		2.00E-03		k	mg/L
²²⁵ Ac	1.55E-06	3,535	2.72E-01				No	pCi/L
²²⁷ Ac	1.89E-04	735	6.17E-02 ^j				No	pCi/L
²⁴¹ Am	6.28E-03	665	1.18E-01				No	pCi/L
²¹⁰ Bi	6.94E-07	4,025	5.30E+00				No	pCi/L
²³⁷ Np	5.27E-02	35	1.29E-01 ^j				No	pCi/L
²³¹ Pa	1.97E-04	735	2.59E-01				No	pCi/L
²³³ Pa	5.27E-02	35	8.23E+00				No	pCi/L
²¹⁰ Pb	6.94E-07	4,025	3.82E-02 ^j				No	pCi/L

Table E1.3. MEPAS Results—Comparison of Estimated Maximum Concentrations of Contaminants in RGA Water at the PGDP Fence Line Originating from Soil and Waste Cells to Residential Preliminary Remediation Goals (Continued)

Preliminary Remediation Goals								
Analyte	Maximum Concentration ^a	Time of Maximum Concentration ^b	ELCR ^c	HI ^d	Regulatory Value ^e	Background/ Exceeded ^f	Criteria Exceeded ^g	Units
²¹⁰ Po	6.94E-07	4,025	1.18E-01				No	pCi/L
²³⁹ Pu	2.66E-02	175	1.22E-01				No	pCi/L
²²³ Ra	1.89E-04	735	1.65E-01				No	pCi/L
²²⁵ Ra	1.55E-06	3,535	2.46E-01				No	pCi/L
²²⁶ Ra	4.00E-02	1,155	1.30E-01 ^j				No	pCi/L
²²² Rn	7.09E-07	4,025	1.03E+00 ^j				No	pCi/L
⁹⁹ Tc	3.46E-02	1,365	2.76E+01				No	pCi/L
²²⁷ Th	1.89E-04	735	9.56E-01				No	pCi/L
²²⁹ Th	1.55E-06	3,535	1.08E-01 ^j				No	pCi/L
²³⁰ Th	1.04E-01	1,085	1.03E+00		1.40E+00		No	pCi/L
²³¹ Th	1.34E-02	665	2.16E+01				No	pCi/L
²³⁴ Th	1.61E-01	665	2.00E+00				No	pCi/L
²³³ U	3.24E-05	315	8.62E-01				No	pCi/L
²³⁴ U	1.51E-01	665	8.70E-01		1.20E+00		No	pCi/L
²³⁵ U	1.34E-02	665	8.21E-01 ^j		1.50E-01		No	pCi/L
²³⁸ U	1.61E-01	665	6.23E-01 ^j		1.10E+00		No	pCi/L

Table taken from DOE 1997.

Blank cells indicate that value is not available or not applicable.

^aMaximum concentration of analyte predicted to be in RGA water at the PGDP security fence by MEPAS. All modeling was performed over a 10,000-year period.

^bTime at which MEPAS predicts maximum concentration will be reached.

^cDirect contact residential use risk-based preliminary remediation goal calculated using 1×10^{-7} as the target excess lifetime cancer risk (ELCR) for chemicals and 1×10^{-6} as the target ELCR for radionuclides.

^dDirect contact residential use risk-based preliminary remediation goal calculated using 0.1 as the target hazard index.

^eThe value reported is the respective analyte's maximum contaminant level (MCL). All MCLs are Primary Drinking Water Standards except where marked with *. Marked values are either proposed Primary Drinking Water Standards or Secondary Drinking Water Standards (SMCLs).

^fConcentration of analyte in uncontaminated media. For all water samples, the background values reported are those for the RGA.

^gPreliminary remediation goals exceeded. In this table, maximum detected concentrations are not directly comparable to preliminary remediation goals because MEPAS only predicts the additional contamination added by migration. However, the difference in magnitude between preliminary remediation goals and the maximum predicted concentrations indicates that contaminants from SWMU 2 are unlikely to contribute significantly to contamination in water at the PGDP security fence over the next 10,000 years.

^hSource term concentration based on maximum undetected concentration.

ⁱMEPAS does not offer both *cis*-1,2-dichloroethene and *trans*-1,2-dichloroethene; therefore, both isomers were modeled as *trans*-1,2-dichloroethene. However, the preliminary remediation goals reported are the lesser of those for the respective isomers.

^jPreliminary remediation goal calculated using the toxicity value (i.e., slope factor) for parent isotope and short-lived daughters.

^kExceeds human health risk based preliminary remediation goal (PRG).

^lExceeds regulatory value (i.e., MCL).

Of the seven chemicals found to have contributed concentrations that exceed screening criteria, the results for four, Aroclor 1221, Aroclor 1232, Aroclor 1242, and 1,1-dichloroethene, should be considered suspect because their contributed concentrations were based on the respective chemical's maximum nondetected concentration. Analytes that were not detected in any sample were retained in the source terms by using these analytes' maximum nondetected value. Although this approach was conservative, this procedure allowed these four chemicals, which have relative high detection limits in relation to their toxicity, to appear to migrate to exposure points at levels that may be of concern. In addition, the significance of arsenic's and vinyl chloride's contributed concentrations in relation to their screening criteria can be questioned because each chemical's contributed concentration is less than its MCL over the time period modeled.

The contributed concentrations of TCE, unlike the other chemicals, exceed both the human health risk-based PRG and the MCL in at least one time period. However, the concentrations of TCE exceed the MCL for only approximately the first 250 years modeled. After this time, the concentrations rapidly fall and are below the human health risk-based PRG based on ELCR prior to model year 500.

A secondary source of TCE may exist in the RGA at SWMU 2. Although the detected maximum concentration of TCE in RGA water is less than the criteria established for secondary sources in DOE (1996) (i.e., 10 mg/L); this concentration is 53.5% of this level. The results of the modeling of the potential secondary source in the RGA to the security fence and plant boundary are shown in Table E.1.4. The results in this table show that by year 105 from present, all contributed concentrations will be below TCE's PRGs at both the security fence and plant boundary. These results also show that the current concentration of TCE at the security fence and plant boundary exceed all of the respective analytes' PRGs and that the potential contributed concentration from SWMU 2 at 35 year is 0.5% and 0.3% respectively, of the current concentration at these locations.

Table E1.4. MEPAS Results—Comparison of Estimated Maximum Concentrations of Contaminants in RGA Water at the PGDP Fence Line Originating from Soil and Waste Cells to Residential Preliminary Remediation Goals

Time (years)	Preliminary Remediation Goals					Criteria Exceeded ^f	Units
	Concentration ^a	ELCR ^b	HI ^c	Regulatory Value ^d	Background ^e		
Results for 1,1-trichloroethene at the security fence							
Present ^g	1.50E+01	2.01E-04	7.86E-03	5.00E-03	None	PR	mg/L
35	6.11E-02					PR	mg/L
105	3.94E-07					No	mg/L
Results for 1,1-trichloroethene at the plant boundary							
Present ^g	1.50E+01	2.01E-04	7.86E-03	5.00E-03	None	PR	mg/L
35	4.51E-02					PR	mg/L
105	3.52E-06					No	mg/L

Note: Table taken from DOE 1997.

^a Present concentrations are measured values; future concentrations are additional materials that will be in addition to materials that will be in addition to materials migrating from other sources (i.e., contributed concentrations).

^b Direct contact residential use risk-based preliminary remediation goal calculated using 1×10^{-7} as the target excess lifetime cancer risk (ELCR) for chemicals.

^c Direct contact residential use risk-based preliminary remediation goal calculated using 0.1 as the target hazard index.

^d The value reported is the respective analyte's maximum contaminant level (MCL). All MCLs are Primary Drinking Water Standards.

^e Concentration of analyte in uncontaminated media. For all water samples, the background values reported are those for the RGA.

^f Summary of preliminary remediation goals exceeded. In this table, maximum detected concentrations are not directly comparable to preliminary remediation goals because MEPAS only predicts the additional contamination added by migration. However, the difference in magnitude between preliminary remediation goals and the maximum predicted concentrations indicates if the preliminary remediation goals may be exceeded. Definitions of codes are:

- P One or both of the residential use human health risk-based preliminary remediation goals are exceeded.
- R The regulatory value is exceeded.
- No No preliminary remediation goals are exceeded.

^g Present concentrations were taken from analyses performed for sample from EW230 taken on 11/28/95.

E1.2.2.2 RESRAD Modeling

The RESRAD (Version 5.6.1) computer code was used to model the specific case of migration of radionuclide contaminants from source areas to the RGA directly under SWMU 2. Therefore, the only exposure point considered using this model was the residential use of RGA groundwater drawn from below SWMU 2. The RESRAD computer code was not used to model transport of contaminants to exposure points at the property boundary and security fence because this code cannot model lateral transport. Tables E1.5 and E1.6 show environmental setting and source term used.

Table E1.5. RESRAD Modeling Parameters: Soil Characteristics, Hydrology, and Hydrogeology

Parameter	Waste			HU2			
	Soil	HU1	HU2A	Conf	HU2B	HU3	RGa
Thickness, m ^a	1.40	5	2.29	2.44	1.83	2.74	---
Soil density ^b , g/cm ³	1.96	1.96	2.24	1.96	2.24	1.963	2.16
Total porosity ^c	0.33	0.33	0.25	0.40	0.25	0.38	0.30
Effective porosity ^d	0.25	0.25	0.13	0.35	0.13	0.30	0.25
Soil-specific b parameter ^e	10.4	10.4	4.05	10.4	4.05	10.4	4.05
Hydraulic conductivity ^f , m/yr	0.032	0.032	1.58	0.025	0.32	0.16	6,508
Hydraulic gradient ^g	---	---	---	---	---	---	0.02
Water table drop rate ^h , m/yr	---	---	---	---	---	---	0.001
Distribution coefficients ⁱ , cm ³ /g							
²⁴¹ Am	200	200	82	1,000	82	1,000	82
²¹⁰ Pb	597	597	234	1,830	234	1,830	234
²³⁷ Np	3	3	3	3	3	3	3
²³⁹ Pu	100	100	10	250	10	250	10
²³¹ Pa	50	50	0	500	0	500	0
²²⁶ Ra	100	100	24.3	124	24.3	124	24.3
⁹⁹ Tc	20	20	3	20	3	20	3
²²⁹ Th	500	500	100	2,700	100	2,700	100
²³⁰ Th	500	500	100	2,700	100	2,700	100
²³³ U	253	253	1,170	3,640	1,170	3,640	66.8
²³⁴ U	253	253	1,170	3,640	1,170	3,640	66.8
²³⁵ U	253	253	1,170	3,640	1,170	3,640	66.8
²³⁸ U	253	253	1,170	3,640	1,170	3,640	66.8

Notes:

Table taken from DOE 1997.

“---” indicates that this parameter is not needed for the respective hydrogeologic unit.

^a Taken from the site conceptual model for SWMU 2.

^b Taken from McConnell (1993).

^c Estimated value.

^d Estimated value.

^e RESRAD default estimated from soil characteristics.

^f Taken from site conceptual model for SWMU 2.

^g Taken from the site conceptual model for SWMU 2.

^h Assumes minimal change due to pumping of residential well.

ⁱ All distribution coefficients, except for those for uranium isotopes, were default values taken from MEPAS. The distribution coefficients for uranium isotopes were those determined during the recently completed field investigation at SWMUs 7 and 30 of WAG 22. When selecting the default values for all other radioisotopes, the description of the soil characteristics of each hydrogeologic unit was used. Note, the soil type for the waste zone was assumed to be that found in HU1. This is a conservative estimate because the recently completed field investigation showed that non-native clay was used to backfill all waste pits.

Table E1.6. RESRAD Modeling Parameters: Initial Source Term Analyte Concentrations

Analyte	Input (pCi/g) ^a							
	Surface	HU1	HU2A	HU2 Confining	HU2B	HU3	RGA	Waste
²⁴¹ Am	---	0.199	0.13	0.12	0.10	0.48	0.103	0.44
²³⁷ Np	0.32	0.032	0.02	0.03	0.12	0.00	0.028	0.15
²³⁹ Pu	7.9	0.032	0.01	0.09	0.02	0.05	0.033	0.07
⁹⁹ Tc	58.0	0.664	0.023	0.29	0.00	0.04	0.017	0.012
²³⁰ Th	14.0	1.25	0.873	0.90	0.76	1.33	0.792	0.41
²³⁴ U	18.0	7.73	0.712	0.93	0.86	1.20	0.735	7.61
²³⁵ U	1.7	1.66	0.083	0.11	0.12	0.07	0.066	7.0E-06
²³⁸ U	69.0	53.70	0.756	0.97	0.93	1.27	0.787	10,200.00

Notes:

Table adapted from DOE 1997.

NA = not applicable.

^a Maximum values selected from SAP database.

Table E1.7 presents the results of the RESRAD modeling. The result of this modeling is the additional contamination that may migrate to the RGA from the sources at SWMU 2. The table shows both the total contributed dose and the contributed dose from each source at SWMU 2.

Table E1.7. RESRAD Results—Contribution of Radionuclides in Soil and Waste Cells to Potential Total Dose to Resident (mrem/year) through Residential Ingestion of RGA Water

Years	Source Area										Total Dose
	RGA ^a	HU3	HU2B	HU2	HU2A	HU2 Total ^b	HU1	Waste Cells	Surface		
50	8.83E-01	3.58E-02	1.42E+00	5.44E-01	0.00E+00	1.96E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	2.88E+00
63	8.72E-01	4.20E-02	8.44E-01	3.54E-01	0.00E+00	1.20E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	2.11E+00
79.37	8.62E-01	4.92E-02	4.41E-01	2.06E-01	3.32E-01	9.79E-01	0.00E+00	0.00E+00	0.00E+00	0.00E+00	1.89E+00
100	8.44E-01	5.71E-02	1.94E-01	1.04E-01	1.72E-01	4.71E-01	5.87E-01	1.44E+00	0.00E+00	0.00E+00	3.40E+00
126	7.95E-01	6.05E-02	6.93E-02	4.42E-02	7.52E-02	1.89E-01	3.84E-01	3.50E-01	0.00E+00	0.00E+00	1.78E+00
158.7	7.35E-01	6.35E-02	1.89E-02	1.50E-02	2.66E-02	6.04E-02	2.25E-01	5.91E-02	8.92E-02	8.92E-02	1.23E+00
200	6.81E-01	6.73E-02	5.24E-03	4.18E-03	7.18E-03	1.66E-02	1.15E-01	6.35E-03	1.31E-04	1.31E-04	8.57E-01
252	6.19E-01	7.17E-02	7.12E-04	9.66E-04	1.43E-03	3.10E-03	4.93E-02	4.65E-04	6.93E-07	6.93E-07	7.44E-01
317.5	5.70E-01	7.69E-02	1.04E-04	2.83E-04	2.39E-04	6.26E-04	1.70E-02	1.09E-04	6.73E-07	6.73E-07	6.64E-01
400	5.39E-01	8.44E-02	5.66E-05	1.44E-04	1.14E-04	3.14E-04	5.25E-03	1.10E-04	7.12E-07	7.12E-07	6.29E-01
504	5.14E-01	9.21E-02	5.22E-05	8.69E-05	7.12E-05	2.10E-04	1.47E-03	1.01E-04	8.00E-07	8.00E-07	6.08E-01
635	4.92E-01	9.79E-02	4.69E-05	5.02E-05	6.44E-05	1.62E-04	5.87E-04	9.17E-05	9.65E-07	9.65E-07	5.91E-01
800	4.78E-01	1.03E-01	3.92E-05	2.74E-05	5.55E-05	1.22E-04	3.43E-04	7.92E-05	1.88E-03	1.88E-03	5.83E-01
1008	4.63E-01	1.07E-01	3.00E-05	1.48E-05	4.39E-05	8.86E-05	2.09E-04	6.37E-05	7.33E-06	7.33E-06	5.70E-01
1270	4.46E-01	1.09E-01	4.65E-03	1.23E-03	3.11E-05	5.91E-03	1.13E-04	4.66E-05	1.17E-06	1.17E-06	5.61E-01
1600	4.26E-01	1.10E-01	9.46E-02	2.01E-02	1.94E-05	1.15E-01	5.40E-05	3.04E-05	1.21E-06	1.21E-06	6.51E-01
2016	3.99E-01	1.09E-01	1.33E-01	3.46E-02	1.03E-05	1.68E-01	2.26E-05	1.75E-05	1.23E-06	1.23E-06	6.75E-01
2540	3.67E-01	1.05E-01	1.69E-01	6.01E-02	5.81E-02	2.87E-01	9.31E-03	2.96E-03	1.28E-06	1.28E-06	7.71E-01
3200	3.31E-01	1.00E-01	1.59E-01	5.93E-02	1.16E-01	3.34E-01	6.08E+02	1.73E-02	1.32E-06	1.32E-06	8.44E-01
4032	2.92E-01	9.55E-02	1.61E-01	5.79E-02	1.36E-01	3.54E-01	8.92E-02	3.28E-02	1.38E-06	1.38E-06	8.64E-01
5080	2.50E-01	9.04E-02	1.60E-01	5.81E-02	1.44E-01	3.62E-01	1.15E-01	5.49E-02	4.77E-02	4.77E-02	9.20E-01
6400	2.07E-01	8.50E-02	1.45E+00	5.96E-02	1.43E-01	3.55E-01	1.15E-01	6.72E-02	1.01E-01	1.01E-01	9.31E-01
8063	1.63E-01	7.89E-02	1.44E-01	6.17E-02	1.40E-01	3.46E-01	1.14E-01	8.77E-02	4.58E-02	4.58E-02	8.35E-01
10160	1.21E-01	7.23E-02	1.35E-01	6.23E-02	1.31E-01	3.29E-01	1.55E-01	1.03E-01	4.35E+01	4.35E+01	8.24E-01
12800	8.31E-02	6.46E-02	1.27E-01	4.66E-02	1.22E-01	2.95E-01	1.14E-01	1.21E-01	4.40E-02	4.40E-02	7.22E-01
Maximum Dose	8.83E-01	1.10E-01	1.42E+00	5.44E-01	3.32E-01	1.96E+00	5.87E-01	1.44E+00	1.01E-01	1.01E-01	3.44E+00
Year of Maximum	50	1,600	50	50	79.37	50	100	100	6,400	6,400	100

Note: Table taken from DOE 1997.

^aThe RGA source area was assumed to consist of hydrogeologic units 4 and 5.

^bThis column contains the sum of the results for HU2A, HU2 confining, and HU2B.

E1.3. MODELING APPEARING IN THE WAG 3 RI REPORT FOR SWMU 4

The conservative modeling in Appendix B of Volume 4 of the WAG 3 RI Report (DOE 2000a) was completed to determine if any contaminants could migrate from source areas at SWMU 4 to POEs at the plant boundary and property boundary at a rate that could result in maximum concentrations greater than risk-based screening levels. This modeling was completed using MEPAS and conservative source term estimates developed using comparisons of sampling results to background concentrations and SSLs for protection of groundwater taken from EPA sources.

The sampling results used in source term development were derived from the WAG 3 RI (DOE 2000a), the Data Gaps Investigation Report (DOE 2000b), and from earlier sampling completed in support of the PGDP Comprehensive Environmental Response, Compensation, and Liability Act (CERCLA) SI performed in the early 1990s (CH2M Hill 1991 and 1992). Source terms developed for SWMU 4 are presented in Table E1.8. As noted in the modeling report, “In all cases, modelers applied conservatism (worst case) in the definition of the extent of the source zones. In all cases, the maximum concentrations were used to develop each contaminant source-term inventory.”

Input parameters used in the MEPAS modeling were based on site-specific data when available. When relevant on-site data were not available, data collected at nearby SWMUs having similar hydrogeologic conditions were used to define the input parameter. If no site-specific data were available, then default values provided by MEPAS were used. In the analysis, all sources were modeled as depleting over time and degrading in the environment. The modeled period was 10,000 years. Modeling inputs for SWMU 4 are presented in Table E1.9. The distribution coefficients (K_d) used were default values taken from MEPAS. These values are presented in Table E1.10.

The results of the MEPAS modeling for SWMU 4, taken from Appendix C of Volume 4 of the WAG 3 RI Report, are in Table E1.11. Interpretations of these results from the Southwest Groundwater Plume SI report (DOE 2006) are shown in Tables E1.12 and E1.13.

Based upon these results, the COCs for SWMU 4 for the plant boundary POE are cobalt; copper; iron; manganese; 1,1-DCE; carbon tetrachloride; TCE; vinyl chloride; ^{237}Np ; ^{239}Pu ; radium-226 (^{226}Ra); ^{99}Tc ; ^{234}U ; ^{235}U ; and ^{238}U . The COCs for SWMU 4 for the property boundary POE are copper; iron; manganese; 1,1-DCE; TCE; vinyl chloride; ^{237}Np ; ^{239}Pu ; ^{99}Tc ; ^{234}U ; ^{235}U ; and ^{238}U .

Table E1.8. Source Term for SWMU 4 Developed in the WAG 3 RI Report MEPAS Modeling^a

Contaminant	Level ^b (mg/kg or pCi/g)	Parallel to Flow Axis (feet)		Perpendicular to Flow Axis (feet)		Thickness (feet)	Volume (ft ³)	Inventory ^c (g or Ci)	Note ^d
		Axis (feet)	Axis (feet)	to Flow Axis (feet)	to Flow Axis (feet)				
<i>Inorganic Chemicals (Metals)</i>									
Aluminum	26,400	470	610	54	15,481,800	2.11E+10	Subsurface		
Chromium	296	370	70	1	25,900	317,000	Surface		
	42.3	NV	NV	1	22,049	38,600	Surface		
Cobalt	77.3	470	365	20	3,431,000	13,700,000	Subsurface		
Copper	31.6	470	610	54	15,481,800	25,200,000	Subsurface		
	19.5	130	180	1	23,400	18,900	Surface		
	30.1	80	165	1	13,200	16,400	Surface		
	1,130	470	610	34	9,747,800	5.68E+08	Subsurface		
Iron	30,700	NV	NV	1	22,049	28,000,000	Surface		
	34,500	470	610	54	15,481,800	2.75E+10	Subsurface		
Lead	62.5	470	175	20	1,645,000	5,300,000	Subsurface		
Lithium	0.148	75	80	10	60,000	458	Subsurface		
Manganese	2,920	470	640	24	7,219,200	1.09E+09	Subsurface		
Nickel	153	NV	NV	1	22,049	139,000	Surface		
Strontium	0.639	NV	NV	NV	NV	1,980	Subsurface		
<i>Organic Compounds</i>									
1,1-DCE	0.340	470	610	45	10,786,500	226,000	Subsurface		
1,2-DCE (mixed)	0.063	80	100	15	120,000	390	Subsurface		
1,2-DCE, <i>cis</i> -	1.5	245	610	24	3,586,800	277,000	Subsurface		
	11	245	610	19	2,839,550	1,610,000	Subsurface		
2-Butanone	0.002	75	130	5	120,000	5.02	Subsurface		
	0.031	65	105	5	34,125	54.5	Subsurface		
2-Propanol	0.100	75	130	5	48,750	251	Subsurface		
4-Methyl-3-penten-2-one	0.180	50	50	1	2,500	18.6	Surface		
	0.67	80	100	5	40,000	1,380	Subsurface		
	2	80	100	29	232,000	23,900	Subsurface		
6-(Acetyloxy)-2-hexanone	2.5	470	610	19	5,447,300	702,000	Subsurface		
PCB-1016	0.8	470	610	19	5,447,300	225,000	Subsurface		
PCB-1248	27	470	610	19	5,447,300	7,580,000	Subsurface		
PCB-1254	0.115	35	600	1	21,000	99.8	Surface		
PCB-1260	0.041	450	35	1	15,750	26.7	Surface		
	0.061	185	205	1	37,925	95.6	Surface		
	500	470	610	19	5,447,300	140,000	Subsurface		

Table E1.8. Source Term for SWMU 4 Developed in the WAG 3 RI Report MEPAS Modeling^a (Continued)

Contaminant	Level ^b		Parallel to Flow		Perpendicular		Volume (ft ³)	Inventory ^c (g or Ci)	Note ^d
	(mg/kg or pCi/g)	Axis (feet)	Axis (feet)	to Flow Axis (feet)	to Flow Axis (feet)	Thickness (feet)			
Bis(2-Methoxyethyl)phthalate	0.45	80	100	100	5	40,000	928	Subsurface	
Carbon tetrachloride	0.170	165	95	95	12	188,100	1,650	Subsurface	
Diethyl ether	0.009	70	45	45	54	170,100	78.9	Subsurface	
Ethanol, 2,2'-oxybis, diacetate	2	80	100	100	5	40,000	4,120	Subsurface	
Octachlorodibenzodioxin	8.2	80	100	100	5	40,000	16.9	Subsurface	
Pentachlorophenol	0.21	80	100	100	5	40,000	433	Subsurface	
TCE	48	470	610	610	45	10,786,500	31,900,000	Subsurface	
Vinyl chloride	0.4	470	610	610	45	10,786,500	266,000	Subsurface	
<i>Radionuclides</i>									
Cesium-137	1.48	210	610	610	10	1,281,000	0.0977	Subsurface	
²³⁷ Np	0.266	80	165	165	1	13,200	0.000145	Surface	
	5.78	470	610	610	10	2,867,000	1.62	Subsurface	
²³⁹ Pu	0.0644	55	445	445	1	24,475	0.0000652	Surface	
	4.17	470	610	610	19	2,867,000	1.17	Subsurface	
²²⁶ Ra	2.51	470	610	610	19	2,867,000	0.705	Subsurface	
⁹⁹ Tc	269	NV	NV	NV	NV	NV	75.5	Subsurface	
²³⁰ Th	68.7	NV	NV	NV	NV	NV	19.3	Subsurface	
²³⁴ U	12.8	35	600	600	1	21,000	0.0111	Surface	
	6.59	450	35	35	1	15,750	0.00429	Surface	
	30.1	185	205	205	1	37,925	0.0472	Surface	
²³⁵ U	69	NV	NV	NV	NV	NV	194	Subsurface	
²³⁸ U	7.2	NV	NV	NV	NV	NV	2.02	Subsurface	
	35.9	35	600	600	1	21,000	0.0312	Surface	
	26.4	450	35	35	1	15,750	0.0172	Surface	
	87.3	185	205	205	1	37,925	0.137	Surface	
Uranium, Total	126	NV	NV	NV	NV	NV	1,760	Subsurface	
	6,260	282	110	110	15	465,300	150	Subsurface	
	6,260	110	292	292	15	481,800	155	Subsurface	

NV = no value reported in the WAG 3 RI Report.

²³⁰Th = Thorium-230

^a Information taken from Table B.5 in Appendix B of Volume 4 of the WAG 3 RI Report.

^b The maximum concentration was used to estimate the contaminant inventory for all contaminants.

^c Calculated using a bulk density of 1.46 g/cm³ for Surface and 1.82 g/cm³ for the Subsurface.

^d Surface assumed to extend to 1 ft bgs. Thickness of subsurface differs with contaminant.

Table E1.9. Modeling Inputs for SWMU 4 MEPAS Modeling in the WAG 3 RI Report^a

Description	Name	Value	Reference
<i>Top Soil Parameters (WT)</i>			
Textural Classification	WT-CLASS	Silt loam	Soil Survey
Sand (%)	WT-SAND	15	Soil Survey
Silt (%)	WT-SILT	80	Maximum for soil type
Clay (%)	WT-CLAY	5	By difference
Organic Matter (%)	WT-OMC	0.05	CH2M Hill 1992
Iron and Aluminum (%)	WT-IRON	4	DOE 1995b
pH of Topsoil	WT-pH	7.32	RI
Vegetative Cover (%)	WT-VEGCOV	100	Description
Topsoil water capacity	WT-AVAILW	0.33	Soil Survey
SCS Curve Number	WT-SCSN	71	MEPAS
<i>Partially Saturated Zone Parameters (WP)</i>			
Thickness	WP-THICK	54	RI
Textural classification	WP-CLASS	Sandy loam	RI
Sand (%)	WP-SAND	38	RI
Silt (%)	WP-SILT	41	RI
Clay (%)	WP-CLAY	21	RI
Organic Matter (%)	WP-OMC	0.05	WAG 6
Iron and Aluminum (%)	WP-IRON	4	DOE 1995b
pH of Pore Water	WP-pH	6.0	DOE 1995b
Bulk Density (g/cm ³)	WP-BULKD	1.82	RI
Total porosity (%)	WP-TOTPOR	31.28	RI
Field capacity (%)	WP-FIELDC	14	MEPAS
Longitudinal dispersivity (ft)	WP-LDISP	0.54	MEPAS
Saturated hydraulic conductivity	WP-CONDUCT	3E-01 ft/day 1.06E-04 cm/sec	RI
Soil Moisture Content (%)	WS-MOISTC	31.28	MEPAS
<i>Saturated Zone Parameters (WZ)</i>			
Textural classification	WZ-CLASS	Loamy sand	RI
Sand (%)	WZ-SAND	74	RI
Silt (%)	WZ-SILT	17	RI
Clay (%)	WZ-CLAY	9	RI
Organic Matter (%)	WZ-OMC	0.02	RI
Iron and Aluminum (%)	WZ-IRON	3	RI
pH of Pore Water	WZ-pH	6.36	RI
Total porosity (%)	WZ-TOTPOR	37	RI
Effective porosity (%)	WZ-EFFPOR	30	MEPAS
Darcy velocity (ft/day)	WZ-PVELOC	0.6	Conductivity = 1500 ft/d Gradient = 0.0004
Thickness	WZ-THICK	45	RI
Bulk Density (g/cm ³)	WZ-BULKD	1.67	RI
Travel Distance (ft)	WZ-DIST	Plant boundary: 890 Property boundary: 2,985	RI
Longitudinal dispersivity (ft)	WZ-LDISP	50	Bioscreen Model
Transverse dispersivity (ft)	WZ-TDISP	5.0	Bioscreen Model
Vertical dispersivity (ft)	WZ-VDISP	0.1	near zero
Total flux to aquifer (%)	WZ-FRACT	100	estimate
Perpendicular to receptor	WZ-YDIST	0	on plume centerline
Vertical to receptor	WZ-AQDEPTH	0	minimum

^a Information taken from Table B.2 in Appendix C of Volume 4 of the WAG 3 RI Report.

Table E1.10. Distribution Coefficients (K_ds) Used for SWMU 4 MEPAS Modeling in the WAG 3 RI Report^a

Contaminant	Surface Soil	Subsurface Soil	RGA
<i>Inorganic Chemicals (metals)</i>			
Aluminum	3,980	35,300	35,300
Chromium	10	565	565
Cobalt	0.2	8.81	8.81
Copper	4.19	92.2	92.2
Iron	10	15	15
Lead	10	597	597
Lithium	0	0.2	0.2
Manganese	1.5	25.3	25.3
Nickel	1.2	58.6	58.6
Strontium	2.34	100	100
<i>Organic Compounds</i>			
1,1-DCE	0.292	0.4	0.171
1,2-DCE (mixed)	0.0432	0.059	0.0253
1,2-DCE, <i>cis</i> -	NV	NV	NV
2-Butanone	NV	NV	NV
2-Propanol	NV	NV	NV
4-Methyl-3-penten-2-one	NV	NV	NV
6-(Acetyloxy)-2-hexanone	NV	NV	NV
PCB-1016	809	1,110	424
PCB-1248	1,250	1,700	729
PCB-1254	2,380	3,260	1,400
PCB-1260	30,100	41,200	17,600
Bis(2-Methoxyethyl)phthalate	NV	NV	NV
Carbon tetrachloride	2.26	3.09	1.32
Diethyl ether	NV	NV	NV
Ethanol, 2,2'-oxybis, diacetate	NV	NV	NV
Octachlorodibenzodioxin	NV	NV	NV
Pentachlorophenol	238	326	140
TCE	0.567	0.775	0.332
Vinyl chloride	0.256	0.35	0.15
<i>Radionuclides</i>			
Cesium-137	10	249	249
²³⁷ Np	3	3	3
²³⁹ Pu	4	100	100
²²⁶ Ra	2.43	100	100
⁹⁹ Tc	1	1	0.1
²³⁰ Th	40	500	500
²³⁴ U	0	50	50
²³⁵ U	0	50	50
²³⁸ U	0	50	50
Uranium, Total	0	50	50

NV = Value not listed in the WAG 3 RI Report.

²³⁰Th = Thorium-230

^a Information taken from Table B.1 in Appendix B of Volume 4 of the WAG 3 RI Report. All values in mL/g.

Table E1.11. MEPAS Results for SWMU 4^a

Source	Contaminant ^b	PGDP Plant Boundary		PGDP Property Boundary		MCL ^d	
		Maximum Concentration ^c	Time of Maximum (Years)	Maximum Concentration ^c	Time of Maximum (Years)		
Surface	Chromium	2.81E-40	10,000	1.95E-52	10,000	1E-01	
Soil	Copper	4.40E-04	8,039	1.40E-04	9,585	1.3E+00	
	Iron	1.97E+00	1,337	6.41E-01	1,525	3E-01*	
	Nickel	2.53E-03	5,044	8.45E-04	6,107	1E-01	
	PCB-1260	0	10,000	0	10,000	5E-04	
	²³⁷ Np	5.33E-02	276	1.64E-02	315	NV	
	²³⁹ Pu	4.16E-04	8,717	1.44E-04	10,260	NV	
	²³⁴ U	1.37E+00	4,355	4.16E-01	5,166	20	
	²³⁸ U	2.67E+00	4,356	8.08E-01	5,167	20	
	Subsurface Soil	Aluminum	0	10,000	0	10,000	2E-01*
		Chromium	1.15E-37	10,000	9.22E-53	10,000	1E-01
Cobalt		3.29E+00	788	6.46E-01	961	NV	
Copper		7.32E+00	7,992	1.46E+00	9,539	1.3E+00	
Iron		1.16E+03	1,738	2.41E+02	2,055	3E-01*	
Lead		8.45E-42	10,000	7.54E-53	10,000	1.5E-02	
Lithium		1.76E-03	30	5.06E-04	36	NV	
Manganese		5.13E+01	2,248	9.46E+00	2,566	5E-02*	
Nickel		1.45E-01	5,019	4.29E-02	6,081	1E-01	
Strontium		2.54E-05	8,661	7.44E-06	10,450	NV	
1,1-DCE		2.57E-01	63	5.38E-02	69	7E-03	
1,2-DCE		2.24E-03	18	6.64E-04	21	7E-02**	
Carbon tetrachloride		5.94E-04	301	1.85E-04	307	5E-03	
PCB-1016		0	10,000	0	10,000	5E-04	
PCB-1248		0	10,000	0	10,000	5E-04	
PCB-1254		0	10,000	0	10,000	5E-04	
PCB-1260		0	10,000	0	10,000	5E-04	
Pentachlorophenol		3.35E-18	10,790	6.06E-19	12,910	1E-03	
TCE		2.26E+01	102	4.70E+00	111	5E-03	
Vinyl chloride		3.31E-01	57	6.90E-02	62	2E-03	
Cesium-137		0	12,920	0	12,920	NV	
²³⁷ Np		4.88E+02	316	9.83E+01	381	NV	
²³⁹ Pu		1.09E+01	8,665	2.05E+00	10,210	NV	
²²⁶ Ra		2.21E-01	8,208	2.16E-02	9,765	5	
⁹⁹ Tc		6.34E+04	111	1.32E+04	113	900	
²³⁰ Th		3.56E-28	10,000	1.30E-43	10,000	NV	
²³⁴ U		4.51E+03	4,329	8.94E+02	5,140	20	
²³⁵ U		4.75E+01	4,330	9.45E+00	5,141	20	
²³⁸ U		8.33E+02	4,330	1.66E+02	5,141	20	
		Total Uranium ^e	6.46E+03	4,330	2.13E+03	5,141	20

²³⁰Th = Thorium-230

^a Information taken from Table B.6 of Appendix B in Volume 4 of the WAG 3 RI Report.

^b Table B.6 includes results for degradation products of radionuclides. These are not included here.

^c Concentrations for chemicals and compounds in mg/L. Concentrations for radionuclides in pCi/L.

^d Maximum contaminant levels (MCLs) taken from PGDP Risk Methods Document. MCLs for chemicals and compounds in mg/L. MCLs for radionuclides in pCi/L. All values except those marked with an asterisk (*) are primary MCLs. Values marked with an asterisk (*) are secondary MCLs. The MCL listed for 1,2-DCE (**) is the primary MCL for *cis*-1,2-DCE. The primary MCL for *trans*-1,2-DCE is 1E-01 mg/L.

^e Modeled as ²³⁸U.

Table E1.12. Estimated HQs for a Resident from Exposure to Maximum Modeled Concentrations from Sources at SWMU 4

Source	Contaminant	PGDP Plant Boundary		PGDP Property Boundary		Risk-based Concentrations ^c
		Maximum Concentration ^a	Hazard Quotient ^b	Maximum Concentration ^a	Hazard Quotient ^b	
Surface	Chromium	2.81E-40	<0.1	1.95E-52	<0.1	1.76E+00
Soil	Copper	4.40E-04	<0.1	1.40E-04	<0.1	5.57E-02
	Iron	1.97E+00	0.4	6.41E-01	0.1	4.49E-01
	Nickel	2.53E-03	<0.1	8.45E-04	<0.1	3.01E-02
	²³⁷ Np	5.33E-02	No value	1.64E-02	No value	No value
	²³⁹ Pu	4.16E-04	No value	1.44E-04	No value	No value
	²³⁴ U	1.37E+00	No value	4.16E-01	No value	No value
	²³⁸ U	2.67E+00	No value	8.08E-01	No value	No value
	Subsurface	Chromium	1.15E-37	<0.1	9.22E-53	<0.1
Soil	Cobalt	3.29E+00	3.6	6.46E-01	0.7	9.06E-02
	Copper	7.32E+00	13.1	1.46E+00	2.6	5.57E-02
	Iron	1.16E+03	258	2.41E+02	53.7	4.49E-01
	Lead	8.45E-42	No value	7.54E-53	No value	1.50E-02*
	Lithium	1.76E-03	<0.1	5.06E-04	<0.1	3.02E-02
	Manganese	5.13E+01	147	9.46E+00	27.0	3.50E-02
	Nickel	1.45E-01	0.5	4.29E-02	0.1	3.01E-02
	Strontium	2.54E-05	<0.1	7.44E-06	<0.1	9.01E-01
	1,1-DCE	2.57E-01	10.4	5.38E-02	2.2	2.46E-03
	1,2-DCE	2.24E-03	0.1	6.64E-04	<0.1	2.47E-03
	Carbon tetrachloride	5.94E-04	0.3	1.85E-04	0.1	1.90E-04
	Pentachlorophenol	3.35E-18	<0.1	6.06E-19	<0.1	2.34E-02
	TCE	2.26E+01	1,410	4.70E+00	294	1.60E-03
	Vinyl chloride	3.31E-01	10.8	6.90E-02	2.3	3.06E-03
	²³⁷ Np	4.88E+02	No value	9.83E+01	No value	No value
	²³⁹ Pu	1.09E+01	No value	2.05E+00	No value	No value
	²²⁶ Ra	2.21E-01	No value	2.16E-02	No value	No value
	⁹⁹ Tc	6.34E+04	No value	1.32E+04	No value	No value
	²³⁰ Th	3.56E-28	No value	1.30E-43	No value	No value
	²³⁴ U	4.51E+03	No value	8.94E+02	No value	No value
	²³⁵ U	4.75E+01	No value	9.45E+00	No value	No value
	²³⁸ U	8.33E+02	No value	1.66E+02	No value	No value
		Total Uranium ^d	6.46E+03	No value	2.13E+03	No value

²³⁰Th = Thorium-230

^a Concentrations for chemicals and compounds in mg/L. Concentrations for radionuclides in pCi/L.

^b Calculated using comparison to risk-based concentration. Contaminants with an HQ greater than 0.1 are considered COCs.

^c Risk-based no action screening value from Appendix A of the Risk Methods Document. In some cases, these updated values differ from those used in calculation in the WAG 3 RI Report. Values for chemicals and components are given in mg/L. Values for radionuclides are given in pCi/L. The value for lead (*) is the MCL.

^d Evaluated as ²³⁸U.

Table E1.13. Estimated Cancer Risks for a Resident from Exposure to Maximum Modeled Concentrations from Sources at SWMU 4

Source	Contaminant	PGDP Plant Boundary		PGDP Property Boundary		Risk-based Concentrations ^c
		Maximum Concentration ^a	Cancer Risk ^b	Maximum Concentration ^a	Cancer Risk ^b	
Surface	Chromium	2.81E-40	No value	1.95E-52	No value	No value
Soil	Copper	4.40E-04	No value	1.40E-04	No value	No value
	Iron	1.97E+00	No value	6.41E-01	No value	No value
	Nickel	2.53E-03	No value	8.45E-04	No value	No value
	²³⁷ Np	5.33E-02	<1.00E-06	1.64E-02	<1.00E-06	5.73E-01
	²³⁹ Pu	4.16E-04	<1.00E-06	1.44E-04	<1.00E-06	2.86E-01
	²³⁴ U	1.37E+00	2.41E-06	4.16E-01	<1.00E-06	5.46E-01
	²³⁸ U	2.67E+00	6.03E-06	8.08E-01	1.82E-06	4.43E-01
	Subsurface	Chromium	1.15E-37	No value	9.22E-53	No value
Soil	Cobalt	3.29E+00	No value	6.46E-01	No value	No value
	Copper	7.32E+00	No value	1.46E+00	No value	No value
	Iron	1.16E+03	No value	2.41E+02	No value	No value
	Lead	8.45E-42	No value	7.54E-53	No value	1.50E-02*
	Lithium	1.76E-03	No value	5.06E-04	No value	No value
	Manganese	5.13E+01	No value	9.46E+00	No value	No value
	Nickel	1.45E-01	No value	4.29E-02	No value	No value
	Strontium	2.54E-05	No value	7.44E-06	No value	No value
	1,1-DCE	2.57E-01	5.47E-03	5.38E-02	1.14E-03	4.70E-05
	1,2-DCE	2.24E-03	No value	6.64E-04	No value	No value
	Carbon tetrachloride	5.94E-04	3.28E-06	1.85E-04	1.02E-06	1.81E-04
	Pentachlorophenol	3.35E-18	<1.00E-06	6.06E-19	<1.00E-06	2.08E-04
	TCE	2.26E+01	1.31E-02	4.70E+00	2.72E-03	1.73E-03
	Vinyl chloride	3.31E-01	9.46E-03	6.90E-02	1.97E-03	3.50E-05
	²³⁷ Np	4.88E+02	8.52E-04	9.83E+01	1.72E-04	5.73E-01
	²³⁹ Pu	1.09E+01	3.81E-05	2.05E+00	7.17E-06	2.86E-01
	²²⁶ Ra	2.21E-01	2.21E-06	2.16E-02	<1.00E-06	1.00E-01
	⁹⁹ Tc	6.34E+04	4.53E-03	1.32E+04	9.43E-04	1.40E+01
	²³⁰ Th	3.56E-28	<1.00E-06	1.30E-43	<1.00E-06	4.24E-01
	²³⁴ U	4.51E+03	8.26E-03	8.94E+02	1.64E-03	5.46E-01
	²³⁵ U	4.75E+01	8.83E-05	9.45E+00	1.76E-05	5.38E-01
	²³⁸ U	8.33E+02	1.88E-03	1.66E+02	3.75E-04	4.43E-01
		Total Uranium ^d	6.46E+03	1.46E-02	2.13E+03	4.81E-03

²³⁰Th = Thorium-230

^a Concentrations for chemicals and compounds in mg/L. Concentrations for radionuclides in pCi/L.

^b Calculated using comparison to risk-based concentration. Contaminants with a cancer risk greater than 1.00E-06 are considered COCs.

^c Risk-based no action screening value from Appendix A of the Risk Methods Document. In some cases, these updated values differ from those used in calculation in the WAG 3 RI Report Values for chemicals and components in mg/L. Values for radionuclides are given in pCi/L. The value for lead (*) is the MCL.

^d Evaluated as ²³⁸U.

E1.4. MODELING APPEARING IN THE WAG 3 RI REPORT FOR SWMU 5

The conservative modeling in Appendix B of Volume 4 of the WAG 3 RI Report (DOE 2000a) was completed to determine if any contaminants could migrate from source areas at SWMU 5 to POEs at the plant boundary and property boundary at a rate that could result in maximum concentrations greater than risk-based screening levels. This modeling was completed using MEPAS and conservative source term estimates developed using comparisons of sampling results to background concentrations and SSLs for protection of groundwater taken from EPA sources. MEPAS transport parameters are given in Table E1.14.

Table E1.14. MEPAS Transport Parameters for SWMU 5

Input Parameter Description	Parameter Name	Value	Reference
<i>Topsoil parameters (wt)</i>			
Textural classification	WT-CLASS	Silt loam	McCracken Co. Soil Survey (USDA 1976)
Percent sand (%)	WT-SAND	15	McCracken Co. Soil Survey: conservative estimate (highest % sand)
Percent silt (%)	WT-SILT	80	Maximum % silt for soil type
Percent clay (%)	WT-CLAY	5	= 100% -% sand - % silt
Percent organic matter (%)	WT-OMC	0.05	CERCLA Phase II Site Investigation (CH2M HILL 1992)
Percent iron and aluminum (%)	WT-IRON	4	Background Concentrations and Human Health Risk-Based Screening Criteria for Metals in Soil at PGDP (DOE 1995b)
pH of topsoil	WT-pH	8.25	WAG 3 RI data
Percent vegetative cover off-site (%)	WT-VEGCOV	100	SWMU Maps
Topsoil water capacity	WT-AVAILW	2.44	McCracken Co. Soil Survey = available water capacity (0.20 in./in.) × root zone depth from Table 2.1 MEPAS Guidance (12.2 in.) × vegetative cover (100%)
SCS curve number	WT-SCSN	71	Antecedent Moisture Condition = II (normal moisture); Group C hydrologic soil group; vegetated surface, well vegetated, 60–100% vegetated
<i>Properties of the partially saturated zones (wp)</i>			
Thickness (ft)	WP-THICK	WP1 39 WP2 20	WP1=1–40 ft (HU 1 + HU 2) WP2=HU 3 Boring logs at SWMU 5
Textural classification	WP-CLASS	WP1 sandy clay loam WP2 clay loam	Boring logs at SWMU 5
Sand (%)	WP-SAND	WP1 = 38 WP2 = 10	Boring logs at SWMU 5
Silt (%)	WP-SILT	WP1 = 27 WP2 = 30	Boring logs at SWMU 5
Clay (%)	WP-CLAY	WP1 = 35 WP2 = 60	Boring logs at SWMU 5

Table E1.14. MEPAS Transport Parameters for SWMU 5 (Continued)

Input Parameter Description	Parameter Name	Value	Reference
Organic matter content in soil (%)	WP-OMC	0.05	WAG 6 geotechnical data
Iron + aluminum in soil (%)	WP-IRON	4	DOE 1995b
pH of pore water in partially saturated zone	WP-pH	WP1 = 6 WP2 = 6.56	DOE 1995b and WAG 3 RI data for WP2
Bulk density(g/cm ³)	WP-BULKD	WP1 = 1.76 WP2 = 2.25	WAG 3 geotechnical data available for WP1; 2.65 × (1-Porosity)
Total porosity (%)	WP-TOTPOR	WP1 = 33.7 WP2 = 15	WAG 3 geotechnical data available for WP1; SWMU 6 boring logs used as estimate for WP2
Field capacity (%)	WP-FIELDC	WP1 = 24 WP2 = 10	Table 2.1 of MEPAS Guidance, based on soil type for WP1; SWMU 5 boring logs used as estimate for WP2
Longitudinal dispersivity (ft)	WP-LDISP	WP1 = 0.39 WP2 = 0.20	Estimated based on MEPAS guidance: D _L = 0.01 × thickness
Saturated hydraulic conductivity (ft/day)	WP-CONDUCT	<u>ft/day</u> 0.3 <u>cm/sec</u> 1.06E-4	WAG 3 Work Plan
Moisture content (%)	WS-MOISTC	WP1 = 33.7 WP2 = 15	Moisture content = total porosity
<i>Properties of the saturated zone (wz)</i>			
Textural classification	WZ-CLASS	Loamy sand	WAG 3 Work Plan
Sand (%)	WZ-SAND	74	WAG 3 Work Plan
Silt (%)	WZ-SILT	17	WAG 3 Work Plan
Clay (%)	WZ-CLAY	9	WAG 3 Work Plan
Organic matter in soil (%)	WZ-OMC	0.02	WAG 3 Work Plan
Iron + aluminum in soil (%)	WZ-IRON	3	WAG 3 Work Plan
pH of pore water in saturated zone	WZ-pH	6.47	WAG 3 RI data
Total porosity (%)	WZ-TOTPOR	37	WAG 3 Work Plan
Effective porosity (%)	WZ-EFFPOR	30	Conservative estimate
Darcy velocity (ft/day)	WZPVELOC	0.6	Conservative estimate; uses conductivity of 1500 ft/day and gradient of 0.0004
Thickness (ft)	WZ-THICK	40	RGA (HU 4 + HU 5) interval: 60–100 ft bgs
Bulk density (ft)	WZ-BULKD	1.67	WAG 3 Work Plan
Travel distance (ft)	WZ-DIST	890 ft to PGDP boundary 2,780 ft to DOE property boundary	Distances measured along the groundwater flow direction from the northern perimeter of the SWMU to the PGDP boundary and to the DOE property boundary
Longitudinal dispersivity (ft)	WZ-LDISP	50.0	Reference: Bioplume groundwater model
Transverse dispersivity (ft)	WZ-TDISP	5.0	Reference: Bioplume groundwater model
Vertical dispersivity (ft)	WZ-VDISP	0.1	Conservative estimate
Percent of total flux to aquifer (%)	WZ-FRACT	100	Conservative estimate
Perpendicular distance from groundwater flow to receptor (ft)	WZ-YDIST	0	(Plume centerline concentrations)
Vertical distance below groundwater table (ft)	WZ-AQDEPTH	0	(Most conservative result)

The sampling results used in source term development were taken from sampling completed as part of the WAG 3 RI (DOE 2000), the Data Gaps Investigation Report (DOE 2000b), and from earlier sampling completed in support of the PGDP CERCLA SI performed in the early 1990s (CH2M HILL 1991 and 1992). Source terms developed for SWMU 5 are presented in Table E1.15. As noted in the modeling report, “In all cases, modelers applied conservatism (worst case) in the definition of the extent of the source zones. In all cases, the maximum concentrations were used to develop each contaminant source-term inventory.”

Three model layers, two partially saturated and one saturated were delineated at SWMU 5. The partially saturated layer includes the loess deposits making up HU 1, the permeable but discontinuous sand and gravel lenses of the UCRS, and a silty clay aquitard, HU 3 (1–60 ft bgs). The saturated layer consists of the RGA and extends from 60 ft to 100 ft bgs. The travel distance from the source to each downgradient exposure point is 890 ft to the PGDP boundary and 2,780 ft to the DOE property boundary.

Surface and subsurface soil data provided by the WAG 3 RI (DOE 2000a) and the PGDP CERCLA SI (CH2M Hill 1991 and 1992) were used to develop the source terms and inventories for the site contaminants. Table E1.15 presents the source terms used in the MEPAS modeling for SWMU 5. Metals, organic compounds, and radionuclides were identified as present above screening levels in surface soils at SWMU 5. Originally identified contaminants that were not referenced in the MEPAS chemical database and, therefore, were not modeled included 3-nitrobenzenamine, benzo(ghi)perylene, and dibenzofuran. The results of the MEPAS modeling for SWMU 5 are presented in Table E1.16. Estimated HQs and cancer risks from these modeling results are not available.

Iron is projected to contribute to the RGA from three distinct sources. Results of modeling the sources to the PGDP boundary are 49.8 mg/L in 1,411 years, 18.8 mg/L in 1,591 years, and 464 mg/L in 1,873 years. At the DOE boundary, concentrations from these sources are 18.4 mg/L in 1,602 years, 6.61 mg/L in 1,871 years, and 82.7 mg/L in 2,069 years. Manganese is projected to contribute to the RGA, resulting in 11.54 mg/L at the PGDP boundary in 2,536 years and 7.53 mg/L at the DOE boundary in 2,952 years. Contributions to the RGA from other constituents are minor. ⁹⁹Tc is projected to contribute to the RGA resulting in 229 pCi/L at the PGDP boundary in 130.1 years and 99.6 pCi/L in 138.6 years.

Table E1.15. Development of Source Terms for SWMU 5

Contaminant	Initial source concentration	Length parallel to flow direction (ft)	Width perpendicular to flow direction (ft)	Thickness (ft)	Contaminant inventory calculation for MEPAS			Notes
					Volume (cm ³)	Bulk density (g/cm ³)	Inventory (g)	
<i>Surface soil</i>								
Aluminum	13,800 mg/kg	235	840	1	5.59E+09	1.46	1.13E+08	Entire SWMU boundary chosen as source term.
2-Methylnaphthalene	150 µg/kg						1.22E+03	
3-Nitrobenzenamine	9,450 µg/kg						7.71E+04	
Acenaphthylene	9,450 µg/kg						7.71E+04	
Benz(a)anthracene	19,000 µg/kg						1.55E+05	
Benzo(a)pyrene	24,800 µg/kg						2.02E+05	
Benzo(b)fluoranthene	49,200 µg/kg						4.02E+05	
Benzo(ghi)perylene	14,600 µg/kg						1.19E+05	
Dibenzofuran	3,520 µg/kg						2.87E+04	
Pentachlorophenol	357 µg/kg						2.91E+03	
PCB-1260	306 µg/kg						2.50E+03	
Phenanthrene	34,600 µg/kg						2.82E+05	
⁹⁹ Tc	5.85 pCi/g	235	775	1	5.16E+09	1.46	4.40E-02	Entire length of SWMU chosen as source term for ⁹⁹ Tc and approximately 3/4 width.
<i>Subsurface soil partially saturated zone WPI (HU1 + HU2 Soils)</i>								
Aluminum	12,400 mg/kg	235	840	39	2.18E+11	1.76	4.76E+09	Entire area and thickness of SWMU used for modeling.
Chromium	296 mg/kg	175	215	14	1.49E+10	1.76	7.77E+06	Metals values detected above screening levels at 005-017 in a 20–23 ft sample.
Iron	29,200 mg/kg						7.67E+08	
Cobalt	24.7 mg/kg	90	240	3	1.83E+09	1.76	7.98E+04	Metals detected above screening levels
Iron	33,100 mg/kg						2.49E+08	at H263 in shallow samples.
Manganese	975 mg/kg						7.35E+06	

*Bulk density of 1.46 used in SWMU 1 MEPAS modeling

Table E1.15. Development of Source Terms for SWMU 5 (Continued)

Contaminant	Initial source concentration	Length parallel to flow direction (ft)	Width perpendicular to flow direction (ft)	Thickness (ft)	Contaminant inventory calculation for MEPAS			Notes
					Volume (cm ³)	Bulk density (g/cm ³)	Inventory (g)	
Benzo(ghi)perylene	260 µg/kg	90	240	7	4.28E+09	1.76	1.96E+03	Semivolatiles detected above
Dibenzofuran	87 µg/kg						6.56E+02	screening levels at H263 in shallow samples.
Phenanthrene	1,300 µg/kg	235	370	20	4.92E+10	1.76	9.80E+03	Just over 1/2 width of SWMU used for source term delineation and entire length.
²²⁶ Ra	2.2 pCi/g						1.91E-01	
²³⁸ U	2 pCi/g	235	370	31	7.63E+10	1.76	2.69E-01	
<i>Subsurface soil partially saturated zone WP2 (HU3 Soils)</i>								
Aluminum	16,400 mg/kg	235	840	20	1.12E+11	2.25	4.13E+09	Entire SWMU boundary chosen as source term.
Cobalt	19.4 mg/kg						4.88E+06	
Iron	29,400 mg/kg						7.40E+09	
Manganese	1,750 mg/kg						4.40E+08	
Toluene	7 µg/kg	110	310	6	5.79E+09	2.25	9.12E+01	Detected only at H002 in 36-42 ft sample.
²²⁶ Ra	1.73 pCi/g	235	370	20	4.92E+10	2.25	1.92E-01	Just over 1/2 width of SWMU used for source term delineation and entire length.
⁹⁹ Tc	3.89 pCi/g	235	370	20	4.92E+10	2.25	4.31E-01	
²³⁸ U	1.71 pCi/g	235	630	20	8.38E+10	2.25	3.23E-01	

Table E1.16. MEPAS Results for SWMU 5

Source	Constituent (Daughter products are denoted with an asterisk)	PGDP boundary			DOE property boundary		
		Potential maximum concentration (mg/L or pCi/L)	Hazard Quotient	Cancer Risk	Potential maximum concentration (mg/L or pCi/L)	Hazard Quotient	Cancer Risk
Surface Soil	Aluminum	0	0.00E+00	No value	0	0.00E+00	No value
	PCB-1260	0	0.00E+00	0.00E+00	0	0.00E+00	0.00E+00
	2-Methylnaphthalene	3.88E-05	1.38E-03	No value	7.00E-06	2.49E-04	No value
	Acenaphthylene	4.35E-03	No value	No value	8.05E-04	No value	No value
	Benz(a)anthracene	0	No value	0.00E+00	0	No value	0.00E+00
	Benzo(a)pyrene	0	No value	0.00E+00	0	No value	0.00E+00
	Benzo(b)fluoranthene	0	No value	0.00E+00	0	No value	0.00E+00
	Pentachlorophenol	1.08E-27	No value	6.00E-30	1.25E-28	No value	6.94E-31
	Phenanthrene	2.62E-03	No value	No value	3.69E-04	No value	No value
⁹⁹ Tc	5.78E+01	No value	3.18E-06	9.65E+00	No value	5.30E-07	
UCRS- WP1	Aluminum	0	0.00E+00	No value	0	0.00E+00	No value
	Chromium	0	0.00E+00	No value	0	0.00E+00	No value
	Cobalt	2.51E-05	1.21E-04	No value	7.61E-06	3.66E-05	No value
	Iron	4.98E+01	1.60E+01	No value	1.84E+01	5.92E+00	No value
	Iron (at H263)	1.88E+01	6.05E+00	No value	6.61E+00	2.13E+00	No value
	Manganese	2.32E-01	1.81E+02	No value	8.44E-02	6.59E+01	No value
	Phenanthrene	6.09E-05	No value	No value	1.64E-05	No value	No value
	²²⁶ Ra	5.59E-03	No value	5.53E-06	1.13E-03	No value	1.12E-06
	²³⁸ U	5.14E-19	No value	8.92E-25	2.13E-19	No value	3.70E-25
	* ²³⁴ Th	5.14E-19	No value	No value	2.13E-19	No value	No value
	* ²³⁴ U	2.81E-20	No value	3.95E-26	1.21E-20	No value	1.70E-26
	* ²³⁰ Th	2.40E-21	No value	2.31E-27	1.07E-21	No value	1.03E-27
	* ²²⁶ Ra	1.93E-21	No value	1.91E-24	8.61E-22	No value	8.52E-25
	* ²²² Rn	1.93E-21	No value	No value	8.61E-22	No value	No value
	* ²¹⁰ Pb	1.92E-21	No value	No value	8.58E-22	No value	No value
	* ²¹⁰ Bi	1.92E-21	No value	No value	8.58E-22	No value	No value
* ²¹⁰ Po	1.92E-21	No value	No value	8.58E-22	No value	No value	
UCRS- WP2	Aluminum	0	0.00E+00	No value	0	0.00E+00	No value
	Cobalt	1.89E-03	9.09E-03	No value	2.81E-04	1.35E-03	No value
	Iron	4.64E+02	1.49E+02	No value	8.27E+01	2.66E+01	No value
	Manganese	1.56E+01	1.22E+04	No value	2.76E+00	2.16E+03	No value
	Toluene	2.78E-05	4.32E-05	No value	1.19E-05	1.85E-05	No value
	⁹⁹ Tc	2.29E+02	No value	1.26E-05	9.96E+01	No value	5.47E-06
	²²⁶ Ra	5.33E-03	No value	5.28E-06	1.04E-03	No value	1.03E-06
	²³⁸ U	9.95E-19	No value	1.73E-24	1.91E-19	No value	3.32E-25
	* ²³⁴ Th	9.95E-19	No value	No value	1.91E-19	No value	No value
	* ²³⁴ U	5.45E-20	No value	7.65E-26	1.09E-20	No value	1.53E-26
	* ²³⁰ Th	4.64E-21	No value	4.46E-27	9.57E-22	No value	9.20E-28
	* ²²⁶ Ra	3.71E-21	No value	3.67E-24	7.72E-22	No value	7.64E-25
	* ²²² Rn	3.71E-21	No value	No value	7.72E-22	No value	No value
	* ²¹⁰ Pb	3.69E-21	No value	No value	7.69E-22	No value	No value
	* ²¹⁰ Bi	3.69E-21	No value	No value	7.69E-22	No value	No value
	* ²¹⁰ Po	3.69E-21	No value	No value	7.69E-22	No value	No value

²¹⁰Bi = Bismuth-210

²¹⁰Pb = Lead-210

²¹⁰Po = Polonium-210

²²²Rn = Radon-222

²³⁰Th = Thorium-230

²³⁴Th = Thorium-234

E1.5. MODELING APPEARING IN THE WAG 3 RI REPORT FOR SWMU 6

The conservative modeling in Appendix B of Volume 4 of the WAG 3 RI Report (DOE 2000a) was completed to determine if any contaminants could migrate from source areas at SWMU 6 to POEs at the plant boundary and property boundary at a rate that could result in maximum concentrations greater than risk-based screening levels. This modeling was completed using MEPAS and conservative source term estimates developed using comparisons of sampling results to background concentrations and SSLs for protection of groundwater taken from EPA sources. MEPAS transport parameters are given in Table E1.17.

The sampling results used in source term development were taken from sampling completed as part of the WAG 3 RI (DOE 2000a), the Data Gaps Investigation Report (DOE 2000b), and from earlier sampling completed in support of the PGDP CERCLA SI (CH2M HILL 1991 and 1992) performed in the early 1990s. Source terms developed for SWMU 6 are presented in Table E1.18. Benzo(ghi)perylene originally was identified as a contaminant to be modeled, but it was not referenced in the MEPAS chemical database and, therefore, could not be modeled. As noted in the modeling report, "In all cases, modelers applied conservatism (worst case) in the definition of the extent of the source zones. In all cases, the maximum concentrations were used to develop each contaminant source-term inventory."

Three model layers (two partially saturated and one saturated) were delineated at SWMU 6 (see Sect. 3.2.3 of Vol. 1). The first partially saturated layer extends to a depth of 40 ft bgs and includes the loess deposits making up HU1 and the HU2; the second partially saturated layer extends to a depth of 60 ft bgs and includes the silty clay aquitard, the HU3. The saturated layer includes the RGA and extends from an average depth of 60 ft to 100 ft bgs.

The travel distances from the source to each downgradient exposure point are 920 ft to the PGDP boundary and 2,820 ft to the DOE property boundary. The direction of groundwater flow in the RGA was assumed to be north, based on potentiometric maps of the area.

The results of the MEPAS modeling conducted for SWMU 6 are presented in Table E1.19. These results indicate that contributions from constituents in surface soil to groundwater in the RGA are negligible. Estimated HQs and cancer risks from these modeling results are not available.

Iron is contributing to the RGA from three distinct sources. Results from the sources to the PGDP boundary are 60.1 mg/L in 1,966 years, 32.8 mg/L in 1,787 years, and 7.77 mg/L in 1,787 years. At the DOE boundary, concentrations from these sources are 21.2 mg/L in 2,171 years, 11.9 mg/L in 2,076 years, and 2.56 mg/L in 2076 years. ⁹⁹Tc contamination from the SWMU 6 waste cell is predicted by the model to reach a maximum activity of 91.5 pCi/L at the PGDP boundary in 118.6 years and 31.8 pCi/L at the DOE property boundary in 120.1 years. Contributions to the RGA from other constituents are minor.

Table E1.17. MEPAS Transport Parameters for SWMU 6

Input Parameter Description	Parameter Name	Value	Reference	
<i>Topsoil parameters (wt)</i>				
Textural classification	WT-CLASS	Silt loam	McCracken Co. Soil Survey (USDA 1976)	
Percent sand (%)	WT-SAND	15	McCracken Co. Soil Survey: conservative estimate (highest % sand)	
Percent silt (%)	WT-SILT	80	Maximum % silt for soil type	
Percent clay (%)	WT-CLAY	5	= 100% -% sand - % silt	
Percent organic matter (%)	WT-OMC	0.05	CERCLA Phase II Site Investigation (CH2M HILL 1992)	
Percent iron and aluminum (%)	WT-IRON	4	Background Concentrations and Human Health Risk-Based Screening Criteria for Metals in Soil at PGDP (DOE 1995b)	
pH of topsoil	WT-pH	7.98	WAG 3 RI Data	
Percent vegetative cover of site (%)	WT-VEGCOV	90	SWMU Maps	
Topsoil water capacity	WT-AVAILW	2.20	McCracken Co. Soil Survey = available water capacity (0.20 in./in.) × root zone depth from Table 2.1 MEPAS Guidance (12.2 in.) × vegetative cover (100%)	
SCS curve number	WT-SCSN	71	Antecedent Moisture Condition = II (normal moisture); Group C hydrologic soil group; vegetated surface, well vegetated, 60–100% vegetated	
<i>Properties of the partially saturated zones (wp)</i>				
Thickness (ft)	WP-THICK	WP1 39 WP2 20	WP1=1–40 ft (HU 1 + HU 2) WP2= HU 3	Boring logs at SWMU 6
Textural classification	WP-CLASS	WP1 sandy clay loam WP2 clay loam	Boring logs at SWMU 6	
Sand (%)	WP-SAND	WP1 = 38 WP2 = 10	Boring logs at SWMU 6	
Silt (%)	WP-SILT	WP1 = 27 WP2 = 30	Boring logs at SWMU 6	
Clay (%)	WP-CLAY	WP1 = 35 WP2 = 60	Boring logs at SWMU 6	
Organic matter content in soil (%)	WP-OMC	0.05	WAG 6 geotechnical data	
Iron + aluminum in soil (%)	WP-IRON	4	DOE 1995b	
pH of pore water in partially saturated zone	WP-pH	WP1 = 6.76 WP2 = 6.29	WAG 3 RI data	
Bulk density(g/cm ³)	WP-BULKD	WP1 = 1.66 WP2 = 2.25	WAG 3 geotechnical data available for WP1; 2.65 × (1-Porosity)	
Total porosity (%)	WP-TOTPOR	WP1 = 37.19 WP2 = 15	WAG 3 geotechnical data available for WP1; SWMU 6 boring logs used as estimate for WP2	
Field capacity (%)	WP-FIELDC	WP1 = 24 WP2 = 10	Table 2.1 of MEPAS Guidance, based on soil type for WP1; SWMU 6 boring logs used as estimate for WP2	

Table E1.17. MEPAS Transport Parameters for SWMU 6 (Continued)

Input Parameter Description	Parameter Name	Value	Reference
Longitudinal dispersivity (ft)	WP-LDISP	WP1 = 0.39 WP2 = 0.20	Estimated based on MEPAS guidance: $D_L = 0.01 \times \text{thickness}$
Saturated hydraulic conductivity (ft/day)	WP-CONDOC	<u>ft/day</u> 0.3 <u>cm/sec</u> 1.06E-4	WAG 3 Work Plan
Moisture content (%)	WS-MOISTC	WP1 = 37.19 WP2 = 15	Moisture content = total porosity
<i>Properties of the saturated zone (wz)</i>			
Textural classification	WZ-CLASS	Loamy sand	WAG 3 Work Plan
Sand (%)	WZ-SAND	74	WAG 3 Work Plan
Silt (%)	WZ-SILT	17	WAG 3 Work Plan
Clay (%)	WZ-CLAY	9	WAG 3 Work Plan
Organic matter in soil (%)	WZ-OMC	0.02	WAG 3 Work Plan
Iron + aluminum in soil (%)	WZ-IRON	3	WAG 3 Work Plan
pH of pore water in saturated zone	WZ-pH	6.275	WAG 3 RI data
Total porosity (%)	WZ-TOTPOR	37	WAG 3 Work Plan
Effective porosity (%)	WZ-EFFPOR	30	Conservative estimate
Darcy velocity (ft/day)	WZ-PVELOC	0.6	Conservative estimate; uses conductivity of 1500 ft/day and gradient of 0.0004
Thickness (ft)	WZ-THICK	40	RGA (HU 4 + HU 5) interval: 60–100 ft bgs
Bulk density (g/cm^3)	WZ-BULKD	1.67	WAG 3 Work Plan
Travel distance (ft)	WZ-DIST	920 ft to PGDP boundary 2820 ft to DOE property boundary	Distances measured along the groundwater flow direction from the northern perimeter of the SWMU to the PGDP boundary and to the DOE property boundary
Longitudinal dispersivity (ft)	WZ-LDISP	50.0	Reference: Bioplume groundwater model
Transverse dispersivity (ft)	WZ-TDISP	5.0	Reference: Bioplume groundwater model
Vertical dispersivity (ft)	WZ-VDISP	0.1	Conservative estimate
Percent of total flux to aquifer (%)	WZ-FRACT	100	Conservative estimate
Perpendicular distance from groundwater flow to receptor (ft)	WZ-YDIST	0	(Plume centerline concentrations)
Vertical distance below groundwater table (ft)	WZ-AQDEPTH	0	(Most conservative result)

Table E1.18. Development of Source Terms for SWMU 6

Contaminant	Initial source concentration	Length parallel to flow direction (ft)	Width perpendicular to flow direction (ft)	Thickness (ft)	Contaminant inventory calculation for MEPAS			Notes
					Volume (cm ³)	Bulk density (g/cm ³)	Inventory (g)	
Surface Soil								
Copper	21.3 mg/kg	65	80	1	1.47E+08	1.46	4.58E+03	Detected above screening levels in 006-001.
Benzo(ghi)perylene	124 µg/kg						2.67E+01	
Phenanthrene	461 µg/kg						9.91E+01	
⁹⁹ Tc	18.8 pCi/g	115	85	1	2.77E+08	1.46	7.60E-03	Area for modeling encompasses 006-016 and 006-017.
SWMU 6 Waste Cells								
Aluminum	18,800 mg/kg	40	100	10	1.13E+09	1.66	3.53E+07	Source term is waste cell "J."
PCB-1016	255 µg/kg						4.79E+02	
²³⁷ Np	0.219 pCi/g						4.12E-04	
⁹⁹ Tc	43.3 pCi/g						8.14E-02	
²³⁸ U	1.52 pCi/g						2.86E-03	
Subsurface soil partially saturated zone WPI (HU1 + HU2 Soils)								
Chromium	116 mg/kg	45	60	17	1.30E+09	1.66	2.50E+05	Area for modeling surrounds 006-027, the area of a relatively high detection of chromium.
							9.01E+05	
Aluminum	56.8 mg/kg	90	150	25	9.56E+09	1.66	2.99E+08	Area for modeling is entire area of SWMU
Cobalt	12,100 mg/kg			39	1.49E+10		1.36E+05	
Copper	17.9 mg/kg			12	4.59E+09		3.71E+05	
Iron	20.9 mg/kg			28	1.07E+10		1.45E+09	
Lead	58,700 mg/kg			39	1.49E+10		2.02E+05	
Manganese	35.4 mg/kg			9	3.44E+09		1.18E+07	
⁹⁹ Tc	1,550 mg/kg			12	4.59E+09		9.90E-03	
²³⁷ Np	8.51 pCi/g	50	45	11	7.01E+08	1.66	1.45E-04	Detected above screening levels at 006-010.
²³⁸ U	0.125 pCi/g						2.00E-03	

Table E1.18. Development of Source Terms for SWMU 6 (Continued)

Contaminant	Initial source concentration		Length parallel to flow direction (ft)	Width perpendicular to flow direction (ft)	Thickness (ft)	Contaminant inventory calculation for MEPAS			Notes
						Volume (cm ³)	Bulk density (g/cm ³)	Inventory (g)	
Subsurface soil partially saturated zone WP2 (HU3 Soils)									
Aluminum	22,500	mg/kg	80	185	20	8.38E+09	2.25	4.24E+08	Area for modeling is entire area of SWMU, less southeastern portion.
Cobalt	156	mg/kg						2.94E+06	
	32,900	mg/kg						6.20E+08	
Iron	25.2	mg/kg						4.75E+05	
Lead	36,900	mg/kg	45	60	20	1.53E+09	2.25	1.27E+08	
Iron									Detected above screening level at 006-027, outside SWMU boundary.

Table E1.19. MEPAS Results for SWMU 6

Source	Constituent (Daughter products are denoted with an asterisk)	PGDP boundary			DOE property boundary		
		Potential maximum concentration (mg/L or pCi/L)	Hazard Quotient	Cancer Risk	Potential maximum concentration (mg/L or pCi/L)	Hazard Quotient	Cancer Risk
Surface Soil	Copper	2.56E-12	6.15E-12	No value	2.11E-14	5.07E-14	No value
	Phenanthrene	9.78E-07	No value	No value	2.71E-07	No value	No value
	⁹⁹ Tc	9.71E+00	No value	5.34E-07	3.15E+00	No value	1.73E-07
UCRS- Waste Cells	Aluminum	0	0.00E+00	No value	0	0.00E+00	No value
	PCB-1016	0	0.00E+00	0.00E+00	0	0.00E+00	0.00E+00
	⁹⁹ Tc	9.15E+01	No value	5.03E-06	3.18E+01	No value	1.75E-06
	²³⁷ Np	1.68E-01	No value	4.20E-07	5.53E-02	No value	1.38E-07
	* ²³³ Pa	1.68E-01	No value	No value	5.53E-02	No value	No value
	* ²³³ U	2.45E-04	No value	No value	9.33E-05	No value	No value
	* ²²⁹ Th	3.99E-06	No value	No value	1.69E-06	No value	No value
	* ²²⁵ Ra	3.99E-06	No value	No value	1.68E-06	No value	No value
	* ²²⁵ Ac	3.99E-06	No value	No value	1.68E-06	No value	No value
	²³⁸ U	4.80E-19	No value	8.33E-25	1.42E-19	No value	2.47E-25
	* ²³⁴ Th	4.80E-19	No value	No value	1.42E-19	No value	No value
	* ²³⁴ U	2.66E-20	No value	3.74E-26	8.12E-21	No value	1.14E-26
	* ²³⁰ Th	2.28E-21	No value	2.19E-27	7.22E-22	No value	6.94E-28
	* ²²⁶ Ra	1.83E-21	No value	1.81E-24	5.84E-22	No value	5.78E-25
	* ²²² Rn	1.83E-21	No value	No value	5.84E-22	No value	No value
	* ²¹⁰ Pb	1.82E-21	No value	No value	5.82E-22	No value	No value
* ²¹⁰ Bi	1.82E-21	No value	No value	5.82E-22	No value	No value	
* ²¹⁰ Po	1.82E-21	No value	No value	5.82E-22	No value	No value	
UCRS- WP1	Aluminum	0	0.00E+00	No value	0	0.00E+00	No value
	Chromium	0	0.00E+00	No value	0	0.00E+00	No value
	Cobalt	8.06E-05	3.88E-04	No value	2.33E-05	1.12E-04	No value
	Copper	3.13E-11	7.52E-11	No value	2.44E-13	5.87E-13	No value
	Iron	6.01E+01	1.93E+01	No value	2.12E+01	6.82E+00	No value
	Lead	0	0.00E+00	No value	0	0.00E+00	No value
	Manganese	4.08E-01	3.19E+02	No value	1.41E-01	1.10E+02	No value
	⁹⁹ Tc	1.16E+01	No value	6.37E-07	3.86E+00	No value	2.12E-07
	²³⁷ Np	5.97E-02	No value	4.26E-08	1.95E-02	No value	1.39E-08
	* ²³³ Pa	5.9E-02	No value	No value	1.95E-02	No value	No value
	* ²³³ U	9.02E-05	No value	No value	3.29E-05	No value	No value
	* ²²⁹ Th	1.47E-06	No value	No value	5.95E-07	No value	No value
	* ²²⁵ Ra	1.47E-06	No value	No value	5.95E-07	No value	No value
	* ²²⁵ Ac	1.47E-06	No value	No value	5.95E-07	No value	No value
	²³⁸ U	3.49E-19	No value	6.06E-25	1.00E-19	No value	1.74E-25
	* ²³⁴ Th	3.49E-19	No value	No value	1.00E-19	No value	No value
	* ²³⁴ U	1.93E-20	No value	2.71E-26	5.75E-21	No value	8.08E-27
	* ²³⁰ Th	1.66E-21	No value	1.60E-27	5.11E-22	No value	4.91E-28
	* ²²⁶ Ra	1.33E-21	No value	1.32E-24	4.13E-22	No value	4.09E-25
	* ²²² Rn	1.33E-21	No value	No value	4.13E-22	No value	No value
* ²¹⁰ Pb	1.33E-21	No value	No value	4.12E-22	No value	No value	
* ²¹⁰ Bi	1.33E-21	No value	No value	4.12E-22	No value	No value	
* ²¹⁰ Po	1.33E-21	No value	No value	4.12E-22	No value	No value	

Table E1.19. MEPAS Results for SWMU 6 (Continued)

Source	Constituent (Daughter products are denoted with an asterisk)	PGDP boundary			DOE property boundary		
		Potential maximum concentration (mg/L or pCi/L)	Hazard Quotient	Cancer Risk	Potential maximum concentration (mg/L or pCi/L)	Hazard Quotient	Cancer Risk
UCRS- WP2	Aluminum	0	0.00E+00	No value	0	0.00E+00	No value
	Cobalt	1.66E-03	7.98E-03	No value	4.96E-04	2.38E-03	No value
	Iron	3.28E+01	1.05E+01	No value	1.19E+01	3.83E+00	No value
	Iron (from 006-027)	7.77E+00	2.50E+00	No value	2.56E+00	8.23E-01	No value
	Lead	0	0.00E+00	No value	0	0.00E+00	No value

²²⁵Ac = Actinium-225 ²³³Pa = Protactinium-233 ²³⁰Th = Thorium-230
²¹⁰Bi = Bismuth-210 ²²²Rn = Radon-222 ²³⁴Th = Thorium-234
²¹⁰Pb = Lead-210 ²²⁵Ra = Radium-225 ²³⁵U = Uranium-233
²¹⁰Po = Polonium-210 ²²⁹Th = Thorium-229

E1.6. MODELING APPEARING IN THE WAG 22, SWMUS 7 AND 30 RI/FS

The conservative modeling in Section 5 and Appendix D of the WAG 22, SWMUs 7 and 30 RI/FS Report (DOE 1998a) was completed to determine if any contaminants could migrate from source areas at SWMUs 7 and 30 to groundwater in the UCRS and RGA at a rate that could result in maximum concentrations greater than risk-based screening levels. The following discussion is taken directly from “*Remedial Investigation Report for Solid Waste Management Units 7 and 30 of Waste Area Group 22 at the Paducah Gaseous Diffusion Plant Paducah, Kentucky,*” DOE/OR/07-1604/V1&D2 (DOE 1998a). Source terms developed for SWMUs 7 and 30 are presented in Tables E1.20 through E1.23. Modeling results are discussed for the four source areas, the UCRS, and the RGA. The use of four source areas (designated Pit A, Pit B/C, F Pits, and areas within SWMUs 7 and 30 outside of Pits A, B, C), the UCRS, and the RGA reflects the distribution of contaminants in the SWMU 7 and 30 source areas and surrounding environmental media. Contaminants disposed of in the three primary source areas of SWMUs 7 and 30, Pit A, Pit B/C Pits, and the F Pits include metals, radionuclides (primarily ⁹⁹Tc and uranium), organic solvents (primarily TCE), and fuel-related VOCs and SVOCs. Of the contaminants disposed of in the source areas, only ⁹⁹Tc and several VOCs were detected in the UCRS and RGA. Metals, other radionuclides, and SVOCs were not detected in either unit.

The complex nature of the hydrogeology and contaminants in SWMUs 7 and 30 preclude development of a single computer model to describe fate and transport of contaminants at this site. Rather, a combination of small-scale analytical groundwater transport models and simple estimates of contaminant attenuation/dilution along specific pathways are combined in the framework of the conceptual model for fate and transport analysis.

The output of the contaminant fate and transport modeling is presented as the expected maximum concentration of modeled contaminants at the receptor locations. These data will allow prediction of the approximate locations of future maximum concentrations resulting from the integration of the contributions from multiple sources and different pathways. For the purpose of this analysis, SWMUs 7 and 30 were divided into four source areas representing the disposal areas: (1) Pit A, (2) Pit B/C, (3) F Pits, and (4) the areas within SWMUs 7 and 30 outside of Pits A, B, C. The quantitative modeling accounted for the following:

- Contents of the source area,
- Presence of DNAPL in the source area,
- Presence or absence of a discrete cap,
- Identifiable geologic strata beneath the source area,
- Thickness of each layer in the vadose zone,
- Vertical permeability of the unsaturated soils,
- Presence of contaminated soils submerged in the groundwater,
- Water table fluctuations, and
- Receptor locations.

Once the leachate modeling for these four source areas was completed, the predicted maximum leachate concentrations were compared against the existing groundwater concentrations in the UCRS. If the predicted leachate concentrations exceeded the concentrations in the UCRS, then the leachate concentrations were compared against their respective groundwater risk-based concentrations (RBCs) that were derived using cancer risk of 1E-6 or hazard index of 1.0. For the remaining constituents, the groundwater concentrations in the UCRS were compared against their respective RBCs. All the constituents that exceeded the groundwater RBCs were selected for vertical transport modeling from the UCRS to the RGA. The constituents with concentrations (both in UCRS groundwater and predicted leachate) below the RBC were eliminated from the list of contaminant migration COPCs (CMCOPCs) and no further evaluations were performed. After performing vertical transport modeling of the CMCOPCs from the UCRS to the RGA, the predicted leachate concentrations were again compared with their respective RBCs, and if the maximum predicted leachate concentration of a COPC exceeded the groundwater RBC, it was selected for horizontal transport modeling using AT123D. If the predicted maximum leachate concentration was below the groundwater RBC, the contaminant was eliminated from the list of CMCOPCs, and no further evaluations were performed.

E1.6.1 SESOIL MODELING

The SESOIL model used for leachate modeling, when applicable, estimates pollutant concentrations in the soil profile following introduction via direct application and/or interaction with other media. The model defines the soil compartment as a soil column extending from the ground surface through the unsaturated zone and to the upper level of the saturated soil zone. Processes simulated in SESOIL are categorized in three cycles - the hydrologic cycle, sediment-cycle, and pollutant cycle. Each cycle is a separate submodule in the SESOIL code. The hydrologic cycle includes rainfall, surface runoff, infiltration, soil-water content, evapo-transpiration, and groundwater recharge. The sediment cycle includes sediment washload as a result of rainstorms (i.e., soil erosion that results from surface runoff). The pollutant cycle includes convective transport, volatilization, adsorption/desorption, and degradation/decay. A contaminant in SESOIL can partition in up to four phases (liquid, adsorbed, air, and pure).

Table E1.20. Initial COPCs Based on Soil Screening from the Burial Pit A, WAG 22

Site Related Contaminant	Exposure Concentration^a	SSL	Is Exposure Concentration > SSL?
<i>Metals and Inorganics^b</i>			
Aluminum	79508.96	22,400	YES
Antimony	7.00	0.3	YES
Arsenic	10.00	1	YES
Barium	386.00	82	YES
Cadmium	10.00	0.4	YES
Chromium	55.00	2	YES
Copper	141.00	21.2	YES
Cyanide	0.64	2	NO
Lead	5,760.06	400	YES
Manganese	5,239.00	511	YES
Mercury	0.60	0.02	YES
Molybdenum	0.02	9.43E+00	NO
Nickel	132.00	7	YES
Selenium	1.00	0.3	YES
Tin	10.80	4,020	NO
Vanadium	40.00	300	NO
Zinc	364.05	620	NO
<i>Volatile Organic Compounds^c</i>			
1,1-Dichloroethane	27.04	1,000	NO
1,2-Dichlorobenzene	17.60	N/A	YES
1,2-Dichloroethane	0.91	1	NO
<i>cis</i> -1,2-DCE	20.73	20	YES
<i>trans</i> -1,2-DCE	0.58	30	NO
1,3-Dichlorobenzene	39.42	1,400	NO
1,4-Dichlorobenzene	26.23	100	NO
4-Methy-2pentanone	7.71	200	NO
Acetone	1.71	800	NO
Benzene	2.20	2	YES
Chlorobenzene	88.31	70	YES
Chloroethene	33.62	235.08	NO
Chloromethane	2.19	0.32	YES
Ethylbenzene	9.83	700	NO
Methylene chloride	5.69	1	YES
Toluene	5.52	600	NO

Table E1.20. Initial COPCs Based on Soil Screening from the Burial Pit A, WAG 22 (Continued)

Site Related Contaminant	Exposure Concentration ^a	SSL	Is Exposure Concentration > SSL?
TCE	2.02	3	NO
Vinyl chloride	3.12	0.7	YES
Xylenes	25.82	1,000	NO
<i>Semivolatile Organic Compounds^c</i>			
1,2,4- Trichlorobenzene	9.31	300	NO
2,4-Dichlorophenol	0.65	50	NO
2,4-Dimethylphenol	56.30	400	NO
2-Methylnaphthalene	6.79	N/A	YES
2-Methylphenol	0.57	8.00E+02	NO
4-Methylphenol	0.40	N/A	YES
Acenaphthene	2,046.00	29,000	NO
Anthracene	2,609.00	590,000	NO
Benzo(a)anthracene	5,616.00	80	YES
Benzo(a)pyrene	5,427.00	400	YES
Benzo(a)fluoranthene	15,000.00	200	YES
Benzo(ghi)perylene	4,400.00	4,280	YES
Chrysene	5,804.00	800	YES
Dibenzofuran	1,172.00	81.6	YES
Bis(2-ethylhexyl)phthalate	37,798.72	180,000	NO
Di-n-octyl phthalate	407,184,020.00	1.00E+07	YES
Diethyl phthalate	0.20	2.30E+04	NO
Fluoranthene	23,000.00	210,000	NO
Fluorene	1,796.00	2,8000	NO
Indeno (1,2,3-c,d)pyrene	2,860.00	700	YES
Naphthalene	923.70	4,000	NO
Phenanthrene	13,190.00	N/A	YES
Pyrene	15,000.00	210,000	NO
<i>Pesticides and PCBs^c</i>			
PCB-1254	165.30	1,000	NO
PCB-1260	3,047.26	1,000	YES
<i>Radionuclides^d</i>			
Gross Alpha	24.00	N/A	YES
Gross Beta	35.00	N/A	YES
²³⁷ Np	1.80	0.033	YES
²³⁹ Pu	5.05	1.46	YES
⁹⁹ Tc	18.00	0.0828	YES
²³⁰ Th	38.61	33.99206	YES
²³⁴ U	562.04	2.2	YES
²³⁵ U	1.01	2.16	NO
^{235/236} U	42.79	2.16	YES
²³⁸ U	686.09	2.29	YES

^a Soil exposure concentrations represent either back-calculated soil concentrations or soil concentrations obtained from soil sampling analysis.

^b Concentrations of all metals and inorganics are expressed in mg/kg.

^c Concentrations of all VOCs, SVOCs, and PCBs are expressed in µg/kg.

^d Concentrations of all radionuclides are expressed in pCi/g.

Table E1.21. Initial COPCs Based on Soil Screening from Burial Pits B/C, WAG 22 (Continued)

Site Related Contaminant	Exposure Concentration ^a	SSL	Is Exposure Concentration > SSL?
<i>Metals and Inorganics^b</i>			
Aluminum	1,950,219.70	22400	YES
Antimony	186.36	0.3	YES
Arsenic	41.00	1	YES
Barium	2,322.52	82	YES
Beryllium	31.21	3.00	YES
Cadmium	165.10	0.4	YES
Chromium	371.08	2	YES
Cobalt	3,861.50	1,180	YES
Copper	31,933.45	21.2	YES
Cyanide	0.77	2	NO
Lead	63,520.67	400	YES
Manganese	40,636.65	511	YES
Mercury	2.04	0.0205	YES
Molybdenum	21.28	9.42756	YES
Nickel	122,168.78	7	YES
Silver	65.86	2	YES
Tin	94.54	4020	NO
Uranium	3,544.37	N/A	YES
Vanadium	1,800.30	300.0	YES
Zinc	114,284.86	620	YES
<i>Volatile Organic Compounds^c</i>			
1, 1- Dichloroethane	5.60	1,000	NO
1, 2-DCE (total)	454.78	51.5	YES
<i>cis</i> -1,2-DCE	0.88	20.0	NO
1, 4-Dichlorobenzene	70.00	100	NO
2-Butanone	29.52	126	NO
4-Methyl-2-Pentanone	35.84	200	NO
Acetone	82.51	800	NO
Benzene	34.73	2	YES
Chloroethane	4.74	235.0855	NO
Ethylbenzene	179.50	700	NO
Methylene Chloride	20.52	1	YES
Toluene	753.40	600	YES
TCE	7.00	3	YES
Vinyl chloride	65.80	7.00E-01	YES
Xylenes, Total	1,065.00	10,000	NO
<i>Semivolatile Organic Compounds^c</i>			
1,2,4-Trichlorobenzene	77.00	300	NO
2-Chlorophenol	39.00	200	NO
2,4,5-Trichlorophenol	35.00	14,000	NO
2,4,6-Trichlorophenol	41.00	8	YES
2,4-Dimethylphenol	5,034.00	400	YES
2-Methylnaphthalene	958.30	N/A	YES
2-Methylphenol	664.40	800	NO
4-Methylphenol	8,129.00	N/A	YES
Acenaphthene	31.00	29,000	NO
Benzo(b)fluoranthene	789.60.	200	YES
bis(2-ethylhexyl)phthalate	947.10	180,000	NO
Chlorobenzene	7.00	70	NO

Table E1.21. Initial COPCs Based on Soil Screening from Burial Pits B/C, WAG 22 (Continued)

Site Related Contaminant	Exposure Concentration ^a	SSL	Is Exposure Concentration > SSL?
Chrysene	590.20	800	NO
Di-n-butylphthalate	91.00	270,000	NO
Dibenzofuran	0.79	81.62258881	NO
Diethyl phthalate	0.57	23,000	NO
Fluoranthene	795.90	210,000	NO
Hexachlorobutadiene	58.00	36.6	YES
Hexachloroethane	34.00	12.4	YES
Naphthalene	8.72	4,000	NO
Phenanthrene	967.20	N/A	YES
Phenol	702.60	5,000	NO
Pyrene	609.40	210,000	NO
<i>Pesticides and PCBs^c</i>			
PCB-1248	720.70	1,000	NO
PCB-1254	7,913.80	1,000	YES
PCB-1260	11,375.23	1,000	YES
<i>Radionuclides^d</i>			
Gross Alpha	37.00	NA	YES
Gross Beta	47.00	NA	YES
²³⁷ Np	19.00	0.033	YES
^{239/240} Pu	72.00	1.46	YES
⁹⁹ Tc	656.50	0.0828	YES
²³⁰ Th	95.70	34	YES
²³⁴ U	362.80	2.2	YES
²³⁵ U	150.00	2.16	YES
^{235/236} U	14.28	2.16	YES
²³⁸ U	2,100.00	2.29	YES

^a Soil exposure concentrations represent either back-calculated soil concentrations or soil concentrations obtained from soil sampling analysis.

^b Concentrations of all metals and inorganics are expressed in mg/kg.

^c Concentrations of all VOCs, SVOCs, and PCBs are expressed in µg/kg.

^d Concentrations of all radionuclides are expressed in pCi/g.

Table E1.22. Initial COPCs Based on Soil Screening from Burial Pit F, WAG 22

Site Related Contaminant	Exposure Concentration ^a	SSL	Is Exposure conc. > SSL?
<i>Metals and Inorganics^b</i>			
Aluminum	615,069.29	22,352.98	YES
Barium	180.00	82	YES
Beryllium	12.00	3.00E+00	YES
Cobalt	247.03	1,177.9812	NO
Lead	5,120.05	400	YES
Manganese	1,200.00	51 I	YES
Molybdenum	175.24	9.42756	YES
Nickel	1,891.06	7	YES
Nitrate	0.52	N/A	YES
Tin	3.80	4,020	NO
Uranium	21,013.03	N/A	YES
Vanadium	700.12	300	YES
Zinc	2,080.27	620	YES
<i>Volatile Organic Compounds^c</i>			
1,1,1- Trichloroethane	5.11	1.00E+02	YES
1,1,2- Trichloro-1,2,2, trifluoroethene	3.79	56,085.504	YES
1,1-Dichloroethane	34.55	1.00E+03	YES
cis-1,2-DCE	0.64	2.00E+01	YES
1,3-Dichlorobenzene	0.90	1,400.137787	YES
Acetone	14.00	800	YES
Benzene	1.46	2	YES
Chloroethane	4.35	235.0855911	YES
Ethylbenzene	34.04	700	YES
Tetrachloroethene	0.76	3	YES
Toluene	4.41	600	YES
TCE	0.65	3.00E+00	YES
Vinyl chloride	3.89	7.00E-01	YES
Xylenes	160.17	10,000	YES
<i>Semivolatile Organic Compounds^c</i>			
1,2,4- Trichlorobenzene	37.00	300	NO
2,4-Dimethylphenol	20.15	400	NO
2-Methylnaphthalene	21.62	N/A	YES
4-Methylphenol	0.75	N/A	YES
4-chloro-3-methylphenol	3.70	N/A	YES
Bis(2-ethylhexyl) phthalate	483.64	1.80E+05	NO
Diethyl phthalate	0.24	2.30E+04	NO
Di-n-octylphthalate	72.00	1.00E+07	NO
Napthalene	9.79	4.00E+03	NO
<i>Pesticides and PCBs^c</i>			
PCB-1016	33.02	N/A	YES
PCB-1248	8,522.20	1,000	YES
PCB-1260	565.19	1.00E+03	NO

Table E1.22. Initial COPCs Based on Soil Screening from Burial Pit F, WAG 22 (Continued)

Site Related Contaminant	Exposure Concentration ^a	SSL	Is Exposure conc. > SSL?
<i>Radionuclides^d</i>			
Gross Alpha	23.00	15.8	YES
Gross Beta	42.00	27.9	YES
²³⁷ Np	0.04	0.033012	YES
^{239/240} Pu	2.06	1.464244	YES
⁹⁹ Tc	4.17	0.0828	YES
²³⁰ Th	9.34	33.99206	NO
²³⁴ U	193.42	2.2	YES
^{235/236} U	29.37	2.16	YES
²³⁸ U	1,243.06	2.29	YES

^a Soil exposure concentrations represent either back-calculated soil concentrations or soil concentrations obtained from soil sampling analysis.

^b Concentrations of all metals and inorganics are expressed in mg/kg.

^c Concentrations of all VOCs, SVOCs, and PCBs are expressed in µg/kg.

^d Concentrations of all radionuclides are expressed in pCi/g.

Table E1.23. Initial COPCs Based on Soil Screening from Subsurface Soils Outside Pits

Site Related Contaminant	Exposure Concentration ^a	SSL	Is Exposure conc. > SSL?
<i>Metals and Inorganics^b</i>			
Aluminum	7,242	22,400	NO
Antimony	4.24	0.3	YES
Arsenic	3.143	1	YES
Barium	104.3	82	YES
Beryllium	0.6603	3'	NO
Cadmium	1.85	0.4	YES
Chromium	21.45	2	YES
Cobalt	8.279	1,180	NO
Copper	16.18	21.2	NO
Cyanide	0.393	2	NO
Lead	12.26	400	NO
Manganese	417.7	511	NO
Nickel	15.99	7	YES
Selenium	0.2117	0.3	NO
Silver	0.9728	2	NO
Thallium	0.9309	0.04	YES
TIN	4.6		NO
Vanadium	23.95	300	NO
Zinc	38.95	620	NO
<i>Volatile Organic Compounds^c</i>			
1,2,4- Trichlorobenzene	33	300	NO
1,4-Dichlorobenzene	25	100	NO
2-Butanone	4	126	NO
2-Chlorophenol	23	200	NO
4-Methyl-2-Pentanone	6	200	NO
Acetone	404.3	800	NO
Carbon Disulfide	4.243	2,000	NO
Methylene Chloride	87.11	1	YES
Tetrachloroethene	2	3	NO
TCE	4.108	3	YES

Table E1.23. Initial COPCs Based on Soil Screening from Subsurface Soils Outside Pits (Continued)

Site Related Contaminant	Exposure Concentration ^a	SSL	Is Exposure conc. > SSL?
<i>Semivolatile Organic Compounds^c</i>			
Acenepthene	17	29,000	NO
Benzo(a)anthracene	56	80	NO
Benzo(a)pyrene	77	400	NO
Benzo(b)fluoroanthene	91	200	NO
Benzo(k)fluoroanthene	81	2,000	NO
bis(2-ethylhexyl)phthalate	235.9	180,000	NO
Chrysene	83	800	NO
Di-n-butylphthalate	181.7	270,000	NO
Fluoroanthene	170	210,000	NO
Indeno (1,2,3-c,d)pyrene	73	700	NO
N-nitrosodiphenylamine	140	26.4	YES
Octachlorodibenzo-p-dioxin	0.6	0.16	YES
Pentachlorophenol	120	5	YES
Phenanthrene	110	1	NO
Pyrene	140		NO
Indeno (1,2,3-c,d)pyrene	73	210,000	NO
<i>Pesticides and PCBs^c</i>			
PCB-1248	27.68	1,000	NO
PCB-1260	55.93	1,000	NO
<i>Radionuclides^d</i>			
Gross Alpha	11.92		
Gross Beta	18.49		
²³⁷ Np	2.362	0.033	YES
²³⁹ Pu	5.986	1.46	YES
⁹⁹ Tc	280	0.0828	YES
²³⁰ Th	22.38	34 2	NO
²³⁴ U	44	.2	YES
²³⁵ U	3.2	2.16	YES
^{235/236} U	0.4	2.16	NO
²³⁸ U	160	2.29	YES

^a Soil exposure concentrations represent either back-calculated soil concentrations or soil concentrations obtained from soil sampling analysis.

^b Concentrations of all metals and inorganics are expressed in mg/kg.

^c Concentrations of all VOCs, SVOCs, and PCBs are expressed in µg/kg.

^d Concentrations of all radionuclides are expressed in pCi/g.

E1.6.1.1 Source Areas

Although 27 constituents from Pit A were identified as the initial COPCs, only 7 of them were selected for SESOIL modeling. Similarly, 14 of the 38 initial COPCs from Pits B/C, 4 of 26 initial COPCs from the F Pits, and 7 of 18 initial COPCs from Subsurface Outside of the Pits were selected for SESOIL Modeling. The model was calibrated against the percolation rate by varying the hydraulic conductivity and the disconnectedness index and keeping all other site-specific geotechnical parameters fixed. The final parameter values used in this modeling are as follows: soil bulk dry density of 1.5 g/cm³, porosity of 0.40, organic carbon content of 0.34%, and volumetric moisture content of 27.5%. Additional parameter values used in the model included a disconnectedness index of 10.0 and an intrinsic permeability of 9.0×10^{-10} cm², which was derived during calibration of the model to a percolation rate of 4.6 inches/year. The percolation rate was derived using water balance data for the site (Geotrans 1992).

The SESOIL model was set up using four layers extending from the ground surface to the average water table surface at 12 ft bgs. The first layer of the model extended from ground surface to 1 ft bgs and corresponds to the observed soil cover over the pits. The second layer extended from 1 ft bgs to 5 ft bgs and corresponds to the sampling interval; therefore, this layer represents the loading zone. The third layer extended from 5 ft bgs to 10 ft bgs. Most of the pit water was collected in this interval, which was used to back-calculate to corresponding soil concentrations; therefore, this layer also represents the loading zone. The fourth layer extended from 10 to 12 ft bgs, formed the leaching zone, and was divided into 5 sub layers for better resolution.

E1.6.1.2 UCRS

SESOIL-predicted maximum leachate concentrations from the individual source areas were compared against the currently observed maximum groundwater concentrations, as stated earlier, and the source term concentrations for transporting the contaminants vertically down to the RGA were developed. However, only 17 of 42 initial COPCs from the UCRS were selected for SESOIL modeling. As before, the model was calibrated against the percolation rate by varying the hydraulic conductivity and the disconnectedness index and keeping all other site-specific geotechnical parameters fixed. The final parameter values used in the modeling from UCRS are as follows: soil bulk dry density of 1.5 g/cm³, porosity of 0.40, organic carbon content of 0.26%, and volumetric moisture content of 29.5%. Additional parameter values used in the model included a disconnectedness index of 10.0 and an intrinsic permeability of 1.65×10^{-10} cm². Of these parameters, porosity, density, and disconnectedness index represent default values for silty-clay, and organic carbon content represents the average measured value. The volumetric moisture content and the intrinsic permeability were derived during calibration of the model to a percolation rate of 4.3 inches/year. The percolation rate was derived using water balance data for the site (Geotrans 1992).

The SESOIL model was set up using three layers extending from the top of the HU2 to the top of the RGA at 45 ft bgs. The first layer of the model extended from top of the HU2 to the top of the confining zone and corresponds to the contaminated zone. This layer was divided into five sublayers and contaminant loading was performed in each of these sub layers which represented the back-calculated soil concentrations. The second layer extended from the top of the confining zone to the top of the RGA and formed the leaching zone. This layer was also divided into five sublayers for better resolutions. The third layer of 0.5 ft was used to read the output concentrations at the water table.

E1.6.1.3 Modeling Results

The results of contaminant fate and transport analysis for individual source areas are summarized below:

E1.6.1.3.1 Pit A

Table E1.24 summarizes the results of fate and transport analyses for Pit A. Presented in this table are the source term concentrations (i.e., either back-calculated soil concentrations or the observed soil concentrations representing the 95% UCL values), the predicted peak contributing concentrations in the UCRS groundwater beneath the source, and the corresponding time for peak concentrations. In addition, this table presents for comparison the current maximum concentrations in the UCRS groundwater and drinking water MCLs or risk-based concentrations (RBCs) (if a MCL is not available). As can be seen from this table, cadmium, chromium, ²³⁷Np, and ⁹⁹Tc were predicted to reach the peak contributing concentrations exceeding groundwater Remedial Goal Options (RGOs). Predicted peak contributing concentrations of methylene chloride, chlorobenzene, and chloromethane among the organics also exceed their respective groundwater RGOs.

Table E1.24. Summary of Leachate Modeling Results for the COPCs¹ from the Burial Pit A

COPCs	Exposure concentration	Predicted C _{gw,max} ² in the UCRS	Predicted T _{max}	Observed C _{gw,max} in the UCRS	Groundwater RGOs	Comment
Metals^a						
Cadmium	10.0	0.375	545	N/A	0.005	M
Chromium	55.0	1.95	415	0.91	0.10	M
Radionuclides^b						
²³⁷ Np	1.8	119	338	0.4	1.31	R
⁹⁹ Tc	18	66,441	5	99	276	R
Volatile Organic Compounds^c						
Methylene chloride	5.69	12.0	4	N/A	5.00	M
Chlorobenzene	88.31	53.4	12	N/A	12.7	R
Chloromethane	2.19	2.27	4	N/A	1.33	R

¹ These COPCs represent the constituents that were selected for SESOIL modeling.

² It should be noted here that the predicted C_{gw, max} in the UCRS represent the peak leachate concentration before reaching the water table based on contaminant leaching from the existing source concentrations.

^a Concentrations of all inorganic compounds are expressed in mg/kg or mg/L.

^b Concentrations of radionuclides are expressed as pCi/g or pCi/L.

^c Concentrations of organic compounds are expressed as µg/g or µg/L.

M = MCL

R = Risk-based

N/A = Not available

E1.6.1.3.2 Pit B/C

The results of fate and transport analyses for Pit B/C are summarized in Table E1.25. Presented in this table are the source term concentrations (i.e., either back-calculated soil concentrations or the observed soil concentrations representing the 95% UCL values), the predicted peak concentrations in the UCRS groundwater beneath the source, and the corresponding time for peak concentrations. In addition, this table presents for comparison the current maximum concentrations in the UCRS groundwater and drinking water MCLs or RBCs (if a MCL is not available). As can be seen from this table, arsenic, barium, cadmium, chromium, copper, mercury, ²³⁷Np, and ⁹⁹Tc, were predicted to reach the peak contributing concentrations exceeding groundwater RGOs. Predicted peak contributing concentrations of benzene; methylene chloride; toluene; and 2,4-dimethylphenol among the organics also exceed their respective groundwater RGOs.

E1.6.1.3.3 F Pits

The results of fate and transport analyses for the F Pits are summarized in Table E1.26. Presented in this table are the source term concentrations (i.e., either back-calculated soil concentrations or the observed

soil concentrations representing the 95% UCL values), the predicted peak concentrations in groundwater beneath the source, and the corresponding time for peak concentrations. In addition, this table presents for comparison the current maximum concentrations in the UCRS groundwater and drinking water MCLs or RBCs (if a MCL is not available). As can be seen from this table, only ⁹⁹Tc and ²³⁷Np were predicted to reach the peak contributing concentrations exceeding groundwater RGOs.

Table E1.25. Summary of Leachate Modeling Results for the COPCs¹ from the Burial Pits B/C

COPCs	Exposure concentration	Predicted C _{gw,max} ² in the UCRS	Predicted T _{max}	Observed C _{gw,max} in the UCRS	Groundwater RGOs	Comment
<i>Metals^a</i>						
Arsenic	41	2.48	254	0.28	0.05	M
Barium	232.22	13.61	784	4.3	2	M
Cadmium	165.1	6.96	524	N/A	0.005	M
Chromium	371.08	19.72	391	0.91	0.1	M
Copper	31,933	634	456	0.46	0.602	R
Mercury	2.04	0.009	114	0.0011	0.002	M
<i>Radionuclides^b</i>						
²³⁷ Np	19.0	879	327	0.4	1.31	R
⁹⁹ Tc	656.5	3,555,651	4	99	276	R
<i>Volatile organic compounds^c</i>						
Benzene	34.7	62.2	6	12	5	M
Methylene chloride	20.52	60.5	4	N/A	5	M
Toluene	753.4	678.2	9	59	1,000	M
<i>Semivolatile organic compounds^d</i>						
2,4,6-Trichlorophenol	41	3.25	215	N/A	3.99	R
2,4-Dimethylphenol	5,034	11,390	10	4.4	230	R
Hexachloroethane	34.0	0.8	585	NA	3.29	R

¹ These COPCs represent the constituents that were selected for SESOIL modeling.

² It should be noted that the predicted C_{gw,max} in the UCRS represent the peak leachate concentration before reaching the water table based on contaminant leaching from the existing source concentrations.

^a Concentrations of all inorganic compounds are expressed as mg/kg or mg/L.

^b Concentrations of radionuclides are expressed as pCi/g or pCi/L.

^c Concentrations of organic compounds are expressed as µg/g or µg/L.

M = MCL

R = Risk-based

N/A = Not available

Table E1.26. Summary of Leachate Modeling Results for the COPCs¹ from the Burial Pit F

COPCs	Exposure concentration	Predicted C _{gw,max} ² in the UCRS	Predicted T _{max}	Observed C _{gw,max} in the UCRS	Groundwater RGOs	Comment
<i>Radionuclides^a</i>						
²³⁷ Np	0.04	1.73	338	0.4	1.31	R
⁹⁹ Tc	4.17	26,430	5	99	276	R
<i>Semivolatile organic compounds^b</i>						
4-Methylphenol	0.75	1.69	8	0.21	N/A	

¹ These COPCs represent the constituents that were selected for SESOIL modeling.

² It should be noted that the predicted C_{gw,max} in the UCRS represent the peak leachate concentration before reaching the water table based on contaminant leaching from the existing source concentrations.

^a Concentrations of radionuclides are expressed as pCi/g or pCi/L.

^b Concentrations of organic compounds are expressed as µg/g or µg/L.

M = MCL

R = Risk-based

N/A = Not available

E1.6.1.3.4 Subsurface Source Outside of the Pits

The results of fate and transport analyses for Subsurface Source Outside of the Pits are summarized in Table E1.27. Presented in this table are the source term concentrations (i.e., either back-calculated soil concentrations or the observed soil concentrations representing the lesser of 95% UCL or maximum values), the predicted peak concentrations in groundwater beneath the source, and the corresponding time for peak concentrations. In addition, this table presents for comparison the current maximum concentrations in the UCRS groundwater and drinking water MCLs or RBCs (if a MCL is not available). As can be seen from this table, arsenic, barium, cadmium, chromium, ⁹⁹Tc, ²³⁷Np, and methylene chloride were predicted to reach the peak contributing concentrations exceeding groundwater RGOs.

Table E1.27. Summary of Leachate Modeling Results for the COPCs¹ from the Subsurface Soil Outside Pits

COPCs	Exposure concentration	Predicted C _{gw,max} ² in the UCRS	Predicted T _{max}	Observed C _{gw,max} in the UCRS	Groundwater RGOs	Comment
<i>Metals^a</i>						
Arsenic	3.143	1.06	265	0.28	0.05	M
Barium	104.3	8.92	827	4.3	2	M
Cadmium	1.85	0.37	557	N/A	0.005	M
Chromium	21.45	2.37	419	0.91	0.1	M
<i>Radionuclides^b</i>						
²³⁷ Np	2.36	428.58	360	0.4	1.31	R
⁹⁹ Tc	280.0	977,625	5	99	276	R
<i>Volatile organic compounds^c</i>						
Methylene chloride	87.1	710.4	4	N/A	5	M

¹ These COPCs represent the constituents that were selected for SESOIL modeling.

² It should be noted that the predicted C_{gw,max} in the UCRS represent the peak leachate concentration before reaching the water table based on contaminant leaching from the existing source concentrations.

^a Concentrations of all inorganic compounds are expressed as mg/kg or mg/L.

^b Concentrations of radionuclides are expressed as pCi/g or pCi/L.

^c Concentrations of organic compounds are expressed as µg/g or µg/L.

M = MCL

R = Risk-based

N/A = Not available

E1.6.1.3.5 UCRS

As discussed in E1.6.1.2, SESOIL modeling for the UCRS used either the maximum leachate concentrations predicted by SESOIL modeling for the four source areas (Pit A, Pit B/C, F Pits, and Subsurface Source Outside of the Pits) or the maximum observed groundwater concentrations to predict the peak leachate concentration in the UCRS. The results of this transport modeling in the UCRS are summarized in Table E1.28. Presented in this table are source concentrations (i.e., either predicted maximum leachate concentrations based on SESOIL modeling of the sources (i.e., Pit A, Pit B/C, F Pits, and the Areas within SWMU 7 and 30 outside of Pits A, B, and C) or the maximum observed groundwater concentrations in the UCRS, whichever is greater), the predicted peak contributing concentrations in the leachate before reaching the RGA, and the corresponding time of peak concentrations. In addition, this table presents for comparison the current maximum concentrations in the RGA groundwater and drinking water MCLs or RBCs (if a MCL is not available). As can be seen from this table, the predicted peak contributing concentrations of 1,2-DCE; *cis*-1,2-DCE; 2,4 dimethylphenol; methylene chloride; TCE; vinyl chloride; ⁹⁹Tc; and mercury exceed their respective groundwater RGOs; therefore, these constituents were considered for lateral transport modeling in the RGA using AT123D (see Section E1.6.2). Mercury was dropped from this list. The maximum concentration of mercury, although slightly higher than its MCL, decreases to lower than its MCL with dilution.

Table E1.28. Summary of Leachate Modeling Results for the COPCs¹ from the UCRS

COPCs	Exposure concentration	Predicted C _{gw,max} ² in the UCRS	Predicted T _{max}	Observed C _{gw,max} in the UCRS	Groundwater RGOs	Comment
<i>Metals^a</i>						
Mercury	0.09	0.0042	405	0.0012	0.002	M
<i>Radionuclides^b</i>						
⁹⁹ Tc	355.6	763,627	14	3,670	276	R
<i>Volatile organic compounds^c</i>						
1,1,1-Trichloroethane	94.77	27.5	26	1.3	200	M
1,2-Dichloroethane	0.59	0.32	14	3	5	M
1,1-DCE	3.89	0.31	16	3.7		
1,2-DCE	1,591.9	281.3	18	110	136	R
<i>cis</i> -1,2-DCE	406.1	133.7	13	140	149	R
4-Methyl 2,2-pentanone	0.09	0.09	10	2.9	N/A	
Acetone	2.54	2.73	9	430	1,510	R
Benzene	1.93	0.48	16	N/A	5	M
Chloromethane	0.03	0.01	10	14	1.33	R
Methylene chloride	18.47	9.40	10	N/A	5	M
TCE	464.4	56.53	20	19,000	5	M
Vinyl chloride	362.7	15.21	12	N/A	2	M
<i>Semivolatile organic compounds^d</i>						
2,4-Dimethylphenol	6,189.3	1,983.42	30	N/A	230	R

¹ These COPCs represent the constituents that were selected for SESOIL modeling.

² It should be noted that the predicted C_{gw,max} in the UCRS represent the peak leachate concentration before reaching the water table based on contaminant leaching from the existing source concentrations.

^a Concentrations of all inorganic compounds are expressed as mg/kg or mg/L.

^b Concentrations of radionuclides are expressed as pCi/g or pCi/L.

^c Concentrations of organic compounds are expressed as µg/g or µg/L.

M = MCL

R = Risk-based

N/A = Not available

E1.6.2 AT123D MODELING

AT123D is an analytical groundwater pollutant fate and transport model chosen to predict the future receptor concentrations for the contaminants. It computes the spatial-temporal concentration distribution of wastes in the aquifer system and predicts the transient spread of a contaminant plume through a groundwater aquifer. The fate and transport processes accounted for in AT123D are advection, dispersion, adsorption/retardation, and decay. This model can be used as a tool for estimating the dissolved concentration of a chemical in three dimensions in the groundwater resulting from a mass release over a source area (point, line, area, or volume source). The model can handle instantaneous as well as continuous source loadings of chemicals of interest at the site. AT123D frequently is used by the scientific and technical community to perform quick and conservative estimates of groundwater plume movement in space and time. In RISKPRO, SESOIL and AT123D are linked so that mass loading to the groundwater predicted by SESOIL can be directly transferred to AT123D.

Six organic compounds and one radionuclide were selected for AT123D modeling in the RGA based on source loading from the UCRS predicted by SESOIL. Maximum concentrations at two receptor locations (PGDP boundary and DOE property boundary) were simulated for these constituents. Maximum concentrations at the end of 30 years of simulation also were predicted for these constituents.

E1.6.2.1 RGA

The results of fate and transport modeling in the RGA based on future contaminant loading from SWMUs 7 and 30 are summarized in Table E1.29. Presented in this table are the predicted peak contributing concentrations in groundwater beneath the source, predicted peak contributing concentrations at 30 years and in 100 years at the PGDP boundary in the direction of flow, and the peak contributing concentrations in 30 and 100 years at the DOE property boundary in the direction of flow. In addition, this table presents for comparison the current maximum concentrations in the RGA groundwater and drinking water MCLs or RBCs (if a MCL is not available). As can be seen from this table, ⁹⁹Tc is predicted to reach the peak contributing concentrations exceeding its groundwater RBC at all locations and for both 30- and 100-year scenarios. None of the organic COPCs are predicted to reach the peak contributing concentrations exceeding their respective groundwater RBCs; however, the predicted results for all the COPCs but ⁹⁹Tc are quite low, when compared to their maximum concentrations currently observed in the RGA. For ⁹⁹Tc, the observed maximum concentrations of 3,670 pCi/L falls within the predicted range of 1,996 pCi/L (30-year peak) to 21,686 pCi/L (100-year peak).

Table E1.29. Summary of Transport Modeling in the RGA Based on Future Contaminant Loading from SWMUs 7 and 30

Constituent	Unit	Predicted maximum concentration beneath the source	Predicted GW concentration at the PGDP boundary in the direction of flow		Predicted GW concentration at the DOE property boundary in the direction of flow	
			30 years	100 years	30 years	100 years
1,2-DCE	µg/L	27	16.8	24.3	5.2	5.2
<i>cis</i> -12-DCE	µg/L	12.3	4.6	11.3	1.5	2.7
TCE	µg/L	5.3	3.8	4.6	1.1	1.1
Methylene chloride	µg/L	0.8	0.14	0.8	0.05	0.19
Vinyl chloride	µg/L	1.0	0.15	0.96	0.06	.23
2,4-Dimethylphenol	µg/L	200	174	1.74	4.1	40.7
⁹⁹ Tc	pCi/L	23,580	1,996	21,686	1,205	5,077

Note: All the constituents that were identified as the initial contaminant migration COPCs in the source areas (i.e., SWMUs 7 and 30, and also the UCRS beneath the site) were modeled to the RGA. However, only the constituents that were predicted to arrive at the RGA with concentrations exceeding their groundwater MCLs/RBCs were modeled to the receptors using AT123D and are shown in this table. All the concentrations shown in this table represent only the contributed concentrations and do not account for the existing concentrations that already have contaminated the RGA groundwater.

Based on fate and transport analyses results, it appears that ⁹⁹Tc is the only constituent that will continue to be a major problem at the receptor locations. Therefore, ⁹⁹Tc was chosen for further fate and transport evaluations in order to address the source units within SWMUs 7 and 30, separately, in terms of contaminant contributions to the receptor locations. Vertical transport of ⁹⁹Tc from the individual source units through the UCRS to the RGA were performed using SESOIL. The results from the SESOIL modeling were used to create input for an AT123D model that was applied for predicting lateral migration of ⁹⁹Tc to the receptor locations. These results are summarized in Table E1.30. Concentration contributions from the individual source units are added to provide the total contributed concentrations based on future loading from SWMUs 7 and 30. The relative (%) contributions of individual source units are also shown (values in parentheses) in this table.

Table E1.30. Results of ⁹⁹Tc Modeling in the RGA Based on Future Contaminant Loading from the Individual Source Units within SWMUs 7 and 30

Constituent	Units	Predicted Maximum Concentration Beneath the Source	Predicted GW Concentration at the PGDP Boundary in the Direction of Flow		Predicted GW Concentration at the DOE Property Boundary in the Direction of Flow	
			30 years	100 years	30 years	100 years
Burial Pit A	pCi/L	60.8 (0.65)	3.1 (0.46)	34.2 (0.47)	1.3 (0.35)	6.3 (0.4)
Burial Pits B/C	pCi/L	3,253.8 (35.02)	165.2 (24.79)	1,822.8 (25.09)	71.2 (19.03)	291.3 (19.1)
Burial Pit F	pCi/L	12.7 (0.14)	0.6 (0.09)	6.5 (0.09)	0.2 (0.05)	1.0 (0.07)
Subsurface soils outside these pits	pCi/L	5,962.5 (64.19)	497.4 (74.66)	5,400.3 (74.35)	301.5 (80.67)	1,228.0 (80.43)
Combined SWMUs 7 and 30	pCi/L	9289.8 (100.00)	666.3 (100.00)	7263.8 (100.00)	347.2 (100.00)	1526.6 (100.00)

Note: All the concentrations shown in this table represent only the contributed concentrations and do not account for the existing concentrations that are already present in the aquifer. Percent of total contribution from SWMUs 7 and 30 combined are shown in parentheses.

E1.7. MODELING APPEARING IN THE SITEWIDE RISK ASSESSMENT MODEL AND ENVIRONMENTAL BASELINE REPORT

Groundwater fate and transport modeling and risk modeling for SWMUs 2, 3, 4, 5, 6, 7, and 30 was performed as part of the PGDP sitewide risk assessment model and environmental baseline (DOE 2003). This section summarizes the results of this modeling for these SWMUs and is taken from *Sitewide Risk Assessment Model and Environmental Baseline for the Paducah Gaseous Diffusion Plant, Paducah, Kentucky* (DOE 2003).

The models selected for groundwater modeling were SESOIL for soil leachability and AT123D for lateral transport in groundwater. The MODFLOW/MODPATH models were used to support the AT123D

modeling. The COPC list for groundwater transport modeling was derived from the full list of significant COPCs. The results of this modeling include the chemical-specific maximum source contributions to cancer risk, hazard and dose for the SWMUs at four integrator points.

Four groundwater integrator points were determined to be relevant to the exposure conceptual site model and endstate goals. At each of these locations, it was assumed that residents could be exposed to contamination originating at PGDP through household use of groundwater, even though no water wells used for this purpose are present at these locations at this time. These integrator points are:

- **Integrator Point 1 (GW-NW-P).** This integrator point is located on the northwest side of the DOE reservation at the center of the current northwest TCE groundwater plume. Sources to this integrator point include SWMUs 2, 3, 5, 6, 7, and 30.
- **Integrator Point 2 (GW-SW-P).** This integrator point is located on the northwest side of the DOE reservation to the west of Integrator Point 1 at a point to which the center of the southwest TCE plume is expected to migrate. (The southwest plume has *not* reached this location.) Sources to this integrator point include SWMU 4.
- **Integrator Point 3 (GW-NE-P).** This integrator point is located on the northeast side of the DOE reservation at the center of the current northeast TCE plume.
- **Integrator Point 4 (GW-North).** This integrator point is located on the north side of the DOE reservation between Integrator Points 1 and 3 and is not associated with any currently identified plume.

E1.7.1 SESOIL MODELING

The SESOIL model is described in Section E1.6.1. The input data are divided into four types: climatic, chemical, soil, and application data. An overview of the parameters used for the sitewide risk assessment modeling follows. Specific parameters used are presented in Appendix E of *Sitewide Risk Assessment Model and Environmental Baseline for the Paducah Gaseous Diffusion Plant, Paducah, Kentucky* (DOE 2003).

E1.7.1.1 Climate Data

The climatic data file of SESOIL consists of an array of values for various climatic parameters. These monthly data span 1 year and are derived from Paducah. The climatic parameters are used within SESOIL to generate the hydrologic model responsible for contaminant transport.

E1.7.1.2 Chemical Data

The pollutant fate cycle of SESOIL uses several chemical transport and transformation processes that occur in the soil zone. The processes of volatilization/diffusion, adsorption/desorption, and biodegradation are used in SESOIL modeling for the purpose of the soil leachability analysis.

E1.7.1.3 Soil Data

The soil data file of SESOIL contains input parameters describing the physical characteristics of the subsurface soil. The parameters include soil bulk density, intrinsic permeability, soil disconnectedness index, Freundlich exponent, total porosity, and organic carbon content. The groundwater recharge rates (infiltration rates) were obtained from the PGDP sitewide groundwater model (DOE 1998b). These

recharge rates were used in SESOIL modeling as a calibration target. The intrinsic permeability for the vadose zone is a critical model parameter in that it permits calibration to the target groundwater recharge rate. The intrinsic permeability was varied in iterative runs until the groundwater recharge rate predicted by SESOIL matched the recharge rate obtained from the groundwater model. Unit-specific data were not available for some of the parameters; therefore, EPA default values were used as input to the model. These parameters included the soil disconnectedness index and Freundlich exponent. There is no measurement method for the soil disconnectedness index (described below), nor is there a measured value of the related Freundlich exponent (used in calculating the adsorbed contaminant concentration). The soil disconnectedness index replaces moisture retention curves (or characteristic curves) used by other unsaturated zone leaching models. The soil disconnectedness index was calibrated for four different soil types ranging from sandy loam to clay (Hetric *et al.* 1986). This parameter has a minor impact upon the groundwater recharge rate and is varied (within the range specified for the corresponding soil type) in the final stages of model calibration. The SESOIL default value was used for the Freundlich exponent.

E1.7.1.4 Application Parameters

The SESOIL application data describe the soil layer configuration and the initial contaminant concentrations in each model layer. The SESOIL model was arranged in layers and sublayers that facilitate contaminant loading at intervals closely approximating the actual waste placement in the disposal facility. They represent constituent loading or leaching zones, as appropriate. The initial loading concentrations (source term) for SESOIL layers are the soil concentrations observed at individual sites. The SESOIL model for any group of SWMUs contained four major layers with multiple sublayers to model leaching through the vadose zone. The thickness of the layers varied depending on depth to the RGA and thickness of the contaminated soil zone within the vadose zone. In these analyses in general, Layers 1 and 2 are assumed for source loading, and the remaining layers are assigned for leaching zones. The last sublayer of Layer 4 represents the interface of the vadose zone and the RGA beneath the site. The predicted leachate concentration is determined in this last sublayer of Layer 4. The simulations using SESOIL were continued until the maximum concentration in leachate beneath the source was attained or a maximum time period of 1,000 years was reached. The maximum predicted leachate concentrations were input into the AT123D model to predict the maximum groundwater concentration at the integrator points.

E1.7.2 AT123D MODELING

The AT123D model is described in Section E1.6.2. For the sitewide risk assessment modeling, AT123D was chosen to predict the future integrator point concentrations for the COPCs. Specific model input parameters for the AT123D are presented in Appendix E of *Sitewide Risk Assessment Model and Environmental Baseline for the Paducah Gaseous Diffusion Plant, Paducah, Kentucky* (DOE 2003).

E1.7.3 MODFLOW AND MODPATH MODELING

The MODFLOW/MODPATH models were used to identify the locations of the integrator points and estimate hydraulic gradients, flow distances, and hydraulic conductivities along SWMU-to-integrator flowpaths. This information was subsequently used to support the AT123D modeling effort. MODFLOW is a three-dimensional, finite-difference model capable of simulating both steady-state and transient head distribution for a saturated groundwater flow field. In contrast, MODPATH is a three-dimensional, particle-tracking model capable of using the steady-state, head distribution generated by MODFLOW to track flowpaths of particles released in the groundwater flow field modeled by MODFLOW.

The MODFLOW model used in this analysis was the sitewide groundwater flow model developed earlier by DOE (1998b). This model covers most of the DOE reservation except that portion above the Porter's Creek Clay terrace. It has been approved by both the PGDP Modeling Steering Committee and the Risk Assessment Working Group; therefore, this model was used without any modification. The parameters used in this model are summarized in the PGDP Quarterly Modeling Report (BJC 2001) and are not discussed further here.

As noted above, the MODPATH model was used to track flowpaths of particles released from a location by using the steady-state head distribution generated by MODFLOW. The key parameter of MODPATH is the particle depth at release. For grouping the SWMUs assigned to different integrator points, the mid-depth of Layer 3 was assumed to represent the average flow condition, and the particles were released from this depth of an aquifer.

E1.7.4 MODELING ASSUMPTIONS

A highly conservative approach was used for this analysis. Several assumptions were made in developing the mathematical models for this analysis. The important assumptions are listed below.

- Infiltration of water through vadose zone soil consists of one-dimensional, steady flow through soil with no dispersion and with uniform average soil properties. More complex flow could either increase or decrease contaminant mobility and transport to the RGA.
- The use of K_d and R_d (retardation factor) to describe the reaction term of the transport equation assumes that an equilibrium relationship exists between the solid- and solution-phase concentrations and that the relationship is linear and reversible.
- Most of the K_d values used in this analysis for the COPCs represent literature-based or calculated values and might not represent the actual site conditions.
- No biodegradation takes place in the vadose zone (i.e., SESOIL modeling assumes no decay) or in the groundwater (i.e., AT123D modeling without accounting for biodegradation).
- In case of radionuclides and organic compounds, ingrowth and decay are not considered.
- Flow and transport in the vadose zone are one-dimensional (i.e., take place in only the vertical direction).
- Initial condition is disregarded in the vadose zone modeling.
- Flow and transport are not affected by density variations.
- Areal distribution of soil contamination is not considered.
- The aquifer is assumed to be homogenous and isotropic.
- The integrator point is located at a nearest downgradient distance.

The inherent uncertainties associated with using these assumptions must be recognized. Because K_d values are highly sensitive to changes in the major chemistry of the solution phase, it is important that the

values be measured or estimated under conditions that will represent, as closely as possible, those of the contaminant plume. It is also important to note that the contaminant plume will change over time and be affected by multiple solutes that are present at the site. Projected organic concentrations in the aquifer are uncertain because of the lack of site-specific data on constituent decay in the vadose zone and groundwater. Use of literature values could produce either over- or underestimation of constituents' concentrations in the aquifer. Deviations from assumed literature values could significantly affect contaminant fate predictions.

E1.7.5 COPC LIST USED IN GROUNDWATER TRANSPORT MODELING

The COPC list for soil and sediment that act as a source for contamination that might arrive at a groundwater integration point (i.e., a location downgradient from the source) was derived from the full list of significant COPCs by deleting those COPCs that could not reasonably be expected to migrate through the groundwater pathway and arrive at an integration point at a concentration greater than the screening levels. The COPCs not expected to migrate through the groundwater pathway were selected by considering each COPC's site-specific K_d and removing those inorganic chemical and radionuclide COPCs with a K_d greater than 500 L/kg and those organic compound COPCs with a K_d greater than 10 L/kg. Additionally, all radionuclides with a half-life less than 6 years (i.e., will pass through five half-lives in a 30-year period) were removed. The COPCs expected to migrate through the groundwater pathway are indicated in Table E1.31.

Table E1.31. COPC List for Groundwater Transport Modeling

Inorganic Chemicals		Organic Compounds		Radionuclides	
Significant COPC	CAS Number	Significant COPC	CAS Number	Significant COPC	CAS Number
Antimony	7440360	Acrylonitrile	107131	Neptunium-237+D	13994202
Arsenic	7440382	Benzene	71432	Strontium-90+D	10098972
Cadmium	7440439	Carbon tetrachloride	56235	Technetium-99	14133767
Chromium III	16065831	Chloroform	67663	Uranium-234	13966295
Chromium VI	18540299	1,1-Dichloroethene	75354	Uranium-235+D	15117961
Chromium (Total)	7440473	1,2-Dichloroethene (mixed)	540590	Uranium-238+D	7440611
Copper	7440508	<i>trans</i> -1,2-Dichloroethene	156605		
Iron	7439896	<i>cis</i> -1,2-Dichloroethene	156592		
Lead	7439921	Ethylbenzene	100414		
Manganese	7439965	Naphthalene	91203		
Mercury	7439976	Tetrachloroethene	127184		
Molybdenum	7439987	Trichloroethene	79016		
Nickel	7440020	Vinyl chloride	75014		
Selenium	7782492	Xylenes (mixture)	1330207		
Silver	7440224				
Thallium	7440280				
Uranium	NA				

Note: Table adapted from DOE 2003.

CAS=Chemical Abstracts Service.

NA = not applicable.

^a The list of significant COPCs matches that presented in Table 3.1 [of DOE 2003], except individual organic compounds comprising the total PAHs, total PCBs, and total dioxins/furans groups are not listed.

^b The COPCs not expected to migrate through the groundwater pathway were selected by considering each COPC's site-specific K_d and removing those inorganic chemical and radionuclide COPCs with a K_d greater than 500 L/kg and those organic compound COPCs with a K_d greater than 10 L/kg. Additionally, all radionuclides with a half-life less than 6 years (i.e., will pass through five half-lives in a 30-year period) were removed.

E1.7.6 MODELING RESULTS

Table E1.32 presents the chemical-specific maximum source contributions to cancer risk, hazard, and dose for SWMUs 2, 3, 4, 5, 6, 7, and 30 at the groundwater integrator points (DOE property boundary). The concentrations used to derive the risks and doses came from the maximum flux contribution information resulting from the groundwater modeling. The table also includes the expected maximum COPC concentrations predicted by the groundwater modeling.

Table E1.32. Maximum Cancer Risk, Hazard, and Dose for Sources Contributing at the DOE Property Boundary

COPC ^a	SWMU						
	2	3	4	5	6	7	30
Concentrations (mg/L or pCi/L)							
Antimony	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Arsenic	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Cadmium	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Chromium	1.80E-02	2.07E-02	1.50E-01	1.77E-01	6.95E-02	4.23E-01	0.00E+00
Copper	1.27E-02	8.28E-03	3.92E-01	7.10E-03	9.05E-03	1.18E+01	2.43E-02
Iron	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Lead	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Manganese	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Mercury	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Molybdenum	2.62E-03	2.11E-03	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Nickel	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Selenium	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	3.20E-10	0.00E+00
Silver	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Thallium	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Uranium	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Benzene	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	5.80E-03	0.00E+00
Carbon tetrachloride	0.00E+00	0.00E+00	2.21E+01	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Chloroform	0.00E+00	0.00E+00	4.90E+02	0.00E+00	0.00E+00	9.86E-04	0.00E+00
Dichloroethene (mixed), 1,2-	0.00E+00	0.00E+00	6.34E+01	0.00E+00	0.00E+00	5.92E-03	0.00E+00
Dichloroethene, 1,1-	0.00E+00	0.00E+00	1.68E+01	2.33E-01	0.00E+00	0.00E+00	0.00E+00
Dichloroethene, <i>cis</i> -1,2-	1.98E+01	0.00E+00	2.15E+04	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Dichloroethene, <i>trans</i> -1,2-	0.00E+00	0.00E+00	9.24E+02	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Ethylbenzene	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	6.41E-03	0.00E+00
Naphthalene	3.94E-03	6.01E-03	0.00E+00	3.50E-01	0.00E+00	6.32E-03	2.11E-02
Tetrachloroethene	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	5.58E-05	0.00E+00
Trichloroethene	8.06E+00	0.00E+00	4.10E+01	5.18E-04	2.19E-04	1.02E-03	5.70E-04
Vinyl chloride	6.08E-02	0.00E+00	2.90E-01	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Xylenes (mixture)	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	3.81E-02	0.00E+00
Neptunium-237+D	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Strontium-90+D	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Technetium-99	4.83E+02	9.69E+01	2.08E+03	6.53E+01	3.60E+02	1.12E+04	4.79E+03
Uranium-234	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Uranium-235+D	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Uranium-238+D	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00

Table E1.32. Maximum Cancer Risk, Hazard, and Dose for Sources Contributing at the DOE Property Boundary (Continued)

COPC ^a	SWMU						
	2	3	4	5	6	7	30
	Cancer Risk						
Antimony	NR	NR	NR	NR	NR	NR	NR
Arsenic	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Cadmium	NR	NR	NR	NR	NR	NR	NR
Chromium	NR	NR	NR	NR	NR	NR	NR
Copper	NR	NR	NR	NR	NR	NR	NR
Iron	NR	NR	NR	NR	NR	NR	NR
Lead	NR	NR	NR	NR	NR	NR	NR
Manganese	NR	NR	NR	NR	NR	NR	NR
Mercury	NR	NR	NR	NR	NR	NR	NR
Molybdenum	NR	NR	NR	NR	NR	NR	NR
Nickel	NR	NR	NR	NR	NR	NR	NR
Selenium	NR	NR	NR	NR	NR	NR	NR
Silver	NR	NR	NR	NR	NR	NR	NR
Thallium	NR	NR	NR	NR	NR	NR	NR
Uranium	NR	NR	NR	NR	NR	NR	NR
Benzene	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	1.51E-05	0.00E+00
Carbon	0.00E+00	0.00E+00	1.22E-01	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Chloroform	0.00E+00	0.00E+00	2.25E+00	0.00E+00	0.00E+00	4.52E-06	0.00E+00
Dichloroethene (mixed), 1,2-	NR	NR	NR	NR	NR	NR	NR
Dichloroethene, 1,1-	0.00E+00	0.00E+00	3.58E-01	4.96E-03	0.00E+00	0.00E+00	0.00E+00
Dichloroethene, <i>cis</i> -1,2-	NR	NR	NR	NR	NR	NR	NR
Dichloroethene, <i>trans</i> -1,2-	NR	NR	NR	NR	NR	NR	NR
Ethylbenzene	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	1.37E-06	0.00E+00
Naphthalene	NR	NR	NR	NR	NR	NR	NR
Tetrachloroethene	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	9.59E-08	0.00E+00
Trichloroethene	4.66E-03	0.00E+00	2.37E-02	3.00E-07	1.27E-07	5.91E-07	3.29E-07
Vinyl	1.74E-03	0.00E+00	8.29E-03	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Xylenes	NR	NR	NR	NR	NR	NR	NR
Neptunium-237+D	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Strontium-90+D	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Technetium-99	3.45E-05	6.92E-06	1.49E-04	4.67E-06	2.57E-05	8.00E-04	3.42E-04
Uranium-234	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Uranium-235+D	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Uranium-238+D	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00

Table E1.32. Maximum Cancer Risk, Hazard, and Dose for Sources Contributing at the DOE Property Boundary (Continued)

COPC ^a	SWMU						
	2	3	4	5	6	7	30
	Hazard Results						
Antimony	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Arsenic	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Cadmium	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Chromium	1.02E-03	1.18E-03	8.54E-03	1.01E-02	3.95E-03	2.40E-02	0.00E+00
Copper	2.28E-02	1.49E-02	7.04E-01	1.27E-02	1.63E-02	2.12E+01	4.37E-02
Iron	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Lead	NR	NR	NR	NR	NR	NR	NR
Manganese	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Mercury	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Molybdenum	3.48E-02	2.80E-02	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Nickel	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Selenium	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	4.25E-09	0.00E+00
Silver	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Thallium	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Uranium	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Benzene	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	1.15E+00	0.00E+00
Carbon tetrachloride	0.00E+00	0.00E+00	1.16E+04	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Chloroform	0.00E+00	0.00E+00	1.71E+06	0.00E+00	0.00E+00	3.44E+00	0.00E+00
Dichloroethene (mixed), 1,2-	0.00E+00	0.00E+00	2.57E+03	0.00E+00	0.00E+00	2.40E-01	0.00E+00
Dichloroethene, 1,1-	0.00E+00	0.00E+00	1.10E+02	1.52E+00	0.00E+00	0.00E+00	0.00E+00
Dichloroethene, <i>cis</i> -1,2-	7.26E+02	0.00E+00	7.89E+05	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Dichloroethene, <i>trans</i> -1,2-	0.00E+00	0.00E+00	1.69E+04	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Ethylbenzene	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	1.14E-02	0.00E+00
Naphthalene	1.38E+00	2.11E+00	0.00E+00	1.23E+02	0.00E+00	2.22E+00	7.42E+00
Tetrachloroethene	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	6.63E-04	0.00E+00
Trichloroethene	5.04E+02	0.00E+00	2.56E+03	3.24E-02	1.37E-02	6.39E-02	3.56E-02
Vinyl chloride	1.99E+00	0.00E+00	9.48E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Xylenes (Mixture)	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	4.11E-01	0.00E+00
Neptunium-237+D	NR	NR	NR	NR	NR	NR	NR
Strontium-90+D	NR	NR	NR	NR	NR	NR	NR
Technetium-99	NR	NR	NR	NR	NR	NR	NR
Uranium-234	NR	NR	NR	NR	NR	NR	NR
Uranium-235+D	NR	NR	NR	NR	NR	NR	NR
Uranium-238+D	NR	NR	NR	NR	NR	NR	NR
	Dose Results^b						
Neptunium-237+D	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Strontium-90+D	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Technetium-99	4.93E-01	9.91E-02	2.13E+00	6.68E-02	3.68E-01	1.14E+01	4.90E+00
Uranium-234	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Uranium-235+D	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00
Uranium-238+D	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00	0.00E+00

Note: Table adapted from DOE 2003.

“NR” = Result is not reported because no screening value is available.

^a Only COPCs identified as being mobile through the groundwater pathway are listed.

^b Does results are only applicable to radionuclides.

Most of the metals and radionuclides are not expected to migrate to the integrator points at any measurable concentrations. The only metal that is predicted to have a higher concentration is chromium. SWMUs 7 and 5, and 4 might be contributing to the chromium problem at Integrator Points 4, 1, and 2, respectively. The only radionuclide that is predicted to have a higher concentration is ⁹⁹Tc. The contributing sources for ⁹⁹Tc are SWMU 4 for Integrator Point 2, and SWMUs 7 and 30 for Integrator Point 1. The chlorinated solvents are expected to migrate to their respective integrator points at significant concentrations from multiple SWMUs, including SWMUs 2 and 5 for Integrator Point 1 and SWMUs 4 and 1 for Integrator Point 2. None of the SWMUs is expected to contribute chlorinated solvents to either Integrator Points 3 or 4. Among the semivolatile organic compounds (SVOCs), major contribution of naphthalene is expected to move from SWMU 30 to Integrator Point 1.

The greatest contributors of the COPCs presenting the maximum potential cancer risks are as follows. (All values reported below are potential cancer risks to the resident that are projected to exist if only the reported source contributed contaminants to groundwater.)

- 1,1-DCE: SWMU 4 (3.58E-01) and SWMU 5 (4.96E-03)
- Carbon tetrachloride: SWMU 4 (1.22E-01)
- Chloroform: SWMU 4 (>1)
- TCE: SWMU 2 (4.66E-03) and SWMU 4 (2.37E-02)
- Vinyl chloride: SWMU 2 (1.74E-03) and SWMU 4 (8.29E-03)
- ⁹⁹Tc: SWMU 2 (3.45E-05), SWMU 3 (6.92E-06), SWMU 4 (1.49E-04), SWMU 5 (4.67E-06), SWMU 6 (2.75E-05), SWMU 7 (8.00E-04), and SWMU 30 (3.23E-04)

The greatest contributors of the COPCs presenting the maximum potential hazards are as follows. (All values reported below are potential hazards to the resident that are projected to exist if only the reported source contributed contaminants to groundwater.)

- *cis*-1,2-DCE: SWMU 2 (726) and SWMU 4 (789,000)
- *trans*-1,2-DCE: SWMU 4 (169,000)
- Carbon tetrachloride: SWMU 4 (11,600)
- Chloroform: SWMU 4 (>1,000,000)
- TCE: SWMU 2 (504)
- Naphthalene: SWMU 2 (1.38), SWMU 3 (2.11), SWMU 5 (123), SWMU 7 (2.22), and SWMU 30 (7.24)
- Copper: SWMU 7 (21.2)

Finally, the greatest contributors of the COPCs presenting the maximum potential dose (in mrem/year) from ⁹⁹Tc are SWMU 4 (2.13), SWMU 7 (11.4), and SWMU 30 (4.90). (As before, the dose values are potential doses to the resident that are projected to exist if only the reported source contributed contaminants to groundwater.)

The assumptions for groundwater modeling were discussed in Section E1.7.4. The inherent uncertainties associated with using the assumptions must be recognized. Because K_d values are highly sensitive to changes in the major chemistry of the solution phase, it is important that the values be measured or estimated under conditions that will represent, as closely as possible, those of a contaminant plume. It is also important to note that any contaminant plume predicted to occur will change over time and be affected by multiple solutes that are currently present or may be present in the future at the site. Projected organic concentrations in the aquifer are overestimated because site-specific data on constituent decay in the vadose zone and in the groundwater is lacking, and zero decay was assumed for all organic compounds.

Neither the SESOIL nor AT123D models account for the in-growth of radionuclides. This may have resulted in an underestimation of representative concentrations of decay products with mobility higher than the starting radionuclide, and an overestimation of representative concentrations of decay products with mobility lower than the starting radionuclide. Similarly, because decay will reduce the concentration of the starting radionuclide in the source, the representative concentrations of the starting radionuclide may be overestimated. Fortunately, most of the radionuclide COPCs included in the modeling either have very short half-lives relative to the time modeled (i.e., ^{90}Sr) or very long half-lives relative to the time modeled (i.e., ^{99}Tc , ^{238}U , etc.). Therefore, in general, the decay and in-growth of radiological constituents should have very little effect on the risk characterization.

The effects of heterogeneity, anisotropy, and spatial distribution of fractures are not addressed in these simulations. The present modeling study using SESOIL and AT123D does not address the effects of flow and contaminant transport across interfaces in a sharply varying heterogeneous media. The migration distance predicted by the model may be uncertain mainly because of homogenous and isotropic assumptions were used in these models whereas site data indicate otherwise.

As such, a conservative approach was used to address the uncertainties. Based upon the data available, the values of the model parameters were selected to ensure that contaminant transport was not underestimated. Such an approach can be expected to lead to overestimates of COPC concentrations in groundwater at the integrator points and, subsequently, to overestimates of the cancer risks, hazards, and doses estimated.

E1.8. REFERENCES

- BJC (Bechtel Jacobs Company LLC) 2001. *Quarterly Modeling Report for the Paducah Gaseous Diffusion Plant, Paducah, Kentucky*, BJC/PAD-198/Revised Final.
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APPENDIX E
ATTACHMENT 2
SOURCE TERM DEVELOPMENT

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E2.1 SOURCE TERM DEVELOPMENT

E2.1.1 INTRODUCTION

This attachment documents the development of Upper Continental Recharge System (UCRS) contaminant source terms for fate and transport modeling in the Burial Grounds Operable Unit (BGOU) Remedial Investigation (RI). The BGOU consists of Solid Waste Management Units (SWMUs) 2, 3, 4, 5, 6, 7, 30, and 145, as shown in Figure E2.1.

It is almost never possible to collect data at all possible locations in an area of interest. Although sampling will provide information at some locations, it still is necessary to use some method to “interpolate” the data that is available to estimate or predict the values at nondata locations. Spatial analysis, specifically geospatial interpolation techniques in Spatial Analysis and Decision Assistance (SADA)(UT 2002) was used to characterize the source zone in the UCRS soils.

The BGOU RI assesses the risk posed by contaminants for each SWMU; consequently fate and transport assessment requires separate models for each SWMU (and each assessed contaminant within each SWMU). Model domains extend from ground surface down to the top of the Regional Gravel Aquifer (RGA), commonly in seven layers that are discretized into rows and columns of uniform spacing. Table E2.1 summarizes the source term model domains.

Table E2.1. Summary of the Source Term Model Domains

SWMU	Modeled Area (acres)	Depth (ft) Simulated in Model	Grid Block Size (ft x ft)	Table of SADA Source Characterization Results^a
2	0.73	64	20 x 20	SWMU 2 results.xls
3	3.86	65	100 x 100	SWMU 3 results.xls
4	6.49	63	20 x 20	SWMU 4 results.xls
5	4.43	60	20 x 20	SWMU 5 results.xls
6	0.19	63	10 x 10	SWMU 6 results.xls
7	2.53	60	20 x 20	SWMU 7 results.xls
30	2.70	61	20 x 20	SWMU 30 results.xls
145	44.40	58	100 x 100	SWMU 145 results.xls

^a These SADA files are contained on the CD under the SADA directory

The discretized rows, columns, and layers defined the soil domain (volume). Observed data within the domain were compiled, and contaminant concentration in every cell of the domain was predicted using geospatial interpolation. Observed data were available at well sample locations scattered throughout the domain, and the interpolation was used to estimate the mass of contaminant between the sample locations.

Source term development consisted of the following steps:

- Initial interpolation runs of the observed data within the model domain
- Visual inspection of the results of interpolation runs
- Selection of an acceptable interpolation
- Final interpolation
- Analysis (post processing) of the final interpolation

The model domain consisted of a surface soil (0–1 ft below surface) layer and six other layers down to the top of the RGA. Only the surface soil (Layer 0) was interpolated separately because the majority of Layer 1 samples were sediments that were different in nature than the subsurface soils.

The geospatial interpolation techniques available in SADA for three-dimensional analyses are as follows:

- Nearest neighbor
- Inverse distance
- Kriging

An example of the geospatial analyses is provided for arsenic, *cis*-1,2-DCE, and TCE at SWMU 2. Figure E.2.1 provides the sample locations for SWMU 2. The measured data at the sample locations were used in SADA to evaluate the predicted mass for the contaminants using the various geospatial interpolation techniques. Three-dimensional fence diagrams of the interpolated arsenic, *cis*-1,2-DCE, and TCE concentrations were constructed using the Mining Visualization System, Version 9.0, by the C Tech Development Corporation. The fence diagrams are shown in Figures E2.2 through E2.4. These diagrams were used for comparison to the SADA interpolation results.

The results of the geospatial analysis using SADA for the three interpolation techniques are provided in Tables E2.2 through E2.10 for arsenic, *cis*-1,2-DCE and TCE.

Table E2.2. Nearest Neighbor Source Term Characteristics Developed by SADA for Arsenic at SWMU 2

SADA Layer	Dept h (ft)	Area (ft²)	Volume (ft³)	Mass (gm)^a
L1	0-1	4.80E+04	4.80E+04	4.39E+04
L2	01-10	3.96E+04	3.56E+05	1.37E+05
L3	10-20	3.96E+04	4.36E+05	1.23E+05
L4	20-30	3.96E+04	4.36E+05	1.04E+05
L5	30-40	3.96E+04	4.36E+05	1.16E+05
L6	40-50	3.96E+04	4.36E+05	1.06E+05
L7	50-64	3.96E+04	3.56E+05	7.26E+04
Total Mass				7.03E+05

Table E2.3. Inverse Distance Source Term Characteristics Developed by SADA for Arsenic at SWMU 2

SADA Layer	Dept h (ft)	Area (ft²)	Volume (ft³)	Mass (gm)^a
L1	0-1	4.80E+04	4.80E+04	0.00E+00
L2	01-10	3.96E+04	3.56E+05	1.65E+05
L3	10-20	3.96E+04	4.36E+05	8.01E+04
L4	20-30	3.96E+04	4.36E+05	5.93E+04
L5	30-40	3.96E+04	4.36E+05	1.06E+05
L6	40-50	3.96E+04	4.36E+05	1.34E+05
L7	50-64	3.96E+04	3.56E+05	3.25E+04
Total Mass				5.78E+05

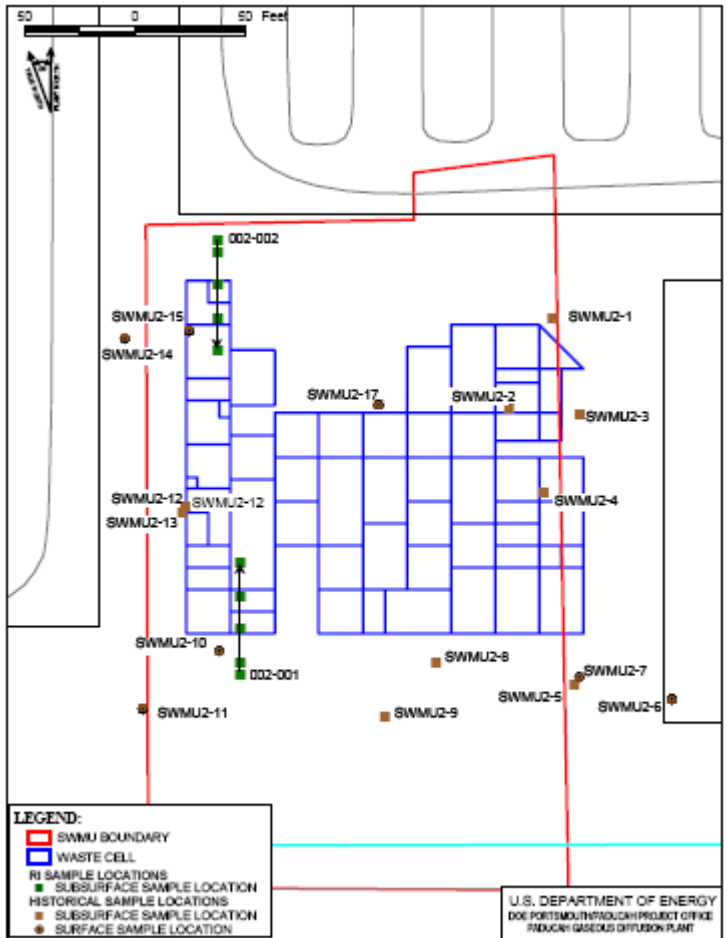
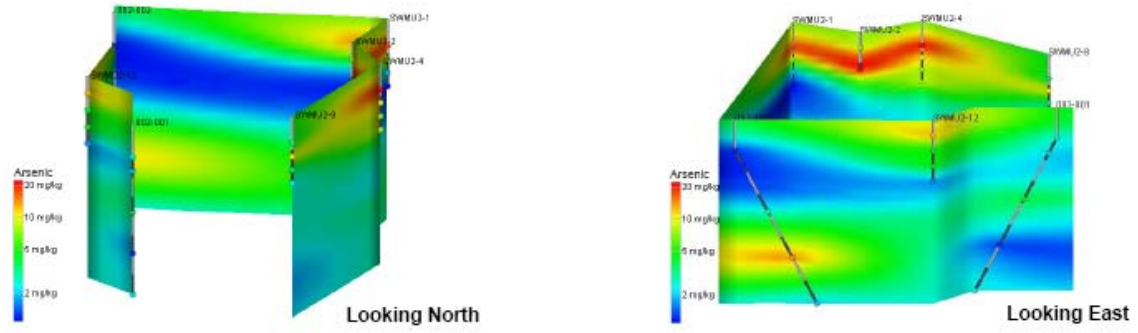


Figure E2.1. Samples Locations at SWMU 2



SWMU 2 Arsenic: 0 – 60 ft bgs
(Vertical Exaggeration = 2)

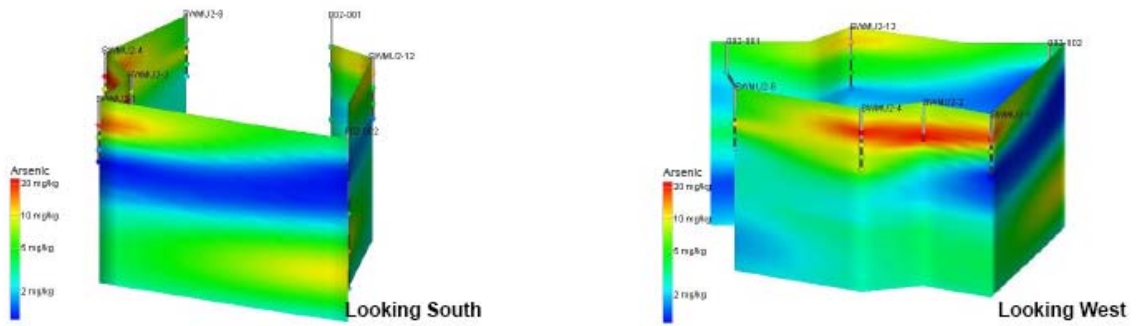
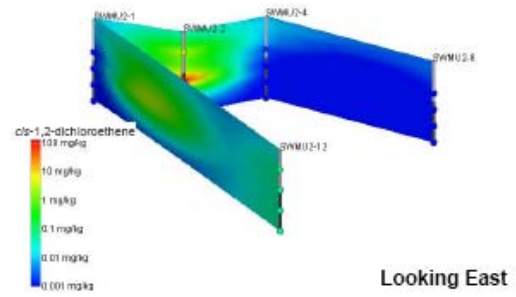
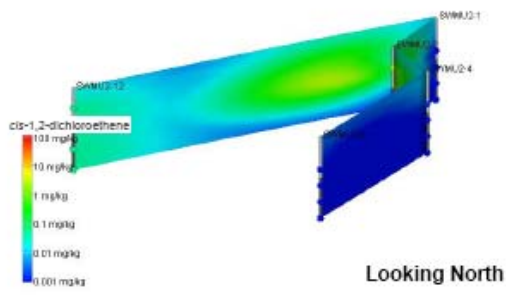


Figure E2.2. Arsenic Concentration Fence Diagrams for SWMU 2



SWMU 2 *cis*-1,2-DCE: 0 – 20 ft bgs
(Vertical Exaggeration = 2)

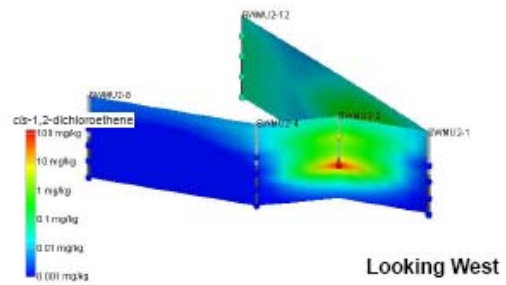
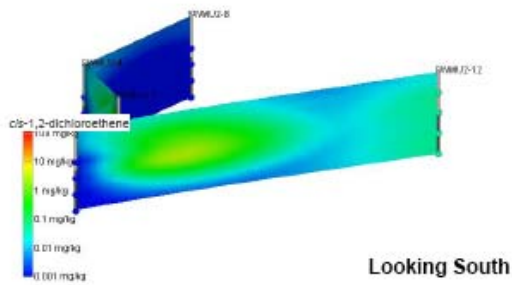
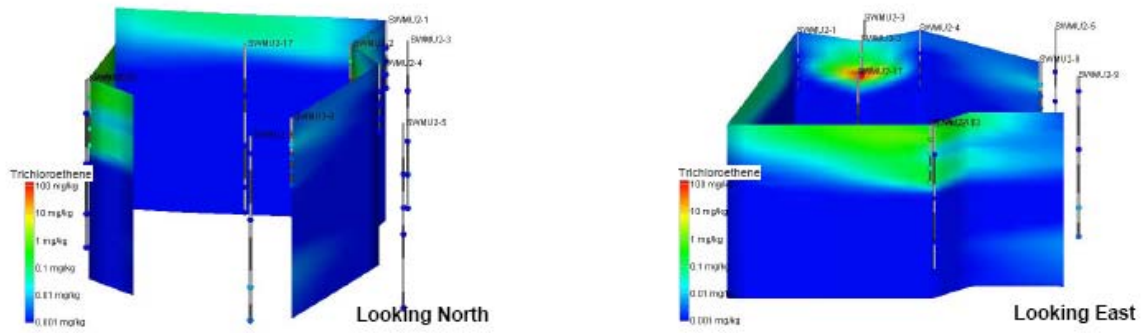


Figure E2.3. *Cis*-1,2-DCE Concentration Fence Diagrams for SWMU 2



SWMU 2 TCE: 0 – 60 ft bgs
(Vertical Exaggeration = 2)

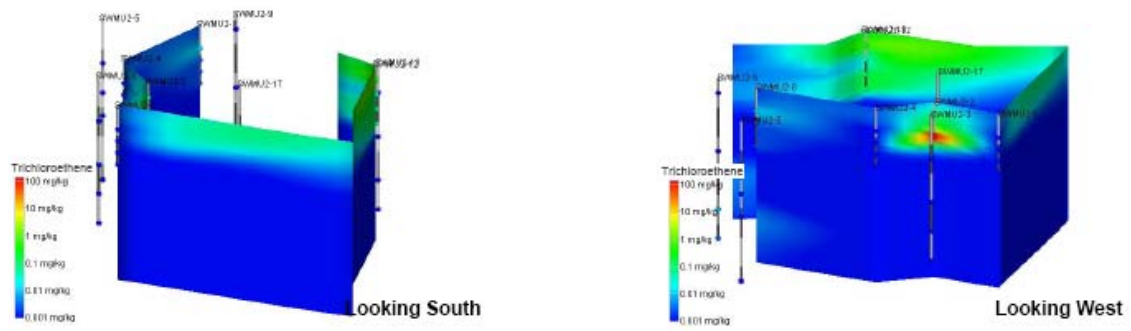


Figure E2.4. TCE Concentration Fence Diagrams for SWMU 2

Table E2.4. Ordinary Kriging Source Term Characteristics Developed by SADA for Arsenic at SWMU 2

SADA Layer	Dept h (ft)	Area (ft²)	Volume (ft³)	Mass (gm)^a
L1	0-1	4.80E+04	4.80E+04	4.38E+04
L2	01-10	3.96E+04	3.56E+05	1.44E+05
L3	10-20	3.96E+04	4.36E+05	8.47E+04
L4	20-30	3.96E+04	4.36E+05	6.19E+04
L5	30-40	3.96E+04	4.36E+05	9.85E+04
L6	40-50	3.96E+04	4.36E+05	1.32E+05
L7	50-64	3.96E+04	3.56E+05	3.25E+04
Total Mass				5.97E+05

Table E2.5. Nearest Neighbor Source Term Characteristics Developed by SADA for *cis*-1,2-DCE at SWMU 2

SADA Layer	Dept h (ft)	Area (ft²)	Volume (ft³)	Mass (gm)^a
L1	0-1	0.00E+00	0.00E+00	0.00E+00
L2	01-10	1.64E+04	1.48E+05	7.17E+03
L3	10-20	6.80E+03	7.48E+04	3.55E+05
L4	20-30	9.60E+03	1.06E+05	3.08E+05
L5	30-40	8.40E+03	9.24E+04	3.08E+05
L6	40-50	1.16E+04	1.28E+05	2.37E+05
L7	50-64	1.12E+04	1.01E+05	1.55E+05
Total Mass				1.37E+06

Table E2.6. Inverse Distance Source Term Characteristics Developed by SADA for *cis*-1,2-DCE at SWMU 2

SADA Layer	Dept h (ft)	Area (ft²)	Volume (ft³)	Mass (gm)^a
L1	0-1	0.00E+00	0.00E+00	0.00E+00
L2	01-10	3.96E+04	3.56E+05	4.18E+04
L3	10-20	3.96E+04	4.36E+05	3.64E+04
L4	20-30	0.00E+00	0.00E+00	0.00E+00
L5	30-40	3.96E+04	4.36E+05	6.67E+02
L6	40-50	3.96E+04	4.36E+05	1.07E+03
L7	50-64	3.96E+04	3.56E+05	1.07E+02
Total Mass				8.00E+04

Table E2.7. Ordinary Kriging Source Term Characteristics Developed by SADA for *cis*-1,2-DCE at SWMU 2

SADA Layer	Dept h (ft)	Area (ft²)	Volume (ft³)	Mass (gm)^a
L1	0-1	0.00E+00	0.00E+00	0.00E+00
L2	01-10	2.52E+04	2.27E+05	7.94E+04
L3	10-20	3.20E+04	3.52E+05	2.33E+05
L4	20-30	0.00E+00	0.00E+00	0.00E+00
L5	30-40	3.84E+04	4.22E+05	6.03E+02
L6	40-50	3.96E+04	4.36E+05	1.09E+03
L7	50-64	3.96E+04	3.56E+05	1.06E+02
Total Mass				3.15E+05

Table E2.8. Nearest Neighbor Source Term Characteristics Developed by SADA for TCE at SWMU 2

SADA Layer	Dept h (ft)	Area (ft²)	Volume (ft³)	Mass (gm)^a
L1	0-1	0.00E+00	0.00E+00	0.00E+00
L2	01-10	9.60E+03	8.64E+04	5.11E+02
L3	10-20	9.20E+03	1.01E+05	1.78E+05
L4	20-30	1.16E+04	1.28E+05	1.28E+05
L5	30-40	1.16E+04	1.28E+05	7.69E+04
L6	40-50	1.28E+04	1.41E+05	5.20E+04
L7	50-64	1.08E+04	9.72E+04	8.06E+02
Total Mass				4.37E+05

Table E2.9. Inverse Distance Source Term Characteristics Developed by SADA for TCE at SWMU 2

SADA Layer	Dept h (ft)	Area (ft²)	Volume (ft³)	Mass (gm)^a
L1	0-1	0.00E+00	0.00E+00	0.00E+00
L2	01-10	4.80E+04	4.32E+05	2.73E+04
L3	10-20	4.80E+04	5.28E+05	3.80E+04
L4	20-30	4.80E+04	5.28E+05	9.20E-01
L5	30-40	4.80E+04	5.28E+05	3.27E+02
L6	40-50	4.80E+04	5.28E+05	2.22E+02
L7	50-64	4.80E+04	4.32E+05	2.85E+03
Total Mass				6.87E+04

Table E2.10. Ordinary Kriging Source Term Characteristics Developed by SADA for TCE at SWMU 2

SADA Layer	Dept h (ft)	Area (ft²)	Volume (ft³)	Mass (gm)^a
L1	0-1	0.00E+00	0.00E+00	0.00E+00
L2	01-10	2.84E+04	2.56E+05	6.07E+04
L3	10-20	2.08E+04	2.29E+05	6.19E+04
L4	20-30	1.56E+04	1.72E+05	9.82E-01
L5	30-40	2.92E+04	3.21E+05	6.02E+02
L6	40-50	4.28E+04	4.71E+05	1.07E+03
L7	50-64	4.80E+04	4.32E+05	1.90E+03
Total Mass				1.26E+05

The nearest neighbor geospatial interpolation provided the maximum total mass of contaminants in the system when compared to the inverse distance and ordinary kriging algorithms. This addresses in part the known low bias in the data caused by the inability to sample the waste. The nearest neighbor interpolation was selected because it provided greater contrast of the interpolated concentrations among the model cells and greater ease of source delineation through visual inspection as shown in Figures E2.5 through E2.7 for TCE. The inverse distance interpolation did not distinctly delineate the modeled contaminant plumes. Kriging interpolations in SADA involved variogram modeling. Modelers concluded that kriging interpolations were not suitable because semi-variogram values for the observed data did not follow monotonically increasing trends as shown in Figure E2.8.

The verification report for SADA (EPA 2000) states that “although geostatistical-based kriging interpolation approaches are more mathematically rigorous than the simple interpolation approaches using nearest neighbor, they are not necessarily better representations of the data. Statistical and geostatistical approaches attempt to minimize the mathematical constraint, similar to a least squares minimization used in curve-fitting of data. While the solution provided is the “best” answer within the mathematical constraints applied to the problem, it is not necessarily the best fit of the data. There are two reasons for this:

“First, in most environmental problems, the data are insufficient to determine the optimum model to use to assess the data. Typically, there are several different models that can provide a defensible assessment of spatial correlation in the data. Each of these models has its own strengths and limitations, and the model choice is subjective” (EPA 2000).

“This conundrum leads to the second reason for the difficulty, if not impossibility, of finding the most appropriate model to use for interpolation—which is that unless the analyst is extremely fortunate, the measured data will not conform to the mathematical model used to represent the data. At best, the interpolation can be reviewed to determine if it is consistent with the data” (EPA 2000).

The nearest neighbor algorithm was used to evaluate the remaining COPCs at all SWMUs. Figure E2.9 shows typical flood contours of the source zone in the UCRS developed from the SADA interpolation in plan view. Three-dimensional figures illustrating the spatial distribution of COCs under each SWMU are provided in Figures E2.10–17. In addition, Figures E2.11 through E2.18 provide examples of the SADA nearest neighbor interpolation results for the layers with the highest total mass for COCs providing the majority of the risk or hazard at the SWMUs. Sample location circles were shaded to match legend scale on right side of

figure. Due to the large number of maps created with SADA (882 total) only one individual soil layer per SWMU with highest total mass was illustrated.

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UT (University of Tennessee) 2002. *Spatial Analysis and Decision Assistance (SADA)*, Version 2.3, User Guide, January. Accessible at <http://www.tiem.utk.edu/~sada/>

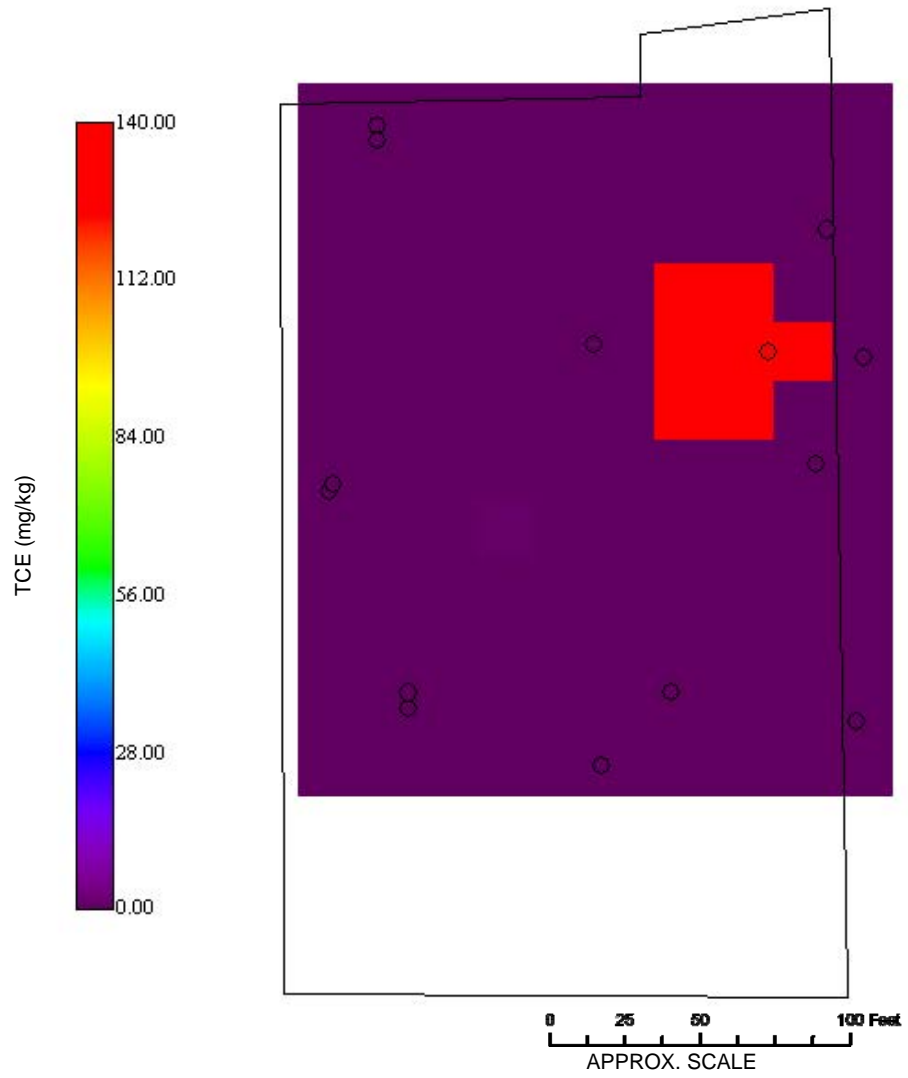


Figure E2.5. Interpolation Nearest Neighbor for SWMU 2–TCE

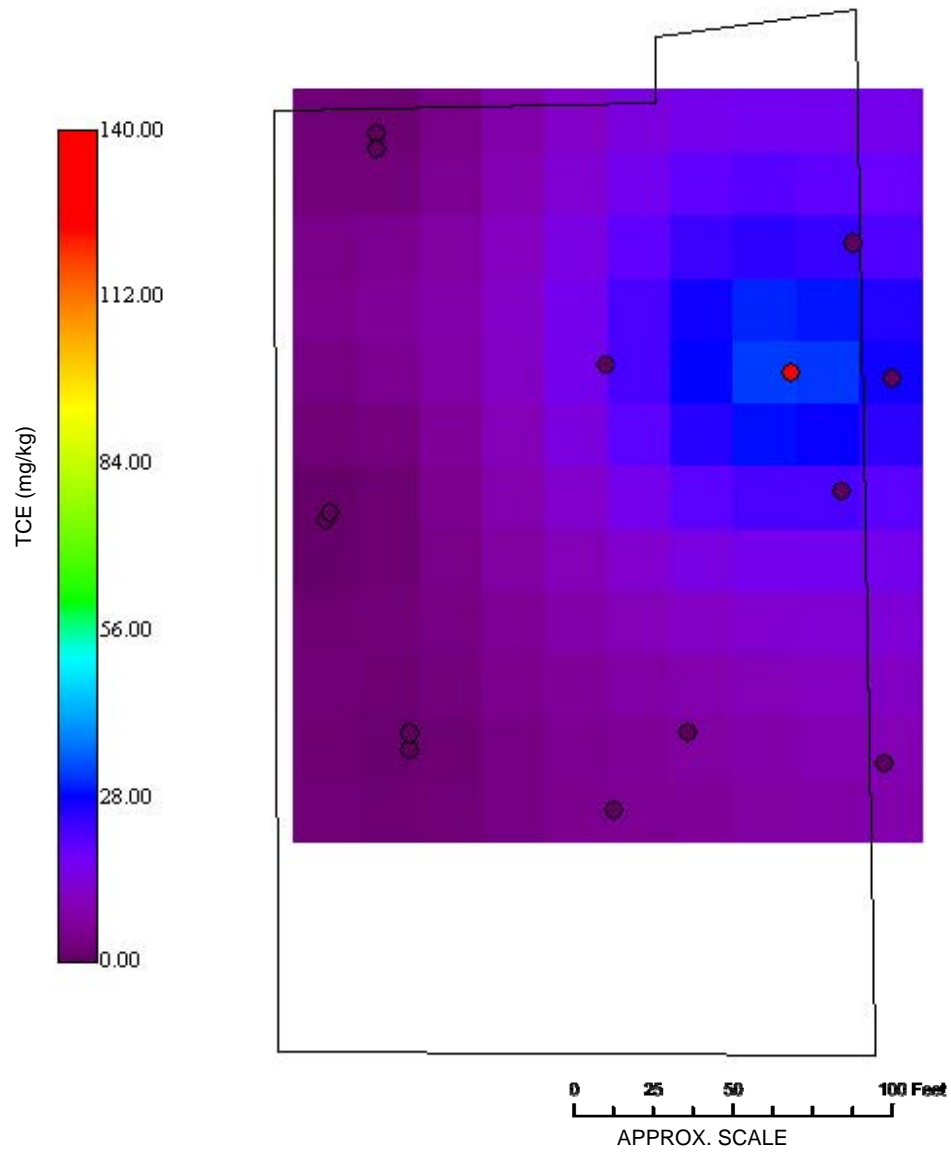


Figure E2.6. Interpolation Inverse Distance for SWMU 2-TCE

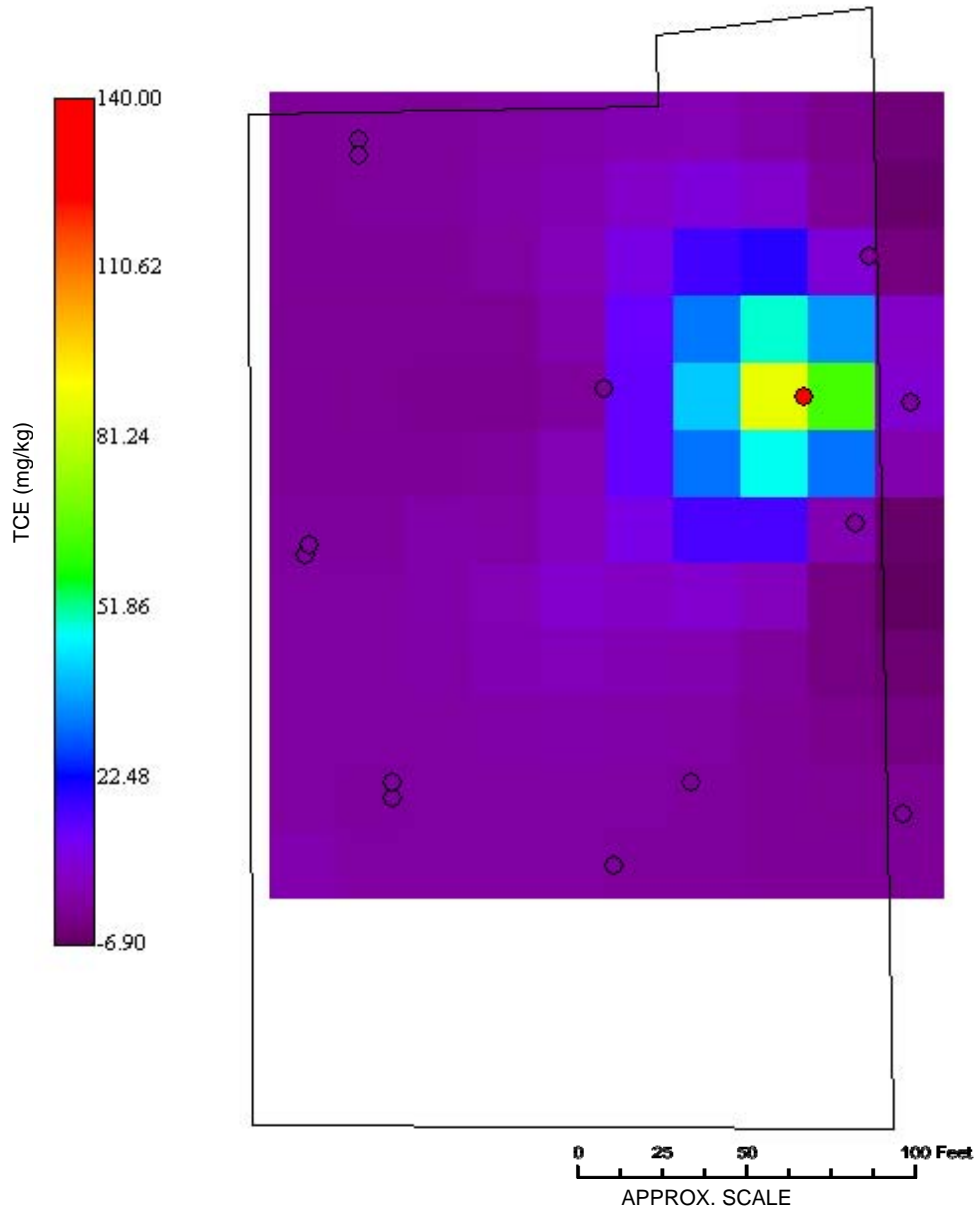


Figure E2.7. Interpolation Ordinary Kriging for SWMU 2-TCE

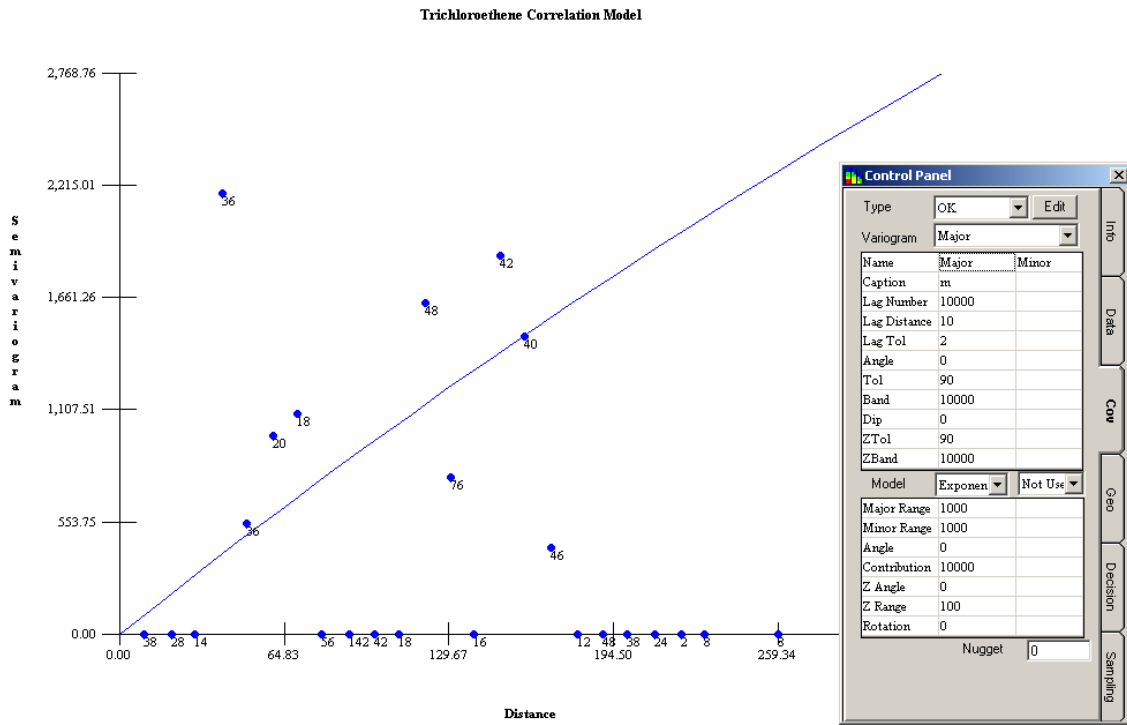


Figure E2.8. Ordinary Kriging Semi-Variogram for SWMU 2-TCE

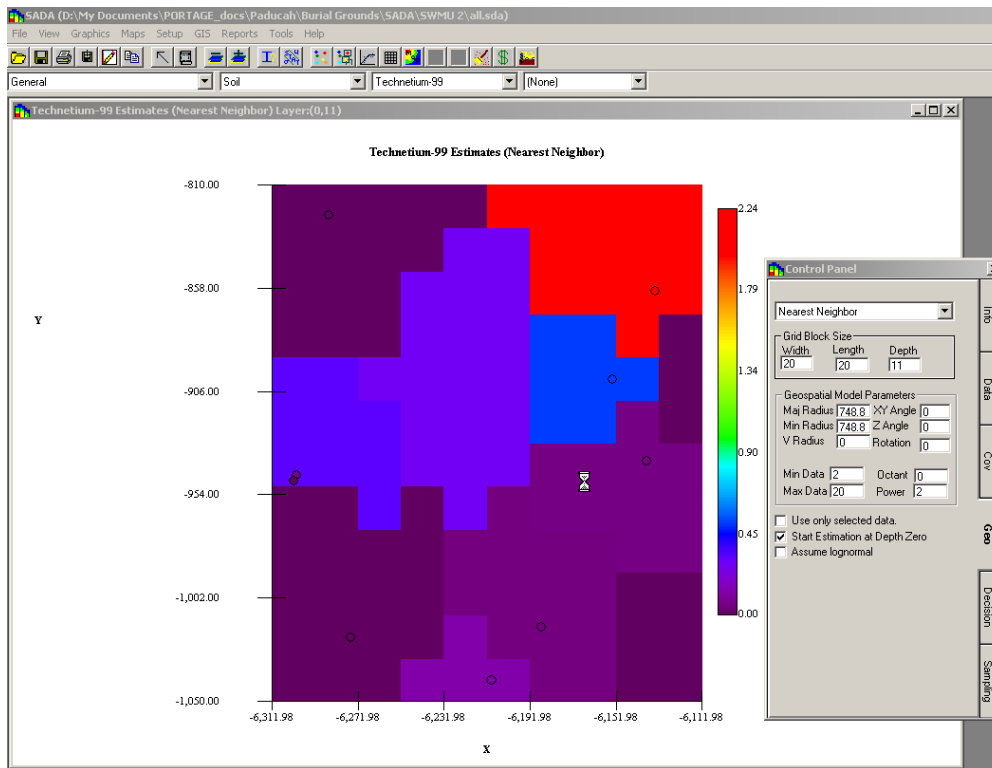


Figure E2.9. Typical SADA Nearest Neighbor Interpolation Map

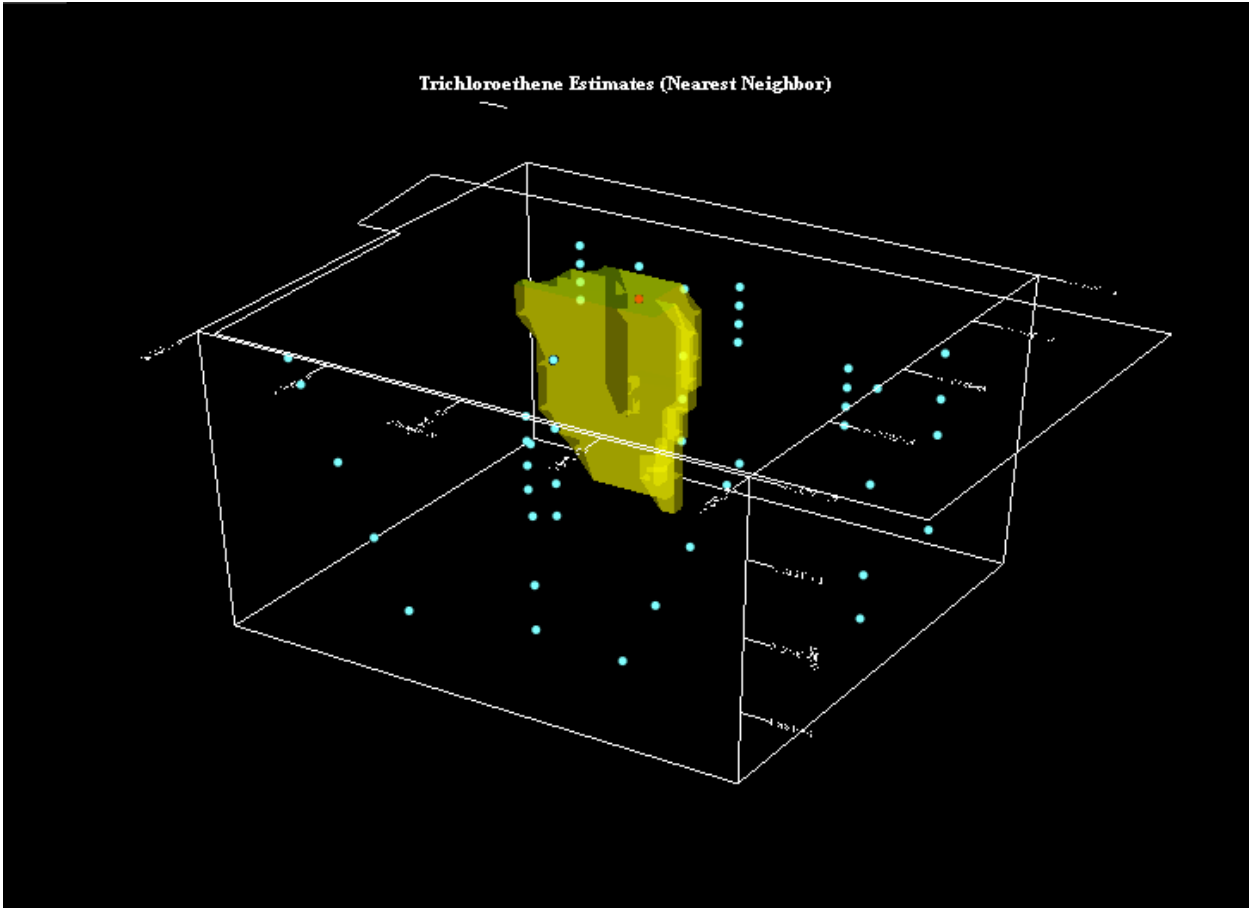


Figure E2.10. SWMU 2 TCE SADA Nearest Neighbor Interpolation Map (3-D)

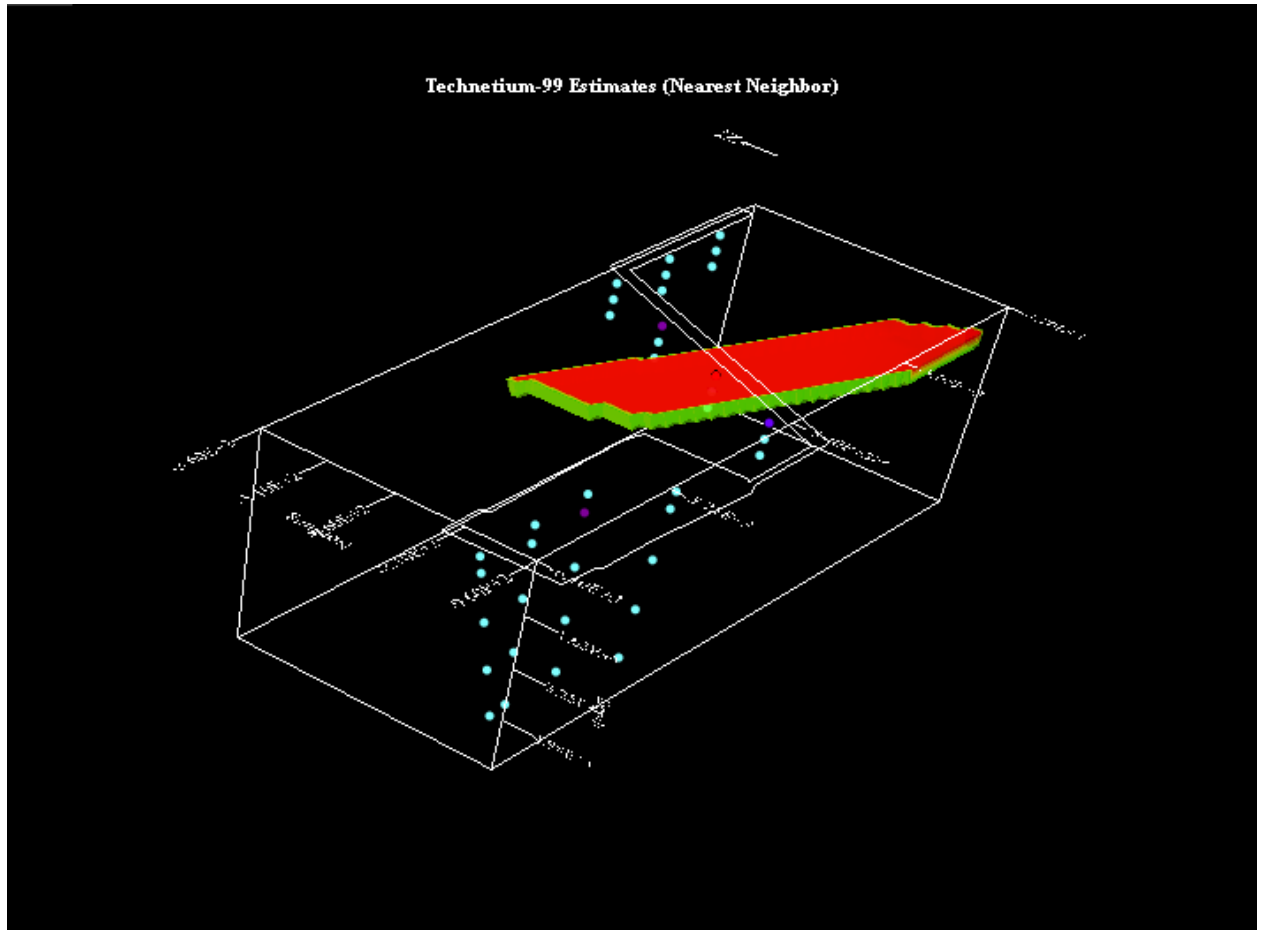


Figure E2.11. SWMU 3 ⁹⁹Tc SADA Nearest Neighbor Interpolation Map (3-D)

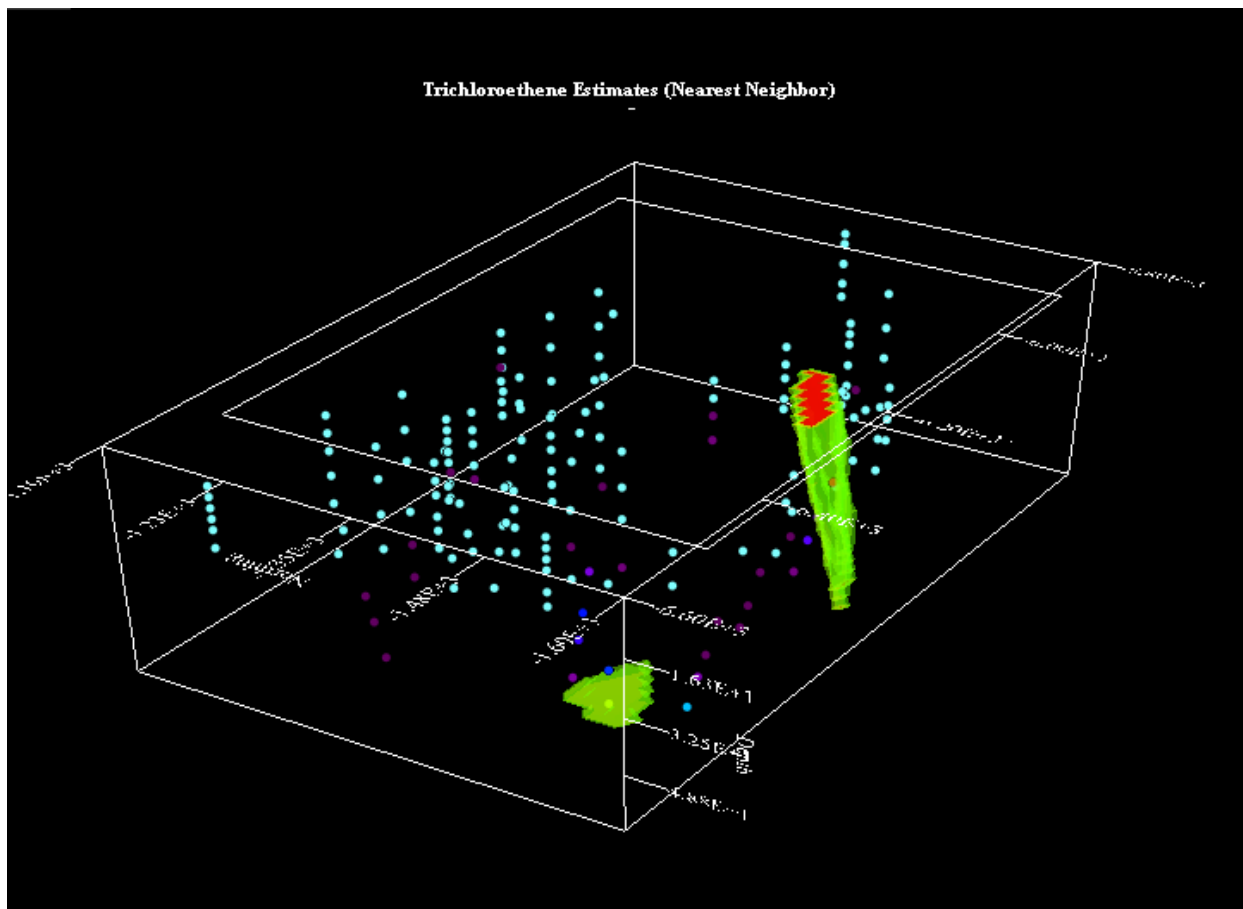


Figure E2.12. SWMU 4 TCE SADA Nearest Neighbor Interpolation Map (3-D)

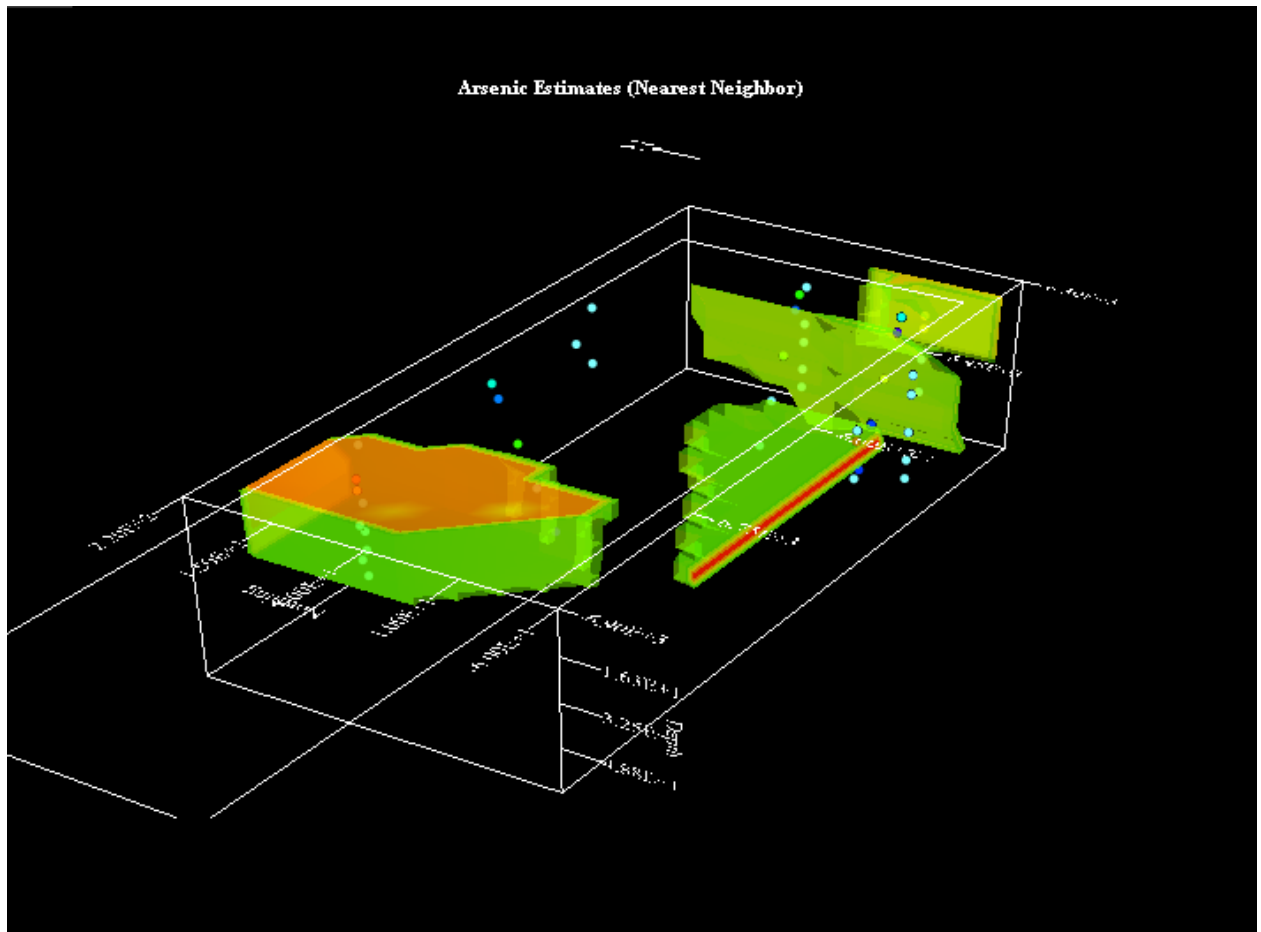


Figure E2.13. SWMU 5 Arsenic SADA Nearest Neighbor Interpolation Map (3-D)

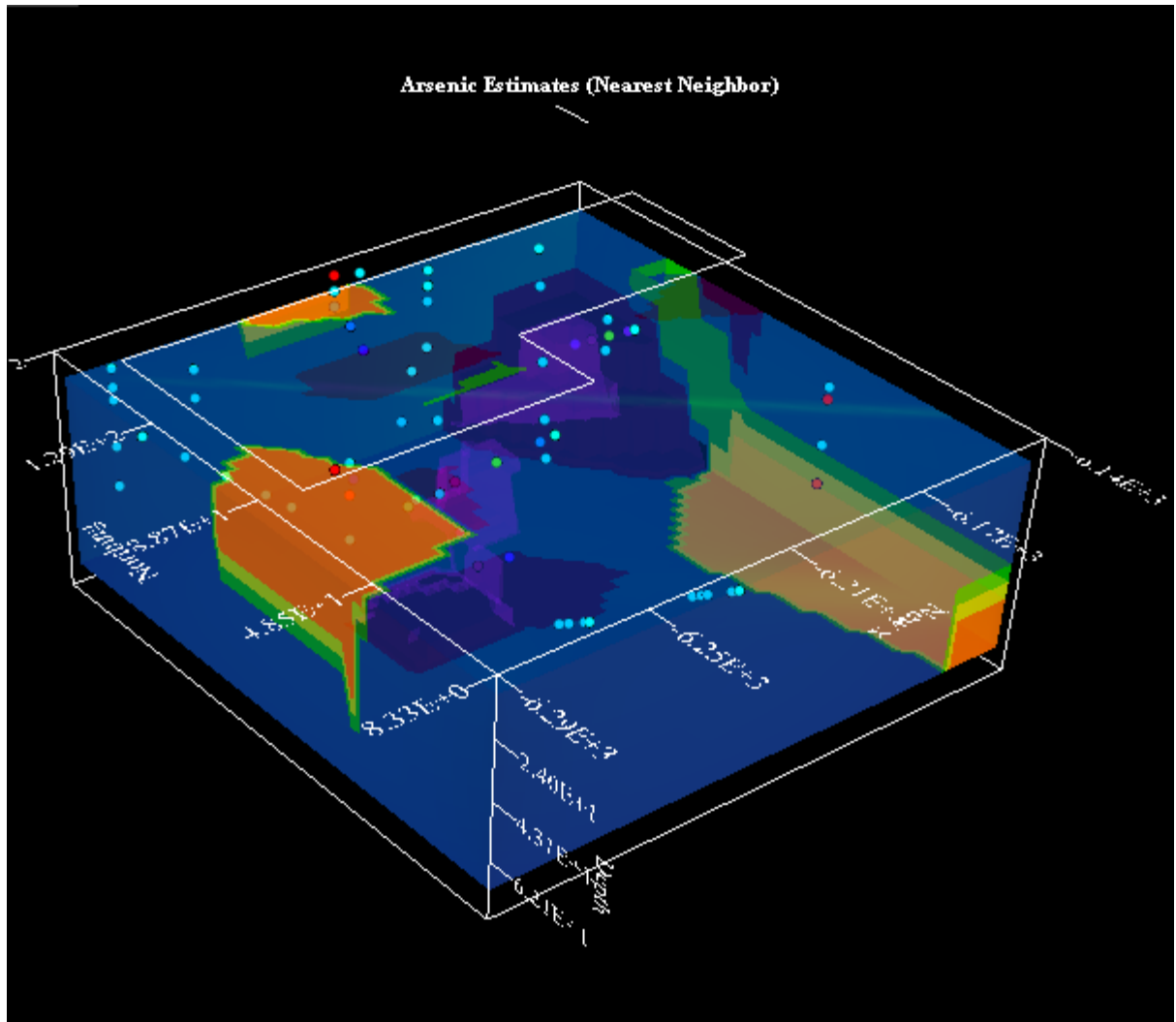


Figure E2.14. SWMU 6 Arsenic SADA Nearest Neighbor Interpolation Map (3-D)

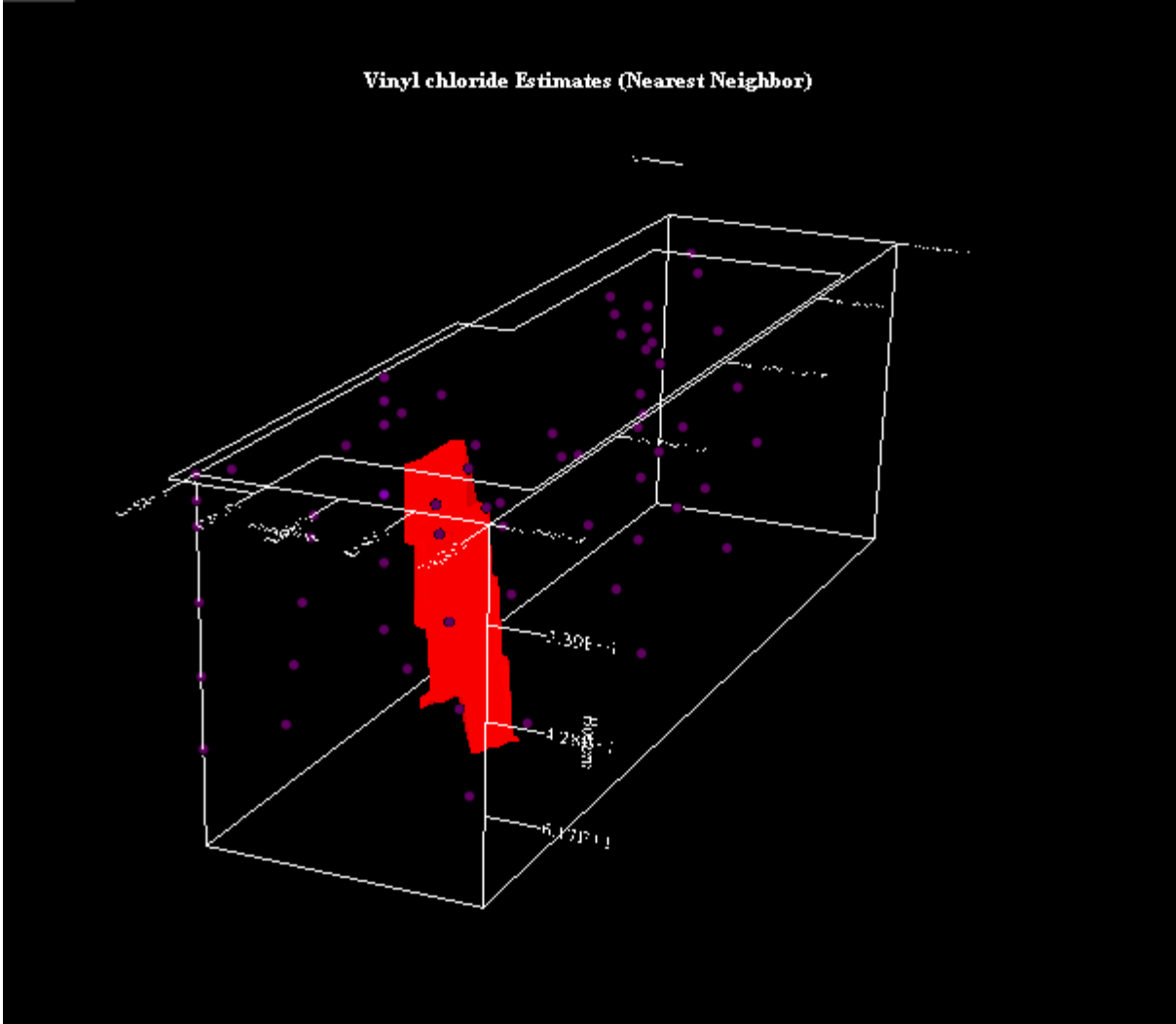


Figure E2.15. SWMU 7 Vinyl Chloride SADA Nearest Neighbor Interpolation Map (3-D)

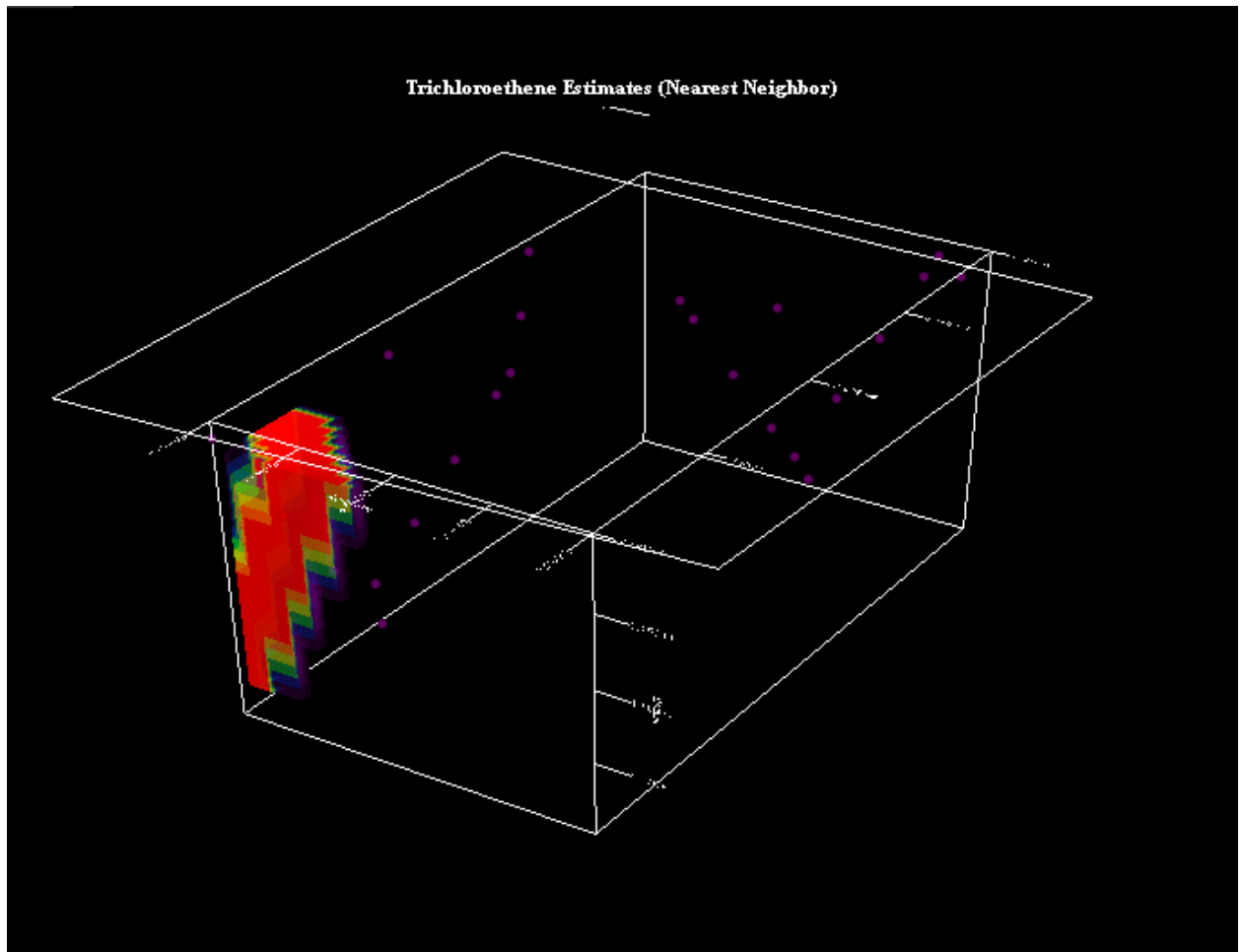


Figure E2.16. SWMU 30 TCE SADA Nearest Neighbor Interpolation Map (3-D)

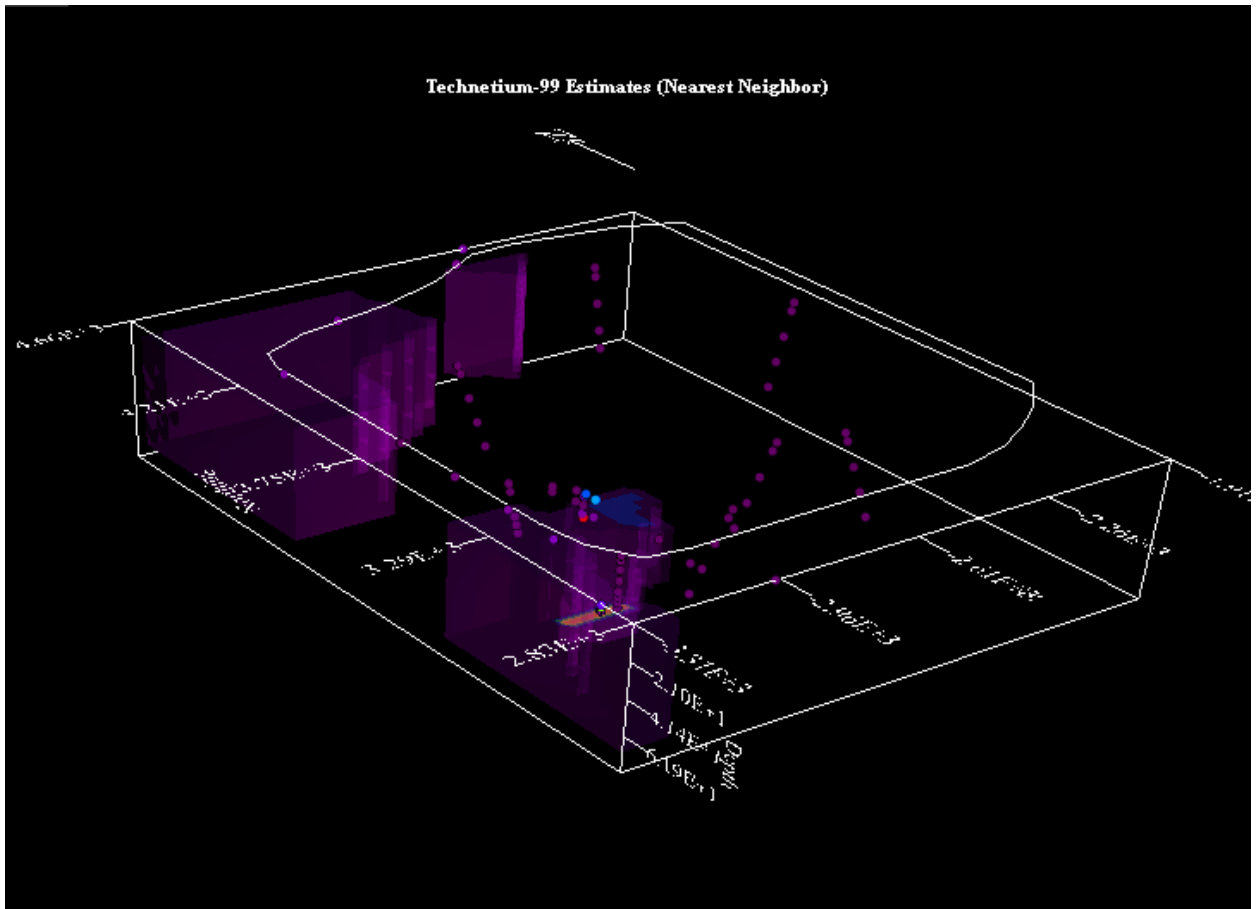


Figure E2.17. SWMU 145 ⁹⁹Tc SADA Nearest Neighbor Interpolation Map (3-D)

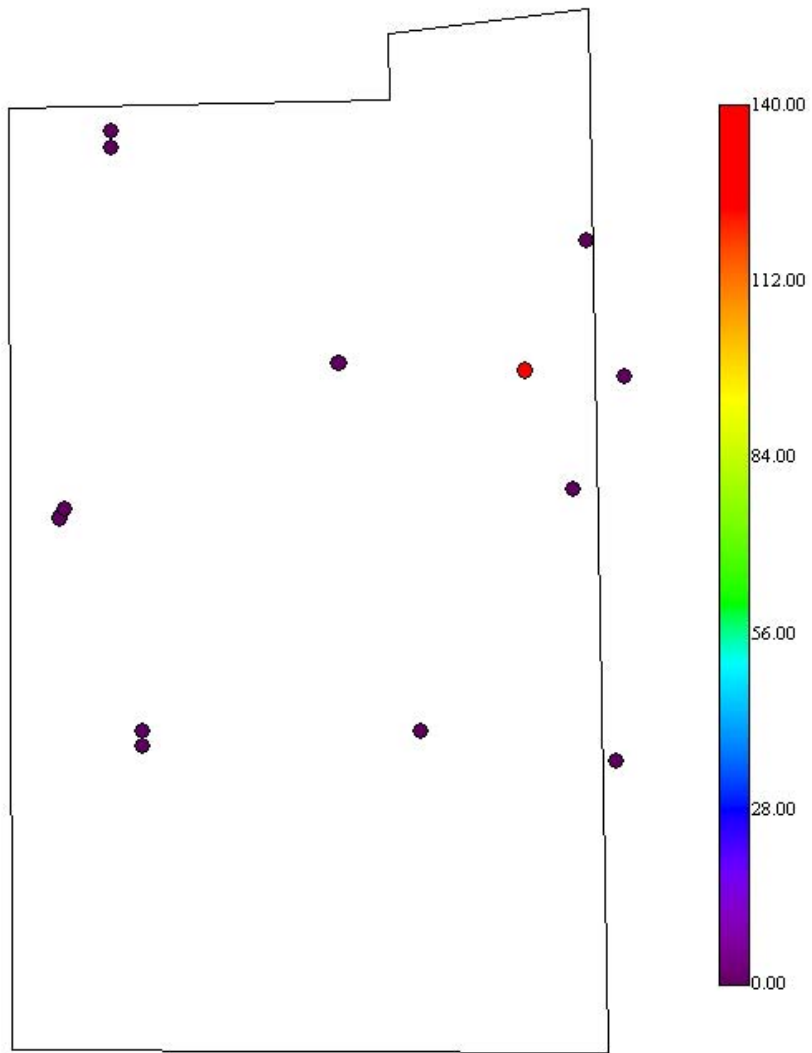


Figure E2.18. SWMU 2, TCE 10-20 ft bgs

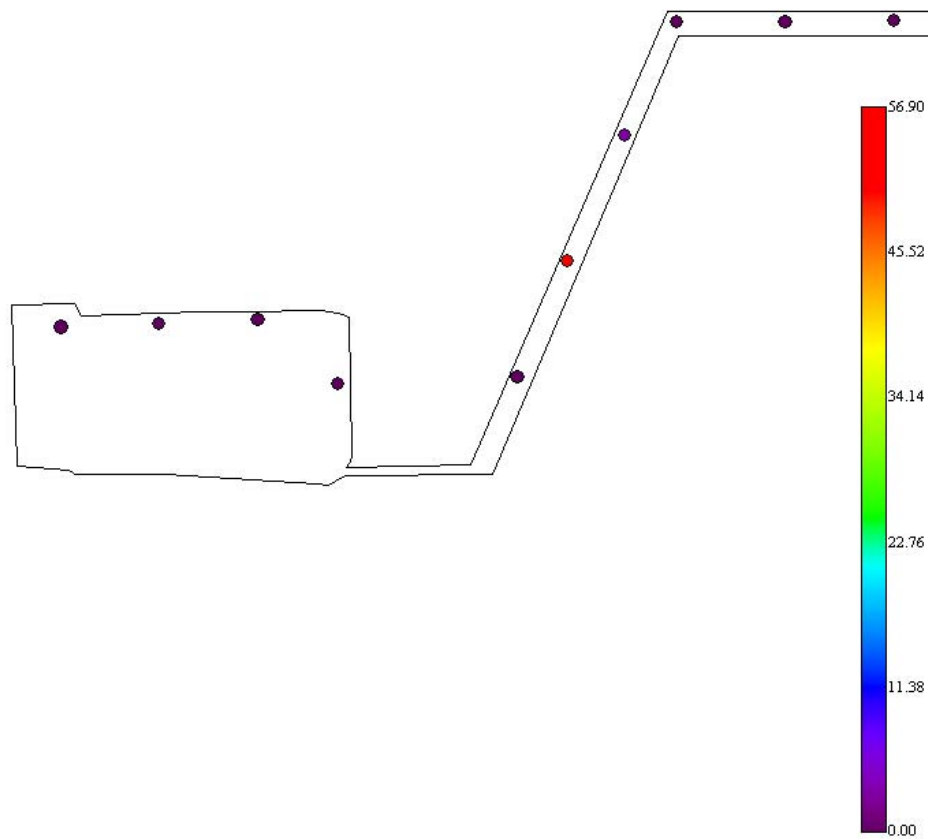


Figure E2.19. SWMU 3, ^{99}Tc 1-10 ft bgs

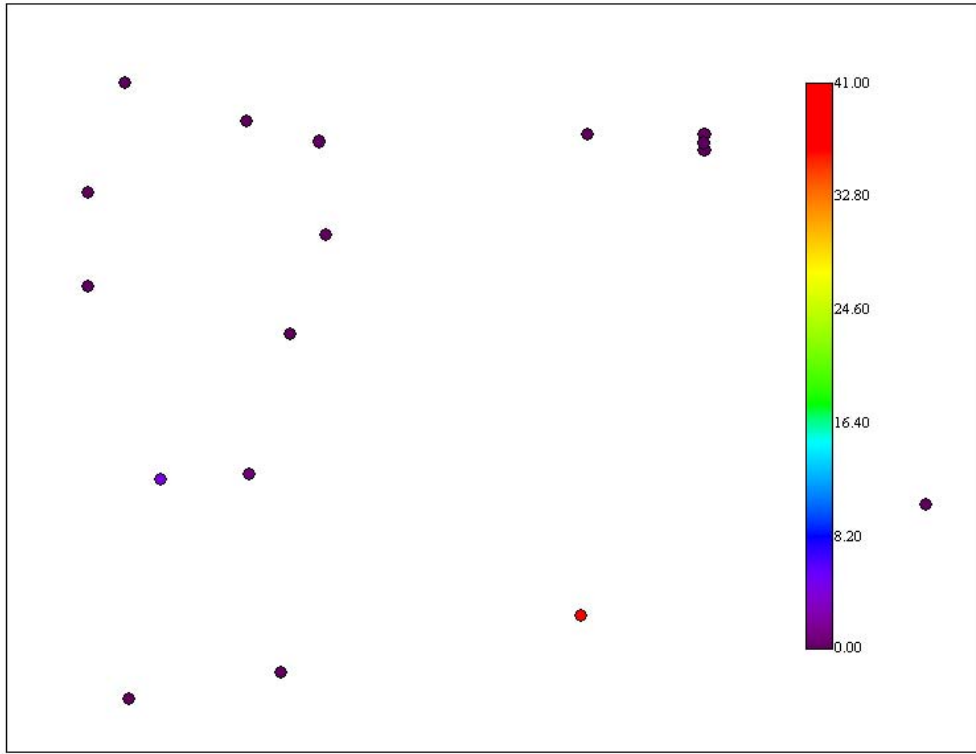


Figure E2.20. SWMU 4, TCE 30-40 ft bgs

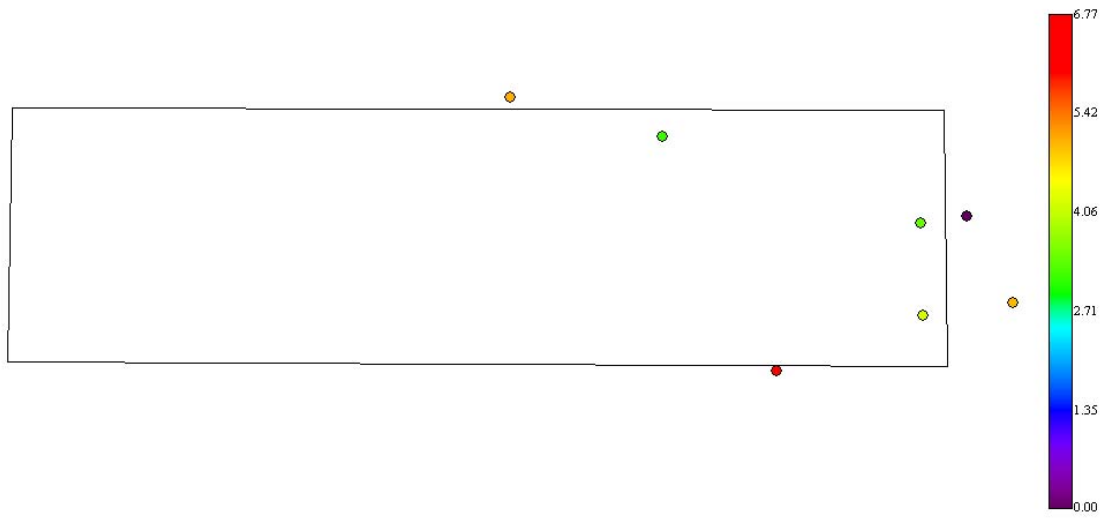


Figure E2.21. SWMU 5, Arsenic 20-30 ft bgs

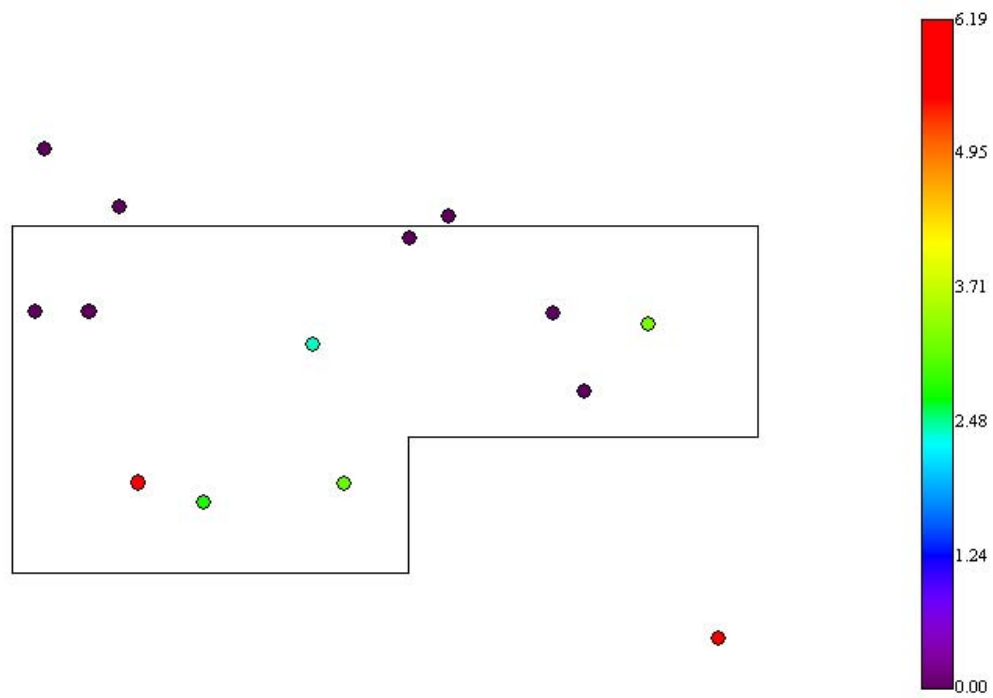


Figure E2.22. SWMU 6, Arsenic 20-30 ft bgs

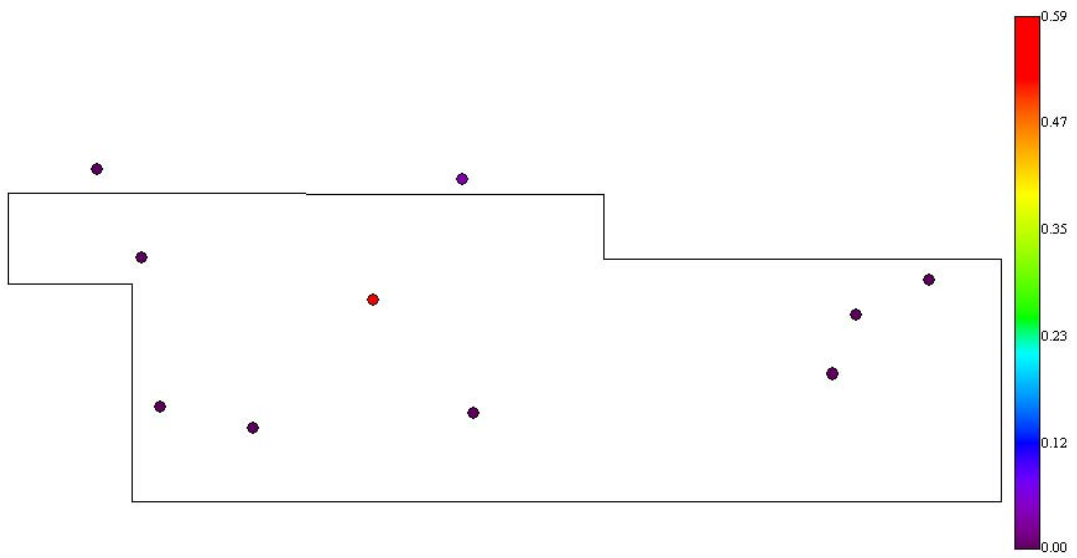


Figure E2.23. SWMU 7, Vinyl Chloride 30-40 ft bgs

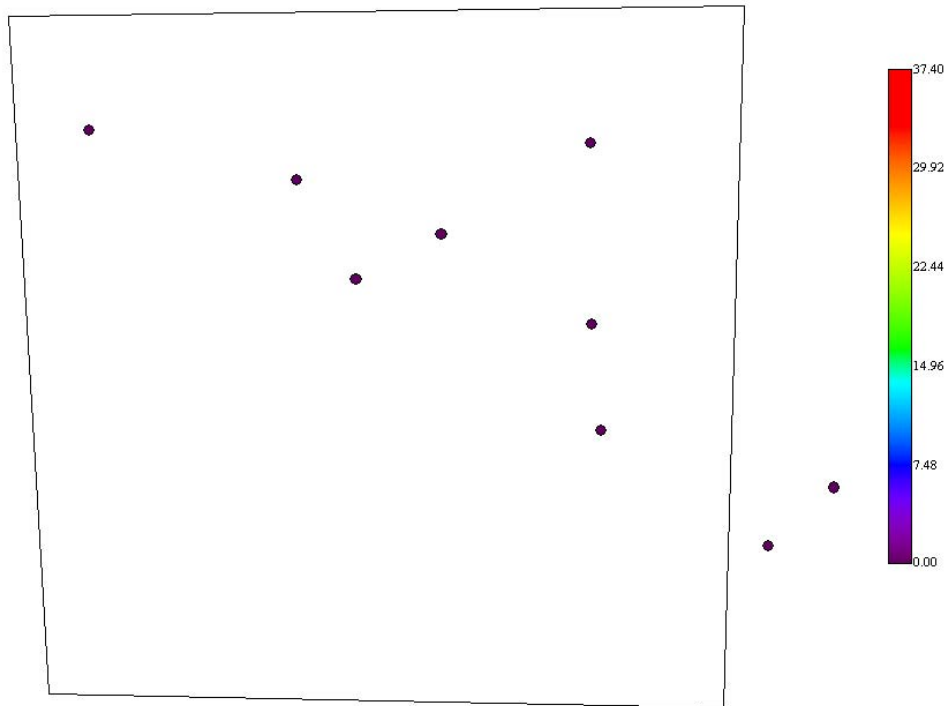


Figure E2.24. SWMU 30, TCE 1-10 ft bgs

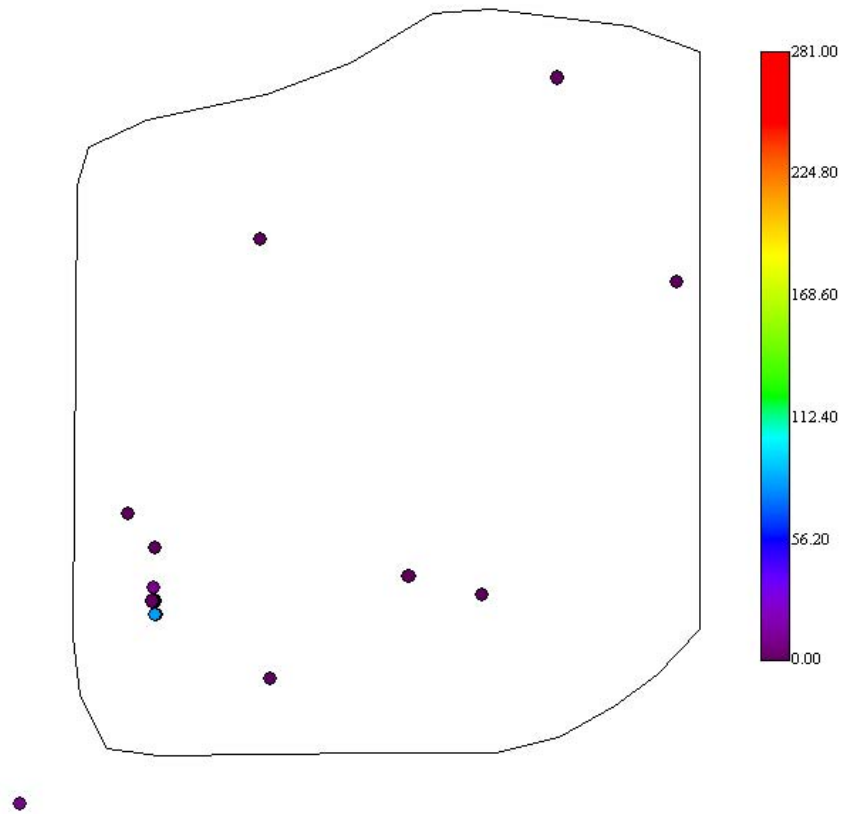


Figure E2.25. SWMU 145, ^{99}Tc 1-10 ft bgs

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**APPENDIX E
ATTACHMENT 3**

GROUNDWATER ANALYTE SCREENING ANALYSIS

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E.3. GROUNDWATER ANALYTE SCREENING ANALYSIS

Tables E3.1 to E3.8 show the comparison of maximum detected concentration of analytes in soil (in any soil horizon) to soil screening levels (SSLs) protective of groundwater and the other criteria described in Section 5 of this Remedial Investigation (RI). These criteria were used to determine which soil analytes were selected to be modeled to groundwater. Modeled groundwater values were compared to residential child groundwater no action levels (NALs) as described in Section 5 to produce the final list of groundwater analytes and their modeled concentrations that were used in the risk assessment in Appendix F.

Table E3.1. Summary of Screening for Detected Analytes–SWMU 2 Groundwater

Analyte	SSL Protective of Groundwater ^A	Exceed SSL	Maximum Detection	Detection Frequency	Included in Groundwater Modeling
Metals (mg/kg)					
Aluminum	NA ^G		1.10E+04	5/5	No
Antimony	2.54E-02	Yes	1.01E+01	2/5	Yes
Arsenic	1.05E-03	Yes	3.00E+01	21/21	Yes
Barium	NA ^G		3.50E+02	21/21	No
Beryllium	2.11E+00	No	1.80E+00	22/23	No
Calcium ^B	NA ^G		2.40E+03	3/3	No
Chromium	3.17E+06	No	3.00E+01	21/21	No
Cobalt	NA ^G		4.74E+00	3/3	No
Copper	NA ^G		7.80E+00	3/3	No
Iron	NA ^G		4.10E+04	3/3	No
Lead	NA ^G		2.90E+01	4/4	No
Magnesium ^B	NA ^G		1.60E+03	3/3	No
Manganese	2.26E+00	Yes	1.20E+03	21/21	Yes
Mercury	2.22E-02	Yes	1.40E-01	1/3	No ^I
Nickel	1.98E+00	Yes	3.70E+01	21/21	Yes
Potassium ^B	NA ^G		9.40E+02	2/2	No
Sodium ^B	NA ^G		2.01E+02	3/3	No
Thallium	NA ^G		4.50E+00	12/21	No
Uranium	NA ^G		1.50E+03	12/50	Yes
Vanadium	9.25E+00	Yes	3.80E+01	21/21	Yes
Zinc	2.78E+01	Yes	1.40E+02	3/3	No ^I
Organic Compounds (mg/kg)					
Total PCB ^C	4.92E-02	Yes	4.35E+00	NA	No ^E
Total PAH ^D	NA ^G		1.67E-01	NA	No ^E
alpha-Chlordane	NA ^G		7.80E-04	1/3	No
Aroclor-1248	NA ^G		4.2E+00	5/20	Yes
Aroclor-1254	NA ^G		1.1E+00	1/20	Yes
Aroclor-1260	NA ^G		5.0E-01	3/20	Yes
delta-BHC	NA ^G		6.50E-03	1/6	No
Gamma-Chlordane	NA ^G		3.00E-03	2/3	No
Benzo(a)pyrene	1.95E-03	Yes	1.40E-01	1/3	Yes
Benzo(b)fluoranthene	NA ^G		2.70E-01	2/3	No
Benzo(ghi)perylene	NA ^G		1.50E-01	1/3	No
Dibenzofuran	NA ^G		1.80E-01	2/3	No
Naphthalene	1.21E-03	Yes	1.80E-01	1/2	Yes
Phenanthrene	NA ^G		5.70E-01	2/3	No
Pyrene	3.81E+00	No	2.00E-01	1/2	No

Table E3.1. Summary of Screening for Detected Analytes–SWMU 2 Groundwater (Continued)

Analyte	SSL Protective of Groundwater ^A	Exceed SSL	Maximum Detection	Detection Frequency	Included in Groundwater Modeling
<i>cis</i> -1,2-DCE	NA ^G		1.30E+02	4/23	Yes
Organic Compounds (mg/kg) (Continued)					
Methylene chloride	NA ^G		1.30E-03	3/4	No
TCE	9.69E-04	Yes	1.40E+02	8/50	Yes
Vinyl Chloride	1.17E-05	Yes	1.40E+00	1/21	No ^H
Radionuclides (pCi/g)					
Americium-241	3.89E+07	No	4.65E+00	54/55	No
Cesium-137 ^F	NA ^G		5.10E+01	16/55	No
Neptunium-237	6.20E+03	No	3.10E-01	51/54	No
Plutonium-239	4.46E+00	Yes	1.61E+01	21/55	Yes
Technetium-99	2.63E+01	No	1.46E+01	54/54	Yes
Uranium-234	1.22E+01	Yes	1.55E+02	55/55	Yes
Uranium-235/236	1.14E+01	Yes	2.58E+01	55/55	Yes
Uranium-238	8.62E+00	Yes	9.47E+02	55/55	Yes

^A Based on a residential child with dilution attenuation factor (DAF) = 1 (DOE 2001)

^B Compound is considered an essential nutrient and has been excluded in accordance with the PGDP Risk Methods Document (2001)

^C Total polychlorinated biphenyl (PCB) maximum concentration reported as actual analytical value and EPC reported as sum of all detected aroclors

^D Total polyaromatic hydrocarbon (PAH) reported as benzo(a)pyrene toxicity equivalence factor (TEF)

^E Groundwater modeling was not performed for Total PCB and Total PAH as they represent groups of compounds. PCBs and PAHs were assessed individually for groundwater modeling.

^F Not applicable as the radionuclide does not reach groundwater within 10,000 years precluding receptor uptake per Section 5 of main text and Appendix E

^G SSL not listed in Table A.7 of PGDP Risk Methods Document (2001)

^H Less than 5% detects, so analyte not retained

^I Maximum detected value less than the soil/sediment no action level (NAL) for a child resident in Table A.17 of PGDP Risk Methods Document (2001), so analyte not retained

Table E3.2. Summary of Screening for Detected Analytes–SWMU 3 Groundwater

Analyte	SSL Protective of Groundwater ^A	Exceed SSL	Maximum Detection	Detection Frequency	Included in Groundwater Modeling
Metals (mg/kg)					
Aluminum	NA ^D		1.18E+04	55/55	No
Antimony	2.54E-02	Yes	1.57E+01	6/55	No
Arsenic	1.05E-03	Yes	1.37E+01	50/55	Yes
Barium	NA ^D		1.27E+02	57/57	No
Beryllium	2.11E+00	No	1.06E+00	5/55	No
Calcium ^B	NA ^D		3.56E+04	56/57	No
Chromium	3.17E+06	No	3.57E+01	57/57	No
Cobalt	NA ^D		2.00E+01	42/58	No
Copper	NA ^D		3.00E+01	57/57	No
Iron	NA ^D		4.20E+04	57/57	No
Lead	NA ^D		2.39E+01	56/56	No
Magnesium ^B	NA ^D		2.50E+03	57/57	No
Manganese	2.26E+00	Yes	6.44E+02	57/57	Yes
Mercury	2.22E-02	Yes	2.40E-02	9/56	Yes
Molybdenum	1.55E-01	Yes	1.80E+01	6/57	Yes
Nickel	NA ^D		1.61E+01	11/56	Yes
Sodium ^B	NA ^D		4.45E+02	1/57	No
Uranium	NA ^D		6.10E-01	1/57	No
Uranium	NA ^D		8.36E+01	19/58	Yes
Vanadium	9.25E+00	Yes	3.37E+01	56/57	Yes
Zinc	2.78E+01	Yes	5.00E+01	31/57	Yes
Organic Compounds (mg/kg)					
TCE	9.69E-04	Yes	4.28E-01	4/56	Yes
Radionuclides (pCi/g)					
Americium-241	3.89E+07	No	8.00E-02	2/57	No
Cesium-137 ^C	NA ^D		4.56E-01	2/55	No
Plutonium-239	4.46E+00	No	5.62E-02	3/57	No
Technetium-99	2.63E+01	Yes	5.69E+01	8/57	Yes
Uranium-234	1.22E+01	No	3.02E+00	25/57	No
Uranium-238	8.62E+00	Yes	2.24E+01	29/57	Yes

^A Based on a residential child with dilution attenuation factor (DAF) = 1 (DOE 2001)

^B Compound is considered an essential nutrient and has been excluded in accordance with the PGDP Risk Methods Document (2001)

^C Not applicable as the radionuclide does not reach groundwater within 10,000 years precluding receptor uptake, per Section 5 of main text and Appendix E

^D SSL not listed in Table A.7 of PGDP Risk Methods Document (2001)

Table E3.3. Summary of Screening for Detected Analytes–SWMU 4 Groundwater

Analyte	SSL Protective of Groundwater ^A	Exceed SSL	Maximum Detection	Detection Frequency	Included in Groundwater Modeling
<i>Inorganic Chemicals (Metals) (mg/kg)</i>					
Aluminum	NA ^G		1.90E+04	139/139	No
Arsenic	1.05E-03	Yes	1.71E+01	16/138	Yes
Barium	NA ^G		3.13E+02	139/139	No
Beryllium	2.11E+00	No	2.02E+00	96/139	No
Calcium ^B	NA ^G		1.31E+05	138/138	No
Chromium	3.17E+06	No	2.96E+02	139/139	No
Cobalt	NA ^G		3.16E+01	137/139	No
Copper	NA ^G		4.64E+01	134/139	No
Iron	NA ^G		4.19E+04	139/139	No
Lead	NA ^G		3.02E+01	4/139	No
Magnesium ^B	NA ^G		2.65E+03	138/139	No
Manganese	2.26E+00	Yes	2.70E+03	138/139	Yes
Mercury	2.22E-02	Yes	4.50E-01	1/139	No ^H
Nickel	1.98E+00	Yes	1.53E+02	95/139	Yes
Potassium ^B	NA ^G		2.39E+03	139/139	No
Sodium ^B	NA ^G		3.15E+03	83/139	No
Uranium ^I	NA ^G		2.05E+04	8/36	Yes
Vanadium	9.25E+00	Yes	7.55E+01	139/139	Yes
Zinc	2.78E+01	Yes	9.37E+01	116/139	Yes

Table E3.3. Summary of Screening for Detected Analytes–SWMU 4 Groundwater (Continued)

Organic Compounds (mg/kg)					
Total PCB ^C	4.92E-02	Yes	2.83E+01	NA	No ^E
Total PAH ^D	NA ^G		4.10E-01	NA	No ^E
Aroclor-1254	7.67E-03	Yes	2.70E+01	7/184	No ^H
Aroclor-1260	2.49E-02	Yes	8.98E-01	7/184	No ^H
Bis(2-ethylhexyl)phthalate	NA ^G		7.47E-01	6/139	No
Diethyl phthalate	NA ^G		2.80E+00	2/139	No
Di-n-butyl phthalate	NA ^G		6.10E+00	22/139	No
1,1,2-TCA	NA ^G		2.10E-02	1/115	No
1,1-DCE	1.95E-05	Yes	1.40E-02	1/339	No ^H
Chloroform	8.32E-06	Yes	1.2E-02	1/113	No ^H
<i>cis</i> -1,2-DCE	7.80E-04	Yes	9.8E+00	23/338	Yes
<i>trans</i> -1,2-DCE	1.86E-03	Yes	4.5E-01	1/339	No ^H
TCE	9.69E-04	Yes	4.10E+01	47/335	Yes
Vinyl chloride	1.17E-05	Yes	2.9E-01	7/339	Yes
Radionuclides (pCi/g)					
Cesium-137 ^F	NA ^G		1.81E+02	8/179	No
Technetium-99	2.63E+01	Yes	2.69E+02	15/200	Yes
Neptunium-237	6.20E+03	No	5.78E+00	13/47	No
Plutonium-239	4.46E+00	Yes	2.71E+01	10/48	Yes
Uranium-234	1.22E+01	Yes	6.90E+01	21/29	Yes
Uranium-235	1.14E+01	No	4.20E+00	1/171	No
Uranium-238	8.62E+00	Yes	1.26E+02	21/29	Yes

^A Based on a residential child with dilution attenuation factor (DAF) = 1 (DOE 2001)

^B Compound is considered an essential nutrient and has been excluded in accordance with the PGDP Risk Methods Document (2001)

^C Total polychlorinated biphenyl (PCB) maximum concentration reported as sum of all detected aroclors

^D Total polyaromatic hydrocarbon (PAH) reported as benzo(a)pyrene toxicity equivalence factor (TEF)

^E Groundwater modeling was not performed for Total PCB and Total PAH as they represent groups of compounds. PCBs and PAHs were assessed individually for groundwater modeling

^F Not applicable as the radionuclide does not reach groundwater within 10,000 years precluding receptor uptake, per Section 5 of main text and Appendix E

^G SSL not listed in Table A.7 of PGDP Risk Methods Document (2001)

^H Less than 5% detects, so analyte not retained

^I Uranium analytical laboratory results were provided in units of pCi/g. The calculations to convert the uranium to units of mg/kg were performed using the specific activity of the nuclides in uranium and the error associated with the sample result. It was assumed, based on information found at www.wise-uranium.org/, that the U-238% of activity is about 85% and U-234 at 14%, with U-235 making up the remainder. U-238 dominates the unit conversion calculation due to its low specific activity which results in the largest estimate of uranium mass; therefore, the unit conversion was conducted using the specific activity of U-238 with the addition of the sample error. This example calculation is provided for clarification: A maximum detection of 6,260 pCi/g and an associated error of 628 pCi/g gives a total concentration of 6,888 pCi/g or 6.888E+06 pCi/kg. The specific activity of U-238 is 3.36E+05 pCi/g or 3.36E+02 pCi/mg; therefore, 6.888E+06 pCi/kg divided by 3.36E+02 pCi/mg equals 2.05E+04 mg/kg of uranium.

Table E3.4. Summary of Screening for Detected Analytes–SWMU 5 Groundwater

Analyte	SSL Protective of Groundwater ^A	Exceed SSL	Maximum Detection	Detection Frequency	Included in Groundwater Modeling
<i>Metals (mg/kg)</i>					
Aluminum	NA ^G		1.64E+04	84/84	No
Arsenic	1.05E-03	Yes	1.22E+01	24/85	Yes
Barium	NA ^G		3.43E+02	85/85	No
Beryllium	2.11E+00	Yes	2.59E+00	41/84	No
Calcium ^B	NA ^G		2.07E+05	84/84	No
Chromium	3.17E+06	No	2.96E+02	84/85	No
Cobalt	NA ^G		2.85E+01	69/84	No
Copper	NA ^G		1.44E+01	82/84	No
Iron	NA ^G		3.29E+04	84/84	No
Lead	NA ^G		3.19E+01	18/85	No
Magnesium ^B	NA ^G		4.78E+03	84/84	No
Manganese	2.26E+00	Yes	1.75E+03	84/84	Yes
Mercury	2.22E-02	Yes	3.60E-02	1/85	No ^H
Nickel	1.98E+00	Yes	1.35E+02	69/85	Yes
Potassium ^B	NA ^G		1.89E+03	55/55	No
Selenium	3.92E-02	Yes	1.23E+00	2/85	Yes
Silver	6.67E-02	Yes	5.14E+00	1/85	No ^H
Sodium ^B	NA ^G		3.89E+02	37/72	No
Uranium	NA ^G		2.79E+02	6/29	Yes
Vanadium	9.25E+00	Yes	5.69E+01	84/84	Yes
Zinc	2.78E+01	Yes	1.63E+02	54/84	Yes
<i>Organic Compounds (mg/kg)</i>					
Total PCB ^C	4.92E-02	Yes	3.06E-01	NA	No ^E
Total PAH ^D	NA ^G		1.15E+02	NA	No ^E
Acenaphthene	1.92E-01	Yes	3.20E+01	18/96	Yes
Acenaphthylene	NA ^G		9.45E+00	2/96	No
Anthracene	4.53E+00	Yes	4.00E+01	20/96	No ^I
Aroclor-1260	2.49E-02	Yes	3.06E-01	6/79	Yes
Benz(a)anthracene	NA ^G		1.30E+02	30/96	No
Benzo(a)pyrene	1.95E-03	Yes	8.00E+01	30/96	Yes
Benzo(b)fluoranthene	NA ^G		1.70E+02	29/96	No
Benzo(ghi)perylene	NA ^G		2.80E+01	26/97	No
Benzo(k)fluoranthene	NA ^G		1.17E+01	6/55	No
Bis(2-ethylhexyl)phthalate	NA ^G		5.70E+00	23/96	No
Carbazole	NA ^G		7.10E+01	10/96	No
Chrysene	NA ^G		9.50E+01	30/96	No

**Table E3.4. Summary of Screening for Detected Analytes–SWMU 5 Groundwater
(Continued)**

Analyte	SSL Protective of Groundwater ^A	Exceed SSL	Maximum Detection	Detection Frequency	Included in Groundwater Modeling
<i>Organic Compounds (mg/kg)</i>					
Dibenz(a,h)anthracene	NA ^G		7.49E-01	2/55	Yes
Dibenzofuran	NA ^G		3.52E+00	3/55	No
Di-n-butyl phthalate	NA ^G		7.30E+00	27/55	No
Indeno(1,2,3-cd)pyrene	NA ^G		3.70E+01	24/96	No
Fluoranthene	4.67E+00	Yes	5.3E+01	3/34	Yes
Fluorene	2.66E-01	Yes	2.80E+01	16/96	Yes
2-Methylnaphthalene	NA ^G		7.30E+00	2/96	No
Naphthalene	1.21E-03	Yes	1.60E+01	3/96	Yes
Phenanthrene	NA ^G		6.40E+01	31/96	No
Pyrene	3.81E+00	Yes	1.50E+02	35/96	Yes
1,1-DCE	1.95E-05	Yes	2.80E+00	1/102	No ^H
TCE	9.69E-04	Yes	5.10E-03	12/102	Yes
<i>Radionuclides (pCi/g)</i>					
Cesium-137 ^F	NA ^G		7.41E-02	23/61	Yes
Technetium-99	2.63E+01	No	1.73E+01	52/90	Yes
Uranium-234	1.22E+01	No	1.47E+00	11/40	No
Uranium-238	8.62E+00	No	2.00E+00	21/40	No

^A Based on a residential child with dilution attenuation factor (DAF) = 1 (DOE 2001)

^B Compound is considered an essential nutrient and has been excluded in accordance with the PGDP Risk Methods Document (2001)

^C Total polychlorinated biphenyl (PCB) maximum concentration reported as sum of all detected aroclors

^D Total polyaromatic hydrocarbon (PAH) reported as benzo(a)pyrene toxicity equivalence factor (TEF)

^E Groundwater modeling was not performed for Total PCB and Total PAH as they represent groups of compounds. PCBs and PAHs were assessed individually for groundwater modeling.

^F Not applicable as the radionuclide does not reach groundwater within 10,000 years precluding receptor uptake, per Section 5 of main text and Appendix E

^G SSL not listed in Table A.7 of PGDP Risk Methods Document (2001)

^H Less than 5% detects, so analyte not retained.

^I Maximum detected value less than the soil/sediment no action level (NAL) for a child resident in Table A.17 of PGDP Risk Methods Document (2001), so analyte not retained

Table E3.5. Summary of Screening for Detected Analytes–SWMU 6 Groundwater

Analyte	SSL Protective of Groundwater ^A	Exceed SSL	Maximum Detection	Detection Frequency	Included in Groundwater Modeling
Metals (mg/kg)					
Aluminum	NA ^F		2.25E+04	85/85	No
Arsenic	1.05E-03	Yes	6.38E+00	25/85	Yes
Barium	NA ^F		1.53E+02	85/85	No
Beryllium	2.11E+00	Yes	3.07E+00	41/85	Yes
Chromium	3.17E+06	No	1.16E+02	85/85	No
Cobalt	NA ^F		1.56E+02	76/85	No
Copper	NA ^F		2.13E+01	82/85	No
Iron	NA ^F		5.87E+04	85/85	No
Lead	NA ^F		3.54E+01	24/85	No
Magnesium ^B	NA ^F		4.41E+03	85/85	No
Manganese ^B	2.26E+00	Yes	1.55E+03	85/85	Yes
Nickel	1.98E+00	Yes	6.86E+01	52/85	Yes
Uranium	NA ^F		1.14E+02	16/33	Yes
Vanadium	9.25E+00	Yes	7.91E+01	84/85	Yes
Zinc	2.78E+01	Yes	1.28E+02	57/85	Yes
Organic Compounds (mg/kg)					
Total PCB ^C	4.92E-02	Yes	6.00E-02	NA	No ^E
Total PAH ^D	NA ^F		4.99E-01	NA	No ^E
Anthracene	4.53E+00	No	1.56E-01	1/62	No
Benz(a)anthracene	NA ^F		2.55E-01	2/62	No
Benzo(a)pyrene	1.95E-03	Yes	4.02E-01	1/62	No ^G
Benzo(b)fluoranthene	NA ^F		5.00E-01	2/62	No
Benzo(ghi)perylene	NA ^F		1.24E-01	2/62	No
Bis(2-ethylhexyl)phthalate	NA ^F		6.00E-01	3/62	No
Fluoranthene	4.67E+00	No	6.36E-01	2/55	No
Pyrene	3.81E+00	No	6.63E-01	2/62	No
Acetone	NA ^F		8.50E-02	6/48	No
TCE	9.69E-04	Yes	1.01E-02	5/115	Yes
Radionuclides (pCi/g)					
Technetium-99	2.63E+01	No	1.88E+01	3/90	Yes

^A Based on a residential child with dilution attenuation factor (DAF) = 1 (DOE 2001)

^B Compound is considered an essential nutrient and has been excluded in accordance with the PGDP Risk Methods Document (2001)

^C Total polychlorinated biphenyl (PCB) maximum concentration reported as method detection limit as there were no detections for Total PCB

^D Total polyaromatic hydrocarbon (PAH) reported as benzo(a)pyrene toxicity equivalence factor (TEF)

^E Groundwater modeling was not performed for Total PCB and Total PAH as they represent groups of compounds. PCBs and PAHs were assessed individually for groundwater modeling.

^F SSL not listed in Table A.7 of PGDP Risk Methods Document (2001)

^G Less than 5% detects, so analyte not retained.

^H Maximum detected value less than the soil/sediment no action level (NAL) for a child resident in Table A.17 of PGDP Risk Methods Document (2001), so analyte not retained

Table E3.6. Summary of Screening for Detected Analytes–SWMU 7 Groundwater

Analyte	SSL Protective of Groundwater ^A	Exceed SSL	Maximum Detection	Detection Frequency	Included in Groundwater Modeling
Metals (mg/kg)					
Aluminum	NA ^G		1.60E+04	83/83	No
Arsenic	1.05E-03	Yes	1.60E+01	72/83	Yes
Barium	NA ^G		1.80E+02	83/83	No
Beryllium	2.11E+00	No	1.55E+00	29/85	No
Cadmium	5.02E-02	Yes	1.30E+00	12/83	Yes
Calcium ^B	NA ^G		2.10E+05	83/83	No
Chromium	3.17E+06	No	5.58E+01	83/85	No
Cobalt	NA ^G		1.77E+01	56/83	No
Copper	NA ^G		9.90E+01	70/83	No
Iron	NA ^G		3.47E+04	83/83	No
Lead	NA ^G		1.20E+02	83/83	No
Magnesium ^B	NA ^G		3.30E+03	83/83	No
Manganese	2.26E+00	Yes	1.20E+03	83/83	Yes
Mercury	2.22E-02	Yes	9.20E-02	22/85	Yes
Nickel	1.98E+00	Yes	1.40E+02	58/83	Yes
Potassium ^B	NA ^G		8.70E+02	20/20	No
Selenium	3.92E-02	Yes	8.80E-01	8/83	Yes
Sodium ^B	NA ^G		4.00E+02	47/83	No
Silver	6.67E-02	Yes	1.90E+00	7/83	No ^I
Thallium	NA ^G		2.00E+00	17/83	No
Tin	NA ^G		8.7E+00	20/20	No
Uranium	NA ^G		1.27E+03	31/85	Yes
Vanadium	9.25E+00	Yes	5.20E+01	76/83	Yes
Zinc	2.78E+01	Yes	2.40E+02	47/83	Yes
Organic Compounds (mg/kg)					
Total PCB ^C	4.92E-02	Yes	1.81E+01	NA	No ^E
Total PAH ^D	NA ^G		6.37E+00	NA	No ^E
Aroclor-1254	7.67E-03	Yes	1.30E-01	4/86	Yes
Aroclor-1260	2.49E-02	Yes	3.1E+00	13/86	Yes
1,2,4-Trichlorobenzene	NA ^G		7.70E-02	2/80	No
1,4-Dichlorobenzene	NA ^G		7.00E-02	2/88	No
2,4,6-Trichlorophenol	NA ^G		4.10E-02	2/160	No
2-Methylphenol	NA ^G		2.00E-02	1/17	No
3-Methylcholanthrene	NA ^G		1.10E-01	1/17	No
4-Methylphenol	NA ^G		1.60E-02	1/17	No
Acenaphthene	1.92E-01	Yes	4.40E-01	2/80	No ^I
Acenaphthylene	NA ^G		1.90E-02	1/80	No
Benz(a)anthracene	NA ^G		4.30E+00	7/80	No
Benzo(a)pyrene	1.95E-03	Yes	4.10E+00	7/80	Yes
Benzo(b)fluoranthene	NA ^G		5.20E+00	7/80	No
Benzo(ghi)perylene	NA ^G		3.80E+00	3/80	No
Benzo(k)fluoranthene	NA ^G		1.70E+00	6/80	No

Table E3.6. Summary of Screening for Detected Analytes–SWMU 7 Groundwater (Continued)

Analyte	SSL Protective of Groundwater ^A	Exceed SSL	Maximum Detection	Detection Frequency	Included in Groundwater Modeling
Organic Compounds (mg/kg)(Continued)					
Bis(2-ethylhexyl)phthalate	NA ^G		3.90E-01	2/80	No
Chrysene	NA ^G		4.20E+00	7/80	No
Dibenz(a,h)anthracene	NA ^G		9.20E-01	1/80	No
Di-n-octylphthalate	NA ^G		7.20E-02	1/80	No
Fluoranthene	4.67E+00	Yes	7.90E+00	8/80	Yes
Fluorene	2.66E-01	Yes	4.10E-01	1/80	No ^I
Hexachloroethane	NA ^G		3.40E-02	1/80	No
Indeno(1,2,3-cd)pyrene	NA ^G		3.80E+00	3/80	No
Naphthalene	1.21E-03	Yes	5.60E-02	1/80	No ^I
Pentachlorophenol	NA ^G		6.90E-02	1/80	No
Pyrene	3.81E+00	Yes	9.00E+00	9/80	Yes
1,1,1-TCA	NA ^G		1.59E-01	2/71	No
1,1,2-TCA	NA ^G		1.49E-01	2/71	No
1,1-DCE	1.95E-05	Yes	1.66E+00	4/71	Yes
1,1-Dichloroethane	NA ^G		3.78E-01	3/72	No
1,2-Dichloroethane	NA ^G		1.63E-02	2/71	No
Acetone	NA ^G		8.37E-02	10/71	No
cis-1,2-DCE	7.80E-04	Yes	6.84E-01	15/71	Yes
Methylene chloride	NA ^G		5.70E+00	5/71	No
Tetrachloroethene	3.38E-04	Yes	6.20E-03	1/71	Yes
TCE	9.69E-04	Yes	2.60E-01	11/71	Yes
Vinyl chloride	1.17E-05	Yes	5.85E-01	5/71	Yes
Radionuclides (pCi/g)					
Americium-241	3.89E+07	No	2.00E+00	3/64	No
Cesium-137 ^F	NA ^G	No	1.83E-01	3/64	No
Neptunium-237	6.20E+03	No	7.20E-01	26/86	Yes
Plutonium-239	4.46E+00	No	6.80E-01	19/86	No
Technetium-99	2.63E+01	Yes	4.06E+02	37/84	Yes
Uranium-234	1.22E+01	Yes	3.18E+02	59/94	Yes
Uranium-235	1.14E+01	No	4.21E+01	22/22	Yes
Uranium-238	8.62E+00	Yes	2.39E+03	53/94	Yes

^A Based on a residential child with dilution attenuation factor (DAF) = 1 (DOE 2001)

^B Compound is considered an essential nutrient and has been excluded in accordance with the PGDP Risk Methods Document (2001)

^C Total polychlorinated biphenyl (PCB) maximum concentration reported as sum of all detected aroclors

^D Total polyaromatic hydrocarbon (PAH) reported as benzo(a)pyrene toxicity equivalence factor (TEF)

^E Groundwater modeling was not performed for Total PCB and Total PAH as they represent groups of compounds. PCBs and PAHs were assessed individually for groundwater modeling.

^F Not applicable as the radionuclide does not reach groundwater within 10,000 years precluding receptor uptake, per Section 5 of main text and Appendix E

^G SSL not listed in Table A.7 of PGDP Risk Methods Document (2001)

^H Less than 5% detects, so analyte not retained

^I Maximum detected value less than the soil/sediment no action level (NAL) for a child resident in Table A.17 of PGDP Risk Methods Document (2001), so analyte not retained

Table E3.7. Summary of Screening for Detected Analytes–SWMU 30 Groundwater

Analyte	SSL Protective of Groundwater ^A	Exceed SSL	Maximum Detection	Detection Frequency	Included in Groundwater Modeling
<i>Inorganic Chemicals (Metals) (mg/kg)</i>					
Aluminum	NA ^F		1.90E+04	33/33	No
Arsenic	1.05E-03	Yes	8.9+00	30/33	Yes
Barium	NA ^F		1.70E+02	35/35	No
Beryllium	2.11E+00	No	1.48E+00	15/33	No
Cadmium	5.02E-02		2.80E+00	6/35	Yes
Calcium ^B	NA ^F		2.40E+04	33/33	No
Chromium	3.17E+06	No	4.90E+01	35/35	No
Cobalt	NA ^F		1.40E+01	25/33	No
Copper	NA ^F		1.70E+02	32/33	No
Iron	NA ^F		2.90E+04	33/33	No
Lead	NA ^F		7.10E+01	33/35	No
Magnesium ^B	NA ^F		2.20E+03	33/33	No
Manganese	2.26E+00	Yes	1.20E+03	33/33	Yes
Mercury	2.22E-02	Yes	1.70E-01	12/35	Yes
Nickel	1.98E+00	Yes	5.70E+02	29/33	Yes
Potassium ^B	NA ^F		1.50E+03	12/12	No
Selenium	3.92E-02	Yes	1.00E+00	28/35	Yes
Sodium ^B	NA ^F		1.87E+02	9/33	No
Thallium	NA ^F		1.80E+00	5/35	No
Uranium	NA ^F		1.40E+03	16/33	Yes
Vanadium	9.25E+00	Yes	4.00E+01	32/33	Yes
Zinc	2.78E+01	Yes	7.50E+02	11/33	Yes
<i>Organic Compounds (mg/kg)</i>					
Total PCB ^C	4.92E-02	Yes	7.60E-01	NA	No ^E
Total PAH ^D	NA ^F		1.25E+01	NA	No ^E
Aroclor-1254	7.67E-03	Yes	2.00E-01	3/36	Yes
Aroclor-1260	1.92E-01	Yes	1.50E+01	5/34	Yes
Acenaphthene	1.92E-01	Yes	1.70E+00	5/34	No ^G
Acenaphthylene	NA ^F		9.10E-02	3/34	No
Anthracene	4.53E+00	No	3.20E+00	3/34	No
Benz(a)anthracene	NA ^F		9.10E+00	10/34	No
Benzo(a)pyrene	1.95E-03	Yes	8.40E+00	7/34	Yes
Benzo(b)fluoranthene	NA ^F		9.60E+00	8/34	No
Benzo(ghi)perylene	NA ^F		5.20E+00	6/34	No
Benzo(k)fluoranthene	NA ^F		4.30E+00	7/34	No
Benzoic acid	NA ^F		5.30E-01	7/34	No
Bis(2-ethylhexyl)phthalate	NA ^F		6.20E-01	2/34	No
2-Chlorophenol	NA ^F		2.30E-02	1/34	No
Chrysene	NA ^F		9.90E+00	9/34	No
Dibenz(a,h)anthracene	NA ^F		1.60E+00	30/34	Yes
Dibenzofuran	NA ^F		8.30E-01	3/34	No
Di-n-butyl phthalate	NA ^F		1.00E-01	2/34	No

Table E3.7. Summary of Screening for Detected Analytes–SWMU 30 Groundwater (Continued)

Analyte	SSL Protective of Groundwater ^A	Exceed SSL	Maximum Detection	Detection Frequency	Included in Groundwater Modeling
Fluoranthene	NA ^F		2.00E+01	3/34	No
Fluorene	NA ^F		1.30E+00	3/34	Yes
Indeno(1,2,3-cd)pyrene	NA ^F		5.40E+00	6/34	No
2-Methylnaphthalene	NA ^F		2.70E-01	2/34	No
Naphthalene	NA ^F		3.10E-01	1/34	Yes
Phenanthrene	NA ^F		1.70E+01	9/34	No
Pyrene	3.81E+00	Yes	2.30E+01	11/34	Yes
1,2,4-Trichlorobenzene	NA ^F		3.30E-02	1/54	No
1,1-DCE	1.95E-05	Yes	5.00E-03	1/28	Yes
1,4-Dichlorobenzene	NA ^F		2.50E-02	1/39	No
Acetone	NA ^F		9.73E-03	3/26	No
TCE	9.69E-04	Yes	3.74E-02	1/28	Yes
Radionuclides (pCi/g)					
Neptunium-237	6.20E+03	No	1.68E+00	9/33	No
Plutonium-239	4.46E+00	No	6.20E-01	11/33	No
Technetium-99	2.63E+01	Yes	3.60E+02	13/34	Yes
Uranium-234	1.22E+01	Yes	1.15E+02	27/34	Yes
Uranium-235	1.14E+01	Yes	1.66E+01	14/34	Yes
Uranium-238	8.62E+00	Yes	5.65E+02	24/36	Yes

^A Based on a residential child with dilution attenuation factor (DAF) = 1 (DOE 2001)

^B Compound is considered an essential nutrient and has been excluded in accordance with the PGDP Risk Methods Document (2001)

^C Total polychlorinated biphenyl (PCB) maximum concentration reported as sum of all detected aroclors

^D Total polyaromatic hydrocarbon (PAH) reported as benzo(a)pyrene toxicity equivalence factor (TEF)

^E Groundwater modeling was not performed for Total PCB and Total PAH as they represent groups of compounds. PCBs and PAHs were assessed individually for groundwater modeling.

^F SSL not listed in Table A.7 of PGDP Risk Methods Document (2001)

^G Maximum detected value less than the soil/sediment no action level (NAL) for a child resident in Table A.17 of PGDP Risk Methods Document (2001), so analyte not retained

Table E3.8. Summary of Screening for Detected Analytes–SWMU 145 Groundwater

Analyte	SSL Protective of Groundwater ^A	Exceed SSL	Maximum Detection	Detection Frequency	Included in Groundwater Modeling
<i>Inorganic Chemicals (Metals)(mg/kg)</i>					
Aluminum	NA ^F		1.62E+04	74/74	No
Antimony	2.54E-02	Yes	2.02E+01	18/74	Yes
Arsenic	1.05E-03	Yes	2.19E+01	52/79	Yes
Barium	NA ^F		3.00E+02	88/88	No
Beryllium	2.11E+00	No	2.08E+00	50/89	No
Cadmium	5.02E-02	Yes	2.47E+00	13/89	Yes
Calcium ^B	NA ^F		8.30E+04	74/74	No
Chromium	3.17E+06	No	1.20E+02	89/89	No
Cobalt	NA ^F		2.05E+01	70/89	No
Copper	NA ^F		1.35E+02	72/74	No
Iron	NA ^F		3.14E+04	74/74	No
Lead	NA ^F		4.67E+01	68/88	No
Magnesium ^B	NA ^F		2.35E+03	74/74	No
Manganese	2.26E+00	Yes	1.90E+03	74/74	Yes
Mercury	2.22E-02	Yes	4.70E-01	31/89	Yes
Molybdenum	NA ^F		3.39E+00	13/32	No
Nickel	1.98E+00	Yes	1.01E+02	65/74	Yes
Potassium ^B	NA ^F		1.30E+03	32/34	No
Selenium	3.92E-02	Yes	1.10E+00	2/86	No ^H
Sodium ^B	NA ^F		6.93E+02	46/70	No
Silver	6.67E-02	Yes	1.96E+01	3/88	No ^H
Thallium	NA ^F		1.60E+00	14/74	No
Uranium	NA ^F		3.11E+02	26/55	No
Vanadium	9.25E+00	Yes	6.52E+01	54/74	Yes
Zinc	2.78E+01	Yes	2.61E+02	52/79	No ^I
<i>Organic Compounds (mg/kg)</i>					
Total PCB ^C	4.92E-02	Yes	1.44E+01	NA	No ^E
Total PAH ^D	NA ^F		5.31E-02	NA	No ^E
Aroclor-1254	7.67E-03	Yes	1.9E+00	4/86	Yes
Aroclor-1260	2.49E-02	Yes	1.25E+01	3/90	Yes
Benz(a)anthracene	NA ^F		5.00E-02	1/14	No
Benzo(a)pyrene	1.95E-03	Yes	4.40E-02	1/13	Yes
Benzo(b)fluoranthene	NA ^F		4.00E-02	1/13	No
Chrysene	NA ^F		6.70E-02	1/13	No
Di-n-butyl phthalate	NA ^F		8.10E-01	1/9	No
Fluoranthene	4.67E+00	No	1.30E-01	1/9	No
Phenanthrene	NA ^F		1.10E-01	1/13	No
Pyrene	3.81E+00	No	8.40E-02	1/13	No
1,2-Dimethylbenzene	NA ^F		3.60E-02	2/62	No
2-Butanone	NA ^F		2.83E-02	3/64	No
4-Methyl-2-pentanone	NA ^F		1.50E-02	1/64	No
Acetone	NA ^F		1.16E-01	20/64	No
Ethylbenzene	4.48E-03	Yes	1.80E-02	3/64	No
m,p-Xylene	6.35E-02	No	0.04	2/62	No
Toluene	NA ^F		0.037	5/63	No
Methylene chloride	NA ^F		3.00E-03	2/64	No

Table E3.8. Summary of Screening for Detected Analytes–SWMU 145 Groundwater

Analyte	SSL Protective of Groundwater ^A	Exceed SSL	Maximum Detection	Detection Frequency	Included in Groundwater Modeling
Radionuclides (pCi/g)					
Americium-241	3.89E+07	No	1.10E+01	13/79	No
Cesium-137 ^G	NA ^F	No	1.05E-00	14/67	No
Neptunium-237	6.20E+03	No	1.22E+00	18/86	No
Plutonium-239	4.46E+00	Yes	1.01E+01	21/75	Yes
Technetium-99	2.63E+01	Yes	2.81E+02	26/79	Yes
Uranium-234	1.22E+01	Yes	2.54E+02	58/86	Yes
Uranium-235	1.14E+01	Yes	2.2E+00	25/71	Yes
Uranium-238	8.62E+00	Yes	3.26E+02	59/86	Yes

^A Based on a residential child with dilution attenuation factor (DAF) = 1 (DOE 2001)

^B Compound is considered an essential nutrient and has been excluded in accordance with the PGDP Risk Methods Document (2001)

^C Total polychlorinated biphenyl (PCB) maximum concentration reported as sum of all detected aroclors

^D Total polyaromatic hydrocarbon (PAH) reported as benzo(a)pyrene toxicity equivalence factor (TEF)

^E Groundwater modeling was not performed for Total PCB and Total PAH as they represent groups of compounds. PCBs and PAHs were assessed individually for groundwater modeling.

^F SSL not listed in Table A.7 of PGDP Risk Methods Document (2001)

^G Not applicable as the radionuclide does not reach groundwater within 10,000 years precluding receptor uptake, per Section 5 of the main text and Appendix E^H Less than 5% detects, so analyte not retained.

^I Maximum detected value less than the soil/sediment no action level (NAL) for a child resident in Table A.17 of PGDP Risk Methods Document (2001), so analyte not retained

APPENDIX F
BASELINE HUMAN HEALTH RISK ASSESSMENT

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ACRONYMS

ABS	dermal absorption factor
ATSDR	Agency for Toxic Substances and Disease Registry
BAF	bioaccumulation factor
BGOU	Burial Grounds Operable Unit
bgs	below ground surface
BHHRA	Baseline Human Health Risk Assessment
CAS	Chemical Abstract Service
CDI	chronic daily intake
CNS	central nervous system
COC	contaminant of concern
COPC	chemical of potential concern
CSM	conceptual site model
DAF	dilution attenuation factor
DCE	dichloroethene
DHHS	Department of Health and Human Services
DNA	deoxyribonucleic acid
DOE	U.S. Department of Energy
DQA	Data Quality Assessment
ELCR	excess lifetime cancer risk
EPA	U. S. Environmental Protection Agency
EPC	exposure point concentration
GI	gastrointestinal tract
HEAST	Health Effects Assessment Summary Tables
HI	hazard index
HQ	hazard quotient
IARC	International Agency for Research on Cancer
IRIS	Integrated Risk Information System
KDEP	Kentucky Department for Environmental Protection
LET	linear energy transfer
MCL	maximum contaminant level
MEPAS	Multimedia Environmental Pollution Assessment System
NAL	no action level
NCEA	National Center for Environmental Assessment
NSDD	North-South Diversion Ditch
OREIS	Oak Ridge Environmental Information System
PAH	polyaromatic hydrocarbon
PCB	polychlorinated biphenyl
PGDP	Paducah Gaseous Diffusion Plant
POC	pathway of concern
POE	point of exposure
PRG	preliminary remediation goal
PVC	polyvinyl chloride
RAGS	Risk Assessment Guidance for Superfund
RAIS	Risk Assessment Information System
RCRA	Resource Conservation and Recovery Act
RDA	recommended dietary allowances
RESRAD	RESidual RADioactive Materials
RfC	reference concentration

RfD	reference dose
RGA	Regional Gravel Aquifer
RGO	remedial goal option
RI	remedial investigation
RME	reasonable maximum exposure
SF	slope factor
SI	site investigation
SQL	sample quantitation limit
SSL	soil screening level
SVOC	semivolatile organic compound
SWMU	solid waste management unit
SWOU	Surface Water Operable Unit
⁹⁹ Tc	technetium-99
TCE	trichloroethene
TEF	toxicity equivalence factor
TVA	Tennessee Valley Authority
VOC	volatile organic compound
WAG	waste area grouping
WKWMA	West Kentucky Wildlife Management Area
WMU	waste management unit

BASELINE HUMAN HEALTH RISK ASSESSMENT

This Baseline Risk Human Health Assessment (BHHRA) utilizes information collected during the recently completed remedial investigation (RI) of eight Burial Grounds Operable Unit (BGOU) Solid Waste Management Units (SWMUs), in addition to information collected during previous investigations (listed in Section F.1), to characterize the baseline risks posed to human health from contact with contaminants in soil and water at these SWMUs and at locations to which contaminants may migrate. The units included are SWMU 2, SWMU 3, SWMU 4, SWMU 5, SWMU 6, SWMU 7, SWMU 30, and SWMU 145 located at the Paducah Gaseous Diffusion Plant (PGDP) in Paducah, Kentucky. A summary of the data used is presented in Attachment F1 to this appendix.

Part of Goal 2 for the BGOU RI, as presented in the BGOU work plan (DOE 2006), was to determine if contaminants at the BGOU units are contributing to groundwater contamination; this risk assessment supports that goal by using modeled concentrations of contaminants to the Regional Gravel Aquifer (RGA) to support the refinement of an assessment of risks to human health and the environment through groundwater. The work plan also specified that the RI should include a risk assessment for residential, industrial, and recreational receptors. Risk assessments for each of those scenarios are presented here.

The methods and presentations used in this BHHRA are consistent with those presented in *Methods for Conducting Risk Assessments and Risk Evaluations at the Paducah Gaseous Diffusion Plant*, DOE/OR/07-1506&D2 (DOE 2001). The Risk Methods Document integrates the human health risk assessment guidance from the U.S. Environmental Protection Agency (EPA) and the Kentucky Department for Environmental Protection (KDEP) and incorporates instructions contained in regulatory agency comments on earlier risk assessments performed for PGDP.

Consistent with the 2001 revision to the Risk Methods Document, this BHHRA is presented in nine sections. The first section reviews the results of previous risk assessments that are useful in understanding the risks posed to human health by contaminants at or migrating from the source areas. Identification of chemicals of potential concern (COPCs) is in the second section. The third section documents the exposure assessment for the sources, including the characterization of the exposure setting, identification of exposure pathways, consideration of land use, determination of potential receptors, delineation of exposure points and routes [including development of the conceptual site model (CSM)], and calculation of chronic daily intakes (CDIs). The fourth section presents the toxicity assessment, including information on the noncarcinogenic (i.e., systemic toxicity or hazard) and carcinogenic effects of the COPCs and the uncertainties in the toxicity information. The fifth section reports the results of the risk characterization for current and future land use and identifies contaminants, pathways, and land use scenarios of concern. The sixth section contains qualitative and quantitative analyses of the uncertainties affecting the results of the BHHRA. The seventh section summarizes the methods used in the BHHRA and presents the BHHRA's conclusions and observations. The eighth section uses the results of the BHHRA to develop site-specific risk-based remedial goal options (RGOs). The ninth section contains references.

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F.1. RESULTS OF PREVIOUS STUDIES

Four previous reports contain risk assessment results for one or more of the burial grounds considered in this RI. The results of these assessments are summarized here and are presented in more detail in Attachment F2 to this appendix. For groundwater, these previous assessments were based on measured groundwater concentrations, while this risk assessment used modeled concentrations. In the previous assessments summarized here, lead had exceedingly high hazard indices (HIs) and was the overwhelming risk driver. This finding may be attributed to the use of a very conservative (1.0E-07 mg/kg-day) reference dose (RfD) value provided by KDEP. That RfD is no longer in use by KDEP; therefore, the contaminants of concern (COCs), risks, and hazards from those reports presented in this appendix are based on the assessment with lead excluded as a COPC, which also was presented in these previous reports. This issue is discussed further in the uncertainty section of this appendix. Reports containing previous assessments and the year the assessment was completed are listed below:

- *Remedial Investigation Report for Waste Area Grouping 3 at the Paducah Gaseous Diffusion Plant, Paducah, Kentucky*, DOE/OR/07-1895/V1-V4&D1, U.S. Department of Energy, Paducah, KY (DOE 2000).
- *Remedial Investigation Report for Solid Waste Management Units 7 and 30 of Waste Area Group 22 at the Paducah Gaseous Diffusion Plant, Paducah, Kentucky*, DOE/OR/07-1604/V1&D2, U.S. Department of Energy, Paducah, KY (DOE 1998a).
- *Remedial Investigation Addendum for Waste Area Grouping 22, Burial Grounds Solid Waste Management Units 2 and 3 at the Paducah Gaseous Diffusion Plant, Paducah, Kentucky*, DOE/OR/07-1141&D2, U.S. Department of Energy, Paducah, KY (DOE 1994).
- *Data Summary and Interpretation Report for Interim Remedial Design at Solid Waste Management Unit 2 of Waste Area Group 22 at the Paducah Gaseous Diffusion Plant, Paducah, Kentucky*, DOE/OR/07-1549&D1, U.S. Department of Energy, Paducah, KY (DOE 1997).

The soil depths used to calculate exposure point concentrations (EPCs) for site receptors is consistent with the Risk Methods Document. Exposure to soil, 0-1 ft bgs was used for current industrial workers, future industrial workers, and future on-site rural residents; and 0-10 ft bgs was used for future excavation workers.

F.1.1 RESULTS OF REMEDIAL INVESTIGATION REPORT FOR WASTE AREA GROUPING 3 AT THE PADUCAH GASEOUS DIFFUSION PLANT, PADUCAH, KENTUCKY

The Waste Area Grouping (WAG) 3 RI report BHHRA contains results for SWMU 4, SWMU 5, and SWMU 6 (DOE 2000). Scenarios assessed included the following:

- **Current Industrial Worker**
 - Incidental ingestion of soil
 - Dermal contact with soil
 - Inhalation of vapors and particulates emitted from soil
 - External exposure to ionizing radiation emitted from soil

- **Future Industrial Worker**
 - Incidental ingestion of soil
 - Dermal contact with soil
 - Inhalation of vapors and particulates emitted from soil
 - External exposure to ionizing radiation emitted from soil
 - Ingestion of groundwater
 - Dermal contact with groundwater while showering
 - Inhalation of vapors emitted by groundwater while showering

- **Future Excavation Worker**
 - Incidental ingestion of soil (soil and waste)
 - Dermal contact with soil (soil and waste)
 - Inhalation of vapors and particulates emitted from soil (soil and waste)
 - External exposure to ionizing radiation emitted from soil (soil and waste)

- **Future Recreational User**
 - Ingestion of venison grazing on vegetation grown in contaminated soil
 - Ingestion of rabbit grazing on vegetation grown in contaminated soil
 - Ingestion of quail grazing on vegetation grown in contaminated soil

- **Future On-site Rural Resident**
 - Incidental ingestion of soil
 - Dermal contact with soil
 - Inhalation of vapors and particulates emitted from soil
 - External exposure to ionizing radiation emitted from soil
 - Ingestion of groundwater
 - Dermal contact with groundwater while showering
 - Inhalation of vapors emitted by groundwater during household use
 - Inhalation of vapors emitted by groundwater while showering
 - Ingestion of vegetables grown in contaminated soil

- **Off-site Rural Resident (at PGDP security fence)**
 - Ingestion of groundwater
 - Dermal contact with groundwater while showering
 - Inhalation of vapors emitted by groundwater during household use
 - Inhalation of vapors emitted by groundwater while showering

For all SWMUs in WAG 3, the cumulative human health systemic toxicity and excess lifetime cancer risk (ELCR) exceeded the accepted standards of KDEP and EPA for one or more land use scenarios when assessed using default exposure parameters. The land use scenarios for which risks exceeded *de minimis* levels [i.e., for KDEP, a cumulative HI of 1 or a cumulative ELCR of 1.0E-06, and for EPA, an HI of 1 and a range of E-04–E-06 for ELCR] are summarized in the WAG 3 report and Tables F2.13 through F2.15 in Attachment F2 of this appendix.

F.1.1.1 Exposure Routes

Dermal contact with soil has been a driving exposure route in previous BHHRA at PGDP, with most of this risk arising from contact with metals. This is a direct result of using dermal absorption factors that exceed gastrointestinal absorption values and may be overly conservative. In such circumstances, risk estimates from the dermal exposure route may be unrealistic and exceed the real risk posed by this route

of exposure. Although chemical-specific dermal absorption factor (ABS) values were used when available, default ABS values were used for most chemicals because chemical-specific values are lacking. Chemical-specific ABS values are available for polychlorinated biphenyls (PCBs) and cadmium and were used in this BHHRA. Remedial decisions based on the dermal contact with soil exposure route should be carefully considered because of the uncertainty associated with risk from this exposure route. While the dermal exposure route may represent an important route of contaminant uptake for persons exposed to soil at WAG 3, ingestion of groundwater appears to represent the most important mechanism of uptake of contaminants from the RGA aquifer and McNairy Formation, with ingestion of groundwater-irrigated vegetables also representing a significant exposure route for the hypothetical on-site resident.

F.1.1.2 Current and Future Industrial Worker

Soil hazards (total HIs) for the current industrial worker exceed *de minimis* levels (HI >1 or ELCR >1.0E-06) at only SWMU 4+ (HI = 3.62) (Exhibit 1.23, p. 1-171, DOE 2000). The contaminants at SWMU 4 contributing more than 10% to total HI are chromium, iron, and vanadium, with dermal contact as the driving exposure route (Exhibit 1.55, p. 1-208, DOE 2000). Soil cancer risks (total ELCRs) for the current industrial worker exceed 1.0E-04 at SWMUs 4, 5, and 6 (Exhibit 1.25, p. 1-172, DOE 2000). The major contaminant in surface soils at these SWMUs is beryllium, with significant contributions from polyaromatic hydrocarbons (PAHs) at SWMUs 5 and 6 (Exhibit 1.26, p. 1-172, DOE 2000). For SWMUs 4, 5, and 6, dermal contact is the driving exposure route (Exhibits 1.55, 1.56, 1.57, pp.1-208 to 1-215, DOE 2000). The future industrial land use scenario is identical to the current industrial land use scenario except that the future industrial land use scenario also evaluates use of groundwater. Groundwater HIs for the future industrial worker exceed *de minimis* levels at SWMUs 4, 5, and 6 (16,000–216,000); however, these hazards are markedly reduced by excluding lead as a COPC (19.1–75.9) (Exhibits 1.27, 1.28, pp. 1-174 to 1-175, DOE 2000). Including lead as a COPC masks the contribution from other COPCs. Iron, manganese, vanadium, and trichloroethene (TCE) contribute more than 10% to total HIs, with ingestion as the driving exposure route (Exhibits 1.55, 1.56, 1.57, pp.1-208 to 1-215, DOE 2000). Iron is both widespread and predominant as a COC, contributing 61–80% to HI, depending on location (Exhibit 1.28, p. 1-175, DOE 2000). Groundwater ELCRs for the future industrial worker exceed 1.0E-04 at SWMUs 4, 5, and 6 (>1.0E-04) (Exhibit 1.35, p. 1-180, DOE 2000). Arsenic, beryllium, TCE, and radium-226 (²²⁶Ra) contribute more than 10% to ELCR, with ingestion as the driving exposure route (Exhibits 1.55, 1.56, 1.57, pp.1-208 to 1-215, DOE 2000).

F.1.1.3 Future Excavation Worker

Total soil and waste HIs for the future excavation worker exceed *de minimis* levels at SWMUs 4, 5, and 6 (2.16–1750), but fall below 3 when lead is excluded as a COPC (Exhibits 1.33, 1.34, p. 1-179, DOE 2000). Including lead as a COPC masks the contribution from other COPCs. Chromium, iron, manganese, and vanadium are the contaminants contributing more than 10% to HI, with dermal contact as the driving exposure route (Exhibits 1.55, 1.56, 1.57, pp.1-208 to 1-215, DOE 2000). Total soil and waste ELCRs for the future excavator exceed 1.0E-04 at SWMUs 4, 5, and 6 (Exhibit 1.41, p. 1-184, DOE 2000). Total Uranium is the major contributor to ELCR at SWMU 4 (83%), with external exposure as the driving exposure route (Exhibit 1.55, p. 1-208, DOE 2000). Beryllium and Total PAHs contribute 10% or more to ELCR at SWMU 5, with dermal contact as the driving exposure route (Exhibit 1.56, p. 1-211, DOE 2000). Beryllium is the major contributor to ELCR at SWMU 6, with dermal contact as the driving exposure route (Exhibit 1.57, p. 1-213, DOE 2000).

F.1.1.4 Future Rural Resident

Soil HIs for the future child on-site rural resident exceed *de minimis* levels at SWMUs 4, 5, and 6, but are less than 100 when lead is excluded as a COPC (Exhibit 1.29, p. 1-176, DOE 2000). Including lead as a

COPC masks the contribution from other COPCs. Aluminum, arsenic, chromium, iron, and nickel contribute more than 10% to total HIs, with dermal contact with soil and ingestion of vegetables raised in soil as the driving exposure routes (Exhibits 1.55, 1.56, 1.57, pp.1-208 to 1-215, DOE 2000). The uncertainty associated with the dermal exposure route has been previously discussed. Exclusion of the vegetable exposure route would reduce soil HIs for the rural resident by as much as 87% (Exhibits 1.55, 1.56, 1.57, pp.1-208 to 1-215, DOE 2000). Soil ELCRs for the future rural resident exceed *de minimis* levels at SWMUs 4, 5, and 6 ($> 1.0E-03$). Beryllium and uranium-238 (^{238}U) contribute 10% or more to ELCR at SWMU 4, with dermal contact as the driving exposure route (Exhibit 1.55, p. 1-208, DOE 2000). Arsenic and Total PAHs contribute 10% or more to ELCR at SWMU 5, with ingestion of vegetables as the driving exposure route (Exhibit 1.56, p. 1-211, DOE 2000). Beryllium and Total PAHs contribute 10% or more to ELCR at SWMU 6, with ingestion of vegetables as the driving exposure route (Exhibit 1.57, p. 1-213, DOE 2000). Exclusion of the vegetable exposure route would reduce soil ELCRs for the rural resident by as much as 90% (Exhibits 1.55, 1.56, 1.57, pp.1-208 to 1-215, DOE 2000).

Groundwater HIs for the future child on-site rural resident exceed *de minimis* levels at SWMUs 4, 5, and 6 (178,000 – 2,390,000), but are reduced by several orders of magnitude, with lead excluded as a COPC (223–798) (Exhibits 1.29, 1.30, p. 1-176 to 1-177, DOE 2000). Including lead as a COPC masks the contribution from other COPCs. Iron, manganese, vanadium, carbon tetrachloride, and TCE contribute 10% or more to total HI, with ingestion of water and ingestion of vegetables irrigated with water as the driving exposure routes (Exhibits 1.55, 1.56, 1.57, pp.1-208 to 1-215, DOE 2000). As with the future industrial worker land use scenario, iron is both widespread and predominant as a COC, contributing 49-77% to HI, depending on location (Exhibits 1.55, 1.56, 1.57, pp.1-208 to 1-215, DOE 2000). Exclusion of the vegetable exposure route would reduce groundwater HIs for the rural resident by as much as 40% (Exhibits 1.55, 1.56, 1.57, pp.1-208 to 1-215, DOE 2000). Groundwater ELCRs for the future rural resident exceed *de minimis* levels at SWMUs 4, 5, and 6 ($> 1.0E-03$). Arsenic, beryllium, 1,1-dichloroethene (DCE), TCE, ^{226}Ra , and technetium-99 (^{99}Tc) contribute more than 10% to ELCR, with ingestion of water and ingestion of vegetables irrigated with water as the driving exposure routes (Exhibits 1.55, 1.56, 1.57, pp.1-208 to 1-215, DOE 2000). Exclusion of the vegetable exposure route would reduce groundwater ELCRs for the rural resident by as much as 46% (Exhibits 1.55, 1.56, 1.57, pp.1-208 to 1-215, DOE 2000).

F.1.1.5 Future Recreational User

The future recreational user scenario is not of concern regarding total soil HI at any WAG 3 SWMU (Exhibit 1.32, p. 1-178, DOE 2000). In terms of cancer risks, total soil ELCR exceeds *de minimis* levels only at SWMU 5 ($1.0E-05$), where PAHs contribute 96% to risk, with ingestion of rabbit as the driving exposure route (Exhibits 1.55, 1.56, 1.57, pp.1-208 to 1-215, DOE 2000).

F.1.1.6 Modeled On-site and Off-site COCs

As noted previously, this baseline human health risk assessment (BHHRA) used results of fate and transport modeling [Multimedia Environmental Pollution Assessment System (MEPAS)] to estimate the baseline risks posed to human health through contact with media impacted by contaminants migrating off-site from the various sources in WAG 3. The following chemicals are “priority COCs” for MEPAS-modeled off-site use of groundwater (i.e., rural residential use in the home). These COCs may migrate from a source at a SWMU in WAG 3 to an off-site location and present a chemical-specific HI or ELCR to the rural resident that is greater than 0.1 or $1.0E-06$, respectively (Section 1.5.5, p. 1-185, DOE 2000).

- SWMU 4—arsenic, cobalt, copper, iron, manganese, nickel, vanadium, 1,1-DCE, 1,2-DCE, carbon tetrachloride, TCE, vinyl chloride, neptunium-237 (^{237}Np), plutonium-239 (^{239}Pu), ^{99}Tc , Total Uranium (assessed as ^{238}U), and ^{238}U

- SWMU 5—iron and manganese
- SWMU 6—iron and manganese

The RESidual RADioactive Materials (RESRAD) model was used to model both dose and excess cancer risk for radionuclides, accounting for in-growth of decay products. The following chemicals are “priority COCs” for modeled on-site soil use (i.e., industrial and excavator) and on-site groundwater use (i.e., rural residential use in the home). These chemicals are radionuclides that, through in-growth of decay products, present a chemical-specific ELCR that exceeds 1.0E-06 from exposure to surface and subsurface soil and waste at SWMUs in WAG 3 and radionuclides that may migrate from a source at a SWMU in WAG 3 to on-site RGA groundwater and present a chemical-specific ELCR to the rural resident that is greater than 1.0E-06 (Section 1.5.6, p. 1-190, DOE 2000):

- SWMU 4—thorium-230 (^{230}Th), Total Uranium (modeled as ^{238}U), and ^{238}U
- SWMU 5— ^{226}Ra and ^{238}U
- SWMU 6— ^{237}Np , ^{99}Tc , and ^{238}U

F.1.2 RESULTS OF REMEDIAL INVESTIGATION REPORT FOR SOLID WASTE MANAGEMENT UNITS 7 AND 30 OF WASTE AREA GROUPING 22 AT THE PADUCAH GASEOUS DIFFUSION PLANT, PADUCAH, KENTUCKY, DOE/OR/07-1604/V1&D2

The following is a summary of the results of the BHHRA found in the RI report for the WAG 22, Burial Grounds Solid Waste Management Units 7 and 30 (DOE 1998a). A complete summary of the BHHRA result can be found in WAG 22 report and in Attachment F2.

- **Current Industrial Worker**
 - Soil ingestion
 - Inhalation of volatile organic compounds (VOCs) or airborne soil particulates
 - Dermal contact
 - External exposure to ionizing radiation
- **Future Industrial Worker**
 - Soil ingestion
 - Inhalation of VOCs or airborne soil particulates
 - Dermal contact
 - External exposure to ionizing radiation
 - Ingestion of groundwater
 - Inhalation of VOCs released from groundwater while showering
 - Dermal contact with groundwater contaminants while showering
- **Future On-site Rural Resident**
 - Soil ingestion
 - Inhalation of VOCs or airborne soil particulates
 - Dermal contact
 - External exposure to ionizing radiation
 - Ingestion of groundwater
 - Inhalation of VOCs released from groundwater while showering
 - Dermal contact with groundwater contaminants while showering
 - Exposure to contaminated biota (i.e., garden vegetables)

- **Future Excavation Worker**
Soil ingestion
Inhalation of VOCs or airborne soil particulates
Dermal contact
External exposure to ionizing radiation
- **Future Recreational User**
Ingestion of game species including deer, rabbit, and quail

The total ELCR values calculated using default exposure factors were $> 1.0E-04$ for all receptor scenarios except the recreational user. The total HI value calculated using default factors was > 1 for all scenarios. Total ELCR values calculated for the recreational user were within the $1.0E-04$ to $1.0E-06$ range.

F.1.2.1 Exposure Routes

The dermal contact with soil exposure route poses considerable risk, and most of this risk comes from contact with metals in soil. In fact, for all land use scenarios evaluated, the systemic toxicity and the ELCR posed through the soil dermal exposure route exceeds that posed by the soil ingestion route. This is a direct result of using ABS values that may be too conservative because they exceed gastrointestinal absorption values. This observation indicates that the risk estimates from the dermal exposure route may be unrealistic and exceed the real risk posed by this route of exposure. Although chemical-specific ABS values were used when available, default ABS values were used for most chemicals because chemical-specific values are lacking. Chemical-specific ABS values were available for dioxins, PCBs, cadmium, and carbon disulfide and are used in this BHHRA. Remedial decisions based on the dermal contact with soil exposure route should be carefully considered because of the uncertainty associated with risk from this exposure route.

F.1.2.2 HI and ELCR Discussions

In the next subsections, soil hazards (HI) and ELCRs are compared to *de minimis* levels for each of the scenarios evaluated. Two types of HIs and ELCRs were calculated for this risk assessment and are summarized here. The first is the total HI/ELCR derived using all default exposure values and will be referred to as the default HI/ELCR. The second is the total site-specific or average HI/ELCR without groundwater, with EPA default dermal values, without lead, and without food exposure routes. These risk and hazard values are referred to as site-specific HI/ELCR. The values that will be discussed are those that exceed *de minimis* levels, $HI > 0.1$ and $ELCR > 1.0E-06$.

F.1.2.2.1 Current Industrial Worker

The default HI of 4,800 for combined exposure to soil and groundwater for SWMU 7 and 3,900 for SWMU 30 exceeded *de minimis* levels. The default ELCR of $4E-03$ for both SWMUs and the site-specific ELCR, $1.0E-05$ for both SWMUs exceeded *de minimis* levels. For both SWMUs 7 and 30, dermal contact with soil is the driving exposure route. The primary contributing COC for ELCR is beryllium, and the primary contributing COC for HI is iron (Sections 1.5.3.1, 1.5.3.2, 1.7.4.3, pp. p.1-101, 1-146, DOE 1998a).

F.1.2.2.2 Future Industrial Worker

The default HI of 53,000 for combined exposure to soil and groundwater for SWMU 7 and 22,000 for SWMU 30 and the site-specific HIs of 0.3 for SWMU 7 and 0.2 for SWMU 30 exceeded *de minimis* levels. The default ELCR of $6.0E-03$ for SWMU 7 and $4.0E-03$ for SWMU 30 and the site-specific

ELCRs of 2.0E-04 for both SWMUs exceeded *de minimis* levels. For SWMU 7, the driving exposure route is dermal contact with soil for ELCR, with beryllium being the primary contributor. The driving exposure route for HI is ingestion of groundwater, and the primary contributor is PCBs. For SWMU 30, the driving exposure route is dermal contact with soil, with the contributing COCs as beryllium for ELCR and TCE for HI (Sections 1.5.4.1, 1.5.4.2, 1.7.4.3, p. 1-102, 1-146, DOE 1998a).

F.1.2.2.3 Future On-site Rural Resident

The default HI of 860,000 for SWMU 7 and 460,000 for SWMU 30 and the site-specific HIs of 2 for both SWMUs exceeded *de minimis* levels. The default ELCR of 5.0E-02 for SWMU 7 and 4.0E-02 for SWMU 30 and the site-specific ELCRs of 1.0E-03 for SWMU 7 and 8.0E-04 for SWMU 30 exceeded *de minimis* levels. The future child rural resident scenario was evaluated to determine the exposure route for HI. The driving exposure route for HI is ingestion of vegetables from groundwater with 1,2-DCE (total) being the primary contributor for SWMU 7. The driving exposure route for HI is ingestion of vegetables from soil, with uranium being the primary contributor for SWMU 30. A combined child and adult rural resident was evaluated to determine the driving exposure routes for ELCR. The driving exposure route also was ingestion of vegetables from soil, and beryllium was the primary contributor for both SWMUs (Sections 1.5.4.1, 1.5.4.2, 1.7.4.3, p. 1-102, 1-146, DOE 1998a).

F.1.2.2.4 Future Excavation Worker

The default HI of 7,100 for SWMU 7 and 5,100 for SWMU 30 exceeded *de minimis* levels. Only the default ELCR of 2.0E-03 for SWMU 7 and 1.0E-03 for SWMU 30 exceeded *de minimis* levels. For both SWMUs 7 and 30, the driving exposure route was determined to be dermal contact with soil. Beryllium was the primary contributor for ELCR, and iron was the primary contributor for HI (Sections 1.5.4.1, 1.5.4.2, 1.7.4.3, p. 1-102, 1-146, DOE 1998a).

F.1.2.2.5 Future Recreational User

The default HI of 3 for SWMU 7 and 2 for SWMU 30 exceeded *de minimis* levels. The default ELCR of 1E-05 for SWMU 7 and 2E-05 for SWMU 30 exceeded *de minimis* levels. (Note: the default ELCR for SWMU 30 is reported incorrectly in Table ES.6 and in Section 1.7.4.3 of DOE 1998a. The default ELCR is reported as 1.50E-05 in Table 1.67, so 2E-05 is shown here.) The future adult recreational user scenario was used to determine the exposure route for ELCR. HI was not evaluated because the exposure pathway HI was less than 0.1. The driving exposure route for both SWMUs is ingestion of rabbit, and dibenz(a,h)anthracene was the primary contributor (Sections 1.5.4.1, 1.5.4.2, 1.7.4.3, p. 1-102, 1-146, DOE 1998a).

F.1.2.3 COCs

The COCs were identified in the BHHRA using the approach set forth in the PGDP Risk Methods Document (DOE 2001). The COCs were identified by applying the *de minimis* thresholds to the quantitative risk results for applicable use scenarios including the industrial worker, excavation worker, and off-site rural residential scenarios. A number of COPCs were not evaluated because no toxicity information was available. These COPCs were not retained as COCs in the BHHRA.

Chemicals of concern identified for SWMUs 7 and 30 include VOCs in groundwater; semivolatile organic compounds (SVOCs) in SWMU 7 soil, SWMU 30 soil, and drainage ditch sediments; PCBs in SWMU 30 soil and drainage ditch sediments; metals in SWMU 7 soil, SWMU 30 soil, and drainage ditch sediments; and radionuclides in SWMU 7 soil, SWMU 30 soil, groundwater, and drainage ditch sediments. No COCs were identified for surface water in the ditches because surface-water flow in the

ditches is intermittent and not available for exposure. The COCs identified are listed below (Section 1.5.7.2, p. 1-110, DOE 1998a):

VOCs: vinyl chloride;

SVOCs: benzo(a)anthracene, benzo(a)pyrene, benzo(b)fluoranthene, dibenzo(a,h)anthracene, indeno(1,2,3-cd)pyrene;

PCBs: Aroclor-1260;

Metals: aluminum, antimony, arsenic, beryllium, cadmium, chromium, copper, iron, nickel, manganese, uranium, vanadium;

Radionuclides: ^{237}Np , ^{239}Pu , ^{99}Tc , uranium-234 (^{234}U), uranium-235 (^{235}U), uranium-235/236 ($^{235/236}\text{U}$), ^{238}U .

COCs were identified as those contaminants that had chemical-specific ELCRs summed over all exposure routes within the use scenario of concern that were greater than or equal to 1×10^{-6} or whose hazard quotients (HQs) summed over all exposure routes within a use scenario of concern were greater than or equal to 0.1. Results from SESOIL modeling and modeling for ^{99}Tc are provided in Appendix E, Section E.1.6.

F.1.3 RESULTS OF REMEDIAL INVESTIGATION ADDENDUM FOR WASTE AREA GROUPING 22, BURIAL GROUNDS SOLID WASTE MANAGEMENT UNITS 2 AND 3 AT THE PADUCAH GASEOUS DIFFUSION PLANT, PADUCAH, KENTUCKY, DOE/OR/07-1141&D2

The following is a summary of the BHHRA found in the RI Addendum for the WAG 22, Burial Grounds Solid Waste Management Units 2 and 3 (DOE 1994). A complete summary of the BHHRA result can be found in WAG 22 report and in Tables F2.1 through F2.12 in Attachment F2.

- **Current Industrial Worker**
 - Soil ingestion
 - Dermal absorption
 - Inhalation
 - External exposure to ionizing radiation
- **Site-specific Industrial Worker**
 - Soil ingestion
 - Dermal absorption
 - Inhalation
 - External exposure to ionizing radiation
- **Future On-site and Off-site Rural Resident**
 - Ingestion of groundwater contaminants from groundwater use
 - Inhalation of VOCs from groundwater use

The future resident scenario exhibited a total ELCR value $> 1.0\text{E-}04$ when calculated using default exposure factors. The future resident scenario also exhibited total HIs $> 1.0\text{E-}04$ calculated using default exposure factors > 1 . Site-specific and current industrial worker scenarios exhibited total ELCR values

that fell between 1.0E-04 and 1.0E-06. The site-specific and current industrial worker scenarios also exhibited HI values < 1 (Sections 2.6.1.4.1, 2.6.1.4.2, pp.2-76 to 2-83, DOE 1994).

F.1.3.1 Exposure Routes

From the evaluation of the chemical ELCR, soil ingestion was the driving exposure route and from the evaluation of the radiological ELCR, external exposure to ionizing radiation was the driving exposure route. Through the HI calculation, dermal absorption was the driving exposure route. The dominant driving exposure route for groundwater was ingestion of groundwater. Remedial decisions based on the dermal absorption exposure route should be carefully considered because of the uncertainty associated with risk from this exposure route.

F.1.3.2 HI and ELCR Discussions

In the next subsections soil hazards (HI) and ELCRs are compared to the *de minimis* ELCR level, ELCR > 1.0E-06, and to HI > 0.1 for each of the scenarios evaluated. The values that are discussed are those that exceed these levels. The chemical ELCR and radiological ELCR are presented separately in the original document and each is compared to the *de minimis* ELCR level of 1.0E-06. The HI is compared to the level of 1.0.

F.1.3.2.1 Current Industrial Worker

The chemical ELCR, 5.0E-06, exceeded *de minimis* levels and arsenic was the primary contributing COC. The driving exposure route was soil ingestion. The radiological ELCR was calculated separately in this assessment, 1.0E-04, exceeded *de minimis* level, with ²³⁵U being the primary contributor. The driving exposure route was external exposure to ionizing radiation. The HI did not exceed 0.1 with a result of 0.007 (Attachment 2-4, 2-5, 2-6, DOE 1994).

F.1.3.2.2 Site-specific Industrial Worker

The chemical ELCR, 5.0E-07, did not exceed *de minimis* level. The radiological ELCR, 1.0E-05, did exceed *de minimis* level, with ²³⁵U being the primary contributor. The driving exposure route was external exposure to ionizing radiation. The HI did not exceed 0.1 at a result of 0.07, and the driving exposure route was dermal absorption (Attachment 2-1, 2-2, 2-3, DOE 1994).

F.1.3.2.3 Future On-site and Off-site Rural Resident

The chemical ELCR of 2.0E-03 exceeded *de minimis*, with N-nitroso-di-n-propylamine being the primary contributor. The radiological ELCR of 1.0E-05 exceeded *de minimis* with the primary contributor as ⁹⁹Tc. The HI exceeded 0.1, and the primary contributor was manganese. The dominant exposure route was ingestion of groundwater (Attachment 2-7, 2-10, 2-12, DOE 1994).

F.1.3.3 COCs

The COCs were identified as those contaminants that had chemical-specific ELCRs summed over all exposure routes within the use scenario of concern that were greater than or equal to 1×10^{-6} or whose HQs summed over all routes within a use scenario of concern were greater than or equal to 0.1 for any of the scenarios evaluated: industrial worker, excavation worker, and off-site rural residential scenarios. A number of COPCs were not evaluated because no toxicity information was available. These COPCs were not retained as COCs in the BHHRA. The COCs identified are as listed below (Table 2-23, DOE 1994):

VOCs: TCE.

Metals: arsenic, barium, beryllium, cadmium, chromium, manganese, nickel, silver, thallium, vanadium, uranium (total).

Radionuclides: ⁹⁹Tc.

F.1.4 DATA SUMMARY AND INTERPRETATION REPORT FOR INTERIM REMEDIAL DESIGN AT SOLID WASTE MANAGEMENT UNIT 2 OF WASTE AREA GROUP 22 AT THE PADUCAH GASEOUS DIFFUSION PLANT, PADUCAH, KENTUCKY, DOE/OR/07-1549&D1

F.1.4.1 Screening Approach

This document presents a screening risk assessment for SWMU 2 based on an updated CSM, but, unlike the other documents summarized here, has no BHHRA. The preliminary remediation goals (PRGs) used for the screening were the residential and industrial PRGs from the 1996 Risk Assessments Methods document (DOE 1996). The PRGs for groundwater were based on residential use. A summary of the COPCs from the comparison to PRGs presented in this document is detailed here.

F.1.4.2 Comparison to Industrial PRGs (based on HQ=1 and risk =1E-04)

Sediment COPCs (Section 5.2.1.1, p. 5-4, DOE 1994):

- Arsenic, iron, and vanadium
- Cesium-137 (¹³⁷Cs), ²³⁵U, ²³⁸U, ²³⁹Pu

Surface soil COPCs (Section 5.2.1.2, p. 5-4, DOE 1994):

- Arsenic, beryllium, manganese, and vanadium
- PCBs
- ²³⁵U and ²³⁸U

F.1.4.3 Residential Comparison to PRGs (Section 5.2.1.7, p. 5-6, DOE 1994)

Groundwater (RGA):

- All inorganic chemicals which were analyzed for (particularly arsenic, beryllium, manganese)
- *trans*-1,2-DCE
- ²³⁸U, ⁹⁹Tc, ²³⁹Pu, ²³⁷Np, and Americium-241 (²⁴¹Am)

Because this was a screening assessment, the process stopped at COPC selection and no additional determination of COCs was made in this document.

F.1.5 COPCs FOR GROUNDWATER IDENTIFIED IN BGOU RI WORK PLAN SCREENING

A screening of measured concentrations in the groundwater against no action levels (NALs) and action levels from the 2001 PGDP Risk Methods Document (DOE 2001) is presented in Appendix E of the BGOU RI Work Plan. The priority COCs identified in the screening of measured groundwater data for each of these SWMUs are listed below:

- SWMU 2—arsenic, barium, beryllium, cadmium, iron, manganese, nickel, uranium, vanadium, 1,1-DCE, *cis*-1,2-DCE, TCE, and vinyl chloride
- SWMU 3—arsenic, uranium, TCE, and ^{234}U
- SWMU 4—aluminum, ammonia, antimony, arsenic, barium, beryllium, boron, cadmium, cobalt, copper, iron, manganese, mercury, nickel, nitrate, vanadium, zinc, acetone, PCB-1254, carbon tetrachloride, chloroform, 1,1-DCE, *cis*-1,2-DCE, *trans*-1,2-DCE, naphthalene, 1,1,2-TCA, TCE, vinyl chloride, and ^{99}Tc
- SWMU 5—aluminum, arsenic, barium, beryllium, cadmium, cobalt, copper, iron, manganese, nickel, vanadium, zinc, TCE, and $^{226}\text{Ra}+\text{D}$
- SWMU 6—aluminum, arsenic, barium, beryllium, cadmium, cobalt, copper, iron, manganese, nickel, uranium, vanadium, PCB-1016, TCE, $^{237}\text{Np}+\text{D}$, ^{99}Tc , and ^{234}U
- SWMUs 7 and 30—aluminum, arsenic, barium, beryllium, cadmium, cobalt, copper, iron, manganese, molybdenum, nickel, uranium, acetone, PCB-1016, PCB-1248, PCB-1254, benzene, carbon tetrachloride, chlorobenzene, chloroform, 1,3-dichlorobenzene, 1,4-dichlorobenzene, *cis*-1,2-DCE, 2,4-dimethylphenol, tetrachloroethene, TCE, vinyl chloride, $^{222}\text{Rn}+\text{D}$, ^{99}Tc , and ^{234}U
- SWMU 145—aluminum, antimony, arsenic, barium, boron, iron, manganese, molybdenum, nickel, uranium, vanadium, white phosphorus, acetone, PCB-1016, benzene, chloroform, m-cresol, TCE, $^{222}\text{Rn}+\text{D}$, and ^{234}U

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F.2. IDENTIFICATION OF COPCs

Soil COPCs for direct contact were determined in the previous studies as outlined in Section F.1. Detected soil analytes were screened as described in Section 6 to determine which analytes would be modeled to determine concentrations in groundwater. Modeled groundwater concentrations were used in lieu of measured groundwater concentrations to examine and segregate potential risk and hazard contributions from each SWMU. The maximum detected soil concentrations (surface to 60 ft bgs) was compared to the child resident soil screening levels¹ (SSLs) with a dilution attenuation factor (DAF)=1 (DOE 2001). Those analytes with a maximum concentration greater than their respective SSLs then were compared to soil/sediment child resident NALs (DOE 2001). Those constituents greater than both the SSL and NAL were retained as COPCs for groundwater modeling. TCE, ⁹⁹Tc, and uranium isotopes were retained in all SWMUs, as they are significant risk contributors or known to be part of the facility's process history. Following the review and screening process, the COPCs retained then were modeled as described in Section 5 and Appendix E of the RI Report. Groundwater concentrations used in the assessment also were modeled as shown in Section 5 to several points of exposure away from the SWMU. These were as follows:

- Plant boundary
- Property boundary
- Little Bayou seeps (when particle modeling showed a contribution to the seep)
- Monitoring well located near the Ohio River

Cancer risk and HIs were calculated for groundwater exposure for the rural resident at both the on-site and off-site locations.

¹ SSLs are risk-based soil concentrations considered to be protective of groundwater (DOE 2001).

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F.3. EXPOSURE ASSESSMENT

This section describes the exposure assessment used to determine the pathways of exposure that were considered for the groundwater risk assessment presented in this appendix. Specifically, the exposure assessment process is delineated, the exposure settings of the BGOU are described, the routes of exposure are outlined, and the daily intakes and doses are derived. The ultimate products presented in this section are the CSM for the BGOU and the CDIs used when calculating ELCR and HI in Section 5.

F.3.1 DESCRIPTION OF THE EXPOSURE ASSESSMENT PROCESS

Exposure is the contact of an organism with a chemical or physical agent. The magnitude of exposure (i.e., dose) is determined by measuring or estimating the amount of an agent available at exchange boundaries (e.g., gut, skin, etc.) during a specified period. Exposure assessment is a process that uses information about the exposure setting and human activities to develop CSMs under current and potential future conditions.

The first step in the exposure assessment is to characterize the exposure setting. This includes describing the activities of the human population (on or near a site) that may affect the extent of exposure and the physical characteristics of the site. During this process, sensitive subpopulations that may be present at the site or that may be exposed to contamination migrating from the site also are considered. Generally, site characterization results in a qualitative evaluation of the site and the surrounding population.

The second step in the exposure assessment is to identify exposure pathways. Exposure pathways describe the path a contaminant travels from its source to an individual. A complete exposure pathway includes all links between the source and the exposed population; therefore, a complete pathway consists of a source of release, a mechanism of release, a transport medium, a point of potential human contact, and an exposure route.

The third step in the exposure assessment is to calculate dose by quantifying the magnitude, frequency, and duration of exposure for the populations for the exposure pathways selected for quantitative evaluation. This step involves using the EPCs developed for each COPC to quantify the pathway-specific CDIs for that COPC.

F.3.2 CHARACTERIZATION OF THE EXPOSURE SETTING

The first step in evaluating exposure is to characterize surface features, meteorology, geology, demography and land use, ecology, hydrology, and hydrogeology of the area inhabited by potential receptors. These aspects are discussed in Chapter 3 of this RI report. Physical descriptions of SWMU 2, SWMU 3, SWMU 4, SWMU 5, SWMU 6, SWMU 7, SWMU 30, and SWMU 145 are summarized within this exposure assessment to support later discussions of the conceptual model and its uncertainties.

F.3.2.1 C-749 Uranium Burial Ground (SWMU 2)

The C-749 Uranium Burial Ground (SWMU 2) is located within the west-central portion of the plant, north of Virginia Avenue. SWMU 2 encompasses an area of approximately 32,000 ft², with approximate dimensions of 160 ft by 200 ft. Records indicate that when the burial ground was in use, pits were

excavated to an estimated depth of 7 to 17 ft. After the burial ground no longer was in use, the area was covered with a 6-inch thick clay cap and an 18-inch thick soil layer covered with vegetation (DOE 1995).

F.3.2.2 C-404 Low-Level Radioactive Waste Burial Ground (SWMU 3)

The C-404 Low-Level Radioactive Waste Burial Ground (SWMU 3) includes 1.2 acres located in the west-central portion of the plant area. The unit originally was constructed as a rectangular, aboveground, surface impoundment measuring 387 ft by 137 ft with a floor area of approximately 53,000 ft². The floor of the surface impoundment was constructed of well-tamped earth and clay dikes to a height of 6 ft. The C-404 impoundment was designed with an overflow weir at its southwest corner. From the weir, the surface impoundment effluent flowed west in a ditch [not the North-South Diversion Ditch (NSDD)] and eventually discharged into Kentucky Pollutant Discharge Elimination System Outfall 015.

In March 2003, an additional 37,000 ft² of area were added to the SWMU when a northeast-southwest ditch just east of SWMU 3 was included as part of the SWMU. This ditch was impacted by the discharge of a now-abandoned pipeline with historic leachate flow into the NSDD (DOE 2003). When the C-404 impoundment was converted into a disposal facility, a sump was installed at the weir. The sump was used to pump leachate into an underground transfer line. The transfer line discharged into a northeast-southwest ditch just east of C-404. From this ditch, the leachate flowed into the NSDD. The date of termination of the leachate discharge from the underground transfer line to the NSDD has not been determined. However, it is known that, prior to landfill closure in 1986, this underground transfer line into the NSDD was not in operation, and leachate from the C-404 Landfill was being collected in the sump for treatment at C-400. The wastewater from the treatment of the leachate was discharged to C-403 and, ultimately, to the NSDD. At some time following closure of C-404 Landfill, treatment of leachate from C-404 at C-400 was discontinued and treatment of the leachate was transferred to the C-752 Remedial Action Waste Holding Facility.

F.3.2.3 C-747 Contaminated Burial Yard and C-748-B Burial Area (SWMU 4)

The C-747 Contaminated Burial Yard and the C-748-B Burial Area (SWMU 4) is located in the western section of the plant area. SWMU 4 (which covers an area of approximately 286,700 ft²) is bounded on the north by Virginia Avenue, on the east by 6th Street, on the west by 4th Street, and on the south by an active railroad spur. This SWMU is an open grass field that, at one time, was used for the burial and disposal of various waste materials in designated burial cells. There have been no permanent structures built on the site. SWMU 4 is bounded on three sides (north, east, and west) by shallow drainage swales that direct surface runoff to the northwest corner of the site. There is an elevation difference of approximately 10 ft between the highest point in the SWMU and the adjacent drainage swales. The entire burial yard was covered with 2 to 3 ft of soil material and a 6-inch clay cap was placed over the area in 1982 (DOE 1998b).

F.3.2.4 C-746-F Burial Yard (SWMU 5)

The C-746-F Burial Yard is located in the northwestern section of PGDP. SWMU 5 (which covers an area of approximately 197,400 ft²) is located adjacent to the C-746-P Scrap Yard to the north and SWMU 6 to the east. Disposal pits were located on a grid system. Documentation of the size of these grids ranges from 10 ft by 10 ft cells to 20 ft by 20 ft cells excavated to a depth of 6 to 15 ft bgs. Waste placed in the yard disposal pits was covered with 2 to 3 ft of soil. SWMU 5 is fenced to limit access to authorized personnel only. The ground surface is covered with short grasses and various flowering herbaceous plants (DOE 1998b). The suspected burial area within this SWMU starts approximately 120 ft east of the western boundary of the SWMU and extends to the eastern boundary of the SWMU.

F.3.2.5 C-747-B Burial Ground (SWMU 6)

The C-747-B Burial Ground is located in the northwestern section of the plant area east of SWMU 5. SWMU 6 was in operation from 1960 to 1976. The entire burial area covers an area of approximately 13,500 ft², which is divided into five separate burial cells (Areas H, I, J, K, and L). The following are the dimensions of each of the cells.

- Area H—This disposal site covers an area of about 12 ft by 15 ft and is about 6 ft deep. A 3 ft cover of soil was placed on top of the buried drums.
- Area I—This discard pit is approximately 8 ft by 35 ft and is about 8 ft deep. The waste was covered with about 5 ft of soil.
- Area J—This burial site is about 4,000 ft² (37 ft by 110 ft) and was excavated to a depth of about 6 ft. The area was covered with about 3 ft of soil.
- Area K—This disposal site consists of an area of about 12 ft by 15 ft and is about 6 ft deep. A 3 ft cover of soil was placed on top of the buried drums.
- Area L—This burial area is about 20 ft by 30 ft and about 6 ft deep. The disposed waste was covered with about 3 ft of soil.

This area is relatively flat and is bounded to the north by the roadbed of an abandoned railroad track, to the east by a 5-ft-wide by 4-ft-deep drainage ditch that drains into Ditch 001, and unnamed gravel roads to the west and south. The ground surface is covered by medium to tall grasses (up to 3 ft high).

F.3.2.6 C-747-A Burial Ground (SWMU 7)

The C-747-A area is located in the northwest corner of PGDP. SWMU 7 comprises the eastern two-thirds of C-747-A. The SWMU is bounded on the north and south sides by perimeter ditches, on the west side by the C-747-A Burn Area (SWMU 30), and on the east side by the C-746-E Contaminated Scrap Yard. SWMU 7 covers approximately 240,900 ft² and includes five discrete burial pit areas (Burial Pits B, C, D, F, and G) (DOE 1998a).

Records indicate the burial pits were excavated to a depth of 6 to 7 ft bgs, filled with wastes, and covered with approximately 3 ft of earth; however, the Phase II Site Investigation (SI) discovered waste to a depth of 10 ft on the west side of Burial Pit B, and borings sampled waste to a minimum depth of 8 ft in Burial Pit C (Union Carbide 1978). A stockpile of radiologically-contaminated scrap drums, locally known as Drum Mountain, formerly was located on the southeast corner.

The land surface slopes within SWMU 7. Burial Pits B and C form a slight hill on the north side of SWMU 7, and Burial Pit F forms a lesser mound on the south side of the SWMU. Pit D underlies a level area north of where Drum Mountain once was located. Shallow drainage swales occur on the west side of Burial Pit B and between Burial Pits C and D. The ground surface is covered by grassy vegetation, except where gravel roads extend through the site.

The surface water that drains from SWMU 7 into the surrounding ditches is carried west through Outfall 001 into Bayou Creek. In 2002, a sedimentation basin was constructed to contain runoff from PGDP scrap yards. Runoff now flows into the sedimentation basin and is released periodically into Outfall 001.

F.3.2.7 C-747-A Burn Area (SWMU 30)

SWMU 30 includes the western one-third of C-747-A. It consists of an historical burn-and-burial pit (Burial Pit A) and the location of a former incinerator. The SWMU is bounded on the north and south sides by ditches, on the west side by an unnamed paved road, and on the east side by the C-747-A Burial Ground (SWMU 7). The unit encompasses approximately 128,000 ft². The pit is reported to have been excavated to a depth of 12 ft and covered with 4 ft of earth. The land surface slopes gently, and a slight mound rises over Burial Pit A. SWMU 30 is bordered by drainage ditches on the north and south side. Grassy vegetation covers the ground, except where gravel roads extend through the site.

F.3.2.8 Area P (SWMU 145)

Area P (SWMU 145) is located north of PGDP and is defined by encompassing SWMUs 9 and 10 (the C-746-S&T Landfills, respectively). The SWMU is approximately 44 acres and began operation in the early 1950s. Currently, the C-746-S&T Landfills are located on top of SWMU 145 (DOE 1999). The boundaries of the area are not well defined outside of the area utilized by the C-746-S&T Landfills.

F.3.2.9 Demography and Land Use

As shown in the physical descriptions presented above, current land use of all sources investigated during the BGOU RI is industrial. Under current use, because of access restrictions, only plant workers and authorized visitors are allowed access to the source areas. As discussed in the PGDP Site Management Plan (DOE 2004a), foreseeable future land use of the area is expected to be industrial as well.

At present, both recreational and residential land uses occur in areas surrounding PGDP. Recreational use occurs in the Western Kentucky Wildlife Management Area (WKWMA). The WKWMA is used primarily for hunting and fishing, but other activities include horseback riding, field trials, hiking, and bird watching. An estimated 5,000 fishermen visit the area annually, according to the Kentucky Department of Fish and Wildlife Resources manager of the WKWMA. Residential use near the plant and in areas to which the groundwater from the BGOU may migrate is rural residential and includes agricultural activities. However, current response actions have eliminated exposure to contaminated groundwater by these rural residents. More urban residential use occurs in the villages of Heath, Grahamville, and Kevil, which are within 3 miles of U.S. Department of Energy (DOE) property boundaries, but outside of the area projected to be potentially impacted by the BGOU. The closest major urban area is the municipality of Paducah, Kentucky, which has a population of approximately 26,000 and is approximately 10 miles from PGDP. Other municipalities in the region near PGDP are Cape Girardeau, Missouri, which is approximately 40 miles west of the plant; and the cities of Metropolis and Joppa, Illinois, which are across the Ohio River from PGDP. Total population within a 50-mile radius of the plant is approximately 732,000 people, with about 88,500 people living within 10 miles. The population of McCracken County, in which PGDP lies, is estimated at 65,000 people.

In the area near PGDP and in western Kentucky, in general, the economy has historically been agriculturally based; however, industry has increased in recent years. PGDP is a major employer with approximately 1,400 workers. Another major employer near PGDP is the Tennessee Valley Authority (TVA) Shawnee Steam Plant, which employs approximately 260 individuals.

F.3.3 IDENTIFICATION OF EXPOSURE PATHWAYS

Exposure pathways describe how a contaminant travels from its source to an individual. A complete exposure pathway includes all links between the source and the exposed population. That is, a complete

pathway consists of the source of release, a mechanism of release, a transport medium, a point of potential human contact, and an exposure route. The following discussions focus on points of potential human contact, types of receptors, and exposure routes that are relevant to exposure to contaminated groundwater evaluated in this BHHRA.

F.3.3.1 Points of Human Contact—Land Use Considerations

As discussed earlier, the potential sources to the BGOU are in an industrial area located within a large industrial facility; therefore, the current land use is industrial. Per KDEP and EPA agreement (Risk Methods Document), industrial land use limits the current exposure scenario to an industrial worker (with exposure to the first foot of surface soil) and an excavation worker (with potential exposure to soil in the 0-10 ft bgs depth). The current scenarios do not include any current use of groundwater drawn from the RGA at the sources.

Also as discussed earlier, the current land use can be expected to continue in the foreseeable future, and the most plausible future land use of the BGOU also is industrial. In the future, the expected exposure frequencies and durations may be higher than duration and frequency of the current exposure. Additionally, use of groundwater drawn from the RGA at the sources is not expected; however, uses of areas surrounding PGDP indicate that it would be prudent to examine a range of land uses to provide managers with estimates of the risk that may be posed to humans under alternate uses, however unlikely. In addition, consideration of a range of land uses is consistent with requirements outlined in the Risk Methods Document. Additional possible future land uses considered in earlier BHHRA of the source areas were recreational and rural residential. Recreational and residential scenarios would include use of groundwater drawn from the RGA and potential exposure to groundwater at the seeps at Little Bayou Creek in addition to soil exposure. Baseline risks under each of these uses are presented for each of the SWMUs in the data summary tables in this appendix, with the following exceptions:

- *SWMUs 2 and 3.* SWMUs 2 and 3 were assessed together in the WAG 22 RI. (DOE 1994) Assessment of future residential and excavation worker exposure to on-site soils was not included in the WAG 22 RI and, therefore, not in the current BGOU D2 RI. The risk associated with both these scenarios would be greater than the risk for the industrial worker because industrial worker soil exposure exhibited risks above *de minimis* levels and indicates a need for action; therefore, assessment of the residential and excavation scenarios is not warranted.
- *SWMU 145.* SWMU 145 is partially located beneath SWMUs 9 and 10 and, as a result, limited soil data are available for SWMU 145. Only groundwater exposure pathways were considered in assessing this SWMU. Uncertainty associated with soil exposure at SWMU 145 is addressed in the BGOU FS.

F.3.3.2 Potential Receptor Populations

As noted above, the potential receptor population under current conditions at the source units are the industrial worker and the excavation worker. The potential receptor populations under future conditions in BGOU areas also include recreational and residential exposures. Residential and recreational exposures would include exposure to groundwater.

F.3.3.3 Delineation of Exposure Point/Exposure Routes

As discussed, human health risks are assessed by determining point of exposure (POEs) and exposure routes. POEs are locations where human receptors can contact contaminated media. Exposure routes are the processes by which human receptors contact contaminated media. The exposure routes considered during the exposure assessment for groundwater in all BHHRA per the Risk Methods Document are listed in the

following paragraphs. This material also presents reasons for selecting or not selecting each exposure route for each of the potentially exposed populations in this BHHRA.

Ingestion Groundwater as a Drinking Water Source. Residential and industrial use of groundwater is common in western Kentucky. Potential receptors for this pathway are rural residents. This exposure route is assessed quantitatively in this BHHRA.

Inhalation of Volatile Constituents Emitted While Using Groundwater. As noted previously, residential and industrial use of groundwater is common in western Kentucky. Rural residents are potential receptors for this exposure route. This exposure route is assessed quantitatively in this BHHRA.

Dermal Contact with Groundwater While Showering. As noted earlier, residential and industrial use of groundwater is common in western Kentucky. Rural residents are potential receptors for this exposure route. This exposure route is assessed quantitatively in this BHHRA.

Inhalation of Vapor Released from the Ground Water Into Home Basements. This exposure route was modeled quantitatively in this BHHRA for rural residents. Potentially, industrial workers also could be exposed through this route. Because the resident has an exposure time of 24 hrs/day for this exposure route, the rural resident quantitative assessment is protective of any potential worker exposure.

Inhalation of Volatile Organic Compounds During Irrigation with Contaminated Groundwater. In the Midwest, irrigation of farmland with groundwater using center pivot irrigation is common. Rural residents are potential receptors for this exposure route. Because only modeled groundwater data are available for this BHHRA in areas where migration may occur in the future and because earlier assessments have shown that risk from this exposure route is minimal, this exposure route is not assessed quantitatively in this BHHRA.

Dermal Contact with Water While Swimming or Wading in Privately Owned Fish Ponds Filled with Groundwater. Contamination found in BGOU soils has a reasonable potential of contaminating surface waters through dissolution into the groundwater. Contaminants also may be contacted through suspension of fine particles in the ponds, also originating from groundwater. Recreational use of these ponds by residents may reasonably be expected to occur. During recreational use (e.g., swimming or wading), dermal contact with water could occur. Rural residents are potential receptors for this exposure route. Because only modeled groundwater data are available for this BHHRA in areas where this activity may occur in the future, this exposure route is not assessed quantitatively in this BHHRA.

Incidental Ingestion of Sediment While Swimming or Wading in Privately Owned Fishponds Filled with Groundwater. The rationale for considering ponds is presented previously. In addition, recreational use of these ponds by residents may reasonably be expected to occur. During recreational activities, incidental ingestion of sediment contaminated by constituents in groundwater is possible. Rural residents are potential receptors for this exposure route. Because only modeled groundwater data are available for this BHHRA in areas where this activity may occur in the future, this exposure route is not assessed quantitatively in this BHHRA.

External Exposure to Ionizing Radiation Emitted by Contaminants in Groundwater While Swimming or Wading in Privately Owned Fish Ponds Filled with Groundwater. The rationale for considering ponds is presented previously. During use of these ponds by residents, exposure to ionizing radiation emitted by radionuclides in water could occur. Rural residents are potential receptors for this exposure route. Because only modeled groundwater data primarily were used this BHHRA in areas where this activity may occur in the future, this exposure route is not assessed quantitatively in this BHHRA.

External Exposure to Ionizing Radiation Emitted by Contaminants in Sediment While Swimming or Wading in Privately Owned Fish Ponds Filled with Groundwater. The rationale for considering ponds is presented previously. During use of these ponds by residents, exposure to ionizing radiation emitted by radionuclides in groundwater and sediment could occur. Rural residents are potential receptors for this exposure route. Because only modeled groundwater data primarily were used for this BHHRA in areas where this activity may occur in the future, this exposure route is not assessed quantitatively in this BHHRA.

Consumption of Fish Raised in Privately Owned Fish Ponds Filled with Groundwater. The fish raised in ponds would be exposed to contaminants in groundwater and may accumulate some contaminants in their edible tissues. These fish, caught in either a “pay-to-fish” or a commercial pond by residents, could reasonably be expected to be consumed. Recreational users (i.e., visitors) and rural residents are potential receptors for this exposure route. Because only modeled groundwater data primarily were used for this BHHRA in areas where this activity may occur in the future, this exposure route is not assessed quantitatively in this BHHRA.

Incidental Ingestion of Surface Water in Creeks or Ponds. Open bodies of water, such as Bayou Creek or settling ponds, are attractive for recreation (e.g., swimming and wading). Although such bodies of water are not included in the assessment of the source areas, contaminants may migrate from the sources to these creeks or ponds. Recreational users and industrial workers are potential receptors for this exposure route. This exposure route is not assessed quantitatively in this BHHRA because earlier BHHRAs have concluded that mixing with surface water results in risks that are insignificant. Additionally, risks associated with surface water will be evaluated as part of the Surface Water Operable Unit (SWOU).

Dermal Contact with Surface Water While Swimming or Wading in Creeks or Ponds. Open bodies of water, such as Bayou Creek or settling ponds, are attractive for recreation (e.g., swimming and wading). Although such bodies of water are not included in this assessment of the source areas, contaminants may migrate from sources to these bodies of water. Recreational users and industrial workers are potential receptors for this exposure route. This exposure route is not assessed quantitatively in this BHHRA because earlier BHHRAs have concluded that mixing with surface water results in risks that are insignificant. Additionally, risks associated with surface water will be evaluated as part of the SWOU.

Incidental Ingestion of Sediment While Swimming or Wading in Creeks or Ponds. Open bodies of water, such as Bayou Creek or settling ponds, are attractive for recreation (e.g., swimming and wading). Although such bodies of water are not included in this assessment of the source areas, contaminants may migrate from sources to these bodies of water. Recreational users and industrial workers are potential receptors for this exposure route. This exposure route is not assessed quantitatively in this BHHRA because earlier BHHRAs have concluded that mixing with surface water results in risks that are insignificant. Additionally, risks associated with surface water will be evaluated as part of the SWOU.

External Exposure to Ionizing Radiation Emitted by Contaminants in Surface Water While Swimming or Wading in Creeks or Ponds. Open bodies of water, such as Bayou Creek or settling ponds, are attractive for recreation (e.g., swimming and wading). Although such bodies of water are not included in this assessment of the source areas, contaminants may migrate from sources to these bodies of water. Recreational users and industrial workers are potential receptors for this exposure route. This exposure route is not assessed quantitatively in this BHHRA because earlier BHHRAs have concluded that mixing with surface water results in risks that are insignificant. Additionally, risks associated with surface water will be evaluated as part of the SWOU.

External Exposure to Ionizing Radiation Emitted by Contaminants in Sediment While Swimming or Wading in Creeks or Ponds. Open bodies of water, such as Bayou Creek or settling ponds, are attractive for recreation (e.g., swimming and wading). Although such bodies of water are not included in this assessment of the source areas, contaminants may migrate from sources to these bodies of water. Recreational users and industrial workers are potential receptors for this exposure route. This exposure route is not assessed quantitatively in this BHHRA because earlier BHHRAs have concluded that mixing with surface water results in risks that are insignificant. Additionally, risks associated with surface water will be evaluated as part of the SWOU.

Consumption of Fish Taken from Creeks and Ponds Containing Contaminated Surface Water. Fish living in Bayou Creek or settling ponds may accumulate contaminants in surface water in their edible tissues. Although such bodies of water are not included in this assessment of the source areas, contaminants may migrate from sources to these bodies of water. Recreational users and residents may catch and consume fish from the potentially impacted surface water bodies. Potential receptors for this route of exposure are recreational users. This exposure route is not assessed quantitatively in this BHHRA because earlier BHHRAs have concluded that mixing with surface water results in risks that are insignificant. Additionally, risks associated with surface water will be evaluated as part of the SWOU.

Consumption of Beef from Cattle Contaminated by Consuming Vegetation (Pasture and Concentrates) Irrigated with Groundwater and Drinking Groundwater. During interviews, Agriculture Extension Agents for Ballard and McCracken counties indicated that small scale cow-calf operations are common in western Kentucky. (See Section 2 of Appendix 5 of the Risk Methods Document.) They further noted that slaughtering feeder cattle for home consumption is common. In the study area, such beef may be contaminated by incidental ingestion of soil while on pasture, by consumption of contaminated vegetation (pasture and concentrate), and by ingestion of contaminated groundwater. Residents may eat this beef; therefore, potential receptors for this route of exposure are rural residents. This exposure route was considered in earlier BHHRAs, but is not reassessed in this BHHRA because new soil data are not available, and only modeled groundwater data are available in this BHHRA in areas where this activity may occur in the future. The potential impact of excluding the contribution from this pathway is addressed qualitatively in the uncertainty section.

Consumption of Dairy Products (i.e., Milk) from Cows Contaminated by Consuming Vegetation (Pasture or Concentrates) Irrigated with Groundwater and Drinking Groundwater. During interviews, Agriculture Extension Agents for Ballard and McCracken counties noted that dairy farming still occurs in their counties. (See Section 2 of Appendix 5 of the Risk Methods Document.) Furthermore, the agents stated that these cattle are fed stored feed and are allowed to graze on pasture. As noted previously, the soil at source units is contaminated, and the vegetation may become contaminated. Dairy cattle raised at the sources after the industrial infrastructure is removed may become contaminated through incidental ingestion of soil while on pasture, consumption of contaminated vegetation, and ingestion of contaminated groundwater. Products made from milk from these cows could, in turn, be consumed by residents; therefore, potential receptors for this route of exposure are rural residents. This exposure route was considered in earlier BHHRAs, but is not reassessed in this BHHRA because new soil data are not available, and only modeled groundwater data are available in this BHHRA in areas where this activity may occur in the future. The potential impact of excluding the contribution from this pathway is addressed qualitatively in the uncertainty section.

Consumption of Poultry Given Groundwater to Drink. During interviews, Agriculture Extension Agents for Ballard and McCracken counties noted that commercial broiler production did occur in their counties, but not near PGDP. (See Section 2 of Appendix 5 of the Risk Methods Document.) (Home flocks for both meat and eggs were noted as being uncommon.) Furthermore, they stated that broilers were fed bought (not locally raised) feed, that normal resident time in poultry houses was 2 months, and

that commercial distribution of the product occurs. The agents did note that the birds are most likely watered with groundwater; therefore, broilers may become contaminated through ingestion of contaminated groundwater. For this exposure assessment, the receptor assumed to consume the contaminated poultry is the rural resident. Because only modeled groundwater data are available for this BHHRA in areas where this activity may occur in the future, this exposure route is not assessed quantitatively in this BHHRA. The potential impact of excluding the contribution from this pathway is addressed qualitatively in the uncertainty section.

Consumption of Pork from Swine Watered with Groundwater. During interviews, Agriculture Extension Agents for Ballard and McCracken counties noted that both large commercial and small hog farms exist in their counties. (See Section 2 of Appendix 5 of the Risk Methods Document.) Furthermore, they indicated that swine on both types of farms were fed locally raised feed and, on the smaller farms, that farm-raised pork was consumed by farmers. Any swine raised may be contaminated through consumption of contaminated feed and groundwater, and this pork may be eaten by rural residents; therefore, rural residents are potential receptors for this pathway. Because only modeled groundwater data are available for this BHHRA in areas where this activity may occur in the future, this exposure route is not assessed quantitatively in this BHHRA. The potential impact of excluding the contribution from this pathway is addressed qualitatively in the uncertainty section.

Out of the potential routes described above, five routes of exposure to groundwater were considered in the BHHRA in this RI for the BGOU:

- Ingestion of water while using groundwater as a drinking water source Table F.50
- Dermal contact with groundwater while showering..... Table F.51
- Inhalation of volatiles in groundwater while showering Table F.52
- Inhalation of volatiles in groundwater during household use..... Table F.53
- Inhalation of volatiles as a result of vapor intrusion into home basements.. Appendix E

CDIs, which are calculated for inorganic and organic constituents, and radionuclide intakes, calculated for radionuclides, represent the exposure to a COPC as mass contacted per unit body weight per unit time for the applicable receptor (EPA 1991). Doses, which apply only to radionuclide COPCs, represent the activity of a COPC in contact with an exchange boundary (EPA 1991). Unless otherwise noted, CDIs, RIs and doses are calculated using the values presented in Tables F.1 through F.4, and are from the 2001 approved version of the Risk Methods Document. Values obtained from the draft 2008 revision of the Risk Methods Document are footnoted (DOE 2008). Values in these tables marked as “chemical-specific” were obtained from tables in Appendix B and Appendix D of the Draft 2008 Risk Methods Document. The ABS factors used are from the Draft 2008 Risk Methods Document as well, because these factors apply only to COPCs evaluated for dermal toxicity.

Table F.1. Ingestion of Groundwater by a Rural Resident

$$\text{Chronic Daily Intake (mg/kg x day)} = \text{CDI} = [C_w * \text{IR} * \text{EF} * \text{ED} / (\text{BW} * \text{AT})]$$

$$\text{Radionuclide Intake (pCi)} = \text{RI} = [A_w * \text{IR} * \text{EF} * \text{ED}]$$

Parameter	Value Used	Units
Chemical concentration in water = C_w	Chemical-specific	mg/L
Radiological activity in water = A_w	Chemical-specific	pCi/L
Ingestion rate = IR	2 (adult)	L/day
	1.5 (child) ^a	L/day
Exposure frequency = EF	350	day/yr
Exposure duration = ED	24 (adult) ^a	day
	6 (child)	day
Body weight = BW	70 (adult)	kg
	15 (child)	kg
Averaging time = AT	70 x 365 (carcinogenic)	yr x day/yr
	ED x 365 (noncarcinogenic)	yr x day/yr

^a Value from 2008 draft revision of the Risk Methods Document (DOE 2008)

Table F.2. Dermal Contact with Water While Showering by a Rural Resident

$$\text{Chronic Daily Intake (mg/kg-day)}_{\text{inorganic}} = \text{CDI}_{\text{inorganic}} = [C_w * \text{SA} * K_p * \text{CF} * \text{EF} * \text{ED} * \text{ET} / (\text{BW} * \text{AT})]$$

$$\text{Chronic Daily Intake (mg/kg-day)}_{\text{organic}} = \text{CDI}_{\text{organic}} = [C_w * \text{DA}_{\text{eventfactor}} * \text{SA} * \text{CF} * \text{CF}_o * \text{ED} * \text{EF} * \text{EV} / (\text{BW} * \text{AT})]$$

Parameter	Value Used	Units
Chemical concentration in water = C_w	Chemical-specific	mg/L
Skin surface area exposed = SA	1.815 ^b	m ²
	0.65 ^{a,b}	m ²
Skin permeability constant = K_p	Chemical-specific ^a	cm/hr
Absorbed dose factor per event = $\text{DA}_{\text{eventfactor}}$	Chemical-specific ^a	L/cm ² -event
Conversion factor = CF	10	(L-m)/(cm-m ³)
Conversion factor (organic) = CF_o	10 ³	cm ² /m ²
Exposure frequency = EF	350	baths/yr
Exposure duration = ED	24 (adult) ^a	day
	6 (child)	day
Exposure time = ET	0.2	hr/bath
Event = EV	1	Event/day
Body weight = BW	70 (adult)	kg
	15 (child)	kg
Averaging time = AT	70 x 365 (carcinogenic)	yr x day/yr
	ED x 365 (noncarcinogenic)	yr x day/yr

^a Value from 2008 draft revision of the Risk Methods Document

^b Entire surface area of body for both adult and child

Table F.3. Inhalation of Volatile Organic Compounds in Water While Showering by a Rural Resident

$$\text{Chronic Daily Intake (mg/kg-day)} = \text{CDI} = [C_{\text{shower}} * \text{IR}_{\text{air}} * \text{EF} * \text{ED} * \text{ET} / (\text{BW} * \text{AT})]$$

$$\text{Radionuclide Intake (pCi)} = \text{RI} = [A_{\text{gw}} * \text{IR}_{\text{air}} * \text{ED} * \text{EF} * \text{IEF}]$$

$$C_{\text{shower}} = [((C_{\text{amax}}/2) * t_1) + (C_{\text{amax}} * t_2)] / (t_1 + t_2)$$

$$C_{\text{amax}} = (C_{\text{gw}} * f * F_w * t_1) / V_a$$

Parameter	Value Used	Units
Chemical concentration in groundwater = C_{gw}	Chemical-specific	mg/L
Radionuclide activity in groundwater = A_{gw}	Chemical-specific	pCi/L
Time-adjusted concentration in shower = C_{shower}	Chemical-specific	mg/m ³
Indoor inhalation rate = IR_{air}	0.833 ^a	m ³ /hr
Exposure frequency = EF	350	day/yr
Exposure duration = ED	24 (adult) ^a	day
	6 (child)	day
Exposure time = ET	0.2	hr/day
Inhalation exposure factor = IEF^b	Chemical-specific ^a	(L-hr)/(m ³ -day)
Maximum air concentration = C_{amax}	Chemical-specific ^a	mg/m ³
Time of shower = t_1	0.1	hr
Time after shower = t_2	0.1	hr
Fraction volatilized = f	0.75	unitless
Water flow rate = F_w	890	L/hr
Bathroom volume = V_a	11	m ³
Body weight = BW	70 (adult)	kg
	15 (child)	kg
Averaging time = AT	70 x 365 (carcinogenic)	yr x day/yr
	ED x 365 (noncarcinogenic)	yr x day/yr

^a Value from 2008 draft revision of the Risk Methods Document

^b Default value is 0. Values for tritium and radon are 0.2064 and 5.6, respectively (DOE 2008)

Table F.4. Inhalation of Volatile Organic Compounds in Water During Household Use by a Rural Resident

$$\text{Chronic Daily Intake (mg/kg-day)} = \text{CDI} = [C_{\text{house}} * \text{IR}_{\text{air}} * \text{EF} * \text{ED} * \text{ET} / (\text{BW} * \text{AT})]$$

$$\text{Radionuclide Intake (pCi)} = \text{RI} = [A_{\text{gw}} * \text{IR}_{\text{air}} * \text{ED} * \text{EF} * \text{IEF}]$$

$$C_{\text{house}} = C_{\text{gw}} * \text{WHF} * f / (\text{HV} * \text{ER} * \text{MC})$$

Parameter	Value Used	Units
Chemical concentration in groundwater = C_{gw}	Chemical-specific	mg/L
Radionuclide activity in groundwater = A_{gw}	Chemical-specific	pCi/L
Concentration in household air = C_{house}	Chemical-specific	mg/m ³
Indoor inhalation rate = IR_{air}	0.833	m ³ /hr
Exposure frequency = EF	350	day/yr
Exposure duration = ED	24 (adult) ^a	day
	6 (child)	day
Exposure time = ET	24 ^a	hr/day
Inhalation exposure factor = IEF^b	Chemical-specific ^a	(L-hr)/(m ³ -day)
Water flow rate = WHF	890	L/day
Fraction volatilized = f	0.75	unitless
House volume = HV	450	m ³ /change
Exchanged rate = ER	10	changes/day
Mixing coefficient = MC	0.5	unitless
Body weight = BW	70 (adult)	kg
	15 (child)	kg
Averaging time = AT	70 x 365 (carcinogenic)	yr x day/yr
	ED x 365 (noncarcinogenic)	yr x day/yr

^a Value from 2008 draft revision of the Risk Methods Document

^b Default value is 0. Values for tritium and radon are 0.2064 and 5.6, respectively (DOE 2008)

F.3.3.4 Development of Conceptual Site Models

Using the information presented in the previous subsections, a CSM was developed for the BGOU. This CSM (Figure F.1) illustrates the sources, pathways of migration, and routes of exposure relevant to this and earlier BHHRA. For this screening and the subsequent BHHRA, surface soil was defined as 0–1 ft bgs and subsurface soil was defined as 0–10 ft bgs. Surface soil was used to evaluate direct exposure for residential, recreational, and industrial receptors. Subsurface soil was used to evaluate direct exposure for excavation worker. Table F.5 shows the media evaluated for each land use scenario for each SWMU.

Table F.5. Land Uses And Media Assessed For Each Source Area in Current and Previous BHHRA for the BGOU

	Location								
	SWMU 2	SWMU 3	SWMU 4	SWMU 5	SWMU 6	SWMU 7	SWMU 30	SWMU 145	
Future On-site Industrial Worker									
Surface Soil	P	P	P	PX	P	P	P	NA	
Surface Water	NA	NA	NA	NA	NA	NA	NA	NA	
Future On-site Excavation Worker									
Subsurface Soil	NA	NA	P	P	P	P	P	NA	
Future Recreational User									
Game (Soil)	P	P	P	P	P	P	P	NA	
Surface Soil	NA	NA	NA	NA	P	P	P	NA	
Surface Water	NA	NA	NA	NA	NA	NA	NA	NA	
Future On-site Rural Resident									
Soil	NA	NA	P	P	P	P	P	NA	
Groundwater ^a	PX	PX	PX	PX	PX	PX	PX	X	
Vapor Intrusion ^c	X	X	X	X	X	X	X	X	
Future Off-site Rural Resident									
Groundwater ^b	X	X	X	X	X	X	X	X	
Vapor Intrusion ^c									
Future On-site Terrestrial Biota			P						
Soil	P	NA	NA	P	P	P	P	NA	
Surface Water	NA	NA		NA	NA	NA	NA	NA	

Notes: Scenarios that were assessed in this RI BHHRA are marked with an X. Scenarios assessed in previous BHHRA are marked with a P. Scenarios assessed in both the past and current RBA are marked with a PX. Scenarios not assessed because the scenario is not applicable, or for which the medium is not present, are marked with an NA.

^a The earlier BHHRA assessed risks from use of water drawn from the RGA separately from use of water drawn from the McNairy Formation. The risks assessed in this RI BHHRA are for use of water drawn from the RGA.

^b Modeling results were used to assess groundwater risk to the off-site rural resident. POEs are at the PGDP plant boundary, at the PGDP property boundary, Little Bayou seeps and in a groundwater well at the Ohio River. These POEs are presented in Figure 5.1.

^c Vapor intrusion was modeled for residential basements.

F.3.4 QUANTIFICATION OF EXPOSURE

F.3.4.1 Calculation of EPCs of COPCs

EPCs for groundwater used to determine potential future risks for residential use of groundwater at the four POEs discussed in Sections 5.1 and 5.3 (i.e., under the edge of the SWMU boundary, plant boundary, property boundary, and either Ohio River or Little Bayou seeps) were developed from

modeling. The modeled concentrations in groundwater over time at the four POEs are provided in the figures in Section 5. Modeled groundwater concentrations for a future on-site residential scenario correspond to a resident using groundwater drawn from under the edge of the SWMU footprint. The maximum modeled groundwater concentration over the 1,000 year time frame (see Appendix E for details of the modeling) at each POE was used as the EPC for calculation of the groundwater CDI.

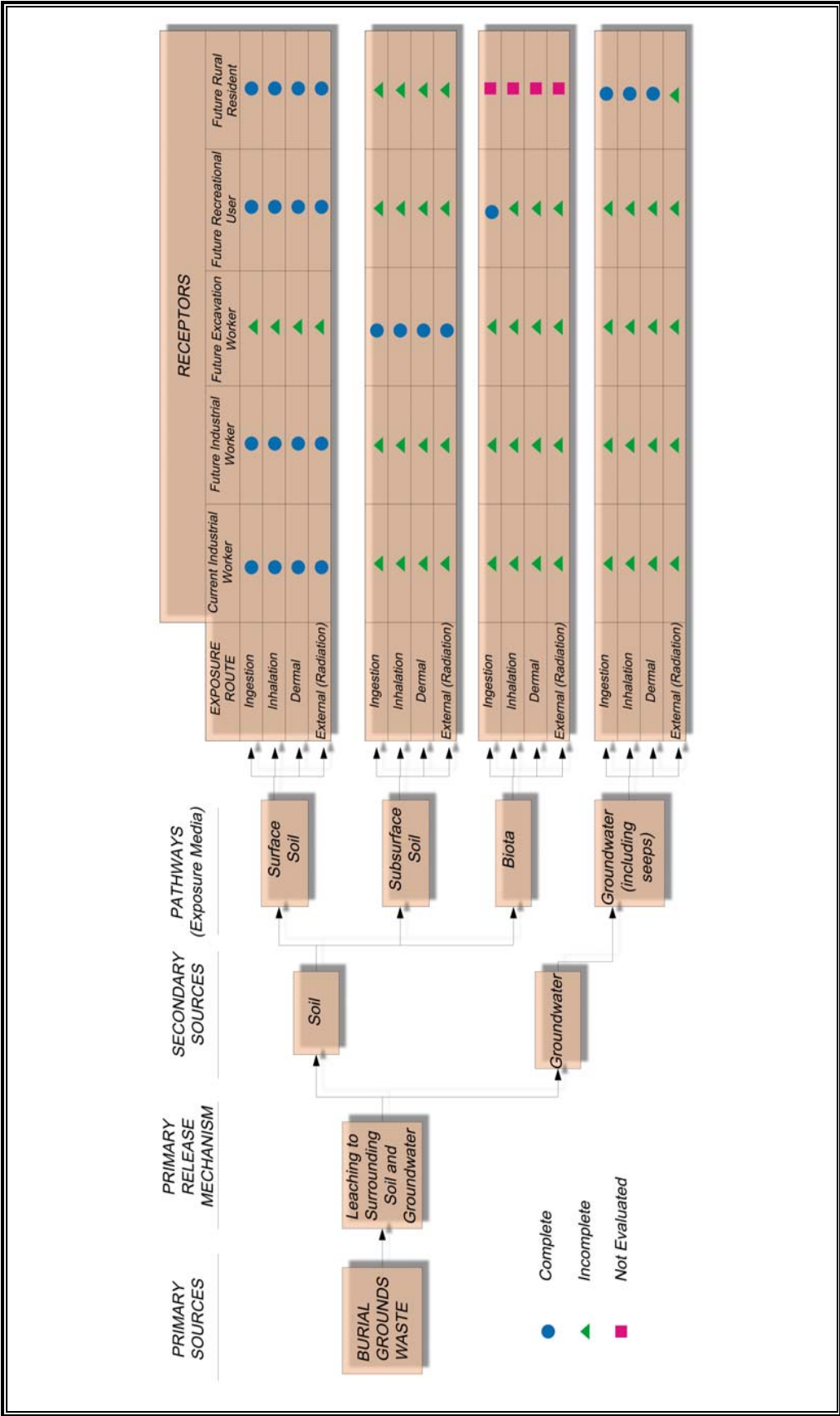


Figure F.1. Conceptual Site Model for BGOU

F.3.4.2 Chronic Daily Intakes

All exposure estimates in this BHHRA represent normalized exposure rates that are evaluated for sources of uncertainty such as variability in data, modeling results, and/or parameter assumptions. Specifically, in this BHHRA, the exposure estimates are an estimation of the reasonable maximum exposure (RME) that can be expected to occur under current or future site conditions. An RME estimate is a conservative estimate of exposure that falls within the upper bound of the range of all possible exposure estimates. In situations where populations are exposed through multiple pathways, RME estimates are calculated for both individual and multiple exposure pathways. Risk estimates for soil and groundwater were calculated separately.

Consistent with the Risk Methods Document, the focus of the exposure assessment for this BHHRA is to determine chronic intake or dose. The chronic exposure estimate is used because it allows for estimation of health consequences that result from long-term or unrestricted exposure to contaminants.

Using the human exposure models for groundwater, the CSM, and the EPCs, the CDIs of groundwater for each of the COPCs were determined. These CDIs are presented in Tables F.6 through F.29. In this presentation, the CDIs used to estimate HI (i.e., noncarcinogenic effects) are presented first, and the values used to estimate ELCR follow.

F.3.5 SUMMARY OF EXPOSURE ASSESSMENT

The receptors evaluated in the current assessment are the adult and child resident for exposure to groundwater.

Table F.6. Chronic Daily Intakes (Non-Carcinogenic) for Child Residential Groundwater User at SWMU 2

COPC	Exposure Point Concentration ^a	Exposure Route-Chronic Daily Intake ^b			
		Ingestion of Water	Dermal Contact	Shower Inhalation	Household Inhalation
<i>Inorganic Compounds</i>					
Arsenic	3.54E-02	3.39E-03	2.94E-06		
Manganese	7.16E-01	6.87E-02	5.95E-05		
Uranium	9.86E-03	9.45E-04	8.19E-07		
<i>Organic Compounds</i>					
<i>cis</i> -1,2-DCE	1.15E+01	1.10E+00	3.89E-02	5.57E-01	4.36E+00
Naphthalene	9.38E-04	8.99E-05	1.63E-05	4.55E-05	3.56E-04
TCE	1.48E+00	1.42E-01	6.64E-03	7.17E-02	5.61E-01

Blank cells indicate that the exposure route is not appropriate to the COPC.

^a Units for metals and organic compounds are mg/L.

^b Units for intakes for metals and organic compounds are mg/(kg-day).

Table F.7. Chronic Daily Intakes (Non-Carcinogenic) for Adult Residential Groundwater User at SWMU 2

COPC	Exposure Point Concentration ^a	Exposure Route-Chronic Daily Intake ^b			
		Ingestion of Water	Dermal Contact	Shower Inhalation	Household Inhalation
<i>Inorganic Compounds</i>					
Arsenic	3.54E-02	9.70E-04	1.76E-06		
Manganese	7.16E-01	1.96E-02	3.56E-05		
Uranium	9.86E-03	2.70E-04	4.90E-07		
<i>Organic Compounds</i>					
cis-1,2-DCE	1.15E+01	3.15E-01	2.33E-02	1.19E-01	9.34E-01
Naphthalene	9.38E-04	2.57E-05	9.76E-06	9.74E-06	7.62E-05
TCE	1.48E+00	4.05E-02	3.97E-03	1.54E-02	1.20E-01

Blank cells indicate that the exposure route is not appropriate to the COPC.

^a Units for metals and organic compounds are mg/L.

^b Units for intakes for metals and organic compounds are mg/(kg-day).

Table F.8. Chronic Daily Intakes (Carcinogenic) for Residential Groundwater User at SWMU 2

COPC	Exposure Point Concentration ^a	Exposure Route-Chronic Daily Intake ^b			
		Ingestion of Water	Dermal Contact	Shower Inhalation	Household Inhalation
<i>Inorganic Compounds</i>					
Arsenic	3.54E-02	6.23E-04	8.56E-07		
<i>Organic Compounds</i>					
TCE	1.48E+00	2.61E-02	1.93E-03	1.14E-02	8.93E-02
<i>Radionuclides</i>					
Technetium-99	1.02E+02	2.03E+06			
Uranium-234	1.58E+00	3.15E+04			
Uranium-238	1.81E+00	3.61E+04			

Blank cells indicate that the exposure route is not appropriate to the COPC.

^a Units for metals and organic compounds are mg/L. Units for radionuclides are pCi/L.

^b Units for intakes for metals and organic compounds are mg/(kg-day). Units for intakes for radionuclides are pCi/(kg-day).

Table F.9. Chronic Daily Intakes (Non-Carcinogenic) for Child Residential Groundwater User at SWMU 3

COPC	Exposure Point Concentration ^a	Exposure Route-Chronic Daily Intake ^b			
		Ingestion of Water	Dermal Contact	Shower Inhalation	Household Inhalation
<i>Inorganic Compounds</i>					
Arsenic	3.29E-02	3.15E-03	2.73E-06		
Manganese	8.95E-01	8.58E-02	7.44E-05		
Uranium	4.89E-02	4.69E-03	4.06E-06		

Blank cells indicate that the exposure route is not appropriate to the COPC.

^a Units for metals and organic compounds are mg/L.

^b Units for intakes for metals and organic compounds are mg/(kg-day).

Table F.10. Chronic Daily Intakes (Non-Carcinogenic) for Adult Residential Groundwater User at SWMU 3

COPC	Exposure Point Concentration ^a	Exposure Route-Chronic Daily Intake ^b			
		Ingestion of Water	Dermal Contact	Shower Inhalation	Household Inhalation
<i>Inorganic Compounds</i>					
Arsenic	3.29E-02	9.01E-04	1.64E-06		
Manganese	8.95E-01	2.45E-02	4.45E-05		
Uranium	4.89E-02	1.34E-03	2.43E-06		

Blank cells indicate that the exposure route is not appropriate to the COPC.

^a Units for metals and organic compounds are mg/L.

^b Units for intakes for metals and organic compounds are mg/(kg-day).

Table F.11. Chronic Daily Intakes (Carcinogenic) for Residential Groundwater User at SWMU 3

COPC	Exposure Point Concentration ^a	Exposure Route-Chronic Daily Intake ^b			
		Ingestion of Water	Dermal Contact	Shower Inhalation	Household Inhalation
<i>Inorganic Compounds</i>					
Arsenic	3.29E-02	5.79E-04	7.95E-07		
<i>Radionuclides</i>					
Technetium-99	5.56E+03	1.11E+08			
Uranium-238	1.59E+01	3.16E+05			

Blank cells indicate that the exposure route is not appropriate to the COPC.

^a Units for metals and organic compounds are mg/L. Units for radionuclides are pCi/L.

^b Units for intakes for metals and organic compounds are mg/(kg-day). Units for intakes for radionuclides are pCi/(kg-day).

Table F.12. Chronic Daily Intakes (Non-Carcinogenic) for Child Residential Groundwater User at SWMU 4

COPC	Exposure Point Concentration ^a	Exposure Route-Chronic Daily Intake ^b			
		Ingestion of Water	Dermal Contact	Shower Inhalation	Household Inhalation
<i>Inorganic Compounds</i>					
Arsenic	1.77E-02	1.70E-03	1.47E-06		
Manganese	5.76E-01	5.52E-02	4.79E-05		
<i>Organic Compounds</i>					
cis-1,2-DCE	6.68E-01	6.41E-02	2.26E-03	3.24E-02	2.53E-01
TCE	1.18E+00	1.13E-01	5.28E-03	5.72E-02	4.47E-01
Vinyl Chloride	2.61E-02	2.50E-03	5.40E-05	2.70E-04	9.90E-03

Blank cells indicate that the exposure route is not appropriate to the COPC.

^a Units for metals and organic compounds are mg/L.

^b Units for intakes for metals and organic compounds are mg/(kg-day).

Table F.13. Chronic Daily Intakes (Non-Carcinogenic) for Adult Residential Groundwater User at SWMU 4

COPC	Exposure Point Concentration ^a	Exposure Route-Chronic Daily Intake ^b			
		Ingestion of Water	Dermal Contact	Shower Inhalation	Household Inhalation
<i>Inorganic Compounds</i>					
Arsenic	1.77E-02	4.85E-04	8.80E-07		
Manganese	5.76E-01	1.58E-02	2.86E-05		
<i>Organic Compounds</i>					
cis-1,2-DCE	6.68E-01	1.83E-02	1.35E-03	6.91E-03	5.43E-02
TCE	1.18E+00	3.23E-02	3.16E-03	1.22E-02	9.59E-02
Vinyl Chloride	2.61E-02	7.15E-04	3.23E-05	2.70E-04	2.12E-03

Blank cells indicate that the exposure route is not appropriate to the COPC.

^a Units for metals and organic compounds are mg/L.

^b Units for intakes for metals and organic compounds are mg/(kg-day).

Table F.14. Chronic Daily Intakes (Carcinogenic) for Residential Groundwater User at SWMU 4

COPC	Exposure Point Concentration ^a	Exposure Route-Chronic Daily Intake ^b			
		Ingestion of Water	Dermal Contact	Shower Inhalation	Household Inhalation
<i>Inorganic Compounds</i>					
Arsenic	1.77E-02	3.12E-04	4.28E-07		
<i>Organic Compounds</i>					
TCE	1.18E+00	2.08E-02	1.54E-03	9.10E-03	7.12E-02
Vinyl Chloride	2.61E-02	4.60E-04	1.05E-02	2.01E-04	1.58E-03
<i>Radionuclides</i>					
Technetium-99	9.01E+03	1.80E+08			

Blank cells indicate that the exposure route is not appropriate to the COPC.

^a Units for metals and organic compounds are mg/L. Units for radionuclides are pCi/L.

^b Units for intakes for metals and organic compounds are mg/(kg-day). Units for intakes for radionuclides are pCi/(kg-day).

Table F.15. Chronic Daily Intakes (Non-Carcinogenic) for Child Residential Groundwater User at SWMU 5

COPC	Exposure Point Concentration ^a	Exposure Route-Chronic Daily Intake ^b			
		Ingestion of Water	Dermal Contact	Shower Inhalation	Household Inhalation
<i>Inorganic Compounds</i>					
Arsenic	9.25E-03	8.87E-04	7.69E-07		
Manganese	1.01E+00	9.68E-02	8.39E-05		
Uranium	4.60E-01	4.41E-02	3.82E-05		
<i>Organic Compounds</i>					
Naphthalene	5.55E-03	5.32E-04	9.66E-05	2.69E-04	2.10E-03

Blank cells indicate that the exposure route is not appropriate to the COPC.

^a Units for metals and organic compounds are mg/L.

^b Units for intakes for metals and organic compounds are mg/(kg-day).

Table F.16. Chronic Daily Intakes (Non-Carcinogenic) for Adult Residential Groundwater User at SWMU 5

COPC	Exposure Point Concentration ^a	Exposure Route-Chronic Daily Intake ^b			
		Ingestion of Water	Dermal Contact	Shower Inhalation	Household Inhalation
<i>Inorganic Compounds</i>					
Arsenic	9.25E-03	2.53E-04	4.60E-07		
Manganese	1.01E+00	2.77E-02	5.02E-05		
Uranium	4.60E-01	1.26E-02	2.29E-05		
<i>Organic Compounds</i>					
Naphthalene	5.55E-03	1.52E-04	5.78E-05	5.76E-05	4.51E-04

Blank cells indicate that the exposure route is not appropriate to the COPC.

^a Units for metals and organic compounds are mg/L.

^b Units for intakes for metals and organic compounds are mg/(kg-day).

Table F.17. Chronic Daily Intakes (Carcinogenic) for Residential Groundwater User at SWMU 5

COPC	Exposure Point Concentration ^a	Exposure Route-Chronic Daily Intake ^b			
		Ingestion of Water	Dermal Contact	Shower Inhalation	Household Inhalation
<i>Inorganic Compounds</i>					
Arsenic	9.25E-03	1.63E-04	2.24E-07		
<i>Radionuclides</i>					
Technetium-99	1.27E+02	2.53E+06			

Blank cells indicate that the exposure route is not appropriate to the COPC.

^a Units for metals and organic compounds are mg/L. Units for radionuclides are pCi/L.

^b Units for intakes for metals and organic compounds are mg/(kg-day). Units for intakes for radionuclides are pCi/(kg-day).

Table F.18. Chronic Daily Intakes (Non-Carcinogenic) for Child Residential Groundwater User at SWMU 6

COPC	Exposure Point Concentration	Exposure Route-Chronic Daily Intake			
		Ingestion of Water	Dermal Contact	Shower Inhalation	Household Inhalation

NO COPCs¹

¹Modeling analysis (Appendix E) did not show any of the identified COPCs at this site as migrating to groundwater.

Table F.19. Chronic Daily Intakes (Non-Carcinogenic) for Adult Residential Groundwater User at SWMU 6

COPC	Exposure Point Concentration	Exposure Route-Chronic Daily Intake			
		Ingestion of Water	Dermal Contact	Shower Inhalation	Household Inhalation

NO COPCS¹

¹Modeling analysis (Appendix E) did not show any of the identified COPCs at this site as migrating to groundwater.

Table F.20. Chronic Daily Intakes (Carcinogenic) for Residential Groundwater User at SWMU 6

COPC	Exposure Point Concentration	Exposure Route-Chronic Daily Intake			
		Ingestion of Water	Dermal Contact	Shower Inhalation	Household Inhalation

NO COPCs¹

¹Modeling analysis (Appendix E) did not show any of the identified COPCs at this site as migrating to groundwater.

Table F.21. Chronic Daily Intakes (Non-Carcinogenic) for Child Residential Groundwater User at SWMU 7

COPC	Exposure Point Concentration ^a	Exposure Route-Chronic Daily Intake ^b			
		Ingestion of Water	Dermal Contact	Shower Inhalation	Household Inhalation
<i>Inorganic Compounds</i>					
Arsenic	1.78E-02	1.71E-03	1.48E-06		
Manganese	3.32E-01	3.18E-02	2.76E-05		
Uranium	3.46E-03	3.32E-04	2.88E-07		
<i>Organic Compounds</i>					
1,1-DCE	8.98E-02	8.61E-03	3.23E-04	4.35E-03	3.40E-02
cis-1,2-DCE	2.35E-02	2.25E-03	7.95E-05	1.14E-03	8.91E-03
Aroclor-1254	5.23E-05	5.02E-06	5.09E-05	2.54E-06	1.98E-05
TCE	1.09E-02	1.05E-03	4.89E-05	5.28E-04	4.13E-03
Vinyl Chloride	1.35E-02	1.29E-03	2.79E-05	6.54E-04	5.12E-03

Blank cells indicate that the exposure route is not appropriate to the COPC.

^aUnits for metals and organic compounds are mg/L.

^bUnits for intakes for metals and organic compounds are mg/(kg-day).

Table F.22. Chronic Daily Intakes (Non-Carcinogenic) for Adult Residential Groundwater User at SWMU 7

COPC	Exposure Point Concentration ^a	Exposure Route-Chronic Daily Intake ^b			
		Ingestion of Water	Dermal Contact	Shower Inhalation	Household Inhalation
<i>Inorganic Compounds</i>					
Arsenic	1.78E-02	4.88E-04	8.85E-07		
Manganese	3.32E-01	9.10E-03	1.65E-05		
Uranium	3.46E-03	9.48E-05	1.72E-07		
<i>Organic Compounds</i>					
1,1-DCE	8.98E-02	2.46E-03	1.93E-04	9.30E-04	7.30E-03
cis-1,2-DCE	2.35E-02	6.44E-04	4.76E-05	2.43E-04	1.91E-03
Aroclor-1254	5.23E-05	1.43E-06	3.05E-05	5.41E-07	4.25E-06
TCE	1.09E02	2.99E-04	2.93E-05	1.13E-04	8.86E-04
Vinyl Chloride	1.35E-02	3.70E-04	1.67E-05	1.40E-04	1.10E-03

Blank cells indicate that the exposure route is not appropriate to the COPC.

^a Units for metals and organic compounds are mg/L.

^b Units for intakes for metals and organic compounds are mg/(kg-day).

Table F.23. Chronic Daily Intakes (Carcinogenic) for Residential Groundwater User at SWMU 7

COPC	Exposure Point Concentration ^a	Exposure Route-Chronic Daily Intake ^b			
		Ingestion of Water	Dermal Contact	Shower Inhalation	Household Inhalation
<i>Inorganic Compounds</i>					
Arsenic	1.78E-02	3.14E-04	4.30E-07		
<i>Organic Compounds</i>					
1,1-DCE	8.98E-02	1.58E-03	9.39E-05	6.93E-04	5.42E-03
Aroclor-1254	5.23E-05	9.21E-07	1.48E-05	4.04E-07	3.16E-06
TCE	1.09E-02	1.92E-04	1.42E-05	8.41E-05	6.58E-04
Vinyl Chloride	1.35E-02	2.38E-04	8.12E-06	1.04E-04	8.15E-04
<i>Radionuclides</i>					
Technetium-99	9.09E+02	1.81E+07			
Uranium-234	7.94E+00	1.58E+05			
Uranium-238	7.59E+00	1.51E+05			

Blank cells indicate that the exposure route is not appropriate to the COPC.

^a Units for metals and organic compounds are mg/L. Units for radionuclides are pCi/L.

^b Units for intakes for metals and organic compounds are mg/(kg-day). Units for intakes for radionuclides are pCi/(kg-day).

Table F.24. Chronic Daily Intakes (Non-Carcinogenic) for Child Residential Groundwater User at SWMU 30

COPC	Exposure Point Concentration ^a	Exposure Route-Chronic Daily Intake ^b			
		Ingestion of Water	Dermal Contact	Shower Inhalation	Household Inhalation
<i>Inorganic Compounds</i>					
Arsenic	1.77E-02	1.70E-03	1.47E-06		
Manganese	3.78E-01	3.62E-02	3.14E-05		
Selenium	1.51E-02	1.45E-03	1.25E-06		
Uranium	8.40E-03	8.05E-04	6.98E-07		
<i>Organic Compounds</i>					
1,1-DCE	6.05E-02	5.80E-03	2.17E-04	2.93E-03	2.29E-02
TCE	9.11E-04	8.74E-05	4.09E-06	4.42E-05	3.45E-04

Blank cells indicate that the exposure route is not appropriate to the COPC.

^a Units for metals and organic compounds are mg/L.

^b Units for intakes for metals and organic compounds are mg/(kg-day).

Table F.25. Chronic Daily Intakes (Non-Carcinogenic) for Adult Residential Groundwater User at SWMU 30

COPC	Exposure Point Concentration ^a	Exposure Route-Chronic Daily Intake ^b			
		Ingestion of Water	Dermal Contact	Shower Inhalation	Household Inhalation
<i>Inorganic Compounds</i>					
Arsenic	1.77E-02	4.85E-04	8.80E-07		
Manganese	3.78E-01	1.04E-02	1.88E-05		
Selenium	1.51E-02	4.14E-04	7.51E-07		
Uranium	8.40E-03	2.30E-04	4.18E-07		
<i>Organic Compounds</i>					
1,1-DCE	6.05E-02	5.00E-02	5.00E-02	6.28E-04	4.92E-03
TCE	9.11E-04	2.50E-05	2.45E-06	9.46E-06	7.40E-05

Blank cells indicate that the exposure route is not appropriate to the COPC.

^a Units for metals and organic compounds are mg/L. Units for radionuclides are pCi/L.

^b Units for intakes for metals and organic compounds are mg/(kg-day). Units for intakes for radionuclides are pCi/(kg-day).

Table F.26. Chronic Daily Intakes (Carcinogenic) for Residential Groundwater User at SWMU 30

COPC	Exposure Point Concentration ^a	Exposure Route-Chronic Daily Intake ^b			
		Ingestion of Water	Dermal Contact	Shower Inhalation	Household Inhalation
<i>Inorganic Compounds</i>					
Arsenic	1.77E-02	3.12E-04	4.28E-07		
<i>Organic Compounds</i>					
1,1- DCE	6.05E-02	1.07E-03	6.32E-05	4.66E-04	3.66E-03
TCE	9.11E-04	1.60E-05	1.19E-06	7.02E-06	5.50E-05
<i>Radionuclides</i>					
Technetium-99	2.87E+02	5.73E+06			
Uranium-234	3.99E+00	7.96E+04			
Uranium-238	5.91E+00	1.18E+05			

Blank cells indicate that the exposure route is not appropriate to the COPC.

^a Units for metals and organic compounds are mg/L. Units for radionuclides are pCi/L.

^b Units for intakes for metals and organic compounds are mg/(kg-day). Units for intakes for radionuclides are pCi/(kg-day).

Table F.27. Chronic Daily Intakes (Non-Carcinogenic) for Child Residential Groundwater User at SWMU 145

COPC	Exposure Point Concentration ^a	Exposure Route-Chronic Daily Intake ^b			
		Ingestion of Water	Dermal Contact	Shower Inhalation	Household Inhalation
<i>Inorganic Compounds</i>					
Antimony	7.99E-02	7.66E-03	6.64E-06		
Arsenic	6.21E-02	5.95E-03	5.16E-06		
Manganese	8.44E-01	8.09E-02	7.01E-05		

Blank cells indicate that the exposure route is not appropriate to the COPC.

^a Units for metals and organic compounds are mg/L.

^b Units for intakes for metals and organic compounds are mg/(kg-day).

Table F.28. Chronic Daily Intakes (Non-Carcinogenic) for Adult Residential Groundwater User at SWMU 145

COPC	Exposure Point Concentration ^a	Exposure Route-Chronic Daily Intake ^b			
		Ingestion of Water	Dermal Contact	Shower Inhalation	Household Inhalation
<i>Inorganic Compounds</i>					
Antimony	7.99E-02	2.19E-03	3.97E-06		
Arsenic	6.21E-02	1.70E-03	3.09E-06		
Manganese	8.44E-01	2.31E-02	4.20E-05		

Blank cells indicate that the exposure route is not appropriate to the COPC.

^a Units for metals and organic compounds are mg/L.

^b Units for intakes for metals and organic compounds are mg/(kg-day).

Table F.29. Chronic Daily Intakes (Carcinogenic) for Residential Groundwater User at SWMU 145

COPC	Exposure Point Concentration ^a	Exposure Route-Chronic Daily Intake ^b			
		Ingestion of Water	Dermal Contact	Shower Inhalation	Household Inhalation
<i>Inorganic Compounds</i>					
Arsenic	6.21E-02	1.09E-03	1.50E-06		
<i>Organic Compounds</i>					
Aroclor-1260	1.92E-03	3.38E-05	7.14E+00	1.48E-05	6.91E-01
<i>Radionuclides</i>					
Technetium-99	1.01E+04	2.02E+08			
Uranium-238	7.67E-02	1.53E+02			

Blank cells indicate that the exposure route is not appropriate to the COPC.

^a Units for metals and organic compounds are mg/L. Units for radionuclides are pCi/L.

^b Units for intakes for metals and organic compounds are mg/(kg-day). Units for intakes for radionuclides are pCi/(kg-day).

F.4. TOXICITY ASSESSMENT

This section summarizes the potential toxicological effects of the COPCs on exposed populations. Many of the toxicological summaries were obtained from the *Risk Assessment Information System* (RAIS) prepared by the Toxicology and Risk Analysis Section of Oak Ridge National Laboratory for DOE (DOE 2004b). This site also lists toxicity values taken from the EPA's Integrated Risk Information System (IRIS) database (EPA 2004a), National Center for Environmental Assessment (NCEA), and Health Effects Assessment Summary Tables (HEAST) database (EPA 1998). This list formed the basis of the toxicity values reported in this section. For those chemicals not profiled in RAIS, a brief summary of information drawn from Agency for Toxic Substances and Disease Registry (ATSDR) or other library research sources is included in this section. The last paragraph of each profile contains the toxicity values used in this BHHRA.

The toxicity information considered in the assessment of potential carcinogenic risks includes (1) a weight-of-evidence classification and (2) a slope factor (SF). The weight-of-evidence classification qualitatively describes the likelihood that an agent is a human carcinogen, based on the available data from animal and human studies. A chemical may be placed in one of three groups to indicate its potential for carcinogenic effects: Group A, a known human carcinogen; Group B, a probable human carcinogen; and Group C, a possible human carcinogen. Group B is divided into Subgroups B1 and B2. Assignment of a chemical to Subgroup B1 indicates that the judgment that the chemical is a probable human carcinogen is based on limited human data, and assignment of a chemical to Subgroup B2 indicates that the judgment that the chemical is a probable human carcinogen is based on animal data because human data are lacking or inadequate. Chemicals that cannot be classified as human carcinogens because of a lack of data are categorized in Group D, and those for which there is evidence of noncarcinogenicity in humans are categorized in Group E.

The slope factor for chemicals is defined as a plausible upperbound estimate of the probability of a response (i.e., development of cancer) per unit intake of a chemical over a lifetime (EPA 1989). Slope factors are specific for each chemical and route of exposure. Slope factors currently are available for ingestion and inhalation pathways. The slope factors used for oral and inhalation routes of exposure for the COPCs considered in this report are shown in Table F.30.

Table F.30. Toxicity Values For Chronic Exposure to Carcinogens Via the Ingestion and Inhalation Exposure Routes

COPC ^a	Class	Oral Slope		Oral Unit Risk ^d	Inhalation		Types of Cancers	
		Factor ^b	Source ^c		Slope Factor ^e	Unit Risk ^f		
<i>Inorganic Chemicals (Metals)</i>								
Arsenic	A	1.50E+00	a	5.00E-02	1.51E+01	a	4.30E-03	Respiratory system tumors
<i>Organic Compounds</i>								
1,1-DCE	C	6.00E-01	b	1.70E-02	1.75E-01	a	5.00E-05	Kidney, adenocarcinoma
Aroclor-1254	B2	4.00E-01	a		3.50E-01	a	5.71E-01	Liver
Aroclor-1260	B2	4.00E-01	a		3.50E-01	a	5.71E-01	Liver

Table F.30. Toxicity Values For Chronic Exposure to Carcinogens Via the Ingestion and Inhalation Exposure Routes (Continued)

COPC ^a	Class	Oral Slope		Oral Unit Risk ^d	Inhalation		Types of Cancers	
		Oral Slope Factor ^b	Factor Source ^c		Slope Factor ^e	Slope Factor Source ^c		Inhalation Unit Risk ^f
TCE ^h	C-B2	3.22E-01	c		3.22E-01	c	1.10E-01	Liver and lung cancer
Vinyl Chloride	A	1.50E+00	a	4.20E-02	3.08E-02	a	8.80E-06	Liver, lung, digestive tract, and brain tumors
<i>Radionuclides</i>								
	ICRP ^g Lung Class							
Technetium-99	M	2.75E-12	d		1.41E-11	d		Various
Uranium-234	M	7.07E-11	d		1.14E-08	d		Various
Uranium-238	M	8.71E-11	d		9.35E-09	d		Various

Note: Blank cells indicate that data are not available or are not appropriate.

^a All groundwater COPCs are listed.

^b The units for the oral slope factors are (mg/kg × day)⁻¹ for nonradionuclides and risk/pCi for radionuclides.

^c Source codes are defined as follows:

a: *Integrated Risk Information System* (IRIS) (EPA 2004a)

b: This value has been withdrawn by EPA's National Center for Environmental Assessment (NCEA) and is being reassessed for IRIS. It is acceptable for use in this risk assessment as no replacement value currently exists (EPA 2000).

c: KDEP

d: *Health Effects Assessment Summary Table* (HEAST) (EPA 1998b)

^d The units for the oral unit risks are (mg/L)⁻¹.

^e The units for the inhalation slope factors are (mg/kg × day)⁻¹ for nonradionuclides and risk/pCi for radionuclides.

^f The units for inhalation unit risks are m³/μg.

^g ICRP Publication 72 is referenced in the HEAST user's guide (ICRP, 1996). Lung class absorption types are defined as follows:

S = slow (particulate)

M = medium (particulate)

F = fast (particulate)

^h Value used is from KDEP (2004) review of TCE slope factors. The slope factors used in previous assessments were 0.052 for the oral slope factor and 0.002 for the inhalation slope factor. This issue is discussed further in the uncertainty section.

Toxicity values used in risk calculations also include the chronic RfD, which is used to estimate the potential for systemic toxicity or noncarcinogenic risk. The chronic RfD is defined as an estimate of a daily exposure level for the human population, including sensitive subpopulations, that is likely to be without an appreciable risk of deleterious effects during a lifetime (EPA 1989). RfD values are specific to the route of exposure. The RfDs used for oral and inhalation routes of exposure for the COPCs considered in this report are presented in Table F.31.

Table F.31. Toxicity Values for Chronic Exposure to Noncarcinogens Via the Ingestion and Inhalation Exposure Routes

COPC ^a	Oral Reference Dose ^b	Oral Reference Dose Source ^c	Inhalation Reference Dose ^d	Inhalation Reference Concentration ^e	Inhalation Reference Concentration Source ^e	RfD basis (vehicle) ^f	Target Organ Critical Effect	Confidence Level ^f	Uncertainty Factor/Modifying Factor ^f
<i>Inorganic Chemicals (Metals)</i>									
Antimony	4.00E-04	a				(O)LOAEL	GI	(O)Low	(O)UF=1,000 (O)MF=1
Arsenic	3.00E-04	a				(O)NOAEL/ LOAEL	Skin	Medium	(O)UF=3 (O)MF=1
Manganese	4.60E-02	a,b	1.43E-05	5.00E-05	a	(O)NOAEL	NA	Low	(I)UF=1,000 (I)MF=1 (O)=100
Selenium	5.00E-03	a				NOAEL/ LOAEL	Lungs (selenosis)	High	(O)UF=3 (O)MF=1
Uranium	6.00E-04	d				LOAEL	Kidney	NA	(O)UF=100 (O)MF=1
<i>Organic Compounds</i>									
1,1-DCE	5.00E-02	a	5.71E-02	2.00E-01	a	LOAEL	Liver	Medium	(O)UF=1,000 (O)MF=1
1,2-DCE, <i>cis</i> -	1.00E-02	x	9.97E-03	3.49E-02	ex	NOAEL	Blood	Low	(O)UF=3,000 (O)MF=1
Aroclor-1254	2.00E-05	a	1.99E-05		ex	(O)LOAEL	Endocrine System	Medium	(O)UF=300 (O)MF=1
TCE	3.00E-04	c	1.14E-02	4.00E-02	c	NA	Liver, kidney, CNS	NA	NA
Vinyl Chloride	3.00E-03	a	2.86E-02	1.00E-01	a	(I)NOAEL/ LOAEL (O)NOAEL/ LOAEL	Liver, kidney, CNS	Medium	(I)UF=30 (I)MF=1 (O)UF=3 (I)MF=1
Naphthalene	2.00E-02	a	8.57E-04	3.00E-03	a	(O)NOAEL (I)LOAEL	Decreased body weight Respiratory	(O)Low (I)Medium	(O)UF=3,000 (O)MF=1 (I)UF=3,000 (I)MF=1

Notes: Blank cells indicate that data are not available or are not appropriate. NA=information not readily available at this time; GI=gastrointestinal; CNS=central nervous system

^a All groundwater COPCs are listed.

^b The units for the oral reference doses are mg/(kg × day).

^c Source codes are defined as follows:

a: *Integrated Risk Information System (IRIS)* (EPA 2004a)

b: IRIS no longer separates manganese values for chronic oral RfDs into water and diet RfDs. IRIS recommends using a modifying factor of 3 to lower the total oral RfD of 1.40E-01 to 4.67E-02. This has been rounded to 4.6E-02 to make the value more conservative.

c: EPA NCEA (EPA 2001)

d: Also see *Soil Screening Guidance for Radionuclides: User's Guide*.

ex: Value is extrapolated from the oral reference dose.

x: A provisional value from EPA National Center for Environmental Assessment (NCEA).

^d The units for the inhalation reference doses are mg/(kg × day).

^e The units for the inhalation reference concentrations are mg/m³.

^f O=oral; I=inhalation; UF=uncertainty factor; MF=modifying factor; NA=not available.

For the dermal routes of exposure (i.e., dermal exposure to contaminated water during swimming or bathing), it is necessary to consider the absorbed dose received by a receptor. This is reflected by the addition of an absorption coefficient in the equations used to calculate the CDI for these pathways. Because the CDI is expressed as an absorbed dose, it is necessary to use RfDs and slope factors that also are expressed in terms of absorbed dose. Currently, EPA has not produced lists of RfDs and slope factors based on absorbed dose but have produced guidance concerning the estimation of absorbed dose RfDs and slope factors from administered dose RfDs and slope factors. This guidance is found in *Risk Assessment Guidance for Superfund, Volume I: Human Health Evaluation Manual (Part E, Supplemental Guidance for Dermal Risk Assessment)* (EPA 2004b) and states, “that to convert an administered dose

slope factor to an absorbed dose slope factor, the administered dose slope factor is divided by the gastrointestinal (GI) absorption efficiency of the contaminant.” Alternatively, to convert an administered dose RfD to an absorbed dose RfD, the administered dose RfD is multiplied by the GI absorption efficiency of the contaminant. The absorbed dose slope factors and RfDs and the information used in their derivation are presented in Tables F.32 and F.33, respectively.

Table F.32. Toxicity Values for Chronic Dermal Contact Exposure to Carcinogens

COPC ^a	Dermal Slope Factor ^b	GI ABS Factor ^c
<i>Inorganic Chemicals (Metals)</i>		
Arsenic	3.66E+00	0.41
<i>Organic Compounds</i>		
1,1-DCE	6.00E-01	1.0
Aroclor-1254	4.44E-01	0.9
Aroclor-1260	4.44E-01	0.9
TCE	2.67E+00	0.15
Vinyl Chloride	1.50E+00	1.0

Note: Blank cells indicate that data are not available or are not appropriate.

^a All groundwater COPCs are listed.

^b The units for these dermal dose slope factors are (mg/kg × d)⁻¹ for nonradionuclides. Absorbed cancer slope factors are calculated by dividing the administered cancer slope factor by GI absorption factor; this value is used in the BHHRA to calculate contribution to cancer risk from dermal exposure.

^c All GI ABS factors from Draft 2008 Risk Methods Document (DOE 2008)

Table F.33. Toxicity Values for Chronic Exposure to Noncarcinogens Via the Dermal Contact Exposure Route

COPC ^a	Dermal Reference Dose ^b	Administered Reference Dose ^c	GI ABS ^d
Antimony	8.00E-06	4.00E-04	0.02
Arsenic	1.23E-04	3.00E-04	0.41
Manganese	1.84E-03	4.60E-02	0.04
Selenium	2.20E-03	5.00E-03	0.44
Uranium ^e	5.10E-04	6.00E-04	0.85
1,1-DCE	5.00E-02	5.00E-02	1
1,2-DCE, <i>cis</i> -	1.00E-02	1.00E-02	1
Aroclor-1254	1.80E-05	2.00E-05	0.9
Naphthalene	1.60E-02	2.00E-02	0.8
TCE	4.50E-05	3.00E-04	0.15
Vinyl Chloride	3.00E-03	3.00E-03	1

Note: Blank cells indicate that data are not available or are not appropriate.

^a All groundwater COPCs are listed except radionuclides because external exposure to water is not assessed.

^b The units for the absorbed doses are mg/(kg × day). All dermal reference doses calculated by multiplying the administered reference dose by the GI absorption factor; this value is used in the BHHRA to calculate contribution to the hazard index from dermal exposure.

^c Administered reference doses are equivalent to the oral reference dose listed in Table F.31. The units are mg/(kg × day).

^d GI absorption factors are from the Draft 2008 Risk Methods Document (DOE 2008) and are unitless

^eUranium Source: 40 *CFR* Part 141 (2000).

F.4.1 INORGANIC COMPOUNDS

F.4.1.1 Antimony (CAS 007440-36-0) (RAIS)

Antimony is a naturally occurring silvery-white metal that is found in the earth's crust. Antimony ores are mined and then mixed with other metals to form antimony alloys or combined with oxygen to form antimony oxide. Little antimony is currently mined in the United States. It is brought into this country from other countries for processing; however, there are companies in the United States that produce antimony as a by-product of smelting lead and other metals. Antimony is used in lead storage batteries, solder, sheet and pipe metal, bearings, castings, and pewter. Antimony oxide is added to textiles and plastics to prevent them from catching fire. It also is used in paints, ceramics, and fireworks, and as enamels for plastics, metal, and glass.

Metallic antimony and a few trivalent antimony compounds are the most significant regarding exposure potential and toxicity. Antimony is a common urban air pollutant, occurring at an average concentration of 0.001 $\mu\text{g}/\text{m}^3$. Exposure to antimony may occur via inhalation oral and dermal routes.

Acute oral and inhalation exposure of humans and animals to high doses of antimony or antimony-containing compounds (antimonials) may cause gastrointestinal disorders (vomiting, diarrhea), respiratory difficulties, and death at extremely high doses. Subchronic and chronic oral exposure may affect hematologic parameters. Long-term oral exposure to high doses of antimony or antimonials has been shown to adversely affect longevity in animals. Long-term occupational exposure of humans has resulted in electrocardiac disorders, respiratory disorders, and possibly increased mortality. Antimony levels for these occupational exposure evaluations ranged from 2.2 to 11.98 $\text{mg Sb}/\text{m}^3$. Based on limited data, occupational exposure of women to metallic antimony and several antimonials has reportedly caused alterations in the menstrual cycle and an increased incidence of spontaneous abortions.

The Department of Health and Human Services (DHHS), the International Agency for Research on Cancer (IARC), and the EPA have not classified antimony as to its human carcinogenicity.

Chronic RfDs for antimony are available in RAIS. The oral RfD used in the BHHRA is 4.00E-04 (mg/kg-day). The absorbed dose RfD is 8.00E-06 (mg/kg-day), which was calculated using a GI absorption factor of 2%, per the Risk Methods Document (DOE 2008).

F.4.1.2 Arsenic (CAS 007440-38-2) (RAIS)

Arsenic is a naturally occurring element widely distributed in the earth's crust. In the environment, arsenic is combined with oxygen, chlorine, and sulfur to form inorganic arsenic compounds. Arsenic in animals and plants combines with carbon and hydrogen to form organic arsenic compounds. Inorganic arsenic compounds are mainly used to preserve wood. Organic arsenic compounds are used as pesticides, primarily on cotton plants. Arsenic cannot be destroyed in the environment. It can only change its form. Arsenic in air will settle to the ground or is washed out of the air by rain. Many arsenic compounds can dissolve in water. Fish and shellfish can accumulate arsenic, but the arsenic in fish is mostly in a form that is not harmful. The toxicity of inorganic arsenic depends on its valence state and also on the physical and chemical properties of the compound in which it occurs.

Water soluble inorganic arsenic compounds are absorbed through the GI tract and lungs; distributed primarily to the liver, kidney, lung, spleen, aorta, and skin; and excreted mainly in the urine at rates as high as 80%. Symptoms of acute inorganic arsenic poisoning in humans are nausea, anorexia, vomiting, epigastric and abdominal pain, and diarrhea. Dermatitis (exfoliative erythroderma), muscle cramps,

cardiac abnormalities, hepatotoxicity, bone marrow suppression and hematologic abnormalities (anemia), vascular lesions, and peripheral neuropathy (motor dysfunction, paresthesia) also have been reported. Oral doses as low as 20-60 µg/kg/day have been reported to cause toxic effects in some individuals. Severe exposures can result in acute encephalopathy, congestive heart failure, stupor, convulsions, paralysis, coma, and death. The acute lethal dose to humans has been estimated to be about 0.6 mg/kg/day.

General symptoms of chronic arsenic poisoning in humans are weakness, general debility and lassitude, loss of appetite and energy, loss of hair, hoarseness of voice, loss of weight, and mental disorders. Primary target organs are the skin (hyperpigmentation and hyperkeratosis), nervous system (peripheral neuropathy), and vascular system. Anemia, leukopenia, hepatomegaly, and portal hypertension also have been reported. In addition, possible reproductive effects include a high male to female birth ratio.

Epidemiological studies have revealed an association between arsenic concentrations in drinking water and increased incidences of skin cancers, as well as cancers of the liver, bladder, respiratory, and GI tracts. Occupational exposure studies have shown a clear correlation between exposure to arsenic and lung cancer mortality. Several studies have shown that inorganic arsenic can increase the risk of lung cancer, skin cancer, bladder cancer, liver cancer, kidney cancer, and prostate cancer. The World Health Organization, the DHHS, and the EPA have determined that inorganic arsenic is a human carcinogen and is classified A, human carcinogen.

Cancer slope factors for arsenic are available from EPA's IRIS. The values used in the BHHRA are 1.50E+00, 1.51E+01, and 3.66E+00 [mg/(kg × day)]⁻¹ for the oral, inhalation, and dermal exposure routes, respectively. The slope factor for the dermal exposure route was calculated by assuming a GI absorption factor of 41% (DOE 2008).

Chronic RfDs for arsenic also are available in RAIS. The values used in the BHHRA were 3.00E-04 and 1.23E-04 mg/(kg × day) for the oral and dermal routes, respectively. The dermal RfD was calculated by assuming a GI absorption factor of 41%.

F.4.1.3 Manganese (CAS 007439-96-5) (RAIS)

Manganese is a silver-colored, naturally occurring metal that is found in many types of rocks and makes up about 0.10% of the earth's crust. Manganese is not found alone, but combines with other substances such as oxygen, sulfur, or chlorine. Manganese also can be combined with carbon to make organic manganese compounds, including pesticides (e.g., maneb or mancozeb) and methylcyclopentadienyl manganese tricarbonyl, a fuel additive in some gasolines. Manganese is an essential trace element and is necessary for good health. Normal nutritional requirements of manganese are satisfied through the diet, which is the normal source of the element, with minor contributions from water and air. The National Research Council recommends a dietary allowance of 2-5 mg/day for a safe and adequate intake of manganese for an adult human. Manganese can be found in several food items, including grains, cereals, and tea.

Manganese can elicit a variety of serious toxic responses upon prolonged exposure to elevated concentrations, either orally or by inhalation. The central nervous system is the primary target. Initial symptoms are headache, insomnia, disorientation, anxiety, lethargy, and memory loss. These symptoms progress with continued exposure and eventually include motor disturbances, tremors, and difficulty in walking, symptoms similar to those seen with Parkinsonism. These motor difficulties are often irreversible. Some individuals exposed to very high levels of manganese for long periods of time at work developed mental and emotional disturbances and slow and clumsy body movements. This combination of symptoms is a disease called "manganism."

There are no human cancer data available for manganese. Manganese has been placed in the EPA weight-of-evidence classification D: not classifiable as to human carcinogenicity. No slope factors, therefore, were used in this BHHRA.

The oral, inhalation, and dermal RfDs from RAIS used in the BHHRA were 4.60E-02, 1.43E-05, and 1.84E-03 mg/(kg × day), respectively. The inhalation RfD was calculated using the inhalation reference concentration (RfC). The dermal RfD was calculated using a GI absorption factor is 4% (DOE 2008).

F.4.1.4 Selenium (CAS 007782-49-2) (RAIS)

Selenium is a metal commonly found in rocks and soil; much of the selenium in rocks is combined with sulfide minerals or with silver, copper, lead, and nickel minerals. Selenium and oxygen combine to form several compounds. Selenium sulfide is a bright red-yellow powder used in anti-dandruff shampoo. Industrially produced hydrogen selenide is a colorless gas with a disagreeable odor. It is probably the only selenium compound that might pose a health concern in the workplace. Selenium dioxide is an industrially produced compound that dissolves in water to form selenious acid. Selenious acid can be found in gun bluing (a solution used to clean the metal parts of a gun). Selenium is an essential trace element important in many biochemical processes that take place in human cells. Recommended human dietary allowances for selenium for adults is about 40-70 µg.

In humans, acute oral exposures can result in excessive salivation, garlic odor to the breath, shallow breathing, diarrhea, pulmonary edema, and death. Other reported signs and symptoms of acute selenosis include tachycardia, nausea, vomiting, abdominal pain, abnormal liver function, muscle aches and pains, irritability, chills, and tremors. The exact levels at which these effects occur are not known. GI absorption in animals and humans of various selenium compounds ranges from about 44% to 95% of the ingested dose. If too much selenium is ingested over long periods of time, brittle hair and deformed nails can develop. Upon contact with skin, selenium compounds have caused rashes, swelling, and pain. Respiratory tract absorption rates of 97% and 94% for aerosols of selenious acid have been reported for dogs and rats, respectively. In humans, inhalation of selenium or selenium compounds primarily affects the respiratory system. Dusts of elemental selenium and selenium dioxide can cause irritation of the skin and mucous membranes of the nose and throat, coughing, nosebleed, loss of sense of smell, dyspnea, bronchial spasms, bronchitis, and chemical pneumonia.

Studies of laboratory animals and humans show that most selenium compounds probably do not cause cancer. In fact, human studies suggest that lower-than-normal selenium levels in the diet might increase the risk of cancer. Other forms of selenium may, however, be carcinogenic according to the DHHS. Selenium sulfide produced a significant increase in the incidence of lung and liver tumors in rats and mice. EPA has placed selenium and selenious acid in Group D, not classifiable as to carcinogenicity in humans, while selenium sulfide is placed in Group B2, probable human carcinogen. Selenium sulfide is very different from the selenium compounds found in foods and in the environment. Selenium sulfide has not caused cancer in animals when it is placed on the skin, and the use of anti-dandruff shampoos containing selenium sulfide is considered safe.

Chronic RfDs from RAIS were available for selenium. The RfDs used in the BHHRA for the oral and dermal routes of exposure were 5.00E-03 and 2.20E-03 mg/(kg × day), respectively. The dermal route RfD was based on a GI absorption factor of 44% (DOE 2008).

F.4.1.5 Uranium (metal and soluble salts) (CAS 007440-61-1)

Uranium is a hard, silvery white amphoteric metal and is a radioactive element. In its natural state it consists of three isotopes: ²³⁴U, ²³⁵U, and ²³⁸U. More than 100 uranium minerals exist; those of commercial importance are the oxides and oxygenous salts. The processing of uranium ore generally

involves extraction then leaching either by an acid or a carbonate method. In addition, the metal may be obtained from its halides by fused salt electrolysis. The primary use of natural uranium is in nuclear energy as a fuel for nuclear reactors, in plutonium production, and as feeds for gaseous diffusion plants; it is also a source of radium salts. Uranium compounds are used in staining glass, glazing ceramics, and enameling; in photographic processes; for alloying steels; and as a catalyst for chemical reactions, radiation shielding, and aircraft counterweights (Sittig 1985).

The primary route of exposure to uranium metals and salts is through dermal contact. Uranium soluble compounds act as a poison to cause kidney damage under acute exposure and pneumoconiosis or pronounced blood changes under chronic exposure conditions. Furthermore, it is difficult to separate the toxic chemical effects of uranium and its compounds from their radiation effects. The chronic radiation effects are similar to those produced by ionizing radiation. Reports now confirm that carcinogenicity is related to dose and exposure time. Cancer of the lung, osteosarcoma, and lymphoma have all been reported (Sittig 1985). An EPA weight-of-evidence classification for uranium metal was not located in the available literature. Slope factors for uranium metal also were not available for use in the BHHRA.

Chronic RfDs from the Risk Methods Document were available for uranium metal (listed as uranium soluble salts). The oral and dermal RfD used in the BHHRA were 6.00E-04 and 5.10E-04 mg/(kg × day), respectively. A GI absorption factor of 85% was used to derive the dermal RfD (DOE 2008).

F.4.2 ORGANIC COMPOUNDS

F.4.2.1 1,1-DCE (CAS 000075-35-4) (RAIS)

1,1-DCE, also known as 1,1-dichloroethene and vinylidene chloride, is a colorless liquid that is used in the production of polyvinylidene chloride (PVC) copolymers and as an intermediate for the synthesis of organic chemicals. The major application of PVC copolymers is the production of flexible films for food packaging.

1,1-DCE is not a naturally-occurring chemical, but is found in the environment due to releases associated with its production and transport and with its polymer production. Because of its high volatility, releases to the atmosphere are the greatest source of ambient 1,1-DCE. Smaller amounts are released to surface waters and soils. Loss of 1,1-DCE from water and soils is primarily due to volatilization. Human exposure to 1,1-DCE is potentially highest in workplace settings and in the vicinity of hazardous waste sites where the compound may contaminate environmental media.

The primary effect of acute exposure to high concentrations (approximately 4,000 ppm) of 1,1-DCE vapor in humans is central nervous system (CNS) depression, which may progress to unconsciousness. Occupational exposure has been reported to cause liver dysfunction in workers. 1,1-DCE is a skin irritant and prolonged contact can cause first degree burns. Direct contact with the eyes may cause conjunctivitis and transient corneal injury. Based on EPA guidelines, 1,1-DCE was assigned to weight-of-evidence group C, possible human carcinogen.

Slope factors for 1,1-DCE are available from RAIS. The values used in the BHHRA for oral, inhalation and dermal slope factors are 6.00E-01, 1.75E-01, and 6.00E-01 mg/(kg × day), respectively. The slope factor for the dermal route was calculated using a GI absorption factor of 100% (DOE 2008).

Chronic RfDs for 1,1-DCE are available from RAIS. The values used in the BHHRA for the oral, inhalation, and dermal routes were 5.00E-02, 5.71E-02, and 5.00E-02 mg/(kg × day), respectively. The inhalation RfD was extrapolated from the inhalation RfC. The dermal RfD was derived using a GI absorption factor of 100% (DOE 2008).

F.4.2.2 *Cis*-1,2-DCE (CAS 000156-59-) (RAIS)

1,2-DCE, also called 1,2-dichloroethene, is a highly flammable, colorless liquid with a sharp, harsh odor. It is used to produce solvents and in chemical mixtures. Very small amounts of 1,2-DCE may be smelled in air (about 17 ppm). There are two forms of 1,2-DCE: *cis*-1,2-DCE and *trans*-1,2-DCE. Sometimes both forms are present as a mixture. Commercial use is not extensive, but mixtures of *cis*- and *trans*-1,2-DCE have been used as intermediates in the production of other chlorinated solvents and compounds, as well as low temperature extraction solvents for dyes, perfumes, and lacquers. Additionally, *cis*- and *trans*-1,2-DCE react violently with potassium hydroxide, sodium, and sodium hydroxide and form shock-sensitive explosives when combined with dinitrogen tetroxide. Both forms of 1,2-DCE are degradation products of TCE.

Humans are exposed to 1,2-DCE primarily by inhalation, but exposure also can occur by oral and dermal routes. Breathing high levels of 1,2-DCE can cause nausea, drowsiness, and tiredness in humans; very high levels can cause death. Animals that ingested extremely high doses of *cis*-1,2-DCE died. Lower doses of *cis*-1,2-DCE caused effects on the blood, such as decreased numbers of red blood cells, and also on the liver.

No cancer bioassays or epidemiological studies were available to assess the carcinogenicity of 1,2-DCE. EPA has placed *cis* -1,2-DCE in weight-of-evidence group D, not classifiable as to human carcinogenicity, based on the lack of or negative human or animal cancer data. No cancer slope factors for *cis* -1,2-DCE are available; therefore, carcinogenicity from exposure could not be quantified in the BHHRA.

The oral, inhalation, and dermal chronic RfDs for *cis*-1,2-DCE used in the BHHRA are 1.00E-02, 9.97E-03, and 1.00E-02 mg/(kg × day), respectively. The inhalation RfD used in the BHHRA was extrapolated from the inhalation RfC. The dermal RfD was derived from the oral toxicity value using a GI absorption factor of 100% (DOE 2001).

F.4.2.3 Aroclor-1254 (CAS 011097-69-1)

Aroclor-1254 is a PCB mixture containing approximately 21% C₁₂H₆Cl₄, 48% C₁₂H₅Cl₅, 23% C₁₂H₄Cl₆, and 6% C₁₂H₃Cl₇, with an average chlorine content of 54%. PCBs are inert, thermally and physically stable, and have dielectric properties. In the environment, the behavior of PCB mixtures is directly correlated to the degree of chlorination. Aroclor is strongly sorbed to soil and remains immobile when leached with water; however, the mixture is highly mobile in the presence of organic solvents. PCBs are resistant to chemical degradation by oxidation or hydrolysis; however, biodegradation, especially of lower chlorinated PCBs, can occur. PCBs have high bioconcentration factors and due to their lipophilicity, especially of highly chlorinated congeners, tend to accumulate in the fat of fish, birds, mammals, and humans.

PCBs are absorbed after oral, inhalation, or dermal exposure and are stored in adipose tissue. The location of the chlorine atoms on the phenyl rings is an important factor in PCB metabolism and excretion. The major route of PCB excretion is in the urine and feces; however, more important is the elimination in human milk. Metabolites are predominately found in urine and bile, while small amounts of the parent compound are found in the feces. Biliary excretion appears to be the source of fecal excretion.

Accidental human poisonings and data from occupational exposure to PCBs suggest initial dermal and mucosal disturbances followed by systemic effects that may manifest themselves several years post-exposure. Initial effects are enlargement and hypersecretion of the Meibomian gland of the eye, swelling of the eyelids, pigmentation of the fingernails and mucous membranes, fatigue, and nausea.

These effects were followed by hyperkeratosis, darkening of the skin, acne form eruptions, edema of the arms and legs, neurological symptoms such as headache and limb numbness, and liver disturbance.

Hepatotoxicity is a prominent effect of Aroclor-1254 that has been well characterized. Effects include hepatic microsomal enzyme induction, increased serum levels of liver-related enzymes indicative of hapocellular damage, liver enlargement, lipid deposition, fibrosis, and necrosis.

Data are suggestive but not conclusive concerning the carcinogenicity of PCBs in humans. The EPA has not determined a weight-of-evidence classification or slope factor for Aroclor-1254 specifically. Hepatocellular carcinomas in rat and mice studies have led EPA to classify PCBs as group B2, probable human carcinogen.

Slope factors for Aroclor-1254 are available from RAIS. The values used in the BHHRA for oral, inhalation and dermal slope factors are 4.00E-01, 3.50E-01, and 4.44E-01 mg/(kg × day), respectively. The slope factor for the dermal route was calculated using a GI absorption factor of 90% (DOE 2008). The value of 4.00E-01 per (mg/kg/day) is the upper bound oral slope factor for the Low Risk and Persistence Tier. Criteria for using this value includes ingestion of water-soluble congeners, inhalation of evaporated congeners, and dermal exposure, if no absorption factor has been applied (DOE 2008).

Chronic RfDs for Aroclor-1254 are available from IRIS. The values used in the BHHRA for the oral, inhalation, and dermal routes were 2.00E-05, 1.99E-05, and 1.80E-05 mg/(kg × day), respectively. The inhalation RfD was extrapolated from the inhalation RfC. The dermal RfD was derived using a GI absorption factor of 90% (DOE 2008).

F.4.2.4 Aroclor-1260 (CAS 11096-82-5)

Aroclor-1260 is a PCB mixture containing approximately 38% C₁₂H₄Cl₆, 41% C₁₂H₃Cl₇, 8% C₁₂H₂Cl₈, and 12% C₁₂H₅Cl₅, with an average chlorine content of 60%. PCBs are inert, thermally and physically stable, and have dielectric properties. In the environment, the behavior of PCB mixtures is directly correlated to the degree of chlorination. Aroclor is strongly sorbed to soil and remains immobile when leached with water; however, the mixture is highly mobile in the presence of organic solvents. PCBs are resistant to chemical degradation by oxidation or hydrolysis; however, biodegradation, especially of lower chlorinated PCBs, can occur. PCBs have high bioconcentration factors and due to lipophilicity, especially of highly chlorinated congeners, tend to accumulate in the fat of fish, birds, mammals, and humans. The use of PCBs in the United States was limited to closed systems in 1974, and in February, 1977, the EPA issued final regulations prohibiting PCB discharge into waterways.

PCBs are absorbed after oral, inhalation, or dermal exposure and are stored in adipose tissue. The location of the chlorine atoms on the phenyl rings is an important factor in PCB metabolism and excretion. The major route of PCB excretion is in the urine and feces; however, of more importance is elimination in human milk. Metabolites are predominately found in urine and bile, while small amounts of parent compound are found in the feces.

Slope factors for Aroclor-1260 are available from IRIS. The values used in the BHHRA for oral, inhalation, and dermal slope factors are 4.00E-01, 3.50E-01, and 4.44E-01 mg/(kg × day), respectively. The slope factor for the dermal route was calculated using a GI absorption factor of 90% (DOE 2008). The value of 4.00E-01 per (mg/kg/day) is the upper-bound Oral Slope Factor for the Low Risk and Persistence Tier. The criteria for using this value includes ingestion of water-soluble congeners, inhalation of evaporated congeners, and dermal exposure, if no absorption factor has been applied (DOE 2008).

No RfD or RfC have been verified for Aroclor-1260. Data are suggestive but not conclusive concerning the carcinogenicity of PCBs in humans. Hepatocellular carcinomas in three strains of rats and two strains of mice have led the EPA to classify PCBs as group B2, probable human carcinogen.

Specific information on the chronic oral toxicity of Aroclor-1260 to humans is not available; however, information from accidental poisonings that occurred in Japan and Taiwan and from occupational exposure to PCBs is available. Oral Exposures Primary target organs are the liver and skin. Hepatotoxicity is a well-characterized effect of Aroclor-1260 and other PCBs. The spectrum of effects include hepatic microsomal enzyme induction, increased serum levels of liver-associated enzymes suggestive of possible liver damage, liver enlargement, lipid deposition, fibrosis and necrosis. Chloracne has been observed in humans and several animal species following PCB exposure.

F.4.2.5 Naphthalene (CAS 000091-20-3)

Naphthalene is a white solid that is found naturally in fossil fuels and that exhibits a typical mothball odor. Naphthalene is a polycyclic aromatic hydrocarbon composed of two fused benzene rings. Burning tobacco or wood produces naphthalene. It occurs in crude oil, from which it may be recovered directly as white flakes; it can also be isolated from cracked petroleum, coke-oven emissions, or from high-temperature carbonization of bituminous coal. The major products made from naphthalene are moth repellents. It is also used for making dyes, resins, leather, tanning agents, and the insecticide carbaryl.

Naphthalene can be absorbed by the oral, inhalation, and dermal routes of exposure and can cross the placenta in amounts sufficient to cause fetal toxicity. Exposure to large amounts of naphthalene may damage or destroy some red blood cells, causing a low level until the body replaces the destroyed cells. People, particularly children, have developed this problem after eating naphthalene-containing mothballs or deodorant blocks. Some of the symptoms of this problem are fatigue, lack of appetite, restlessness, and pale skin. Exposure to large amounts of naphthalene may also cause neurotoxic effects (confusion, lethargy, listlessness, vertigo), gastrointestinal distress, hepatic effects (jaundice, hepatomegaly, elevated serum enzyme levels), renal effects, and ocular effects (cataracts, optical atrophy). The estimated lethal dose of naphthalene is 5-15 g for adults and 2-3 g for children. Animals sometimes develop cloudiness in their eyes after swallowing naphthalene. It is not clear if this also develops in people. When mice were repeatedly exposed to naphthalene vapors for 2 years, their noses and lungs became inflamed and irritated.

Available cancer bioassays were insufficient to assess the carcinogenicity of naphthalene. Using EPA's 1996 Proposed Guidelines for Carcinogen Risk Assessment, the human carcinogenic potential of naphthalene via the oral or inhalation routes "cannot be determined" at this time based on human and animal data. There is suggestive evidence (observations of benign respiratory tumors and one carcinoma in female mice only exposed to naphthalene by inhalation) that naphthalene may cause cancer. Additional support includes increase in respiratory tumors associated with exposure to 1-methylnaphthalene.

Chronic RfDs for naphthalene are available from RAIS. The values used in the BHHRA for the oral, inhalation, and dermal routes were 2.00E-02, 8.57E-04, and 1.60E-02 mg/(kg × day). The inhalation RfD was extrapolated from the inhalation RfC. The dermal RfD was derived using a GI absorption factor of 80% (DOE 2008).

F.4.2.6 TCE (CAS 000079-01-6) (RAIS)

TCE, also known as trichloroethylene, is a colorless, highly volatile liquid that is miscible with water and a number of organic solvents. TCE is a man-made chemical and is not known to occur naturally. It is mainly used as a solvent in industrial degreasing and cleaning of metals, but it also is used as a solvent for waxes, fats, resins, oils, and in numerous other applications. Prior to 1977, TCE had been used as an

anesthetic, grain fumigant, disinfectant, and extractant of spice oleoresins in food and of caffeine in the production of decaffeinated coffee. The evaluation of the toxicity of TCE is complicated by the presence or absence of other chemicals. Industrial grade TCE usually contains stabilizers that are known to be toxic such as triethylamine, triethanolamine, epichlorohydrin, or stearates. In the absence of stabilizers, TCE readily decomposes. These decomposition products also are toxic.

Human and animal data indicate that exposure to TCE can result in toxic effects on a number of organs and systems, including the liver, kidney, blood, skin, immune system, reproductive system, nervous system, and cardiovascular system. Breathing small amounts of TCE may cause headaches, lung irritation, dizziness, poor coordination, and difficulty concentrating. Breathing large amounts of TCE may cause impaired heart function, unconsciousness, and death. Breathing it for long periods may cause nerve, kidney, and liver damage. Drinking large amounts of TCE may cause nausea, liver damage, unconsciousness, impaired heart function, or death. Drinking small amounts of TCE for long periods may cause liver and kidney damage, impaired immune system function, and impaired fetal development in pregnant women, although the extent of some of these effects is not yet clear. Skin contact with TCE for short periods may cause skin rashes.

Epidemiologic studies have been inadequate to determine if a correlation exists between exposure to TCE and increased cancer risk in humans. Some human studies with exposure over long periods to high levels of TCE in drinking water or in workplace air have found evidence of increased cancer; however, these results are inconclusive because the cancer could have been caused by other chemicals. Some studies with mice and rats have suggested that high levels of TCE may cause liver or lung cancer. Although EPA's Science Advisory Board recommended a weight-of-evidence classification of C-B2 continuum (C = possible human carcinogen; B2 = probable human carcinogen), the agency has not adopted a current position on the weight-of-evidence classification. In an earlier evaluation, TCE was assigned to weight-of-evidence Group B2, probable human carcinogen. The IARC has determined that TCE is not classifiable as to human carcinogenicity.

Cancer slope factors for TCE are available from RAIS. The slope factors from EPA for the oral, inhalation, and dermal exposure routes are 4.00E-01, 4.00E-01, and 2.67E+00 [mg/(kg × day)]⁻¹, respectively. The slope factor for the dermal exposure route was calculated using the EPA oral slope factor by assuming a GI absorption factor of 15% (DOE 2008). Cancer slope factors also are available from the review done by KDEP (KDEP 2004). The slope factor from KDEP for the oral exposure route is 3.22E-01 (mg/kg × day)⁻¹. Following guidance in the draft revised Risk Methods Document, the KDEP oral slope factor was used as the slope factor for both the oral and inhalation routes in this BHHRA. Uncertainties related to the selection of toxicity values among the 2001 Risk Methods Document, the EPA value, and the KDEP value are discussed in the uncertainty section.

Chronic RfDs for TCE are available from RAIS. The values used in the BHHRA for the oral, dermal, and inhalation routes were 3.00E-04, 4.50E-05, and 1.14E-02 mg/(kg × day). The inhalation RfD was extrapolated from the inhalation RfC. The dermal RfD was derived using a GI absorption factor of 15% (DOE 2008).

F.4.2.7 Vinyl Chloride (CAS 000075-01-4) (RAIS)

Vinyl chloride, also known as chloroethene, is a halogenated aliphatic hydrocarbon. It is a colorless gas with a mild sweetish odor that is slightly soluble in water and soluble in hydrocarbons, oil, alcohol, chlorinated solvents, and most common organic liquids. Vinyl chloride is produced by thermal cracking of ethylene chloride and does not occur naturally. It is used primarily as an intermediate in the manufacture of PVC; limited quantities are used as a refrigerant and as an intermediate in the production of chlorinated compounds. It is a biodegradation product of TCE, tetrachloroethylene, and 1,1,1-TCA. Vinyl chloride may leach into groundwater from spills, landfills, and industrial sources.

Vinyl chloride is rapidly absorbed from the digestive tract and lungs. Breathing high levels of vinyl chloride can cause dizziness or sleepiness. Breathing very high levels can cause passing out, and breathing extremely high levels can cause death. Humans exposed to vinyl chloride in air for long periods of time can develop changes to the structure of their livers. Workers exposed to vinyl chloride have developed nerve damage and immune reactions. Other workers have developed problems with the blood flow in their hands: the tips of their fingers turn white and hurt when they are in cold temperatures. Sometimes, the bones in the tips of their fingers have broken down. The effects of drinking high levels of vinyl chloride are unknown. If vinyl chloride is spilled on skin, numbness, redness, and blisters may occur. Animal studies have shown that long-term (365 days or longer) exposure to vinyl chloride can damage the sperm and testes. It has not been proven that vinyl chloride causes birth defects in humans, but animal studies have shown that breathing vinyl chloride can harm unborn offspring and also may cause increases in early miscarriages.

Studies show that vinyl chloride causes liver cancer in humans. On the basis of sufficient evidence for carcinogenicity in human epidemiology studies, vinyl chloride is considered to best fit the weight-of-evidence Category “A,” according to current EPA Risk Assessment Guidelines. Agents classified into this category are considered known human carcinogens. This classification is supported by positive evidence for carcinogenicity in animal bioassays including several species and strains, and strong evidence for genotoxicity.

Cancer slope factors for vinyl chloride are available from EPA’s IRIS. The slope factors used in the BHHRA for the oral, inhalation, and dermal exposure routes are and 1.50E+00, 3.08E-02, and 1.50E+00 [mg/(kg × day)]⁻¹, respectively. The slope factor for the dermal exposure route was calculated by assuming a GI absorption factor of 100%.

Chronic RfDs for vinyl chloride are available from RAIS. The values used in the BHHRA for the oral, inhalation, and dermal routes were 3.00E-03, 2.86E-02, and 3.00E-03 mg/(kg × day), respectively. The inhalation RfD was extrapolated from the inhalation RfC. The dermal RfD was derived using a GI absorption factor of 100% (DOE 2008).

F.4.3 RADIONUCLIDES

Radionuclides are unstable atoms of chemical elements that will emit charged particles or energy or both to achieve a more stable state. These charged particles are termed “alpha and beta radiation”; energy is termed “neutral gamma rays.” Interaction of these charged particles (and gamma rays) with matter will produce ionization events, or radiation, which may cause living cell tissue damage. Because the deposition of energy by ionizing radiation is a random process, sufficient energy may be deposited (in a critical volume) within a cell and result in cell modification or death. In addition, ionizing radiation has sufficient energy that interactions with matter will produce an ejected electron and a positively charged ion (known as free radicals) that are highly reactive and may combine with other elements, or compounds within a cell, to produce toxins or otherwise disrupt the overall chemical balance of the cell. These free radicals also can react with deoxyribonucleic acid (DNA), causing genetic damage, cancer induction, or even cell death.

Radionuclides are characterized by the type and energy level of the radiation emitted. Radiation emissions fall into two major categories: particulate (electrons, alpha particles, beta particles, and protons) or electromagnetic radiation (gamma and x-rays). Therefore, all radionuclides are classified by the EPA as Group A carcinogens based on their property of emitting ionizing radiation and on the extensive weight of evidence provided by epidemiological studies of humans with cancers induced by high doses of radiation. Alpha particles are emitted at a characteristic energy level for differing radionuclides. The alpha particle has a charge of +2 and a comparably large size. Alpha particles have the ability to react (and/or ionize)

with other molecules, but they have very little penetrating power and lack the ability to pass through a piece of paper or human skin. However, alpha-emitting radionuclides are of concern when there is a potential for inhalation or ingestion of the radionuclide. Alpha particles are directly ionizing and deposit their energy in dense concentrations [termed high linear energy transfer (LET)], resulting in short paths of highly localized ionization reactions. The probability of cell damage increases as a result of the increase in ionization events occurring in smaller areas; this also may be the reason for increased cancer incidence caused by inhalation of radon gas. In addition, the cancer incidence in smokers may be directly attributed to the naturally occurring alpha emitter, polonium-210, in common tobacco products.

Beta emissions generally refer to beta negative particle emissions. Radionuclides with an excess of neutrons achieve stability by beta decay. Beta radiation, like alpha radiation, is directly ionizing but, unlike alpha activity, beta particles deposit their energy along a longer track length (low LET), resulting in more space between ionization events. Beta-emitting radionuclides can cause injury to the skin and superficial body tissue, but are most destructive when inhaled or ingested. Many beta emitters are similar chemically to naturally occurring essential nutrients and will, therefore, tend to accumulate in certain specific tissues. For example, strontium-90 is chemically similar to calcium and, as a result, accumulates in the bones, where it causes continuous exposure. The health effects of beta particle emissions depend upon the target organ. Those seeking the bones would cause a prolonged exposure to the bone marrow and affect blood cell formation, possibly resulting in leukemia, other blood disorders, or bone cancers. Those seeking the liver would result in liver diseases or cancer, while those seeking the thyroid would cause thyroid and metabolic disorders. In addition, beta radiation may lead to damage of genetic material (DNA), causing hereditary defects.

Gamma emissions are the energy that has been released from transformations of the atomic nucleus. Gamma emitters and x-rays behave similarly, but differ in their origin: gamma emissions originate in nuclear transformations, and x-rays result from changes in the orbiting electron structure. Radionuclides that emit gamma radiation can induce internal and external effects. Gamma rays have high penetrating ability in living tissue and are capable of reaching all internal body organs. Without such sufficient shielding as lead, concrete, or steel, gamma radiation can penetrate the body from the outside and does not require ingestion or inhalation to penetrate sensitive organs. Gamma rays are characterized as low-LET radiation, as is beta radiation; however, the behavior of beta radiation differs from that of gamma radiation in that beta particles deposit most of their energy in the medium through which they pass, while gamma rays often escape the medium because of higher energies, thereby creating difficulties in determining actual internal exposure. For this reason, direct whole-body measurements are necessary to detect gamma radiation, while urine/fecal analyses are usually effective in detecting beta radiation.

People receive gamma radiation continuously from naturally occurring radioactive decay processes going on in the earth's surface, from radiation naturally occurring inside their bodies, from the atmosphere as fallout from nuclear testing or explosions, and from space or cosmic sources. Cesium-137 (from nuclear fallout) decays to barium-137, the highest contributor to fallout-induced gamma radiation. Beta radiation from the soil is a less penetrating form of radiation, but has many contributing sources. Potassium-40, cesium-137, lead-214, and bismuth-214 are among the most common environmental beta emitters. Tritium is also a beta emitter but contributes little to the soil beta radiation because of the low energy of its emission and its low concentration in the atmosphere. Alpha radiation also is emitted by the soil, but is not measurable more than a few centimeters from the ground surface. The majority of alpha emissions are attributable to radon-222 and radon-220 and their decay products. This contributes to what is called background exposure to radiation.

The general health effects of radiation can be divided into stochastic (related to dose) and nonstochastic (not related to dose) effects. The risk of development of cancer from exposure to radiation is a stochastic effect. Examples of nonstochastic effects include acute radiation syndrome and cataract formation, which occur only at high levels of exposures.

Radiation can damage cells in different ways. It can cause damage to DNA within the cell, and the cell either may not be able to recover from this type of damage or may survive but function abnormally. If an abnormally functioning cell divides and reproduces, a tumor or mutation in the tissue may develop. The rapidly dividing cells that line the intestines and stomach and the blood cells in bone marrow are extremely sensitive to this damage. Organ damage results from the damage caused to the individual cells. This type of damage has been reported with doses of 10 to 500 rads (0.1 to 5.0 gray, in SI units). Acute radiation sickness is seen only after doses of >50 rads (0.5 gray), which is a dose rate usually achieved only in a nuclear accident.

When the radiation-damaged cells are reproductive cells, genetic damage can occur in the offspring of the person exposed. The developing fetus is especially sensitive to radiation. The type of malformation that may occur is related to the stage of fetal development and the cells that are differentiating at the time of exposure. Radiation damage to children exposed in the womb is related to the dose the pregnant mother receives. Mental retardation is a possible effect of fetal radiation exposure.

The most widely studied population that has had known exposure to radiation is the atomic bomb survivors of Hiroshima and Nagasaki, Japan. Data indicate an increase in the rate of leukemia and cancers in this population. However, the rate at which cancer incidence is significantly affected by low radiation exposures, such as results of exposure to natural background and industrially contaminated sites, is still undergoing study and is uncertain. In studies conducted to determine the rate of cancer and leukemia increase, as well as genetic defects, several radionuclides must be considered.

F.4.3.1 Technetium-99 (CAS 014133-76-7) (EPA)

Technetium is a radioactive element that occurs in a number of isotopic forms. Technetium is found in some extraterrestrial material (i.e., stars); however, no appreciable amounts have been found in nature due to the relatively short half-lives of its radioactive isotopes (Kutegov *et al.* 1968). While no isotopes of technetium are stable, the existence of three technetium isotopes is well established. Two common forms of technetium, ⁹⁷Tc and ⁹⁸Tc, have half-lives of 2.6×10^6 and 1.5×10^6 years, respectively. The third isotope, ⁹⁹Tc, has a half-life of 2.12×10^5 years. None, however, possesses a half-life sufficiently long to allow technetium to occur naturally (Boyd 1959). Technetium is made artificially for industrial use, and natural technetium, particularly ⁹⁹Tc, has been identified and isolated from the spontaneous fission of uranium, as well as other fissionable material or via the irradiation of molybdenum (Venugopal and Luckey 1978; Clarke and Podbielski 1988).

Technetium is an emitter of beta particles of low specific activity (Boyd 1959). It does not release nuclear energy at a rate sufficient to make the element attractive for the conventional applications of radioactivity (Boyd 1959). ⁹⁹Tc is the only long-lived isotope that is readily available and is the isotope on which most of the chemistry of technetium is based. Although gamma radiation has not been associated with ⁹⁹Tc, the secondary X rays may become important with larger amounts of the element.

Oral and inhalation cancer slope factors used in the BHHRA for ⁹⁹Tc are 2.75E-12 and 1.41E-11 risk/pCi, respectively. Dermal and external exposure cancer slope factors were not calculated because these routes of exposure are not evaluated for groundwater in the BHHRA. Oral, dermal, and inhalation RfDs are not available for this element; therefore, systemic toxicity due to exposure to ⁹⁹Tc is not quantified in the BHHRA.

F.4.3.2 Uranium (CAS 007440-62-2 for Metal, CAS 013966-29-5 for Uranium-234, and CAS 007440-61-1 for Uranium-238) (ATSDR)

Uranium is a mildly radioactive element that occurs widely in the earth's crust. It is found in all soils, most rocks, and, in lesser concentrations, in water, vegetation, and animals, including humans. Uranium

emits a low level of alpha particles and a much lower level of gamma rays. Alpha particles are unable to penetrate skin, but can travel short distances in the body if ingested or inhaled. Consequently, uranium represents a significant carcinogenic hazard only when taken into the body, where alpha particle energy is absorbed by small volumes of tissue. Although the penetrating (gamma) radiation of uranium is not considered to be significant (ATSDR 1989), one of its daughter radionuclides is a strong gamma emitter; therefore, gamma radiation may be a concern in areas containing uranium.

Natural uranium contains the uranium isotopes ^{238}U (which averages 99.27% of total uranium mass), ^{235}U (0.725), and ^{234}U (0.0056%), each of which undergoes radioactive decay. Natural uranium, therefore, contains the radionuclide daughter products from the decay of ^{238}U and ^{235}U (Bowen 1979; ATSDR 1989). The half-lives of the isotopes are 200,000, 700 million, and 5 billion years for ^{234}U , ^{235}U , and ^{238}U , respectively.

Uranium is a radioactive element, but it also is a metallic element. Toxicological effects from the ingestion of uranium are the result of the action of uranium as a metal and its radioactive properties. The primary toxic chemical effect of uranium is seen in kidney damage. Studies in rabbits, mice, and dogs showed effects on the kidney to be dose-related. Fetal skeletal abnormalities and fetal death were found in pregnant mice exposed to 6 mg/kg or uranyl acetate dihydrate.

The primary human exposure studies to uranium have been studies of uranium miners or uranium factory workers. These studies have shown an increase in lung cancer deaths among these workers, which may be attributable to the decay of uranium into radon and its daughters. These workers are exposed to high levels of uranium dust and fumes and other radioactive elements in confined conditions (ATSDR 1989).

Oral cancer slope factors used in the BHHRA for ^{234}U are 7.07E-11 risk/pCi. Oral cancer slope factors used in the BHHRA for ^{238}U are 8.71E-11 risk/pCi. The slope factors for ^{238}U include ingrowth of short-lived degradation products. Inhalation slope factors for ^{234}U and ^{238}U are 1.14E-08 risk/pCi and 9.35E-09 risk/pCi, respectively. Dermal and external exposure cancer slope factors were not calculated for the uranium isotopes because these routes of exposure are not considered significant for radionuclides in groundwater and are not evaluated in the BHHRA. Oral, dermal, and inhalation RfDs are available for uranium and are listed earlier in this section.

F.4.4 CHEMICALS FOR WHICH NO EPA TOXICITY VALUES ARE AVAILABLE

Over all COPCs identified for RGA groundwater associated with the BGOU, oral RfD values exist for all of the inorganic chemical COPCs. Oral RfDs exist for all of the organic COPCs included.

All the inorganic chemical COPCs, except manganese lack inhalation RfD values. Absorbed dose RfD values exist for all of the inorganic and organic chemical COPCs included in the BHHRA.

Arsenic is the only inorganic chemical COPC with an oral slope factor. The organic compound COPCs without an oral slope factor are *cis*-1,2-DCE and naphthalene.

EPA-approved inhalation slope factors are available for only a few of the COPCs. The only inorganic chemical COPC with an inhalation slope factor is arsenic. Most organic compound COPCs have an approved inhalation slope factor. Those without an inhalation slope factor are *cis*-1,2-DCE and naphthalene.

COPCs with absorbed dose slope factors mirror those with oral slope factors. The COPCs without absorbed dose slope factors are all of the inorganic chemicals, except arsenic, and 1,1-DCE, *cis*-1,2-DCE, and naphthalene. All radionuclide COPCs have both oral and inhalation slope factors.

F.4.5 UNCERTAINTIES RELATED TO TOXICITY INFORMATION

Standard EPA RfDs and slope factors were used to estimate potential noncarcinogenic and carcinogenic health effects from exposure to detected chemical contaminants. Considerable uncertainty is associated with the methodology applied to derive slope factors and RfDs. EPA working groups review all relevant human and animal studies for each compound and select the studies pertinent to the derivation of the specific RfD and slope factor. These studies often involve data from experimental studies in animals, high exposure levels, and exposures under acute or occupational conditions. Extrapolation of these data to humans under low-dose, chronic conditions introduces uncertainties. The magnitude of these uncertainties is addressed by applying uncertainty factors to the dose response data for each applicable uncertainty. These factors are incorporated to provide a margin of safety for use in human health assessments. For TCE, there is currently no IRIS slope factor, but several draft slope factors are available. The oral slope factor from the EPA draft reassessment is $4.00\text{E-}01 \text{ (mg/kg} \times \text{day)}^{-1}$ and the KDEP oral slope factor is $3.22\text{E-}01 \text{ (mg/kg} \times \text{day)}^{-1}$. These slope factors are significantly higher than the ones used in previous BHHRA for PGDP. The KDEP oral slope factor was used in this BHHRA, but neither that value nor the EPA one has received final approval.

The dose-response relationship between cancer and ionizing radiation has been evaluated in many reports. Risk factors are extrapolated from the cancer risk established using the Japanese Atomic Bomb Survivors database and a relative risk projection model. EPA's methodology for estimating radionuclide carcinogenic risks currently is being reevaluated.

F.4.6 SUMMARY OF TOXICITY ASSESSMENT

A breakdown of the groundwater COPCs and their available toxicity information by SWMU is provided in the following subsections. These COCs relate to the soil samples that were used for determining the concentrations used for groundwater modeling.

F.4.6.1 SWMU 2 COPC Toxicity Summary

Ten COPCs were retained in groundwater for SWMU 2. Three are inorganic chemicals, all of which have toxicity information; four are organic compounds, all of which have toxicity information; and three are radionuclides, all of which have toxicity information.

F.4.6.2 SWMU 3 COPC Toxicity Summary

Five COPCs were retained in groundwater for SWMU 3. Three are inorganic chemicals, all of which have toxicity information; and two are radionuclides, both of which have toxicity information.

F.4.6.3 SWMU 4 COPC Toxicity Summary

Six COPCs were retained in groundwater for SWMU 4. Two are inorganic chemicals, both of which have toxicity information; three are organic compounds, for which the individual compounds all have toxicity information; and one is a radionuclide that has toxicity information.

F.4.6.4 SWMU 5 COPC Toxicity Summary

Four COPCs were retained in groundwater for SWMU 5. Two are inorganic chemicals, both of which have toxicity information; one is an organic compound, that has toxicity information; and one is a radionuclide, that has toxicity information.

F.4.6.5 SWMU 6 COPC Toxicity Summary

One COPC, manganese, was retained in groundwater for SWMU 6.

F.4.6.6 SWMU 7 COPC Toxicity Summary

Eleven COPCs were retained in groundwater for SWMU 7. Three are inorganic chemicals, all of which have toxicity information; five are organic compounds, for which the individual components all have toxicity information; and three are radionuclides, all of which have toxicity information.

F.4.6.7 SWMU 30 COPC Toxicity Summary

Nine COPCs were retained in groundwater for SWMU 30. Four are inorganic chemicals, all of which have toxicity information; two are organic compounds, for which the individual components all have toxicity information; and three are radionuclides, all of which have toxicity information.

F.4.6.8 SWMU 145 COPC Toxicity Summary

Six COPCs were retained in groundwater for SWMU 145. Three are inorganic chemicals, one is an organic chemical, that has toxicity information; and two are radionuclides, both of which have toxicity information.

F.5. RISK CHARACTERIZATION

Risk characterization is the final step in the risk assessment process. In this step, the information from the exposure and toxicity assessments is integrated to quantitatively estimate both carcinogenic health risks and noncarcinogenic hazard potential. For this assessment, risk is defined as both the lifetime probability of excess cancer incidence for carcinogens and the estimate of daily intake exceeding intake that may lead to toxic effects for noncarcinogens.

F.5.1 DETERMINATION OF POTENTIAL FOR NONCANCER EFFECTS

In this BHHRA, the numeric estimate of the potential for noncancer effects posed by a single chemical within one pathway of exposure is derived as the ratio of the CDI of a chemical, from a single pathway to the appropriate RfD. This ratio also is referred to as a HQ. This value is calculated as shown in the following equation:

$$HQ = \frac{CDI}{RfD}$$

Where:

HQ is the hazard quotient, dimensionless

CDI is the chronic daily intake of a particular chemical, mg/(kg × day)

RfD is the chronic reference dose for a particular chemical and pathway, mg/(kg × day)

When performing this calculation, the proper RfD was used for each CDI. For CDIs that reflect ingestion, the RfD used was that for administered dose. For CDIs that reflect absorption, as in dermal contact, the RfD used was that for absorbed dose. Finally, for CDIs that reflect inhalation exposure, the RfD used was that for inhalation. Similarly, the RfD that was appropriate for the duration of exposure was used. For all adult exposures, the period of exposure was greater than 7 years; therefore, the chronic RfD was used. For all exposures to children, regardless of duration, the chronic RfD was used (Risk Methods Document).

If several chemicals may reach a receptor through a common pathway, guidance (RAGS, Risk Methods Document) recommends adding the HQs of all chemicals reaching the receptor through the common pathway to calculate a pathway HI. This can be represented by the following equation:

$$\text{Pathway HI} = HQ_1 + HQ_2 + HQ_3 + \dots + HQ_n$$

Where:

Pathway HI is the sum of the individual chemical HQs, dimensionless

HQ₁ to HQ_n are the individual chemical hazard quotients relevant to the pathway, dimensionless

Similarly, guidance (Risk Methods Document) recommends summing the pathway HIs for all pathways relevant to an individual receptor to develop a total HI. The total HI is not an estimate of the systemic toxicity posed by all contaminants that may reach the receptor, but can be used to estimate if a toxic effect may result if all contaminants reaching the receptor have additive effects over all pathways. This can be represented as in the following equation:

$$\text{Total HI} = HI_1 + HI_2 + HI_3 + \dots + HI_n$$

Where:

Total HI is the sum of all pathways relevant to a single receptor, dimensionless
HI₁ to HI_n are the individual pathway HIs

Note that the HQ, the pathway HI, and the total HI do not define a dose-response relationship. That is, the magnitude of the HQ or HI does not represent a statistical probability of incurring an adverse effect. If the HQ is less than 1, the estimated exposure to a substance may be judged to be below a level that could present a toxic effect. If the HQ is greater than 1, a toxic effect may or may not result depending on the assumptions used to develop the CDI and assumptions used in deriving the RfD. Similarly, if the pathway HI is less than 1, then the estimated exposure to multiple chemicals contributing to the pathway HI should not be expected to present a toxic effect. If the pathway HI is greater than 1, then exposure may or may not result in a toxic effect depending on what assumptions were used to develop the pathway and how the chemicals included in the pathway interact. Finally, if the total HI is less than 1, then the estimated exposure to multiple chemicals over multiple pathways should not be expected to result in a toxic effect. If the total HI is greater than 1, then a toxic effect may or may not result depending on the rigor used to develop the CSM for all pathways and the interaction between pathways and individual chemicals.

F.5.2 DETERMINATION OF EXCESS LIFETIME CANCER RISK

Estimates of the potential for cancer induction are measured by calculating estimates of ELCR. Generally, ELCR can be defined as the incremental increase in the probability that a receptor may develop cancer if the receptor is exposed to chemicals or radionuclides or both. ELCRs are specific to the CSM used to define the routes and magnitude of exposure. The magnitude of the ELCRs could vary markedly if the exposure assumptions used to develop the CSM are varied.

F.5.2.1 Chemical Excess Cancer Risk

The numeric estimate of the ELCR resulting from exposure to a single chemical carcinogen is derived by multiplying the CDI through a particular pathway by the slope factor appropriate to that pathway. The resulting value is referred to as a chemical-specific ELCR. This value is calculated as shown in the following equation:

$$\text{Chemical specific ELCR} = \text{CDI} \times \text{SF}$$

Where:

Chemical specific ELCR is an estimate of the excess lifetime probability of developing cancer that results because of exposure to the specific chemical, dimensionless

CDI is the chronic daily intake of the chemical [mg/(kg × day)]

SF is the slope factor for the specific chemical [(mg/(kg × day))⁻¹]

As with the calculation used to derive HQs, the proper slope factor was used for each CDI when performing this calculation. For CDIs that reflect ingestion, the slope factor was that for an administered dose. For CDIs that reflect absorption, the slope factor was that for absorbed dose. Finally, for CDIs that reflect inhalation exposure, the slope factor was that for inhalation.

If several chemicals may reach a receptor through a common pathway, the chemical specific ELCRs of all chemicals reaching the receptor through the common pathway are summed to calculate a pathway ELCR. This can be represented by the following equation:

$$\text{Pathway ELCR} = \text{ELCR}_1 + \text{ELCR}_2 + \text{ELCR}_3 + \dots + \text{ELCR}_n$$

Where:

Pathway ELCR is the sum of the chemical-specific ELCRs, dimensionless
ELCR₁ to ELCR_n are the chemical-specific ELCRs relevant to the pathway; dimensionless

Similarly, the pathway ELCRs for all pathways relevant to an individual receptor are summed to develop a total ELCR. The total ELCR is not an actuarial estimate of an individual developing cancer, but can be used to estimate the total ELCR that may result if all contaminants reaching the receptor have additive effects over all pathways. This can be represented as in the following equation:

$$\text{Total ELCR} = \text{ELCR}_{p1} + \text{ELCR}_{p2} + \text{ELCR}_{p3} + \dots + \text{ELCR}_{pn}$$

Where:

Total ELCR is the sum of all pathways relevant to a single receptor, dimensionless
ELCR_{p1} to ELCR_{p2} is the individual pathway ELCRs

Unlike the HQ, the pathway HI, and the total HI; the chemical-specific ELCR, the pathway ELCR, and total ELCR define a dose-response relationship. That is, the ELCRs represent a statistical probability of the increased risk of developing cancer that exists in receptors exposed under the assumptions used in the calculation of the CDI.

F.5.2.2 Radionuclide Excess Cancer Risk

Calculation of cancer risk due to exposure to radionuclides through ingestion or inhalation is conceptually similar to calculation of risks for chemical carcinogens. In performing this calculation, ELCR due to exposure to a particular radionuclide within a specific pathway is calculated by multiplying the intake of the radionuclide by the route-specific cancer slope factor. This can be represented by the following equation:

$$\text{Radionuclide - specific ELCR} = \text{CDI} \times \text{SF}$$

Where:

Radionuclide specific ELCR is an estimate of the excess lifetime probability of developing cancer that results because of exposure to the specific radionuclide, dimensionless
CDI is the ingestion and inhalation chronic daily intake of the radionuclide, pCi
SF is the ingestion and inhalation slope factor for the specific radionuclide, risk/pCi
(Note: For external exposure, the units for CDI and SF are pCi-year/g and risk-g/pCi-year, respectively.)

As with the calculation used to derive chemical-specific ELCRs, the proper slope factor was used for each CDI when performing this calculation. For CDIs that reflect ingestion, the slope factor was that for ingestion. Similarly, for CDIs that reflect inhalation exposure, the slope factor was that for inhalation.

Both the pathway ELCR for radionuclides and the total ELCR from exposure to multiple radionuclides within a pathway and over multiple pathways, respectively, are calculated as illustrated for chemical carcinogens in Subsection F.5.2.1. These equations will not be presented in this risk assessment. The uncertainties related to this method of determining ELCR from exposure to radionuclides is discussed in detail in Section F.6.

In this risk assessment, ELCRs from exposure to chemicals and radionuclides were summed within pathways and over all pathways to indicate the potential health risk to a receptor that may be exposed to radionuclides and chemicals over all pathways. The uncertainties associated with combining radionuclide and chemical ELCRs are discussed in detail in Section F.6.

F.5.3 RISK CHARACTERIZATION FOR SOIL

For the risk characterization for soil, the results of the previous risk assessments described in Section F.1 are used. For most of the SWMUs, no new surface data have been collected since the previous risk assessments were performed. The soils at these units are outside the scope of the BGOU as noted in the approved work plan; therefore, a new quantitative risk assessment was not performed for soils. Vapor intrusion modeling was performed to examine potential risks from vapors from soil contaminants, intruding into basements. The percent contribution listed for each soil COC is listed with the same accuracy (i.e., 2% or 2.0%) as is in the original document from which the value was taken.

F.5.3.1 Systemic Toxicity and Excess Lifetime Cancer Risk (Direct Exposure to Soil)

Results of previous risk assessments are available for SWMUs 2, 3, 4, 5, 6, 7, and 30 (DOE 1994; DOE 1998b; DOE 2000). The results for systemic toxicity (HI) and ELCR for soil exposure are discussed in Section F.1 and presented in Attachment F2 of this appendix.

F.5.3.2 Vapor Intrusion into Basements from Soil

Exposure of on-site residents to vapors from soil contaminants intruding into basements is a potential pathway. To examine potential risks and hazards, vapor intrusion modeling was completed and examined for three POEs: the property boundary, the plant boundary, and a future on-site resident. The HQs and ELCRs for the modeled vapor concentrations are presented in Table E.3.35 of Appendix E. Modeled concentrations for the on-site POE showed an HQ greater than 0.1 for vapor intrusion from TCE, *cis*-1,2-DCE, 1,1-DCE, vinyl chloride, or mercury intrusion for the following:

- SWMU 2: TCE, and *cis*-1,2-DCE
- SWMU 3: mercury
- SWMU 4: TCE, *cis*-1,2-DCE, and vinyl chloride
- SWMU 7: 1,1-DCE, mercury, and vinyl chloride
- SWMU 30: mercury, 1,1-DCE, and TCE
- SWMU 145: mercury

ELCRs for the on-site POE were greater than 1E-06 for several SWMUs based on modeled contaminant concentrations. The following summarizes those SWMUs exhibiting elevated risks based on modeled soil concentrations.

- SWMUs 2: TCE
- SWMU 3: TCE
- SWMU 4: TCE and vinyl chloride
- SWMU 7: TCE, vinyl chloride, and 1,1-DCE
- SWMU 30: TCE and 1,1-DCE

Vapor intrusion into basements also was modeled at the plant boundary and property boundary. At the plant boundary all HIs were below 0.1. ELCRs were below 1E-06 for all SWMUs except for SWMUs 2, 4, 7, and 30. The following lists the risk driver for each SWMU:

- SWMU 2: TCE
- SWMU 4: TCE and vinyl chloride
- SWMU 7: 1,1-DCE
- SWMU 30: TCE and 1,1-DCE

At the property boundary all HIs were below 0.1. The ELCR for TCE exceeded *de minimis* risk levels at the property boundary for TCE at SWMUs 2, 4, and 30. All other risks/hazards were below *de minimis* levels at the property boundary. The quantitative assessment of potential risks and hazards due to exposure to vapor intrusion is summarized in Table 5.14.

F.5.3.3 Vapor Intrusion for the Future Industrial Worker

Vapor intrusion for the future industrial worker was examined at the SWMU boundary. The HI exceeded 0.1 at SWMU 2 due primarily to *cis*-1,2 DCE. Also, the HI was greater than 0.1 at SWMU 30 due primarily to TCE. Risks exceeded 1E-06 at SWMU 2 due to TCE, SWMU 30 due to TCE, and SWMU 7 due to 1,1-DCE. The following tables (Tables F.34-F.40) show the chemicals and associated HQ and risk with an HI and total risk by SWMU.

Table F.34. HQ and Total Risk for SWMU 2

COPC	HQ	% Contribution to HQ	Risk	% Contribution to Risk
Trichloroethene	4.82E-01	11.2%	6.33E-04	100.0%
<i>cis</i> -1,2 dichloroethene	3.83E+00	89.1%	NA	
Naphthalene	6.17E-05	0.0%	NA	
Total	4.3		6.33E-04	

Table F.35. HQ and Total Risk for SWMU 3

COPC	HQ	% Contribution to HQ	Risk	% Contribution to Risk
Trichloroethene	2.79E-04	100.0%	3.66E-07	100.0%
Total	2.79E-04		3.66E-07	

Table F.36. HQ and Total Risk for SWMU 5

COPC	HQ	% Contribution to HQ	Risk	% Contribution to Risk
Trichloroethene	9.29E-05	9.7%	3.65E-07	100.0%
Acenaphthene	6.69E-07	0.1%	NA	
Fluorene	2.52E-07	0.0%	NA	
Naphthalene	8.68E-04	90.2%	NA	
Pyrene	1.49E-08	0.0%	NA	
Total	9.62E-04		3.65E-07	

Table F.37. HQ and Total Risk for SWMU 6

COPC	HQ	% Contribution to HQ	Risk	% Contribution to Risk
Trichloroethene	1.60E-04	100.0%	2.10E-07	100.0%
Total	1.60E-04		2.10E-07	

Table F.38. HQ and Total Risk for SWMU 7

COPC	HQ	% Contribution to HQ	Risk	% Contribution to Risk
Trichloroethene	1.48E-03	1.7%	5.83E-06	1.3%
<i>cis</i> -1,2 dichloroethene	4.18E-03	4.7%	NA	
vinyl chloride	8.39E-02	93.7%	7.92E-05	17.1%
1,1-DCE	3.52E-02	39.3%	3.77E-04	81.6%
Mercury	2.30E-02	25.7%	NA	
Pyrene	5.01E-08	0.0%	NA	
Tetrachloroethene	2.29E-05	0.0%	8.38E-09	0.0%
Total	8.95E-02		4.62E-04	

Table F.39. HQ and Total Risk for SWMU 30

COPC	HQ	% Contribution to HQ	Risk	% Contribution to Risk
Trichloroethene	1.21E-03	0.1%	1.52E-06	78.7%
1,1-DCE	1.20E-04	0.0%	4.11E-07	21.3%
Acenaphthene	9.45E-08	0.0%	NA	
Fluorene	2.00E-08	0.0%	NA	
Mercury	3.94E-02	2.8%	NA	
Naphthalene	7.37E-05	0.0%	NA	
Pyrene	4.46E-09	0.0%	NA	
	1.4		1.93E-06	

Table F.40. HQ and Total Risk for SWMU 145

COPC	HQ	% Contribution to HQ	Risk	% Contribution to Risk
Mercury	3.25E-02	100.0%	NA	
Total	3.25E-02			

F.5.4 RISK CHARACTERIZATION FOR RESIDENTIAL USE OF GROUNDWATER DRAWN FROM THE RGA

This subsection presents the risk for residential use of groundwater drawn from the RGA. Tables and discussion in this subsection provide the total HI or ELCR for the each source area and list the major exposure routes and COPCs contributing to the total HI or ELCR. Environmental data for each source area was used to model groundwater concentrations at the POEs (see Section 5 and Appendix E for details of the groundwater modeling). The EPCs for groundwater are equal to the maximum concentration over the 1,000 year period over which groundwater was modeled. The groundwater assessment is conducted only for the residential scenario, but was conducted for all SWMUs including SWMU 145. Characterization of risks from groundwater at off-site POEs (plant boundary, property boundary, and Ohio River) are discussed in Section F.5.5.

F.5.4.1 Systemic Toxicity (Groundwater Use)

Tables F.41 through F.56 summarize the HIs for the modeled groundwater concentrations at each SWMU for the child and adult resident. As shown in these tables, the total scenario HIs are greater than 1 for the all of the SWMUs except SWMU 6 for both the child and adult resident. The source with the greatest HI for the child receptor is SWMU 2, which has a HI=1300, with the major contribution coming from ingestion of water containing TCE (52.1%) and *cis*-1,2-DCE (46.8%). The source with the greatest HI for the adult is SWMU 2, which has a HI=379, with the major contribution coming from ingestion of water (45%) and household inhalation (27.5%). The major contributors are TCE (62.1%) and *cis*-1,2-DCE (36.8%).

Table F.41. HI Child Residential Groundwater Use at SWMU 2

	Exposure Point Concentration ^a	Ingestion of Water	Dermal Contact	Inhalation while showering	Household Inhalation	Total Hazard ^b	Percent of Total Hazard
<u>Inorganic Compounds</u>							
Arsenic	3.54E-02	1.13E+01	2.39E-02			1.13E+01	0.9%
Manganese	7.16E-01	1.49E+00	3.23E-02			1.52E+00	0.1%
Uranium	9.86E-03	1.58E+00	1.68E-03			1.58E+00	0.1%
<u>Organic Compounds</u>							
<i>cis</i> -1,2-DCE	1.15E+01	1.10E+02	3.89E+00	5.59E+01	4.37E+02	6.07E+02	46.8%
Naphthalene	9.38E-04	4.50E-03	1.02E-03	5.31E-02	4.15E-01	4.74E-01	0.0%
TCE	1.48E+00	4.73E+02	1.48E+02	6.29E+00	4.92E+01	6.76E+02	52.1%
Total Hazard		5.98E+02	1.52E+02	6.22E+01	4.87E+02	1.30E+03	100.00%
% of Total Hazard		46.0%	11.7%	4.8%	37.5%		

Blank cells indicate that the exposure route is not appropriate to the COPC.

^a Units for metals and organic compounds are mg/L.

^b Only COPCs relevant to the endpoint are included in the table.

Table F.42. HI Child Residential Groundwater Use at SWMU 3

COPC	Exposure Point Concentration ^a	Ingestion of Water	Dermal Contact	Inhalation while showering	Household Inhalation	Total Hazard ^b	Percent of Total Hazard
<i>Inorganic Compounds</i>							
Arsenic	3.29E-02	1.05E+01	2.22E-02			1.05E+01	4751.9%
Manganese	8.95E-01	1.87E+00	7.75E-02			1.94E+00	9.6%
Uranium	4.89E-02	7.82E+00	7.97E-03			7.82E+00	38.6%
Total Hazard		2.19E02E+01	1.08E-01	0.00E+00	0.00E+00	2.03E+01	100.0%
% of Total Hazard		99.5%	0.5%	0.0%	0.0%		

Blank cells indicate that the exposure route is not appropriate to the COPC.

^a Units for metals and organic compounds are mg/L.

^b Only COPCs relevant to the endpoint are included in the table.

Table F.43. HI Child Residential Groundwater Use at SWMU 4

COPC	Exposure Point Concentration ^a	Ingestion of Water	Dermal Contact	Inhalation while showering	Household Inhalation	Total Hazard ^b	Percent of Total Hazard
<i>Inorganic Compounds</i>							
Arsenic	1.77E-02	5.66E+00	1.20E-02			5.67E+00	1.0%
Manganese	5.76E-01	1.20E+00	2.60E-02			1.23E+00	0.2%
<i>Organic Compounds</i>							
<i>cis</i> -1,2-DCE	6.68E-01	6.41E+00	2.26E-01	3.25E+00	2.54E+01	3.53E+01	6.1%
TCE	1.18E+00	3.77E+02	1.17E+02	5.02E+00	3.92E+01	5.39E+02	92.5%
Vinyl Chloride	2.61E-02	8.34E-01	1.80E-02	9.45E-03	3.46E-01	1.21E+00	0.2%
Total Hazard		3.91E+02	1.18E+02	8.27E+00	6.50E+01	5.82E+02	100.0%
% of Total Hazard		67.2%	20.2%	1.4%	11.2%		

Blank cells indicate that the exposure route is not appropriate to the COPC.

^a Units for metals and organic compounds are mg/L.

^b Only COPCs relevant to the endpoint are included in the table.

Table F.44. HI Child Residential Groundwater Use at SWMU 5

COPC	Exposure Point Concentration ^a	Ingestion of Water	Dermal Contact	Inhalation while showering	Household Inhalation	Total Hazard ^b	Percent of Total Hazard
<i>Inorganic Compounds</i>							
Uranium	4.60E-01	7.35E+01	7.50E-02			7.36E+01	90.3%
Arsenic	9.25E-03	12.96E+00	6.25E-03			12.96E+00	3.6%
Manganese	1.01E+00	12.11E+00	4.56E-02			2.15E+00	2.6%
<i>Organic Compounds</i>							
Naphthalene	5.55E-03	2.66E-02	6.04E-03	3.14E-01	2.45E+00	2.80E+00	3.4%
Total Hazard		7.86E+01	1.33E-01	3.14E-01	2.45E+00	8.15E+01	100.0%
% of Total Hazard		96.4%	0.2%	0.4%	3.0%		

Blank cells indicate that the exposure route is not appropriate to the COPC.

^a Units for metals and organic compounds are mg/L.

^b Only COPCs relevant to the endpoint are included in the table.

Table F.45. HI Child Residential Groundwater Use at SWMU 6

COPC	Exposure Point Concentration	Ingestion of Water	Dermal Contact	Inhalation while showering	Household Inhalation	Total Hazard	Percent of Total Hazard
Manganese	8.32E-02	1.73E-01	3.76E-03			1.77E-01	100%
Total Hazard		1.73E-01	3.76E-03			1.77E-01	100%
% of Total Hazard		97.9%	2.1%				

Table F.46. HI Child Residential Groundwater Use at SWMU 7

COPC	Exposure Point Concentration ^a	Ingestion of Water	Dermal Contact	Inhalation while showering	Household Inhalation	Total Hazard ^b	Percent of Total Hazard
<i>Inorganic Compounds</i>							
Arsenic	1.78E-02	5.69E+00	1.20E-02			5.70E+00	30.2%
Manganese	3.32E-01	6.92E-01	1.50E-02			7.07E-01	3.7%
Uranium	3.46E-03	5.53E-01	5.64E-04			5.54E-01	2.9%
<i>Organic Compounds</i>							
1,1-DCE	8.98E-02	1.72E-01	6.45E-03	7.62E-02	5.96E-01	8.51E-01	4.5%
cis-1,2-DCE	2.35E-02	2.25E-01	7.95E-03	1.14E-01	8.94E-01	1.24E+00	6.6%
Aroclor-1254	5.23E-05	2.51E-01	2.83E+00	1.27E-01	9.94E-01	4.20E+00	22.3%
TCE	1.09E-02	3.48E+00	1.09E+00	4.63E-02	3.63E-01	4.98E+00	26.4%
Vinyl Chloride	1.35E-02	4.32E-01	9.30E-03	2.29E-02	1.79E-01	6.43E-01	3.4%
Total Hazard		1.15E+01	3.97E+00	3.87E-01	3.03E+00	1.89E+01	100.0%
% of Total Hazard		60.9%	21.0%	2.0%	16.0%		

Blank cells indicate that the exposure route is not appropriate to the COPC.

^a Units for metals and organic compounds are mg/L.

^b Only COPCs relevant to the endpoint are included in the table.

Table F.47. HI Child Residential Groundwater Use at SWMU 30

COPC	Exposure Point Concentration ^a	Ingestion of Water	Dermal Contact	Inhalation while showering	Household Inhalation	Total Hazard ^b	Percent of Total Hazard
<i>Inorganic Compounds</i>							
Arsenic	1.77E-02	5.66E+00	1.20E-02			5.67E+00	63.8%
Manganese	3.78E-01	7.88E-01	1.71E-02			8.05E-01	8.8%
Selenium	1.51E-02	2.90E-01	5.70E-04			2.90E-01	3.2%
Uranium	8.40E-03	1.34E+00	1.37E-03			1.34E-01	14.7%
<i>Organic Compounds</i>							
1,1-DCE	8.18E-05	1.57E-04	5.88E-06	5.13E-02	4.01E-01	4.53E-01	5.0%
TCE	9.11E-04	2.91E-01	9.08E-02	3.88E-03	3.03E-02	4.16E-01	4.6%
Total Hazard		8.53E+00	1.22E-01	5.52E-02	4.31E-01	9.14E+00	100.0%
% of Total Hazard		93.3%	1.3%	0.6%	4.7%		

Blank cells indicate that the exposure route is not appropriate to the COPC.

^a Units for metals and organic compounds are mg/L.

^b Only COPCs relevant to the endpoint are included in the table.

Table F.48. HI Child Residential Groundwater Use at SWMU 145

COPC	Exposure Point Concentration ^a	Ingestion of Water	Dermal Contact	Inhalation while showering	Household Inhalation	Total Hazard ^b	Percent of Total Hazard
<i>Inorganic Compounds</i>							
Antimony	7.99E-02	1.92E+01	8.30E-01			2.00E+01	48.0%
Arsenic	6.21E-02	1.98E+01	4.20E-02			1.99E+01	47.7%
Manganese	8.44E-01	1.76E+00	3.81E-02			1.80E+00	4.3%
Total Hazard		4.08E+01	9.10E-01	0.00E+00	0.00E+00	4.17E+01	100.0%
% of Total Hazard		97.8%	2.2%	0.0%	0.0%		

Blank cells indicate that the exposure route is not appropriate to the COPC.

^a Units for metals and organic compounds are mg/L.

^b Only COPCs relevant to the endpoint are included in the table.

Table F.49. HI Adult Residential Groundwater Use at SWMU 2

COPC	Exposure Point Concentration ^a	Ingestion of Water	Dermal Contact	Inhalation while showering	Household Inhalation	Total Hazard ^b	Percent of Total Hazard
<u>Inorganic Compounds</u>							
Arsenic	3.54E-02	3.23E+00	1.43E-02			3.25E+00	0.9%
Manganese	7.16E-01	4.26E-01	1.93E-02			4.46E-01	0.1%
Uranium	9.86E-03	4.50E-01	9.61E-04			4.51E-01	0.1%
<u>Organic Compounds</u>							
<i>cis</i> -1,2-DCE	1.15E+01	3.15E+01	2.33E+00	1.20E+01	9.37E+01	1.40E+02	36.8%
Naphthalene	9.38E-04	1.28E-03	6.10E-04	1.14E-02	8.89E-02	1.02E-01	0%
TCE	1.48E+00	1.35E+02	8.83E+01	1.35E+00	1.05E+01	2.35E+02	62.1%
Total Hazard		1.71E+02	9.07E+01	1.33E+01	1.04E+02	3.79E+02	100.00%
% of Total Hazard		45.0%	23.9%	3.5%	27.5%		

Blank cells indicate that the exposure route is not appropriate to the COPC.

^a Units for metals and organic compounds are mg/L.

^b Only COPCs relevant to the endpoint are included in the table.

Table F.50. HI Adult Residential Groundwater Use at SWMU 3

COPC	Exposure Point Concentration ^a	Ingestion of Water	Dermal Contact	Inhalation while showering	Household Inhalation	Total Hazard ^b	Percent of Total Hazard
<u>Inorganic Compounds</u>							
Arsenic	3.29E-02	3.00E+00	1.33E-02	NA	NA	3.02E+00	51.7%
Manganese	8.95E-01	5.33E-01	4.64E-02	NA	NA	5.79E-01	9.9%
Uranium	4.89E-02	2.23E+00	4.77E-03	NA	NA	2.24E+00	38.3%
Total Hazard		5.77E+00	6.44E-02	0.00E+00	0.00E+00	5.83E+00	100.0%
% of Total Hazard		98.9%	1.1%	0.0%	0.0%		

Blank cells indicate that the exposure route is not appropriate to the COPC.

^a Units for metals and organic compounds are mg/L.

^b Only COPCs relevant to the endpoint are included in the table.

Table F.51. HI Adult Residential Groundwater Use at SWMU 4

COPC	Exposure Point Concentration ^a	Ingestion of Water	Dermal Contact	Inhalation while showering	Household Inhalation	Total Hazard ^b	Percent of Total Hazard
<u>Inorganic Compounds</u>							
Arsenic	1.77E-02	1.62E+00	7.16E-03			1.62E+00	0.8%
Manganese	5.76E-01	3.43E-01	1.56E-02			3.59E-01	0.2%
<u>Organic Compounds</u>							
<i>cis</i> -1,2-DCE	6.68E-01	1.83E+00	1.35E-01	6.94E-01	5.44E+00	8.10E+00	4.1%
TCE	1.18E+00	1.08E+02	7.02E+01	1.07E+00	8.41E+00	1.87E+02	94.7%
Vinyl Chloride	2.61E-02	2.38E-01	1.08E-02	9.45E-03	7.41E-02	3.33E-01	0.2%
Total Hazard		1.12E+02	7.03E+01	1.77E+00	1.39E+01	1.98E+02	100.0%
% of Total Hazard		56.5%	35.6%	0.9%	7.0%		

Blank cells indicate that the exposure route is not appropriate to the COPC.

^a Units for metals and organic compounds are mg/L.

^b Only COPCs relevant to the endpoint are included in the table.

Table F.52. HI Adult Residential Groundwater Use at SWMU 5

COPC	Exposure Point Concentration ^a	Ingestion of Water	Dermal Contact	Inhalation while showering	Household Inhalation	Total Hazard ^b	Percent of Total Hazard
<i>Inorganic Compounds</i>							
Arsenic	9.25E-03	8.45E-01	3.74E-03			8.48E-01	3.7%
Manganese	1.01E+00	6.02E-01	2.73E-02			6.29E-01	2.7%
Uranium	4.60E-01	2.10E+01	4.49E-02	NA	NA	2.10E+01	91.0%
<i>Organic Compounds</i>							
Naphthalene	5.55E-03	7.60E-03	3.61E-03	6.72E-02	5.26E-01	6.05E-01	2.6%
Total Hazard		2.25E+01	7.95E-02	6.72E-02	5.26E-01	2.31E+01	100.0%
% of Total Hazard		97.1%	0.3%	0.3%	2.3%		

Blank cells indicate that the exposure route is not appropriate to the COPC.

^a Units for metals and organic compounds are mg/L.

^b Only COPCs relevant to the endpoint are included in the table.

Table F.53. HI Adult Residential Groundwater Use at SWMU 6

COPC	Exposure Point Concentration	Ingestion of Water	Dermal Contact	Inhalation while showering	Household Inhalation	Total Hazard	Percent of Total Hazard
Manganese	8.32E-02	4.96E-02	2.25E-03	NA	NA	5.18E-02	100%
Total Hazard		4.96E-02	2.25E-03	0.00E+00	0.00E+00	5.18E-02	100%
% of Total Hazard		95.7%	4.3%	0.0%	0.0%		

Table F.54. HI Adult Residential Groundwater Use at SWMU 7

COPC	Exposure Point Concentration ^a	Ingestion of Water	Dermal Contact	Inhalation while showering	Household Inhalation	Total Hazard ^b	Percent of Total Hazard
<i>Inorganic Compounds</i>							
Arsenic	1.78E-02	1.63E+00	7.20E-03			1.63E+00	25.5%
Manganese	3.32E-01	1.98E-01	8.97E-03			2.07E-01	3.2%
Uranium	3.46E-03	1.58E-01	3.37E-04			1.58E-01	2.5%
<i>Organic Compounds</i>							
1,1-DCE	8.98E-02	4.92E-02	3.86E-03	1.63E-02	1.28E-01	1.97E-01	3.1%
cis-1,2-DCE	2.35E-02	6.44E-02	4.76E-03	2.44E-02	1.92E-01	2.85E-01	4.5%
Aroclor-1254	5.23E-05	7.16E-02	1.69E+00	2.71E-02	2.13E-01	2.01E+00	31.4%
TCE	1.09E-02	9.95E-01	6.50E-01	9.90E-03	7.77E-02	1.73E+00	27.1%
Vinyl Chloride	1.35E-02	1.23E-01	5.57E-03	4.89E-03	3.84E-02	1.72E-01	2.7%
Total Hazard		3.29E+00	2.37E+00	8.26E-02	6.48E-01	6.39E+00	100.0%
% of Total Hazard		51.4%	37.2%	1.3%	10.1%		

Blank cells indicate that the exposure route is not appropriate to the COPC.

^a Units for metals and organic compounds are mg/L.

^b Only COPCs relevant to the endpoint are included in the table.

Table F.55. HI Adult Residential Groundwater Use at SWMU 30

COPC	Exposure Point Concentration ^a	Ingestion of Water	Dermal Contact	Inhalation while showering	Household Inhalation	Total Hazard ^b	Percent of Total Hazard
<i>Inorganic Compounds</i>							
Arsenic	1.77E-02	1.62E+00	7.16E-03			1.62E+00	50.5%
Manganese	3.78E-01	2.25E-01	1.02E-02			2.35E-01	7.1%
Selenium	1.51E-02	8.27E-02	3.41E-04			8.31E-02	2.5%
Uranium	8.40E-03	3.84E-01	8.19E-04			3.84E-01	11.6%
<i>Organic Compounds</i>							
1,1-DCE	8.18E-05	5.00E-01	2.50E-01	4.39E-03	3.44E-02	7.89E-01	23.9%
TCE	9.11E-04	8.32E-02	5.44E-02	8.30E-04	6.49E-03	1.45E-01	4.4%
Total Hazard		2.94E+00	3.23E-01	5.22E-03	4.09E-02	3.31E+00	100.0%
% of Total Hazard		88.8%	9.8%	0.2%	1.2%		

Blank cells indicate that the exposure route is not appropriate to the COPC.

^a Units for metals and organic compounds are mg/L.

^b Only COPCs relevant to the endpoint are included in the table.

Table F.56. HI Adult Residential Groundwater Use at SWMU 145

COPC	Exposure Point Concentration ^a	Ingestion of Water	Dermal Contact	Inhalation while showering	Household Inhalation	Total Hazard ^b	Percent of Total Hazard
<i>Inorganic Compounds</i>							
Antimony	7.99E-02	5.47E+00	4.97E-01			5.97E+00	49.0%
Arsenic	6.21E-02	5.67E+00	2.51E-02			5.70E+00	46.7%
Manganese	8.44E-01	5.03E-01	2.28E-02			5.25E-01	4.3%
Total Hazard		1.16E+01	5.45E-01	0.00E+00	0.00E+00	1.22E+01	100.0%
% of Total Hazard		95.5%	4.5%	0.0%	0.0%		

Blank cells indicate that the exposure route is not appropriate to the COPC.

^a Units for metals and organic compounds are mg/L.

^b Only COPCs relevant to the endpoint are included in the table.

F.5.4.2 Excess Lifetime Cancer Risk (Groundwater Use)

Tables F.57 through F.64 summarize the ELCRs for the modeled groundwater exposure above each SWMU for the rural resident over a lifetime. As shown in these tables, the total ELCRs (bold value in “Total Risk” column) are greater than both 1×10^{-6} and 1×10^{-4} for all of the SWMUs, except SWMU 6, which has no groundwater COCs, with a carcinogenic endpoint. The source with the greatest ELCR is SWMU 4, which has an ELCR of 5.41×10^{-2} . The major pathway is inhalation of vapor during household water use (42.4%). The major contribution is from TCE (67.7%) and vinyl chloride (30.5%).

Table F.57. ELCR Residential Groundwater Use at SWMU 2

COPC	Exposure Point Concentration ^a	Ingestion of Water	Dermal Contact	Inhalation while showering	Household Inhalation	Total Risk ^b	Percent of Total Risk
<u>Inorganic Compounds</u>							
Arsenic	3.54E-02	9.35E-04	3.13E-06			9.38E-04	2.0%
<u>Organic Compounds</u>							
Aroclor-1248	1.54E-03	1.08E-05	1.22E-04	4.16E-06	3.25E-05	1.70E-04	0.4%
Aroclor-1260	8.73E-05	6.15E-07	6.68E-05	2.36E-07	1.84E-06	6.95E-05	0.1%
TCE	1.48E+00	8.39E-03	5.16E-03	3.68E-03	2.88E-02	4.60E-02	97.5 %
<u>Radionuclides</u>							
Technetium-99	1.02E+02	5.60E-06				5.60E-06	0.0%
Uranium-234	1.58E+00	2.23E-06				2.23E-06	0.0%
Uranium-238	1.81E+00	3.15E-06				3.15E-06	0.0%
Total Risk		9.35E-03	5.35E-03	3.68E-03	2.88E-02	4.72E-02	100.0%
% of Total Risk		19.8%	11.3%	7.8%	61.0%		

Blank cells indicate that the exposure route is not appropriate to the COPC.

^a Units for metals and organic compounds are mg/L. Units for radionuclides are pCi/L.

^b Only COPCs relevant to the endpoint are included in the table.

Table F.58. ELCR Residential Groundwater Use at SWMU 3

COPC	Exposure Point Concentration ^a	Ingestion of Water	Dermal Contact	Inhalation while showering	Household Inhalation	Total Risk ^b	Percent of Total Risk
<u>Inorganic Compounds</u>							
Arsenic	3.29E-02	8.69E-04	2.91E-06			8.72E-04	72.4%
<u>Radionuclides</u>							
Technetium-99	5.56E+03	3.05E-04				3.05E-04	25.3%
Uranium-238	1.59E+01	2.76E-05				2.76E-05	2.3%
Total Risk		1.20E-03	2.91E-06	0.00E+00	0.00E+00	1.20E-03	100.0%
% of Total Risk		99.8%	0.2%	0.0%	0.0%		

Blank cells indicate that the exposure route is not appropriate to the COPC.

^a Units for metals and organic compounds are mg/L. Units for radionuclides are pCi/L.

^b Only COPCs relevant to the endpoint are included in the table.

Table F.59. ELCR Residential Groundwater Use at SWMU 4

COPC	Exposure Point Concentration ^a	Ingestion of Water	Dermal Contact	Inhalation while showering	Household Inhalation	Total Risk ^b	Percent of Total Risk
<u>Inorganic Compounds</u>							
Arsenic	1.77E-02	4.68E-04	1.57E-06			4.69E-04	0.9%
<u>Organic Compounds</u>							
TCE	1.18E+00	6.69E-03	4.10E-03	2.93E-03	2.29E-02	3.67E-02	67.7%
Vinyl Chloride	2.61E-02	6.90E-04	1.58E-02	6.20E-06	4.85E-05	1.65E-02	30.5%
<u>Radionuclides</u>							
Technetium-99	9.01E+03	4.94E-04				4.94E-04	0.9%
Total Risk		8.34E-03	1.99E-02	2.94E-03	2.30E-02	5.41E-02	100.0%
% of Total Risk		15.4%	36.7%	5.4%	42.4%		

Blank cells indicate that the exposure route is not appropriate to the COPC.

^a Units for metals and organic compounds are mg/L. Units for radionuclides are pCi/L.

^b Only COPCs relevant to the endpoint are included in the table.

Table F.60. ELCR Residential Groundwater Use at SWMU 5

COPC	Exposure Point Concentration ^a	Ingestion of Water	Dermal Contact	Inhalation while showering	Household Inhalation	Total Risk ^b	Percent of Total Risk
<u>Inorganic Compounds</u>							
Arsenic	9.25E-03	2.44E-04	8.18E-07			2.45E-04	97.2%
<u>Radionuclides</u>							
Technetium-99	1.27E+02	6.97E-06				6.97E-06	2.8%
Total Risk		2.51E-04	8.18E-07	0.00E+00	0.00E+00	2.52E-04	100.0%
% of Total Risk		99.7%	0.3%	0.0%	0.0%		

Blank cells indicate that the exposure route is not appropriate to the COPC.

^a Units for metals and organic compounds are mg/L. Units for radionuclides are pCi/L.

^b Only COPCs relevant to the endpoint are included in the table.

Table F.61. ELCR Residential Groundwater Use at SWMU 6

COPC	Exposure Point Concentration ^a	Ingestion of Water	Dermal Contact	Inhalation while showering	Household Inhalation	Total Risk ^b	Percent of Total Risk
NO COPCs¹							

¹ Modeling analysis (Appendix E) did not show any of the identified carcinogenic COPCs at this site as migrating to groundwater.

Table F.62. ELCR Residential Groundwater Use at SWMU 7

COPC	Exposure Point Concentration ^a	Ingestion of Water	Dermal Contact	Inhalation while showering	Household Inhalation	Total Risk ^b	Percent of Total Risk
<u>Inorganic Compounds</u>							
Arsenic	1.78E-02	4.70E-04	1.57E-06			4.72E-04	15.1%
<u>Organic Compounds</u>							
1,1-DCE	8.98E-02	9.49E-04	5.63E-05	1.21E-04	9.48E-04	2.08E-03	66.4%
Aroclor-1254	5.23E-05	3.68E-07	6.58E-06	1.41E-07		7.09E-06	0.2%
TCE	1.09E-02	6.18E-05	3.80E-05	2.71E-05		1.27E-04	4.1%
Vinyl Chloride	1.35E-02	3.57E-04	1.22E-05	3.21E-06		3.72E-04	11.9%
<u>Radionuclides</u>							
Technetium-99	9.09E+02	4.99E-05				4.99E-05	1.6%
Uranium-234	7.94E+00	1.12E-05				1.12E-05	0.4%
Uranium-238	7.59E+00	1.32E-05				1.32E-05	0.4%
Total Risk		1.91E-03	1.15E-04	1.52E-04	9.48E-04	3.13E-03	100.0%
% of Total Risk		61.2%	3.7%	4.9%	30.3%		

Blank cells indicate that the exposure route is not appropriate to the COPC.

^a Units for metals and organic compounds are mg/L. Units for radionuclides are pCi/L.

^b Only COPCs relevant to the endpoint are included in the table.

Table F.63. ELCR Residential Groundwater Use at SWMU 30

COPC	Exposure Point Concentration ^a	Ingestion of Water	Dermal Contact	Inhalation while showering	Household Inhalation	Total Risk ^b	Percent of Total Risk
<u>Inorganic Compounds</u>							
Arsenic	1.77E-02	4.68E-04	1.57E-06			4.69E-04	88.6%
<u>Organic Compounds</u>							
1,1-DCE	8.18E-05	8.64E-07	5.13E-08	1.19E-07	8.64E-07	1.90E-06	0.3%
TCE	9.11E-04	5.17E-06	3.17E-06	2.26E-06	1.77E-05	2.83E-05	5.2%
<u>Radionuclides</u>							
Technetium-99	2.87E+02	1.57E-05				1.57E-05	2.9%
Uranium-234	3.99E+00	5.63E-06				5.63E-06	1.0%
Uranium-238	5.91E+00	1.03E-05				1.03E-05	1.3%
Total Risk		5.18E-04	4.84E-06	2.38E-06	1.86E-05	5.44E-04	100.0%
% of Total Risk		95.3%	0.9%	0.4%	3.4%		

Blank cells indicate that the exposure route is not appropriate to the COPC.

^a Units for metals and organic compounds are mg/L. Units for radionuclides are pCi/L.

^b Only COPCs relevant to the endpoint are included in the table.

Table F.64. ELCR Residential Groundwater Use at SWMU 145

COPC	Exposure Point Concentration ^a	Ingestion of Water	Dermal Contact	Inhalation while showering	Household Inhalation	Total Risk ^b	Percent of Total Risk
<u>Inorganic Compounds</u>							
Arsenic	6.21E-02	1.64E-03	2.28E-05			1.66E-03	5.1%
<u>Organic Compounds</u>							
Aroclor-1260	1.92E-03	6.76E-05	3.04E-02			3.05E-02	93.2%
<u>Radionuclides</u>							
Technetium-99	1.01E+04	5.54E-04				5.54E-04	1.7%
Uranium-238	7.67E-02	1.10E-07				1.10E-07	0.0%
Total Risk		2.26E-03	3.04E-02	0.0%	0.0%	3.27E-02	100.0%
% of Total Risk		6.9%	93.1%	0.0%	0.0%		

Blank cells indicate that the exposure route is not appropriate to the COPC.

^a Units for metals and organic compounds are mg/L. Units for radionuclides are pCi/L.

^b Only COPCs relevant to the endpoint are included in the table.

F.5.5 RISK CHARACTERIZATION FOR RESIDENTIAL USE OF GROUNDWATER AT FUTURE MODELED CONCENTRATIONS AT BOUNDARY AND RIVER POEs

This subsection discusses the potential future risks to a hypothetical resident using RGA groundwater contaminated by migration of COPCs from the SWMU 2, SWMU 3, SWMU 4, SWMU 5, SWMU 6, SWMU 7, SWMU 30, and SWMU 145 sources. As discussed in Section 2 of this BHHRA, the POEs to which contaminants were modeled were the PGDP plant boundary, PGDP property boundary, and near either the Little Bayou Seeps or the Ohio River. Information about the methods used in the model is provided in Section 5 and Appendix E of this RI.

Table F.65 presents the chemical-specific HIs for the child and adult rural residents from exposure to the modeled peak concentration over the 1,000 year time frame of the COPCs in the RGA at the POEs based on household use of groundwater. The major contributors are arsenic, TCE, and *cis*-1,2-DCE. Table F.66 presents the chemical-specific ELCRs for a rural resident from exposure to maximum modeled concentrations over the 1,000 year time frame of contaminants in the RGA groundwater at the POEs based on household use of water. The major contributors to ELCR are TCE, vinyl chloride, and ⁹⁹Tc. Table F.67 presents the chemical-specific ELCRs for a rural resident from exposure to maximum modeled concentrations of contaminants in the RGA at the seeps at Little Bayou Creek. Peak concentrations for contaminants in groundwater emerging at the seeps were modeled based on contaminants migrating from SWMUs 3, 7, and 30.

Tables F.65 through F.67 show the HIs and ELCRs for the predicted maximum (peak) concentrations for individual contaminants over the 1,000 year time frame of the model. Different contaminants migrate at different rates; therefore, the total HI or ELCR in groundwater at a given time may be less than the sum of the maximum risks of individual COCs. Figures F.2 to F.16 show the total hazard and total risks from the predicted concentrations of all COCs at each time step in the model for each SWMU except SWMU 6 (which had no modeled COCs at locations other than below the SWMU). These hazards and risks are calculated using the NALs (DOE 2001) for residential use of groundwater. The values for total hazard or risk for all COCs calculated from the NALs for residential groundwater use and the time at which the total risk or hazard peaks are provided in Tables F.68 and F.69. These figures and tables show that at all SWMUs except SWMU 7, risk and hazard associated with modeled groundwater contaminants at the SWMU unit peak at the same time. For SWMUs 2, 4, and 30, these peaks occur in within the first twenty years of the model run. For SWMUs 3, 5, and 145, these peaks occur at the 1,000 year boundary of the model. For SWMU 7, the risk peaks very early in the model run, but the hazard value continues increasing to the end of the 1,000 year time frame. For SWMUs with contaminants that may migrate to other POEs such as the plant boundary and property boundary, the same pattern of peak times is seen as for the SWMU unit boundary except at SWMU 3. At SWMU 3, the ELCR at the unit boundary increases over the time frame of the model, but risk at the plant boundary and property boundary peak early because a different contaminant acts as the risk driver. These results indicate that some contaminants in the soil column at the SWMUs, which are not currently significant contributors to risk and hazard in groundwater, may become contributors in the future. Section 5 and Appendix E of the RI provide a discussion of the rates of migration for individual contaminants.

Table F.65. HIs for Peak Modeled Water Concentrations at the Plant Boundary, Property Boundary, and Near the Ohio River for Household Use of Groundwater Water Contaminated by COPC Migration from the BGOU SWMUs

COPC ^a	HI (child) at POE			HI (adult) at POE		
	Plant Boundary	Property Boundary	Near Ohio River	Plant Boundary	Property Boundary	Near Ohio River
SWMU 2						
Arsenic	9.32E-01	2.67E-06	NA	2.67E-01	7.66E-07	NA
Manganese	4.04E-05	NA	NA	1.20E-05	NA	NA
Uranium	1.35E-01	NA	NA	3.81E-06	NA	NA
<i>cis</i> -1,2-DCE	9.19E+01	4.53E+01	1.79E+01	1.60E+01	1.04E+01	4.10E+00
Naphthalene	1.43E-01	4.18E-02	1.73E-02	3.08E-02	9.01E-03	3.73E-03
TCE	9.91E+01	5.03E+01	4.61E+00	3.45E+01	1.75E+01	2.59E+00
SWMU 3						
Arsenic	3.98E-01	NA	--	1.12E-01	NA	--
Manganese	8.69E-10	NA	--	2.54E-10	NA	--
Uranium	3.63E-11	NA	--	1.04E-11	NA	--
SWMU 4						
Arsenic	8.65E-01	1.57E-03	NA	2.48E-01	4.49E-04	NA
Manganese	1.07E-02	NA	NA	3.12E-03	NA	NA
<i>cis</i> -1,2-DCE	1.04E+01	4.72E+00	5.70E-01	2.38E+00	1.08E+00	3.84E-01
TCE	1.93E+02	9.77E+01	3.27E+01	6.70E+01	3.40E+01	1.22E+01
Vinyl Chloride	2.83E-01	1.20E-01	2.77E-02	7.58E-02	3.23E-02	9.97E-03
SWMU 5						
Arsenic	5.70E-01	4.07E-02	NA	1.63E-01	1.16E-02	NA
Manganese	1.85E-01	4.90E-11	NA	5.41E-02	1.43E-11	NA
Naphthalene	4.95E-01	1.88E-01	5.45E-02	1.07E-01	4.05E-02	1.18E-02
Uranium	5.31E+00	7.44E-09	NA	1.52E+00	2.13E-09	NA
SWMU 6^a						
NA	NA	NA	NA	NA	NA	--
SWMU 7						
Arsenic	4.04E+00	7.53E-01	--	1.16E+00	2.16E-01	--
Manganese	5.13E-01	2.24E-06	--	1.50E-01	6.54E-07	--
Uranium	4.05E-01	4.29E-07	--	1.16E-01	1.23E-07	--
1,1-DCE	7.81E-01	1.04E-01	--	1.81E-01	2.41E-02	--
<i>cis</i> -1,2-DCE	1.14E+00	1.65E-01	--	2.61E-01	3.80E-02	--
Aroclor-1254	2.48E+00	2.45E-01	--	1.18E+00	1.17E-01	--
TCE	4.51E+00	6.49E-01	--	1.57E+00	2.26E-01	--
Vinyl Chloride	5.90E-01	5.76E-02	--	1.58E-01	1.54E-02	--
SWMU 30						
Arsenic	3.75E+00	7.50E-01	--	1.07E+00	2.15E-01	--
Manganese	5.35E-01	6.07E-04	--	1.56E-01	1.77E-04	--
Selenium	1.59E-01	1.77E-02	--	4.57E-02	5.07E-03	--
Uranium	7.70E-01	3.86E-04	--	2.20E-01	1.10E-04	--
1,1-DCE	7.84E-03	6.29E-04	--	7.67E-03	6.93E-04	--
TCE	7.93E-01	7.10E-02	--	5.59E-01	5.49E-02	--
SWMU 145						
Antimony	--- ^b	3.78E-04	NA	--- ^b	1.13E-04	NA
Arsenic	--- ^b	5.16E-01	NA	--- ^b	1.48E-01	NA

-- = not a POE for groundwater from this SWMU.

NA = not applicable. Modeling results indicate that the constituent does not contribute significantly to groundwater at this point within the 1,000 year modeling time period, and therefore is insignificant at these POEs.

^aNone of the modeled constituents migrated from SWMU 6 at concentrations with a significant HI.

^bExposure point not modeled because SWMU 145 lies outside the plant boundary.

Table F.66. ELCRs for Peak Modeled Water Concentrations at the Plant Boundary, Property Boundary, and Near the Ohio River for Household Use of Groundwater Water Contaminated by COPC Migration from BGOU SWMUs

COPC	ELCR at POE		
	Plant Boundary	Property Boundary	Near Ohio River
SWMU 2			
Arsenic	7.71E-05	2.21E-10	NA
Aroclor-1248	1.41E-10	NA	NA
TCE	6.74E-03	3.42E-03	1.28E-03
Technetium-99	8.72E-07	4.42E-07	1.71E-07
Uranium-234	2.47E-11	NA	NA
Uranium-238	3.53E-11	NA	NA
SWMU3			
Arsenic	3.23E-05	NA	NA
Technetium-99	9.92E-05	7.46E-05	NA
Uranium-238	1.27E-16	NA	NA
SWMU 4			
Arsenic	7.15E-05	1.30E-07	NA
TCE	1.99E-02	6.65E-03	2.38E-03
Vinyl Chloride	1.92E-04	7.44E-05	2.30E-05
Technetium-99	1.37E-04	6.58E-05	2.08E-05
SWMU 5			
Arsenic	4.72E-05	3.37E-06	NA
Technetium-99	2.74E-06	1.45E-06	4.78E-07
SWMU 6			
	NA ^a	NA	NA
SWMU 7			
Arsenic	3.34E-04	6.23E-05	NA
1,1-DCE	1.90E-03	2.54E-04	NA
Aroclor-1254	4.84E-06	4.78E-07	NA
TCE	3.07E-04	4.38E-05	NA
Vinyl Chloride	3.65E-04	3.56E-05	NA
Technetium-99	4.52E-05	1.48E-05	NA
Uranium-234	8.17E-06	1.30E-12	NA
Uranium-238	9.69E-06	1.02E-11	NA
SWMU 30			
Arsenic	3.10E-04	6.20E-05	NA
1,1-DCE	1.77E-06	1.42E-07	NA
TCE	2.67E-05	2.39E-06	NA
Technetium-99	1.45E-05	3.88E-06	NA
Uranium-234	3.88E-06	2.03E-09	NA
Uranium-238	7.07E-06	3.44E-09	NA
SWMU 145			
Arsenic	--- ^b	4.27E-05	NA
Technetium-99	--- ^b	1.01E-04	5.29E-05

NA = not applicable. Modeling results indicate that the constituent does not contribute significantly to groundwater at this point within the 1,000-year modeling time period, and therefore is insignificant at these POEs.

^a Results for SWMU 6, none of the constituents modeled migrated at concentrations with a significant ELCR.

^b Exposure point not modeled because SWMU 145 lies outside the plant boundary.

Table F.67. HIs and ELCRs for Residential Groundwater Use at the Little Bayou Creek Seeps

	COPC	HI (child)	HI (adult)	ELCR
SWMU 3 Seep	Technetium-99	NA	NA	4.41E-05
SWMU 7	1,1-DCE	3.81E-02	8.82E-03	9.29E-05
	<i>cis</i> -1,2-DCE	8.45E-02	1.91E-02	NA
	Aroclor-1254	1.06E-07	5.06E-08	2.07E-13
	TCE	2.31E-01	8.05E-02	1.57E-05
	Vinyl Chloride	1.97E-02	5.26E-03	1.21E-05
	Technetium-99	NA	NA	7.26E-06
SWMU 30 Seep	Selenium	6.05E-03	1.73E-03	NA
	1,1-DCE	1.91E-04	2.10E-04	4.38E-08
	TCE	1.30E-02	7.23E-03	8.08E-07
	Technetium-99	NA	NA	1.60E-06

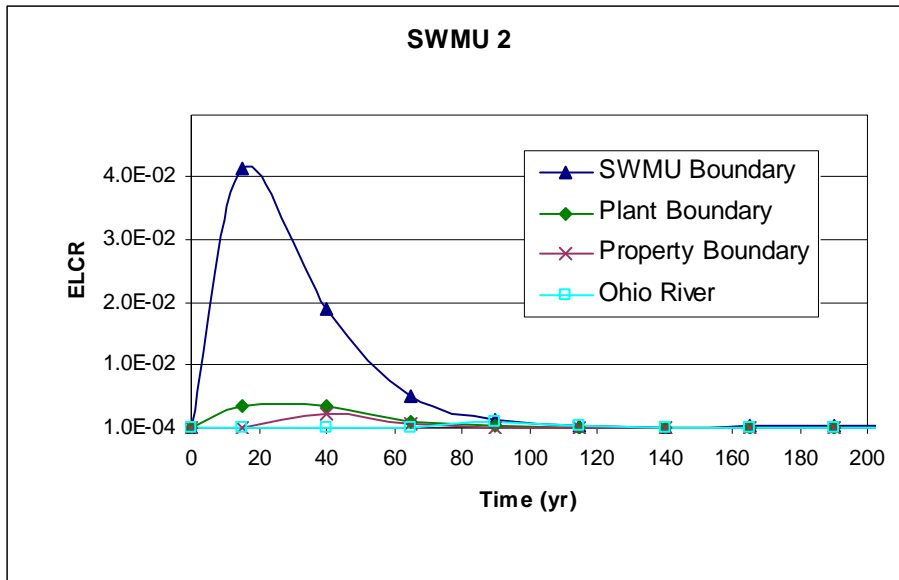


Figure F.2. Total ELCR from All Carcinogenic COCs at SWMU 2

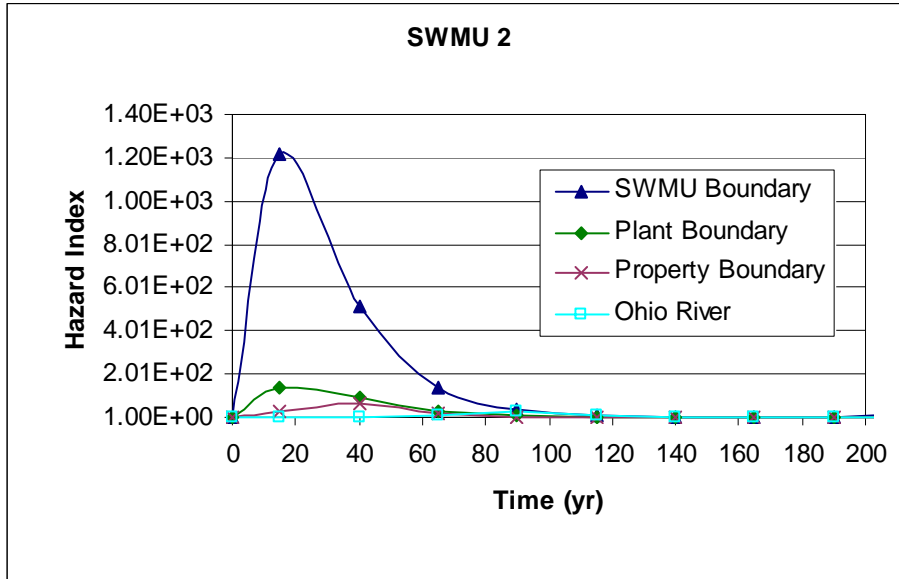


Figure F.3. Total HI from All Noncarcinogenic COCs at SWMU 2

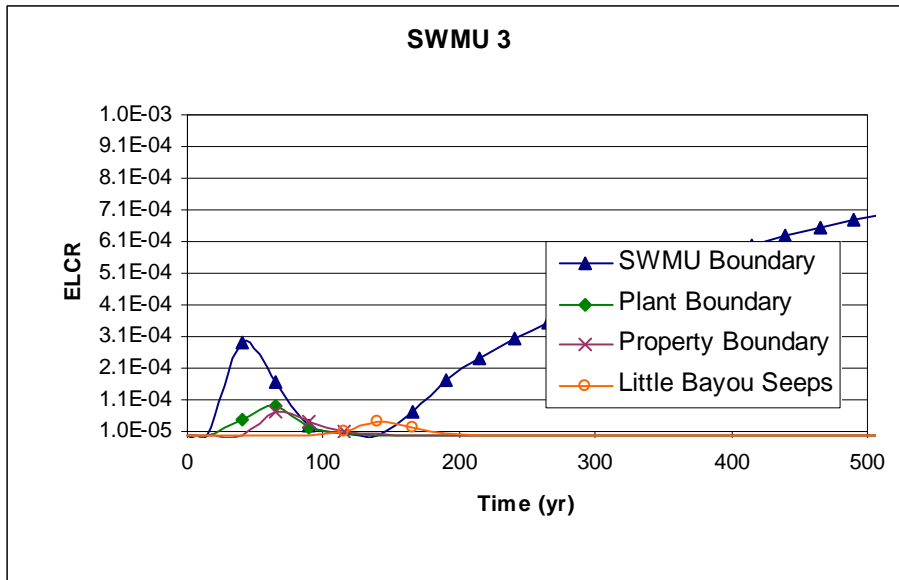


Figure F.4. Total ELCR from All Carcinogenic COCs at SWMU 3

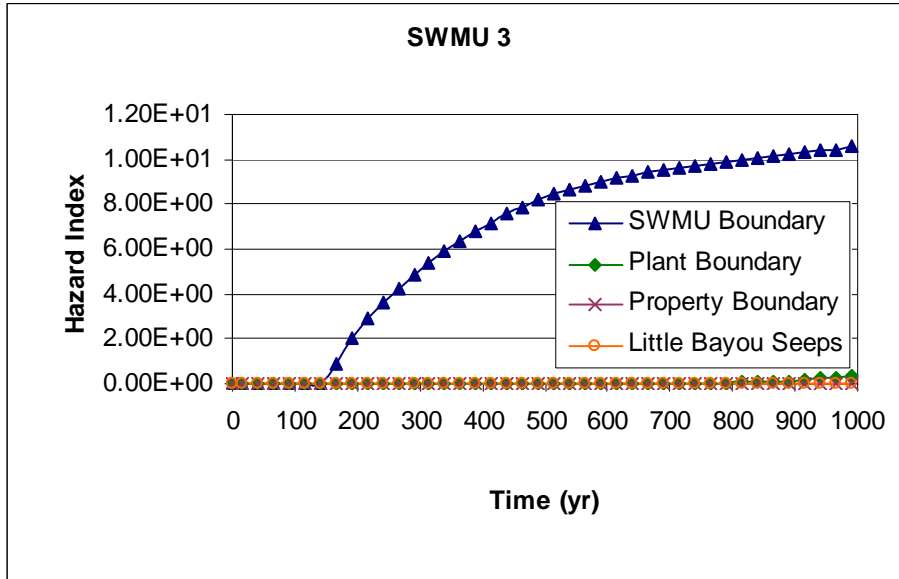


Figure F.5. Total HI from All Noncarcinogenic COCs at SWMU 3

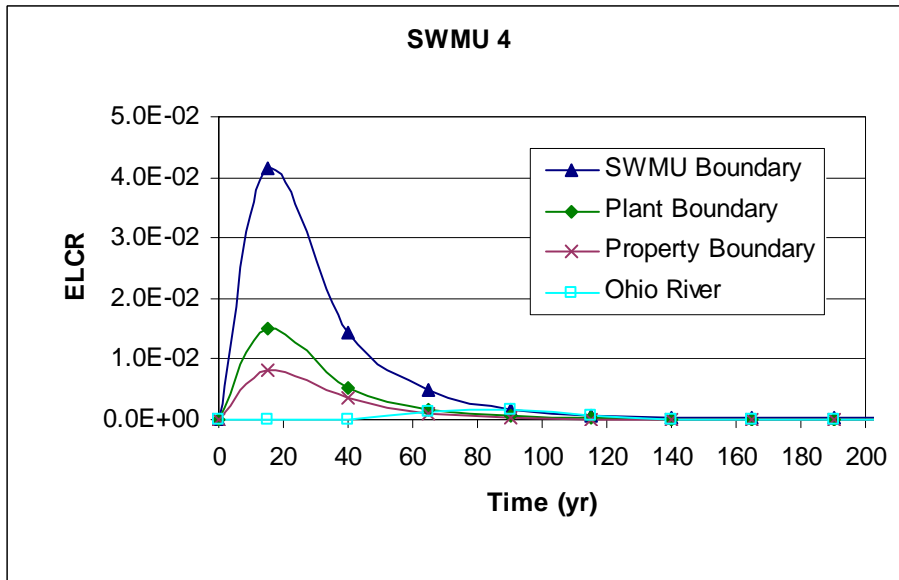


Figure F.6. Total ELCR from All Carcinogenic COCs at SWMU 4

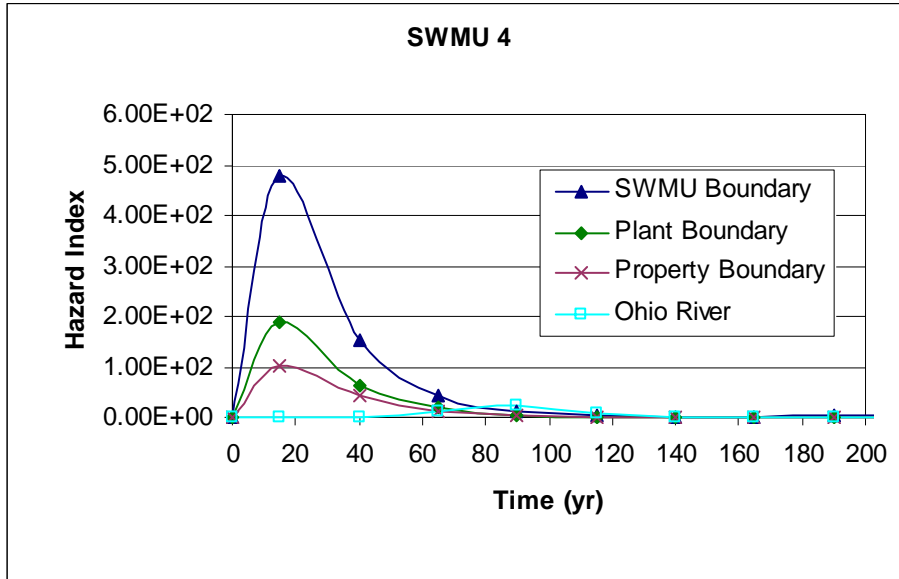


Figure F.7. Total HI from All Noncarcinogenic COCs at SWMU 4

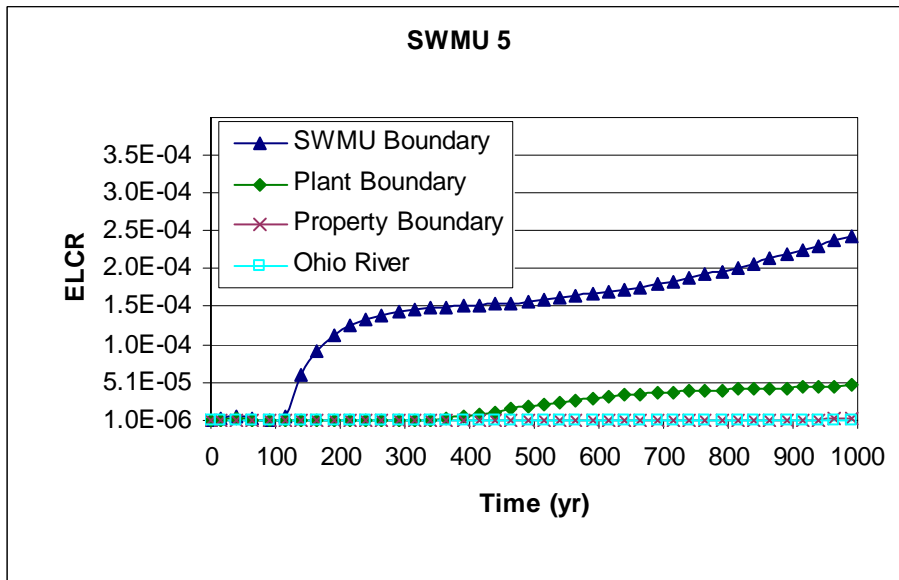


Figure F.8. Total ELCR from All Carcinogenic COCs at SWMU 5

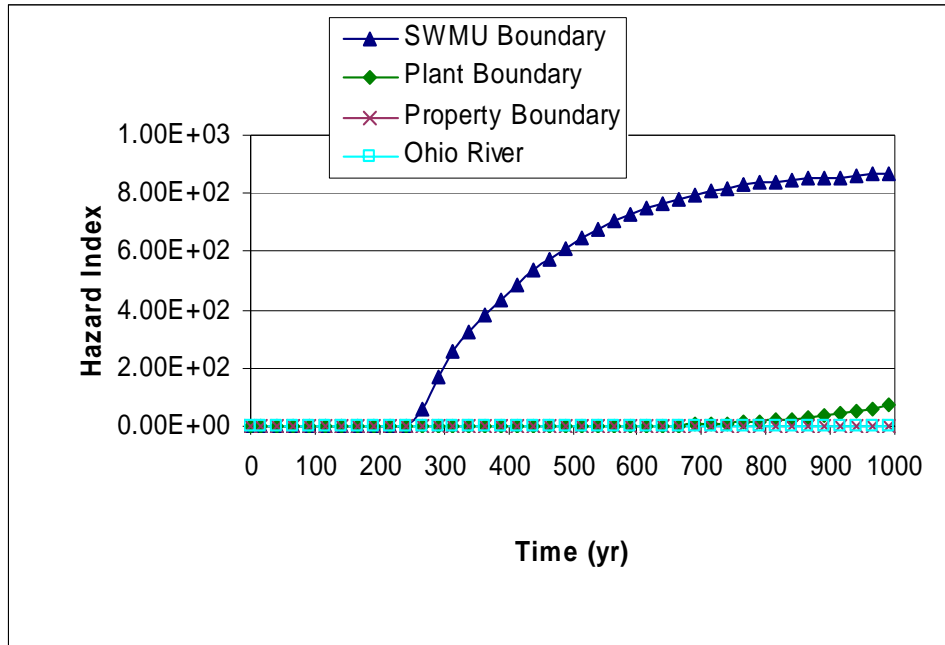


Figure F.9. Total HI from All Noncarcinogenic COCs at SWMU 5

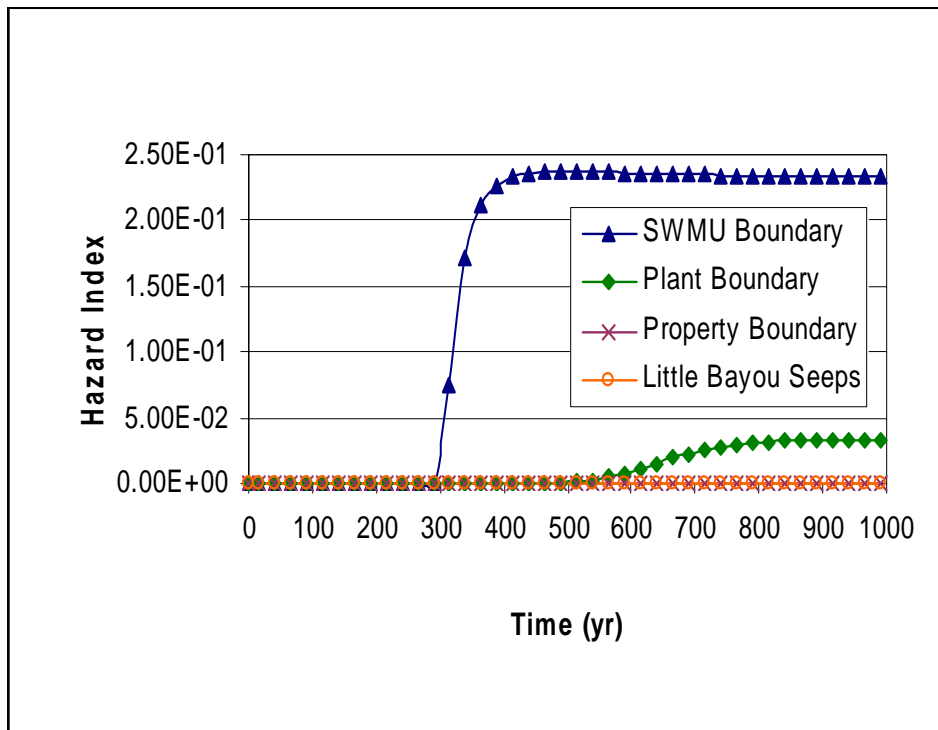


Figure F.10. Total HI from All Noncarcinogenic COCs at SWMU 6

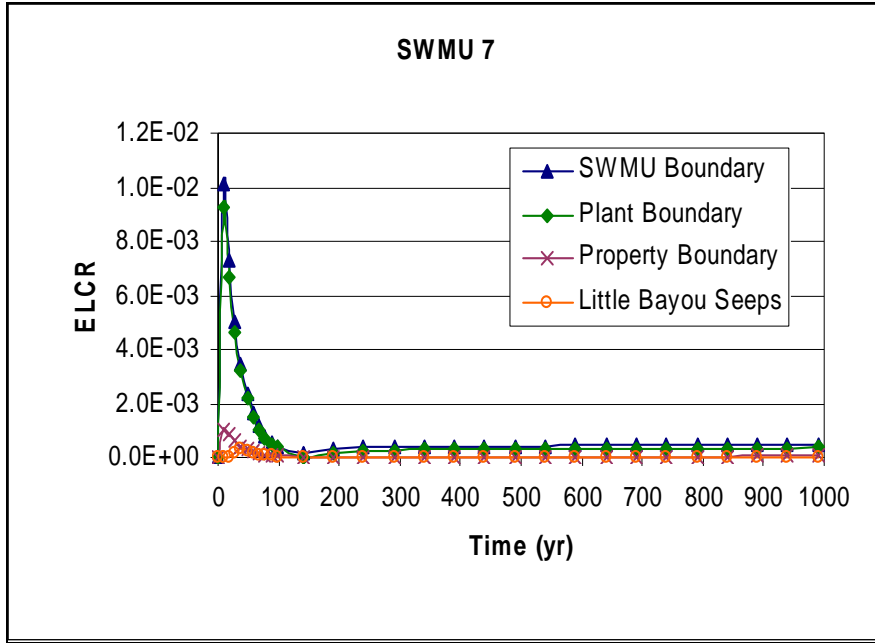


Figure F.11. Total ELCR from All Carcinogenic COCs at SWMU 7

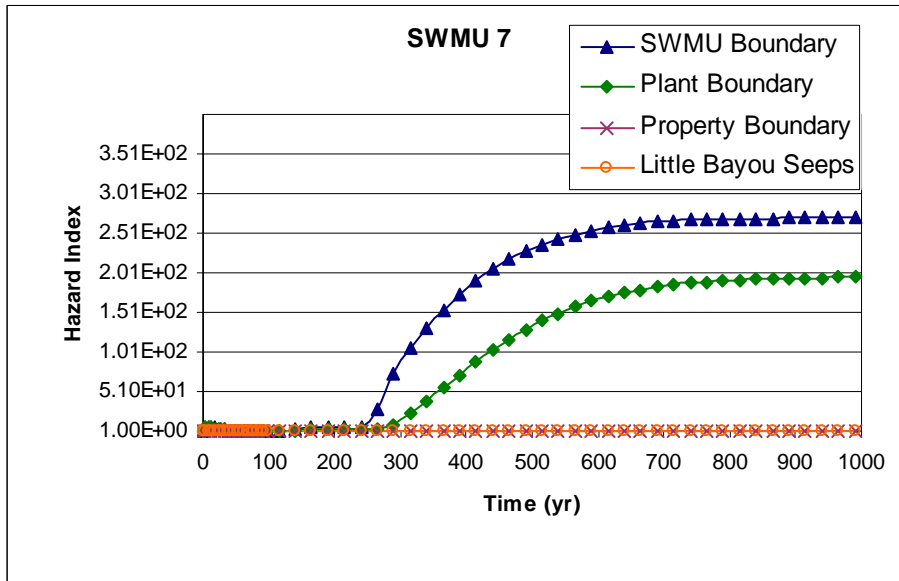


Figure F.12. Total HI from All Noncarcinogenic COCs at SWMU 7

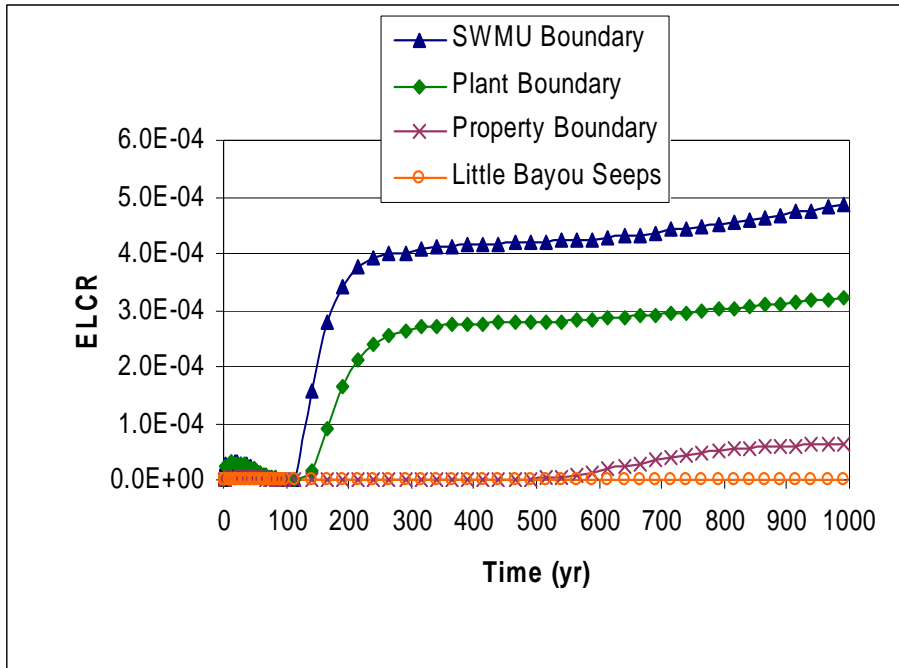


Figure F.13. Total ELCR from All Carcinogenic COCs at SWMU 30

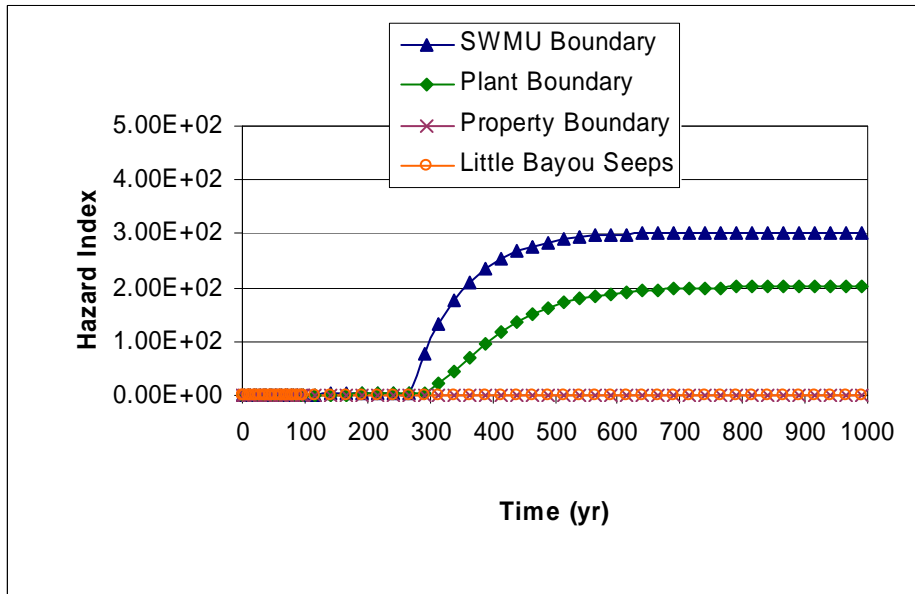


Figure F.14. Total HI from All Noncarcinogenic COCs at SWMU 30

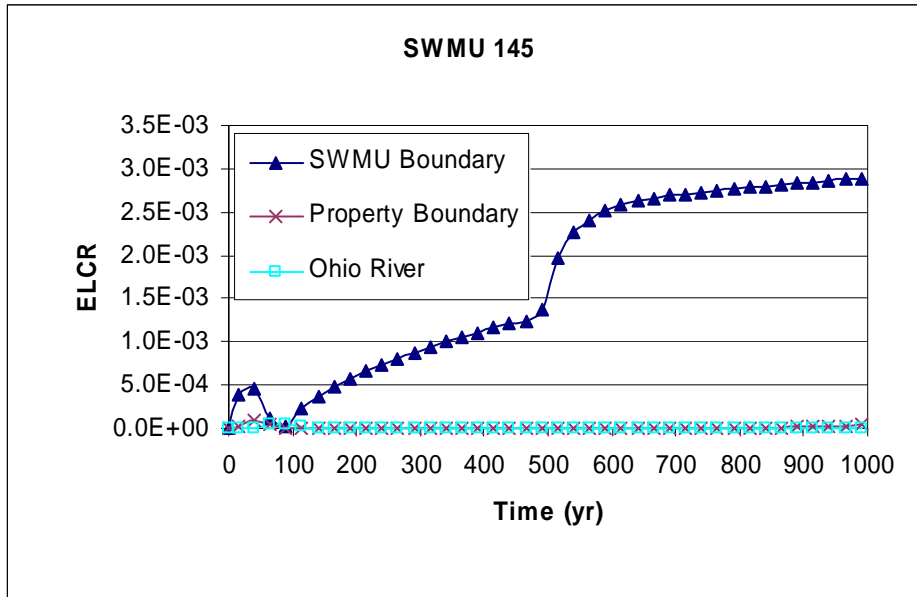


Figure F.15. Total ELCR from All Carcinogenic COCs at SWMU 145

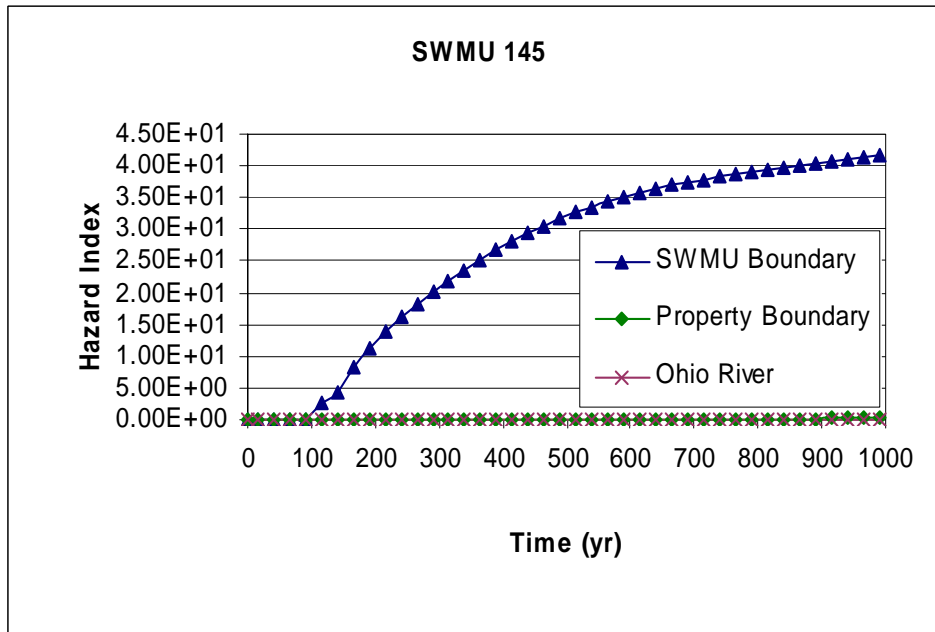


Figure F.16. Total HI from All Noncarcinogenic COCs at SWMU 145

The values for total hazard or risk for all COCs calculated from the NALs for residential groundwater use and the time at which the total risk or hazard peaks are provided in Tables F.61 and F.62.

As shown in Table F.68, the total HI for the child exceeded one for all SWMUs at the SWMU boundary. The total HI for the child for migration from the SWMU 2, SWMU 4, SWMU 5, SWMU 7, and SWMU 30 sources exceeds a HI of 1 at the plant boundary. The total HI for the child for migration from SWMU 2, SWMU 4, and SWMU 30 exceeds an HI of 1 at the property boundary. The total HI for the child for

migration from SWMU 2 and SWMU 4 exceeds an HI of 1 at the Ohio River. The total HI for the child for migration from SWMU 7 and SWMU 30 exceeds an HI of 1 at the Ohio River.

As shown in Table F.69, the total ELCRs resulting from COPC migration are above or equal to 1×10^{-6} at the SWMU boundary, the plant boundary, and the PGDP property boundary POEs for all sources. The total ELCRs were greater than 1×10^{-6} at the Little Bayou Seeps for SWMUs 3, 7, and 30. In addition, total ELCRs were greater than 1×10^{-6} at the Ohio River for SWMU 2, SWMU 4, and SWMU 145.

F.5.6 IDENTIFICATION OF LAND USE SCENARIOS, PATHWAYS, MEDIA, AND CONTAMINANTS OF CONCERN

This subsection outlines land use scenarios, exposure pathways, media, and COCs for each source area. As discussed in Section F.5.3, the results of the previous risk assessments for SWMUs 2, 3, 4, 5, 6, 7, and 30 are used for the risk characterization for soil. (The results of these previous risk assessments are discussed in Section F.1 and presented in Attachment F2 of this appendix.) Section F.8 presents the RGOs for each location and land use scenario.

F.5.6.1 Land Use Scenarios of Concern

To determine whether a land use scenario is of concern, quantitative risk and hazard results were compared to risk and hazard benchmarks for each land use scenario. The benchmarks used for this comparison were a) 1 for HI and b) 1×10^{-6} for ELCR. Land use scenarios with total HIs exceeding the benchmark of 1 are deemed land use scenarios of concern for non-cancer hazard. Land use scenarios with a total ELCR exceeding the benchmark of 1×10^{-6} are deemed land use scenarios of concern for cancer risk. These criteria were used in the previous risk assessments for SWMUs 4, 5, 6, 7, and 30 (DOE 1998a; DOE 2000). For the risk characterization of soil for SWMUs 2 and 3, land use scenarios of concern were determined by using EPA guidance and policy in effect at the time of the risk assessment (DOE 1994). The following are land uses of concern for BGOU at the SWMUs indicated.

- Industrial: SWMUs 2, 3, 4, 5, 6, 7, and 30
- Excavation: SWMUs 4, 5, 6, 7, and 30
- Recreational: SWMUs 5, 7, and 30
- On-Site Residential: SWMUs 2, 3, 4, 5, 6, 7, 30, and 145
- Off-Site Residential: SWMUs 2, 3, 4, 5, 7, 30, and 145

Table F.70 outlines all land use scenarios for all SWMUs that exceed *de minimis* risk or hazard levels.

Table F.68. Maximum Total HIs for Residential Groundwater Used at Each POE

SWMU	SWMU Boundary		Plant Boundary		Property Boundary		Little Bayou Seeps		Ohio River	
	Time (yr)	Peak Hazard	Time (yr)	Peak Hazard	Time (yr)	Peak Hazard	Time (yr)	Peak Hazard	Time (yr)	Peak Hazard
2	20	1.24E+03	20	1.87E+02	30	9.30E+01	NA	NA	80	3.46E+01
3	1,000	10.5	1,000	0.391	0	0	0	0	NA	NA
4	5	565	10	201	15	101	NA	NA	75	36.7
5	1,000	865	1,000	73.7	1,000	4.07E-02	NA	NA	0	0
7	1,000	270	1,000	195	1,000	0.998	36	3.46E-01	NA	NA
30	1,000	302	1,000	201	1,000	26.8	1,000	5.91E-03	NA	NA
145	1,000	19.9	NA	NA	1,000	5.16E-01	NA	NA	0	0

Table F.69. Maximum Total ELCRs for Residential Groundwater Use at Each POE

SWMU	SWMU Boundary		Plant Boundary		Property Boundary		Little Bayou Seeps		Ohio River	
	Time (yr)	Peak Risk	Time (yr)	Peak Risk	Time (yr)	Peak Risk	Time (yr)	Peak Risk	Time (yr)	Peak Risk
2	20	4.60E-02	25	6.74E-03	30	3.42E-03	NA	NA	85	1.28E-03
3	1,000	8.75E-04	55	9.94E-05	70	7.49E-05	140	4.42E-05	NA	NA
4	5	5.32E-02	10	1.69E-02	15	8.25E-03	NA	NA	75	2.81E-03
5	1,000	2.46E-04	1,000	4.73E-05	1,000	3.38E-06	NA	NA	155	4.79E-07
7	3	1.05E-02	4	9.65E-03	9	1.02E-03	34	3.56E-04	NA	NA
30	1,000	4.87E-04	1,000	2.19E-02	1,000	6.22E-05	1,000	1.71E-06	NA	NA
145	1,000	1.65E-03	NA	NA	40	1.01E-04	NA	NA	80	5.30E-05

F.5.6.2 Contaminants of Concern (Soil)

To make a determination about whether contaminants are of concern in soil, quantitative risk and hazard results over all pathways from the previous risk assessments for SWMUs 2, 3, 4, 5, 6, 7, and 30 (DOE 1994; DOE 1998a; DOE 2000) were compared to risk and hazard benchmarks for land use scenarios of concern. The benchmarks used for this comparison were a) 0.1 for HI and b) 1×10^{-6} for ELCR. For SWMUs 2 and 3, the WAG 22 RI calculated risks and hazards for soil for the two SWMUs as a combined unit; therefore, the soil entries for those two SWMUs are identical in the summary tables. The tables present risk and hazard values from the previous risk assessments calculated excluding consideration of lead because, at the time those assessments were written, a now withdrawn RfD for lead from Kentucky still was in use.

Contaminants with chemical-specific HIs or ELCRs exceeding these benchmarks are deemed COCs. Priority COCs are contaminants whose chemical-specific HI is greater than 1 or whose ELCR is greater than 1×10^{-4} for one or more scenarios. The following are priority COCs found in soil at individual SWMUs.

- SWMU 2—none
- SWMU 3—none
- SWMU 4—barium, beryllium, cadmium, chromium, iron, nickel, uranium, vanadium, Total dioxins/furans, Total PCBs, ^{234}U , and ^{238}U
- SWMU 5—aluminum, arsenic, beryllium, chromium, nickel, Total PAHs, and Total PCBs
- SWMU 6—beryllium, chromium, nickel, and Total PAHs
- SWMU 7—aluminum, antimony, arsenic, barium, beryllium, cadmium, chromium, copper, iron, manganese, nickel, uranium, vanadium, benzo(a)pyrene, dibenzo(a,h)anthracene, Aroclor-1254, Aroclor-1260, ^{239}Pu , ^{234}U , ^{235}U , $^{235/236}\text{U}$, and ^{238}U
- SWMU 30—aluminum, antimony, arsenic, barium, beryllium, cadmium, chromium, copper, iron, manganese, mercury, nickel, uranium, vanadium, benzo(a)anthracene, benzo(a)pyrene, benzo(b)fluoranthene, dibenzo(a,h)anthracene, indeno(1,2,3-cd)pyrene, Aroclor-1254, Aroclor-1260, ^{234}U , $^{235/236}\text{U}$, and ^{238}U

Table F.70. Scenarios for Which Human Health Risk Exceeds *De Minimis* Levels^a

Scenario	Location							
	SWMU 2	SWMU 3	SWMU 4	SWMU 5	SWMU 6	SWMU 7	SWMU 30	SWMU 145
Results for excess lifetime cancer risk:								
Current On-site Industrial Worker Exposure to Soil	X	X	X	X	X	X	X	NA
Future On-site Industrial Worker Exposure to Soil	X	X	X	X	X	X	X	NA
Exposure to Surface Water	NA	NA	NA	NA	NA	NA	NA	NA
Future On-site Excavation Worker Exposure to Soil/Soil and Waste	NA	NA	X	X	X	X	X	NA
Future On-site Recreational User Exposure to Game	NA	NA	---	X	---	X	X	NA
Exposure to Soil	NA	NA	NA	NA	NA	NA	NA	NA
Future On-site Rural Resident Exposure to Soil	NA	NA	X	X	X	X	X	NA
Exposure to Groundwater ^b	X	X	X	X	---	X	X	X
Vapor Intrusion ^c	X	X	X	---	---	X	X	---
Future Off-site Rural Resident Exposure to Groundwater ^b	X	X	X	X	---	X	X	X
Vapor Intrusion ^c	X	---	X	---	---	---	X	---
Result for Systematic Toxicity^b								
Current On-site Industrial Worker Exposure to Soil	---	---	X	---	---	X	X	NA
Future On-site Industrial Worker Exposure to Soil	---	---	X	---	---	X	X	NA
Future On-site Excavation Worker Exposure to Soil/Soil and Waste	NA	NA	X	X	X	X	X	NA
Future On-site Recreational User Exposure to Game	NA	NA	---	---	---	---	---	NA
Exposure to Soil	NA	NA	NA	NA	NA	NA	NA	NA
Exposure to Surface Water	NA	NA	NA	NA	NA	NA	NA	NA

Table F.70. Scenarios for Which Human Health Risk Exceeds *De Minimis* Levels^a (Continued)

Result for Systematic Toxicity^b									
Future On-site Rural Resident									
Exposure to Soil	NA	NA	X	X	X	X	X	X	NA
Exposure to Groundwater ^b	X	X	X	X	---	X	X	X	X
Vapor Intrusion ^c	X	X	X	---	---	X	X	X	X
Future Off-site Rural Resident									
Exposure to Groundwater ^b	X	---	X	---	---	X	X	X	---
Vapor Intrusion ^c	---	---	---	---	---	---	---	---	---

Notes: Scenarios where risk exceeds *de minimis* levels are marked with an X. Scenarios where risk did not exceed *de minimis* levels are marked with a ---. NA indicates that the scenario/land use combination was not assessed.

^a Consistent with the PGDP Risk Methods Document (DOE 2001), the *de minimis* levels used are a cumulative ELCR of 1×10^{-6} and a total HI of 1.

^b Systemic toxicity results summarized here for the resident and recreational user are for the child. The off-site POE considered is the property boundary.

^c Based on results of preliminary deterministic contaminant transport modeling. The POE is the property boundary. X indicates that the location contains a source of unacceptable off-site contamination, and --- indicates that the location is not a source of off-site contamination

F.5.6.3 Contaminants of Concern (Groundwater–Modeled from Soil)

Similarly for groundwater, to determine whether contaminants are of concern, quantitative risk, and hazard results over all pathways were compared to risk and hazard benchmarks for land use scenarios of concern. The benchmarks used for this comparison were a) 0.1 for HI and b) 1×10^{-6} for ELCR.

Contaminants with chemical-specific HIs or ELCRs exceeding these benchmarks were deemed COCs. Priority COCs are contaminants whose chemical-specific HI is greater than 1 or whose ELCR is greater than 1×10^{-4} for one or more scenarios. The following presents priority COCs found in groundwater at individual SWMUs.

- SWMU 2—arsenic; Aroclor-1248, manganese; uranium; *cis*-1,2-DCE; and TCE
- SWMU 3—arsenic, manganese, uranium, and ⁹⁹Tc
- SWMU 4—arsenic; manganese; *cis*-1,2-DCE; TCE; vinyl chloride; and ⁹⁹Tc
- SWMU 5—arsenic, manganese, uranium, and naphthalene
- SWMU 6—none
- SWMU 7—arsenic; 1,1-DCE; *cis*-1,2-DCE; Aroclor-1254; TCE; and vinyl chloride
- SWMU 30—arsenic
- SWMU 145—antimony, arsenic, manganese, Aroclor-1260, and ⁹⁹Tc

“Priority COCs” are identified in this section as an aid to risk managers during decision making. Table F.71 summarizes the COCs for both soil and groundwater.

F.5.6.4 Pathways of Concern

To determine whether pathways are of concern, the quantitative risks and hazards for each exposure route are summed over all contaminants and compared to benchmarks for land use scenarios of concern. The benchmarks used for this comparison were a) 0.1 for HI and b) 1×10^{-6} for ELCR. For soil, the quantitative risk and hazard results from the previous risk assessments for SWMUs 2, 3, 4, 5, 6, 7, and 30 (DOE 1994, DOE 1998a; DOE 2000) were used in the comparison. Exposure routes with HIs and ELCRs exceeding these benchmarks are considered POCs. These POCs are shown by SWMU in Table F.80. Each of the pathways included in the BHHRA is a POC for at least one SWMU.

F.5.6.5 Media of Concern

Media of concern are those media that appear in at least one POC. Because they contribute to at least one POC, soil and RGA groundwater are media of concern for all eight SWMUs. Table F.81 provides specific information concerning how each media contributes to risks and hazards for BGOU.

F.5.6.6 Summary of Risk Characterization

Tables F.72 to F.79 present summaries of the risk characterization by location considered in the BHHRA. They present land use scenarios of concern, COCs, and POCs. In addition, each table lists the following:

- Receptor risks for each land use scenario of concern;
- Percent contribution by pathway to the total risk; and
- Percent contribution each COC contributes to the total risk.

Table F.71. COCs and Exposure Medium for Residential Receptor at Each SWMU^a

COC ^b	SWMU 2 ^c			SWMU 3 ^c			SWMU 4 ^d			SWMU 5 ^d			SWMU 6 ^d			SWMU 7 ^e			SWMU 30 ^c			SWMU 145 ^f				
	Total Hazard Child	ELCR	Total	Total Hazard Child	ELCR	Total	Total Hazard Child	ELCR	Total	Total Hazard Child	ELCR	Total	Total Hazard Child	ELCR	Total	Total Hazard Child	ELCR	Total	Total Hazard Child	ELCR	Total	Total Hazard Child	ELCR			
<i>Inorganic Compounds</i>																										
Aluminum				S																						
Antimony																										
Arsenic																										
Barium																										
Beryllium																										
Cadmium																										
Chromium																										
Cobalt																										
Copper																										
Iron																										
Manganese																										
Mercury																										
Nickel																										
Selenium																										
Uranium																										
Vanadium																										
Zinc																										
<i>Organic Compounds</i>																										
1,1-DCE																										
cis-1,2-DCE																										
Benzo(a)anthracene																										
Benzo(a)pyrene																										
Benzo(b)fluoranthene																										
Benzo(k)fluoranthene																										
Bis(2-ethylhexyl)phthalate																										
Chrysene																										
Dibenzo(a,h)anthracene																										
Indeno(1,2,3-cd)pyrene																										
Naphthalene																										

Table F.71. COCs and Exposure Medium for Residential Receptor at Each SWMU (Continued)

COC ^b	SWMU 2 ^c			SWMU 3 ^c			SWMU 4 ^d			SWMU 5 ^d			SWMU 6 ^d			SWMU 7 ^c			SWMU 30 ^c			SWMU 145 ^f			
	Total Hazard Child	ELCR	GW	Total Hazard Child	ELC R	GW	Total Hazard Child	ELCR	GW	Total Hazard Child	ELCR	S	Total Hazard Child	ELCR	GW	Total Hazard Child	ELCR	GW	Total Hazard Child	ELCR	GW	Total Hazard Child	ELCR	GW	
TCE	GW			GW			GW																		
Vinyl Chloride				GW			GW																		
Total PAHs				GW			GW			S															
Aroclor-1248				GW																					
Aroclor-1254																									
Aroclor-1260				GW																					
Total PCBs				GW						S															
<i>Radionuclides^b</i>																									
Neptunium-237																									
Plutonium-129 ^b																									
Technetium-99				GW																					
Uranium-234				GW																					
Uranium-235				GW																					
Uranium-235/236				GW																					
Uranium-238				GW																					

^a Only COPCs that exceed a chemical-specific HI of 0.1 or a chemical-specific ELCR of 1×10^{-6} for a scenario with total HI > 1 are for the child resident and total ELCR > 1×10^{-6} for the resident are listed as COCs.

^b Plutonium-239/240 included with Plutonium-239.

^c Soil was not evaluated for residential receptor at SWMUs 2 and 3 (DOE 1994).

^d From DOE 2000.

^e From DOE 1998a.

^f A previous risk assessment was not performed for SWMU 145 and this site is covered by a cap; therefore, this SWMU was not evaluated for soil exposure.

"S" indicates this chemical is a COC for exposure to soil.

"GW" indicates this chemical is a COC for exposure to groundwater.

Table F.72. Summary of Risk Characterization for SWMU 2

Receptor	Total ELCR ^a	COCs	% Total ELCR	POCs	% Total ELCR	Total HI ^a	COCs	% Total HI	POCs	% Total HI
Current industrial worker/intruder at current concentrations (soil) (from WAG 22 RI Addendum ^b)	1.2E-05	²³⁵ U + daughters ²³⁸ U + daughters	83.8 10.7	External exposure	94.7	6.8E-03	*No COCs	*No COCs		
Future industrial worker at current concentrations (soil) (from WAG 22 RI Addendum ^b)	1.2E-04	Arsenic ²³⁵ U + daughters ²³⁸ U + daughters	2.8 83.9 10.7	Ingestion External exposure	4.7 94.7	7.0E-02	*No COCs		*No COCs	
Future child rural resident at current concentrations (RGA groundwater only)	NA	NA	NA	NA	NA	1.30E+0 3	Arsenic Manganese Uranium <i>cis</i> -1,2-DCE Naphthalene TCE	0.9 0.1 0.1 46.8 0.0 52.1	Ingestion Dermal Inhalation while showering Household inhalation	46.0 11.7 4.8 37.5

Table F.72. Summary of Risk Characterization for SWMU 2 (Continued)

Receptor	Total ELCR ^a	COCs	% Total ELCR	POCs	% Total ELCR	Total HI ^b	COCs	% Total HI	POCs	Total HI
Future adult rural resident at current concentrations (RGA groundwater only)	4.72E-02	Arsenic Aroclor-1248 Aroclor-1260 TCE ⁹⁹ Tc ²³⁴ U ²³⁸ U	2.0 0.4 0.1 98.0 0.0 0.0 0.0	Ingestion Dermal Inhalation while showering Household inhalation	19.9 11.0 7.8 61.3	3.79E+02	Arsenic Manganese Uranium <i>cis</i> -1,2-DCE Naphthalene TCE	0.9 0.1 0.1 36.8 0.0 62.1	Ingestion Dermal Inhalation while showering Household inhalation	45.0 23.9 3.5 27.5
Future child rural resident at modeled concentrations (RGA groundwater drawn at plant boundary)	NA	NA	NA	NA	NA	1.92E+02	Arsenic <i>cis</i> -1,2-DCE Naphthalene TCE	0.5 48 0.1 52	Ingestion Dermal Inhalation while showering Household inhalation	45 12.4 5.4 38
Future adult rural resident at modeled concentrations (RGA groundwater drawn at plant boundary)	6.82E-03	Arsenic TCE	1.1 98.9	Ingestion Dermal Inhalation while showering Household inhalation	19.2 11.1 7.9 61.8	5.08E+01	Arsenic <i>cis</i> -1,2-DCE Naphthalene TCE	0.5 16.2 0.1 83.1	Ingestion Dermal Inhalation while showering Household inhalation	60 32 1 7.2
Future child rural resident at modeled concentrations (RGA groundwater drawn at property boundary)	NA	NA	NA	NA	NA	9.56E+01	<i>cis</i> -1,2-DCE TCE	47.4 52.6	Ingestion Dermal Inhalation while showering Household inhalation	45.4 11.8 4.9 38.0
Future adult rural resident at modeled concentrations (RGA groundwater drawn at property boundary)	3.42E-03	TCE	100	Ingestion Dermal Inhalation while showering Household inhalation	18.3 11.2 8.0 62.5	2.79E+01	<i>cis</i> -1,2-DCE TCE	37.3 62.7	Ingestion Dermal Inhalation while showering Household inhalation	44.4 24.1 3.6 27.9
Future child rural resident at modeled concentrations (RGA groundwater drawn at Ohio River)	NA	NA	NA	NA	NA	2.25E+01	<i>cis</i> -1,2-DCE TCE	79.4 20.5	Ingestion Dermal Inhalation while showering Household inhalation	16.2 18.8 7.4 57.7
Future adult rural resident at modeled concentrations (RGA groundwater drawn at Ohio River)	1.28E-03	TCE	100	Ingestion Dermal Inhalation while showering Household inhalation	18.3 11.2 8.0 62.5	6.7E+00	<i>cis</i> -1,2-DCE TCE	61.2 38.7	Ingestion Dermal Inhalation while showering Household inhalation	15.5 37.7 5.3 41.5

ELCR = excess lifetime cancer risk; HI = hazard index; POC = point of contact; COC = contaminant of concern; Note: NA = ELCR not applicable to child and teen cohorts. ELCR for adult is for lifetime exposure and takes into account exposure as child and teen. .
^aNo COCs = there are no COCs or POCs at this SWMU for this endpoint (may apply to ELCR or HI).^bTotal ELCR and total HI represent total risk or hazard summed across all POCs for all COCs. ^b Remedial Investigation Addendum for Waste Area Grouping 22, Burial Grounds, at the Paducah Gaseous Diffusion Plant, Paducah, Kentucky, DOE/OR/07-1141&D2, September 1994 (DOE 1994), Attachment 2-1 through 2-6. This risk assessment combined SWMU 2 and 3.

Table F.73. Summary of Risk Characterization for SWMU 3

Receptor	Total ELCR ^a	COCs	% Total ELCR	POCs	% Total ELCR	Total HI ^c	COCs	% Total HI	POCs	% Total HI
Current industrial worker/intruder at current concentrations (soil) (from WAG 22 RI Addendum ^b)	1.2E-05	²³⁵ U + daughters ²³⁸ U + daughters	83.8 10.7	External exposure	94.7	6.8E-03	*No COCs		*No COCs	NE
Future industrial worker at current concentrations (soil) (from WAG 22 RI Addendum ^b)	1.2E-04	Arsenic ²³⁵ U + daughters ²³⁸ U + daughters	2.8 83.9 10.7	Ingestion External exposure	4.7 94.7	7.0E-02	*No COCs		*No COCs	NE
Future child rural resident at current concentrations (RGA groundwater only)	NA	NA	NA	NA	NA	2.03E+01	Arsenic Manganese Uranium	51.9 9.6 38.6	Ingestion Dermal	99.5 0.5
Future adult rural resident at current concentrations (RGA groundwater only)	1.20E-03	Arsenic ⁹⁹ Tc ²³⁸ U	72.4 25.3 2.3	Ingestion Dermal	99.8 0.2	5.83E+00	Arsenic Manganese Uranium	51.7 9.9 38.3	Ingestion Dermal	98.9 1.1
Future child rural resident at modeled concentrations (RGA groundwater drawn at plant boundary)	NA	NA	NA	NA	NA	3.98E-01	Arsenic	100	Ingestion	97.9
Future adult rural resident at modeled concentrations (RGA groundwater drawn at plant boundary)	1.32E-04	Arsenic ⁹⁹ Tc	24.6 75.4	Ingestion	99.9	1.12E-01	Arsenic	100	Ingestion	99.6
Future child rural resident at modeled concentrations (RGA groundwater drawn at property boundary)	NA	NA	NA	NA	NA		*No COCs		*No COCs	
Future adult rural resident at modeled concentrations (RGA groundwater drawn at property boundary)	7.46E-05	⁹⁹ Tc	100	Ingestion	100		*No COCs		*No COCs	

Table F.73. Summary of Risk Characterization for SWMU 3 (Continued)

Receptor	Total ELCR ^a	COCs	% Total ELCR	POCs	% Total ELCR	Total HI ^b	COCs	% Total HI	POCs	% Total HI
Future child rural resident at modeled concentrations (RGA groundwater drawn at Little Bayou Seeps)	NA	NA	NA	NA	NA	NA	*No COCs		*No COCs	
Future adult rural resident at modeled concentrations (RGA groundwater drawn at Little Bayou Seeps)	4.41E-05	⁹⁹ Tc	100.0				*No COCs		*No COCs	
Future child rural resident at modeled concentrations (RGA groundwater drawn at Ohio River)		Not a POE for groundwater from this SWMU.								
Future adult rural resident at modeled concentrations (RGA groundwater drawn at Ohio River)		Not a POE for groundwater from this SWMU.								

ELCR = excess lifetime cancer risk; HI = hazard index; POC = point of contact; COC = contaminant of concern

Note: NA = ELCR not applicable to child and teen cohorts. ELCR for adult is for lifetime exposure and takes into account exposure as child and teen.

* No COCs = there are no COCs or POCs.

^a Total ELCR and total HI represent total risk or hazard summed across all POCs for all COCs.

^b Remedial Investigation Addendum for Waste Area Grouping 22: Burial Grounds, at the Paducah Gaseous Diffusion Plant, Paducah, Kentucky, DOE/OR-07-1141&D2, September 1994 (DOE 1994), Attachment 2-1 through 2-6. This risk assessment combined SWMU 2 and 3.

Table F.74. Summary of Risk Characterization for SWMU 4

Receptor	Total ELCR ^a	COCs	% Total ELCR	POCs	% Total ELCR	Total HI ^c	COCs	% Total HI	POCs	% Total HI						
Current industrial worker at current concentrations (soil) (WAG 3 RI ^b)	5.4E-04	Beryllium ²³⁸ U	97	Dermal	97	3.62E+00	Beryllium	5	Dermal	5						
			2	External exposure	2		Chromium	45	Iron	24	Vanadium	24	Barium	2		
Future industrial worker at current concentrations (soil) (WAG 3 RI ^b)	5.4E-04	Beryllium ²³⁸ U	97	Dermal	97	3.62E+00	Beryllium	5	Dermal	5						
			2	External exposure	2		Chromium	45	Iron	24	Vanadium	24	Barium	2		
Future child rural resident at current concentrations (soil) (WAG 3 RI ^b)	NA	NA	NA	NA	NA	9.82E+01	Barium	2	Ingestion	1						
			NA	NA	NA		Beryllium	2	Dermal	21	Cadmium	2	Ingestion of vegetables	78		
Future adult rural resident at current concentrations (soil) (WAG 3 RI ^b)	4.3E-03	Beryllium Total PCB ²³⁴ U ²³⁸ U	72	Dermal	36	2.84E+01	Barium	9	Dermal	14						
			5	External exposure	2		Beryllium	2	Ingestion of vegetables	85	Cadmium	2	Chromium	22	Iron	63
Future child rural resident at current concentrations (RGA groundwater only)	NA	NA	NA	NA	NA	5.82E+02	Arsenic	1.0	Ingestion	67.2						
			NA	NA	NA		Manganese	0.2	Dermal	20.2	<i>cis</i> -1,2-DCE	6.1	Inhalation while showering	1.4	TCE	92.5
Future adult rural resident at current concentrations (RGA groundwater only)	5.41E-02	Arsenic TCE Vinyl chloride ⁹⁹ Tc	0.9	Ingestion	15.4	1.98E+02	Arsenic	0.8	Ingestion	56.5						
			67.7	Dermal	36.7		Manganese	0.2	Dermal	35.6	<i>cis</i> -1,2-DCE	4.1	Inhalation while showering	0.9	TCE	94.7
Future child rural resident at modeled concentrations (RGA groundwater drawn at plant boundary)	NA	NA	NA	NA	NA	2.04E+02	Arsenic	0.4	Ingestion	67.5						
			NA	NA	NA		<i>cis</i> -1,2-DCE	4.6	Dermal	20.6	TCE	94.4	Inhalation while showering	1.4	Vinyl chloride	0.1

Table F.74. Summary of Risk Characterization for SWMU 4 (Continued)

Receptor	Total ELCR ^a	COCs	% Total ELCR	POCs	% Total ELCR	Total HI ^a	COCs	% Total HI	POCs	% Total HI
Future adult rural resident at modeled concentrations (RGA groundwater drawn at plant boundary)	2.03E-02	Arsenic TCE Vinyl chloride ⁹⁹ Tc	0.4 98.0 0.9 0.7	Ingestion Dermal Inhalation while showering Household inhalation	13.6 7.2 5.2 74.0	6.97E+01	Arsenic <i>cis</i> -1,2-DCE TCE	0.4 3.0 96.6	Ingestion Dermal Inhalation while showering Household inhalation	56.5 36.1 0.8 6.6
Future child rural resident at modeled concentrations (RGA groundwater drawn at property boundary)	NA	NA	NA	NA	NA	1.03E+02	<i>cis</i> -1,2-DCE TCE Vinyl chloride	4.6 95.3 0.1	Ingestion Dermal Inhalation while showering Household inhalation	67.6 20.8 1.3 10.3
Future adult rural resident at modeled concentrations (RGA groundwater drawn at property boundary)	6.79E-03	TCE Vinyl chloride ⁹⁹ Tc	97.9 1.1 1.0	Ingestion Dermal Inhalation while showering Household inhalation	19.8 11.0 7.8 61.3	3.51E+01	<i>cis</i> -1,2-DCE TCE	3.1 96.8	Ingestion Dermal Inhalation while showering Household inhalation	56.4 36.3 0.8 6.4
Future child rural resident at modeled concentrations (RGA groundwater drawn at Ohio River)	NA	NA	NA	NA	NA	3.33E+01	<i>cis</i> -1,2-DCE TCE	1.7 98.2	Ingestion Dermal Inhalation while showering Household inhalation	74.6 22.9 1.4 1.0
Future adult rural resident at modeled concentrations (RGA groundwater drawn at Ohio River)	2.43E-03	TCE Vinyl chloride ⁹⁹ Tc	98.2 0.9 0.9	Ingestion Dermal Inhalation while showering Household inhalation	19.6 11.0 7.9 61.5	1.26E+01	<i>cis</i> -1,2-DCE TCE	3.0 96.9	Ingestion Dermal Inhalation while showering Household inhalation	56.4 36.3 0.8 6.4
Future child recreational user at current concentrations (soil) (WAG 3 RI ^b)	NA	NA	NA	NA	NA	< 1	*No COCs		*No COCs	
Future teen recreational user at current concentrations (soil) (WAG 3 RI ^b)	NA	NA	NA	NA	NA	< 1	*No COCs		*No COCs	
Future adult recreational user at current concentrations (soil) (WAG 3 RI ^b)	< 1.0E-06	*No COCs		*No COCs		< 1	*No COCs		*No COCs	

Table F.74. Summary of Risk Characterization for SWMU 4 (Continued)

Receptor	Total ELCR ^a	COCs	% Total ELCR	POCs	% Total ELCR	Total HI ^b	COCs	% Total HI	POCs	% Total HI
Future excavation worker at current concentrations (soil and waste) (WAG 3 RI ^b)	2.7E-03	Arsenic	1	Ingestion	37	2.61E+00	Aluminum	8	Ingestion	13
		Beryllium	7	Dermal	10		Arsenic	4	Dermal	87
		Total dioxins/furans	4	External exposure	54		Barium	2		
		Total PCB	2				Beryllium	2		
		²²⁶ Ra	2				Cadmium	1		
		Total uranium ^c	83				Chromium	24		
		²³⁸ U	1							
							Iron	24		
							Manganese	14		
							Vanadium	20		

ELCR = excess lifetime cancer risk; HI = hazard index; POC = point of contact; COC = contaminant of concern

Note: NA = ELCR not applicable to child and teen cohorts. ELCR for adult is for lifetime exposure and takes into account exposure as child and teen.

* No COCs = There are no COCs or POCs.

^a Total ELCR and total HI represent total risk or hazard summed across all POCs for all COCs.

^b Remedial Investigation Report for Waste Area Grouping 3 at the Paducah Gaseous Diffusion Plant, Paducah, Kentucky, DOE/OR/07-1895&DJ, September 2000 (DOE 2000), Table 1.55. In this table, lead has been excluded as a COC.

^c Risk associated with total uranium at SWMU 4 was calculated using a total uranium analytical result in pCi/g units and toxicity information for ²³⁸U. Individual isotopes also were included in the risk calculation, resulting in a double-counting of risk due to uranium isotopes. This approach likely accounts for the discrepancy between risk related to total uranium and ²³⁸U.

Table F.75. Summary of Risk Characterization for SWMU 5

Receptor	Total ELCR ^a	COCs	% Total ELCR	POCs	% Total ELCR	Total HI ^c	COCs	% Total HI	POCs	% Total HI
Current industrial worker at current concentrations (soil) (WAG 3 R) ^b	4.1E-04	Arsenic Beryllium Total PAH	6 49 45	Ingestion Dermal	2 98	< 1	*No COCs	*No COCs		
Future industrial worker at current concentrations (soil) (WAG 3 R) ^b	4.1E-04	Arsenic Beryllium Total PAH	6 49 45	Ingestion Dermal	2 98	< 1	*No COCs	*No COCs		
Future child rural resident at current concentrations (soil) (WAG 3 R) ^b	NA	NA	NA	NA	NA	4.62E+01	Aluminum Arsenic Beryllium Chromium Nickel Zinc	24 53 1 17 3 1	Ingestion Dermal Ingestion of vegetables	1 12 87
Future adult rural resident at current concentrations (soil) (WAG 3 R) ^b	>1.0E-02*	Arsenic Beryllium Total PAH Total PCB	21 9 68 2	Dermal Ingestion of vegetables	9 90	1.39E+01	Aluminum Arsenic Beryllium Chromium Nickel Zinc	24 55 1 15 3 1	Dermal Ingestion of vegetables	8 92
Future child rural resident at current concentrations (RGA groundwater only)	NA	NA	NA	NA	NA	8.15E+01	Uranium Arsenic Manganese Naphthalene	90.3 3.6 2.6 3.4	Ingestion Dermal Inhalation while showering Household inhalation	96.4 0.2 0.4 3.0
Future adult rural resident at current concentrations (RGA groundwater only)	2.52E-04	Arsenic ⁹⁹ Tc	97.2 2.8	Ingestion Dermal	99.7 0.3	2.31E+01	Uranium Arsenic Manganese Naphthalene	91.0 3.7 2.7 2.6	Ingestion Dermal Inhalation while showering Household inhalation	97.1 0.3 0.3 2.3
Future child rural resident at modeled concentrations (RGA groundwater drawn at plant boundary)	NA	NA	NA	NA	NA	6.56E+00	Uranium Arsenic Manganese Naphthalene	80.9 8.7 2.8 7.5	Ingestion Household inhalation	92.4 6.6
Future adult rural resident at modeled concentrations (RGA groundwater drawn at plant boundary)	4.99E-05	Arsenic ⁹⁹ Tc	94.5 5.5	Ingestion	99.7	1.84E+00	Uranium Arsenic Naphthalene Manganese	82.4 8.9 5.8 2.1	Ingestion Household inhalation	93.9 5.1

Table F.75. Summary of Risk Characterization for SWMU 5 (Continued)

Receptor	Total ELCR ^a	COCs	% Total ELCR	POCs	% Total ELCR	Total HI ^a	COCs	% Total HI	POCs	% Total HI
Future child rural resident at modeled concentrations (RGA groundwater drawn at property boundary)	NA	NA	NA	NA	NA	2.28E-01	Naphthalene	82.2	Household inhalation	72.0
Future adult rural resident at modeled concentrations (RGA groundwater drawn at property boundary)	4.81E-06	Arsenic ^{99m} Tc	69.9 30.1	Ingestion	99.8	<0.1	*No COCs		*No COCs	
Future child rural resident at modeled concentrations (RGA groundwater drawn at property boundary)	NA	NA	NA	NA	NA		*No COCs		*No COCs	
Future adult rural resident at modeled concentrations (RGA groundwater drawn at Ohio River)	NA	*No COCs		*No COCs			*No COCs		*No COCs	
Future child recreational user at current concentrations (soil) (WAG 3 RI ^b)	NA	NA	NA	NA	NA	<1	*No COCs		*No COCs	
Future teen recreational user at current concentrations (soil) (WAG 3 RI ^b)	NA	NA	NA	NA	NA	<1	*No COCs		*No COCs	
Future adult recreational user at current concentrations (soil) (WAG 3 RI ^b)	1.0E-05	Arsenic Total PAH Total PCB	2 96 2	Ingestion of venison Ingestion of rabbit Ingestion of quail	16 63 21	<1	*No COCs		*No COCs	
Future excavation worker at current concentrations (soil and waste) (WAG 3 RI ^b)	2.9E-04	Arsenic Beryllium Total PAH Total PCB	8 62 28 1	Ingestion Dermal	13 87	2.16E+00	Aluminum Arsenic Barium Beryllium Chromium Iron Manganese	9 7 2 3 18 38 22	Ingestion Dermal	18 82

ELCR = excess lifetime cancer risk; HI = hazard index; POC = point of contact; COC = contaminant of concern
 Note: NA = ELCR not applicable to child and teen cohorts. ELCR for adult is for lifetime exposure and takes into account exposure as child and teen.
 * No COCs = There are no COCs or POCs.

* = The ELCR is approximate because the linearized multistage model returns imprecise values at risks > 1.0E-02.

^a Total ELCR and total HI represent total risk or hazard summed across all POCs for all COCs.

^b Remedial Investigation Report for Waste Area Grouping 3 at the Paducah Gaseous Diffusion Plant, Paducah, Kentucky, DOE/OR/07-1895&D1, September 2000 (DOE 2000), Table 1.56. In this table, lead has been excluded as a COC.

Table F.76. Summary of Risk Characterization for SWMU 6

Receptor	Total ELCR ^a	COCs	% Total ELCR	POCs	% Total ELCR	Total HI ^a	COCs	% Total HI	POCs	% Total HI
Current industrial worker at current concentrations (soil) (WAG 3 RI ^b)	2.4E-04	Beryllium Total PAH	90	Dermal	99	<1	*No COCs		*No COCs	
Future industrial worker at current concentrations (soil) (WAG 3 RI ^b)	2.4E-04	Beryllium Total PAH	90	Dermal	99	<1	*No COCs		*No COCs	
Future child rural resident at current concentrations (soil) (WAG 3 RI ^b)	NA	NA	NA	NA	NA	9.38E+00	Beryllium Chromium Nickel Zinc	8	Dermal	34
Future adult rural resident at current concentrations (soil) (WAG 3 RI ^b)	2.4E-03	Beryllium Total PAH	54	Dermal	69	2.57E+00	Beryllium Chromium Nickel Zinc Manganese	72	Ingestion of vegetables	65
Future child rural resident at current concentrations (RGA groundwater only)		*No COCs		*No COCs		1.77E-01		5		
Future adult rural resident at current concentrations (RGA groundwater only)		*No COCs		*No COCs		5.18E-02		7	Dermal	24
Future child rural resident at modeled concentrations (RGA groundwater drawn at plant boundary)		*No COCs		*No COCs				17	Ingestion of vegetables	75
Future adult rural resident at modeled concentrations (RGA groundwater drawn at plant boundary)		*No COCs		*No COCs				6		
Future child rural resident at modeled concentrations (RGA groundwater drawn at property boundary)		*No COCs		*No COCs				100	Ingestion Dermal	97.9
Future adult rural resident at modeled concentrations (RGA groundwater drawn at property boundary)		*No COCs		*No COCs						2.1
Future child rural resident at modeled concentrations (RGA groundwater drawn at property boundary)		*No COCs		*No COCs						95.7
Future adult rural resident at modeled concentrations (RGA groundwater drawn at property boundary)		*No COCs		*No COCs						4.3
Future child rural resident at modeled concentrations (RGA groundwater drawn at Ohio River)		*No COCs		*No COCs						
Future adult rural resident at modeled concentrations (RGA groundwater drawn at Ohio River)		*No COCs		*No COCs						

Table F.76. Summary of Risk Characterization for SWMU 6 (Continued)

Receptor	Total ELCR ^a	COCs	% Total ELCR	POCs	% Total ELCR	Total HI ^a	COCs	% Total HI	POCs	% Total HI
Future child recreational user at current concentrations (soil) (WAG 3 RI ^b)	NA	NA	NA	NA	NA	<1	*No COCs		*No COCs	
Future teen recreational user at current concentrations (soil) (WAG 3 RI ^b)	NA	NA	NA	NA	NA	<1	*No COCs		*No COCs	
Future adult recreational user at current concentrations (soil) (WAG 3 RI ^b)	< 1.0E-06	*No COCs		*No COCs		<1	*No COCs		*No COCs	
Future excavation worker at current concentrations (soil and waste) WAG 3 RI ^b)	2.3E-04	Beryllium Total PAH	90 9	Ingestion Dermal	5 95	2.44E+00	Aluminum Barium Beryllium Chromium Iron Manganese Vanadium	8 2 3 15 32 15 26	Ingestion Dermal	12 88

ELCR = excess lifetime cancer risk; HI = hazard index; POC = point of contact; COC = contaminant of concern

Note: NA = ELCR not applicable to child and teen cohorts. ELCR for adult is for lifetime exposure and takes into account exposure as child and teen.

- = There are no COCs or POCs.

^a Total ELCR and total HI represent total risk or hazard summed across all POCs for all COCs.

^b Remedial Investigation Report for Waste Area Grouping 3 at the Paducah Gaseous Diffusion Plant, Paducah, Kentucky, DOE/OR/07-1895&D1, September 2000 (DOE 2000), Table 1.57. In this table, lead has been excluded as a COC.

Table F.77. Summary of Risk Characterization for SWMU 7

Receptor	Total ELCR ^a	COCs	% Total ELCR	POCs	% Total ELCR	Total HI ^b	COCs	% Total HI	POCs	% Total HI
Current industrial worker at current concentrations (soil) (from WAG 22 RI ^b)	3.8E-03	Arsenic	0.6	Ingestion	0.5	5.0E+00	Aluminum	4.1	Ingestion	4.1
		Beryllium	97.6	Dermal	97.4		Antimony	4.4	Dermal	4.4
		Benzo(a)anthracene	<0.1	External exposure	2.5		Arsenic	2.6		
		Benzo(a)pyrene	0.3				Beryllium	9.6		
		Benzo(b)fluoranthene	<0.1				Chromium	13.6		
		Dibenzo(a,h)anthracene	0.4				Iron	20.6		
		Indeno(1,2,3-cd)pyrene	0.1				Manganese	10.7		
		Indeno(1,2,3-cd)pyrene	<0.1				Uranium	13.7		
		²³⁷ Np	<0.1				Vanadium	17.7		
		²³⁴ U	<0.1							
		²³⁵ U	0.2							
		^{235,236} U	0.3							
		²³⁸ U	2.1							
		Future industrial worker at current concentrations (soil) (from WAG 22 RI ^b)	3.9E-03	Arsenic	0.6		Ingestion	0.5	5.0E+00	Aluminum
Beryllium	96.0			Dermal	97.1	Antimony	4.4	Dermal		4.4
Benzo(a)anthracene	<0.1			External exposure	2.4	Arsenic	2.6			
Benzo(a)pyrene	0.3					Beryllium	9.6			
Benzo(b)fluoranthene	<0.1					Chromium	13.6			
Dibenzo(a,h)anthracene	0.4					Iron	20.6			
Indeno(1,2,3-cd)pyrene	0.1					Manganese	10.7			
Indeno(1,2,3-cd)pyrene	<0.1					Uranium	13.7			
²³⁷ Np	<0.1					Vanadium	17.7			
²³⁴ U	<0.1									
²³⁵ U	0.2									
^{235,236} U	0.3									
²³⁸ U	2.1									
Future child rural resident at current concentrations (soil) (from WAG 22 RI ^b)	NA			NA	NA	NA	NA	3.7E+02		Aluminum
		NA	NA	NA	Antimony	0.9	Dermal		7.7	
		NA	NA	NA	Arsenic	6.2	Ingestion of vegetables from soil		90.9	
		NA	NA	NA	Barium	0.3				
		NA	NA	NA	Beryllium	1.3				
		NA	NA	NA	Cadmium	0.8				
		NA	NA	NA	Chromium	2.7				
		NA	NA	NA	Cobalt	0.1				
		NA	NA	NA	Copper	0.3				
		NA	NA	NA	Iron	19.7				
		NA	NA	NA	Manganese	1.9				
		NA	NA	NA	Nickel	0.4				
		NA	NA	NA	Uranium	58.4				
		NA	NA	NA	Vanadium	2.4				
NA	NA	NA	Zinc	0.2						
NA	NA	NA	Aroclor-1254	1.7						

Table F.77. Summary of Risk Characterization for SWMU 7 (Continued)

Receptor	Total ELCR ^a	COCs	% Total ELCR	POCs	% Total ELCR	Total HI ^b	COCs	% Total HI	POCs	% Total HI
Future adult rural resident at current concentrations (soil) (from WAG 22 RI ^b)	3.4E-02	Arsenic	7.3	Ingestion Dermal	0.5	1.1E+02	Aluminum	2.7	Ingestion	0.5
		Beryllium	65.4	External exposure	33.0		Antimony	0.8	Dermal	5.0
		Aroclor-1254	0.2	Ingestion of vegetables from soil	1.9		Arsenic	6.5	Ingestion of vegetables from soil	94.6
		Aroclor-1260	0.4		64.6		Barium	0.3		
		Benzo(a)anthracene	0.2				Beryllium	1.1		
		Benzo(a)pyrene	1.7				Cadmium	0.8		
		Benzo(b)fluoranthene	0.2				Chromium	2.3		
		Benzo(k)fluoranthene	<0.1				Copper	0.3		
		Dibenzo(a,h)anthracene	1.9				Iron	19.8		
		Indeno(1,2,3-cd)pyrene	0.3				Manganese	1.6		
		²³⁷ Np	0.2				Nickel	0.4		
		²³⁹ Pu	0.4				Uranium	59.5		
		²³⁴ U	3.3				Vanadium	2.0		
		²³⁵ U	0.3				Zinc	0.2		
		^{235/236} U	0.5				Aroclor-1254	1.7		
		²³⁸ U	17.6							
		Future child rural resident at current concentrations (RGA groundwater only)	NA	NA	NA	NA	NA	1.89E+01	Arsenic	30.2
Future adult rural resident at current concentrations (RGA groundwater only)	3.13E-03	Arsenic	15.1	Ingestion	61.2	6.39E+00	Arsenic	25.5	Ingestion	51.4
		1,1-DCE	66.4	Dermal contact	3.7		Manganese	3.2	Dermal contact	37.2
		Total PCBs	0.2	Inhalation while showering	4.9		Uranium	2.5	Inhalation while showering	1.3
		TCE	4.1	Inhalation during household use	30.3		1,1-DCE	3.1	Inhalation household use	10.1
		Vinyl chloride	11.9				<i>cis</i> -1,2-DCE	4.5		
		⁹⁹ Tc	1.6				Total PCBs	31.4		
		²³⁴ U	0.4				TCE	27.1		
		²³⁸ U	0.4				Vinyl chloride	2.7		

Table F.77. Summary of Risk Characterization for SWMU 7 (Continued)

Receptor	Total ELCR ^a	COCs	% Total ELCR	POCs	Total ELCR	Total HI ^b	COCs	% Total HI	POCs	% Total HI
Future child rural resident at modeled concentrations (RGA groundwater drawn at plant boundary)	NA	NA	NA	NA	NA	1.45E+01	Arsenic Manganese Uranium 1,1-DCE <i>cis</i> -1,2-DCE Total PCBs TCE	27.9 3.6 2.8 5.4 7.9 17.2 31.2	Ingestion Dermal contact Inhalation while showering Inhalation household use	62.3 18.7 2.2 16.9
Future adult rural resident at modeled concentrations (RGA groundwater drawn at plant boundary)	2.98E-03	Arsenic 1,1-DCE Total PCBs TCE Vinyl chloride ⁹⁹ Tc ²³⁴ U ²³⁸ U	11.2 63.9 0.2 10.3 12.3 1.5 0.3 0.3	Ingestion Dermal contact Inhalation while showering Inhalation during household use	55.4 3.4 4.7	4.78E+00	Arsenic Manganese Uranium 1,1-DCE <i>cis</i> -1,2-DCE Total PCBs TCE Vinyl chloride	24.2 3.1 2.4 3.8 5.5 24.8 32.9 3.3	Ingestion of groundwater Dermal contact Inhalation household use	53.8 33.8 11.0
Future child rural resident at modeled concentrations (RGA groundwater drawn at property boundary)	NA	NA	NA	NA	NA	1.97E+00	Arsenic 1,1-DCE <i>cis</i> -1,2-DCE Total PCBs	38.1 5.3 8.4 12.4	Ingestion Dermal contact Inhalation household use	66.3 15.8 15.9
Future adult rural resident at modeled concentrations (RGA groundwater drawn at property boundary)	4.11E-04	Arsenic 1,1-DCE TCE Vinyl chloride ⁹⁹ Tc	15.1 61.8 10.7 8.7 3.6	Ingestion Dermal contact Inhalation while showering Inhalation during household use	56.7 3.2 4.5	6.36E-01	Arsenic Total PCBs TCE	32.9 33.9 18.4 35.5	Ingestion Dermal contact	58.8 29.3
Future child rural resident at modeled concentrations (RGA groundwater drawn at Little Bayou Seeps)	NA	NA	NA	NA	NA	3.373E-01	TCE	61.0	Ingestion Inhalation household use	52.5 30.0
Future adult rural resident at modeled concentrations (RGA groundwater drawn at Little Bayou Seeps)	1.28E-04	1,1-DCE TCE Vinyl chloride ⁹⁹ Tc	72.6 12.3 9.5 5.7	Ingestion Dermal contact Inhalation while showering Inhalation during household use	49.6 3.6 5.3 41.4	1.15E-01	*No COCs		*No COCs	

Table F.77. Summary of Risk Characterization for SWMU 7 (Continued)

Receptor	Total ELCR ^a	COCs	% Total ELCR	POCs	% Total ELCR	Total HI ^c	COCs	% Total HI	POCs	% Total HI
Future child rural resident at modeled concentrations (RGA groundwater drawn at Ohio River)		Not a POE for groundwater from this SWMU.								
Future adult rural resident at modeled concentrations (RGA groundwater drawn at Ohio River)		Not a POE for groundwater from this SWMU.								
Future child recreational user at current concentrations (from WAG 22 RI ^b)	NA	NA	NA	NA	NA	7.3E-02	*No COCs		*No COCs	
Future teen recreational user at current concentrations (from WAG 22 RI ^b)	NA	NA	NA	NA	NA	6.4E-02	*No COCs		*No COCs	
Future adult recreational user at current concentrations (from WAG 22 RI ^b)	1.1E-05	Aroclor-1260 Benzo(a)pyrene Dibenzo(a,h)anthracene	18.6 9.5 42.5 15.7	Ingestion of deer Ingestion of rabbit Ingestion of quail	10.0 70.9 21.8	7.5E-02	*No COCs		*No COCs	
Future excavation worker at current concentrations (soil) (from WAG 22 RI ^b)	1.6E-03	Arsenic Beryllium Benzo(a)pyrene Dibenzo(a,h)anthracene	1.8 42.2 0.1 1.7	Ingestion Dermal External exposure	25.6 43.8 32.5	5.4E+00	Aluminum Antimony Arsenic Chromium Copper Iron Manganese Nickel Uranium Vanadium	5.0 11.3 3.4 17.6 2.9 21.3 11.0 3.9 7.5 10.9	Ingestion Dermal	18.4 81.5

ELCR = excess lifetime cancer risk; HI = hazard index; POC = point of contact; COC = contaminant of concern
 Note: NA = ELCR not applicable to child and teen cohorts. ELCR for adult is for lifetime exposure and takes into account exposure as child and teen.
^a No COCs = There are no COCs or POCs.
^b Total ELCR and total HI represent total risk or hazard summed across all POCs for all COCs.
^c Remedial Investigation Report for Solid Waste Management Units 7 and 30 of Waste Area Group 22 at the Paducah Gaseous Diffusion Plant, Paducah, Kentucky, DOE/OR/07-1604-V1&D2, January 1998 (DOE 1998b), Tables 1.59 through 1.68, excluding lead as a COC.

Table F.78. Summary of Risk Characterization for SWMU 30

Receptor	Total ELCR ^a	COCs	% Total ELCR	POCs	Total ELCR	Total HI ^a	COCs	% Total HI	POCs	% Total HI
Current industrial worker at current concentrations (soil) (from WAG 22 RI ^b)	3.7E-03	Arsenic	0.5	Ingestion	0.5	4.4E+00	Aluminum	5.1	Ingestion	5.1
		Beryllium	97.5	Dermal			Antimony	3.7		
		Aroclor-1260	0.1	External exposure			Arsenic	2.7		
		Benzo(a)anthracene	0.1				Beryllium	10.8		
		Benzo(a)pyrene	0.8				Cadmium	3.5		
		Benzo(b)fluoranthene	0.1				Chromium	13.5		
		Dibenzo(a,h)anthracene	0.3				Iron	19.8		
		Indeno(1,2,3-cd)pyrene	0.1				Manganese	11.3		
		²³⁷ Np	<0.1				Uranium	9.0		
		²³⁴ U	<0.1				Vanadium	17.6		
		²³⁵ U	0.2							
		^{235/236} U	0.3							
		²³⁸ U	1.4							
		Future industrial worker at current concentrations (soil) (from WAG 22 RI ^b)	3.8E-03	Arsenic			0.5	Ingestion	0.5	
Beryllium	96.2			Dermal	Antimony	3.7				
Aroclor-1260	0.1			External exposure	Arsenic	2.7				
Benzo(a)anthracene	0.1				Beryllium	10.8				
Benzo(a)pyrene	0.8				Cadmium	3.5				
Benzo(b)fluoranthene	0.1				Chromium	13.5				
Dibenzo(a,h)anthracene	0.3				Iron	19.8				
Indeno(1,2,3-cd)pyrene	0.1				Manganese	11.3				
²³⁷ Np	<0.1				Uranium	9.0				
³⁴ U	<0.1				Vanadium	17.6				
³⁸ U	0.2									
^{235/236} U	0.3									
²³⁸ U	1.4									
Future child rural resident at current concentrations (soil) (from WAG 22 RI ^b)	NA			NA	NA	NA	NA	2.6E+02		Aluminum
		NA	NA	Antimony	0.9					
		NA	NA	Arsenic	7.5					
		NA	NA	Barium	0.6					
		NA	NA	Beryllium	1.8					
		NA	NA	Cadmium	2.2					
		NA	NA	Chromium	3.2					
		NA	NA	Copper	0.6					
		NA	NA	Iron	22.6					
		NA	NA	Manganese	2.5					
		NA	NA	Mercury	0.7					
		NA	NA	Nickel	0.8					
		NA	NA	Uranium	46.8					
		NA	NA	Vanadium	3.0					
NA	NA	Zinc	0.2							
NA	NA	Aroclor-1254	2.6							
				Ingestion of vegetables from soil	89.3					

Table F. 78. Summary of Risk Characterization for SWMU 30 (Continued)

Receptor	Total ELCR ^a	COCs	% Total ELCR	POCs	% Total ELCR	Total HI ^b	COCs	% Total HI	POCs	% Total HI
Future adult rural resident at current concentrations (soil) (from WAG 22 RI ^b)	3.2E-02	Arsenic	6.8	Ingestion	0.5	7.9E+01	Aluminum	4.1	Ingestion	0.5
		Beryllium	66.7	Dermal	35.4		Antimony	0.8	Dermal	6.1
		Aroclor-1254	0.2	External exposure	1.3		Arsenic	7.9	Ingestion of vegetables from soil	93.4
		Aroclor-1260	1.8	Ingestion of vegetables from soil	62.8		Barium	0.6		
		Benzo(a)anthracene	0.4				Beryllium	1.5		
		Benzo(b)fluoranthene	4.4				Cadmium	2.2		
		Benzo(k)fluoranthene	0.5				Chromium	2.9		
		bis(2-ethylhexyl)phthalate	<0.1				Copper	0.6		
		Chrysene	<0.1				Iron	22.8		
		Dibenz(a,h)anthracene	1.7				Manganese	2.1		
		Indeno(1,2,3-cd)pyrene	0.4				Mercury	0.7		
		²³⁷ Np	0.2				Nickel	0.9		
		²³⁴ U	4.5				Uranium	47.5		
		²³⁵ U	0.3				Vanadium	2.4		
^{235/238} U	0.6			Zinc	0.2					
²³⁸ U	11.5			Aroclor-1254	2.7					
Future child rural resident at current concentrations (RGA groundwater only)	NA	NA	NA	NA	NA	9.14E+00	Arsenic	63.8	Ingestion	93.3
Future adult rural resident at current concentrations (RGA groundwater only)	5.44E-04	1,1-DCE	88.6	Ingestion	95.3	3.31E+00	Manganese	7.1	Dermal contact	8.8
		TCE	0.3	Dermal contact	0.9		Selenium	2.5	Inhalation while showering	0.2
		⁹⁹ Tc	5.2	Inhalation while showering	0.4		Uranium	11.6	Inhalation household use	1.2
		²³⁴ U	2.9	Inhalation household use	3.4		1,1-DCE	23.9		
		²³⁸ U	1.0				TCE	4.4		
			1.93				Arsenic	63.1	Ingestion of groundwater	91.1
			NA	NA	NA		Manganese	8.7	Dermal contact	1.7
			NA	NA	NA		Selenium	2.6	Inhalation while showering	0.1
			NA	NA	NA		1,1-DCE	12.3	Inhalation household use	7.1
			NA	NA	NA		TCE	0.1		
Future adult rural resident at modeled concentrations (RGA groundwater drawn at plant boundary)	3.75E-04	Arsenic	85.6	Ingestion	93.6	2.10E+00	Arsenic	52.9	Ingestion	76.1
		1,1-DCE	0.5	Dermal contact	1.1		Manganese	7.4	Dermal contact	3
		TCE	7.1	Inhalation while showering	0.6		Selenium	2.2	Inhalation while showering	0
		⁹⁹ Tc	3.9	Inhalation household use	4.7		Uranium	10.5	Inhalation household use	20.8
		²³⁴ U	1				1,1-DCE	0.4		
		²³⁸ U	1.9				TCE	26.6		

Table F.78. Summary of Risk Characterization for SWMU 30 (Continued)

Receptor	Total ELCR ^a	COCs	% Total ELCR	POCs	% Total ELCR	Total HI ^r	COCs	% Total HI	POCs	% Total HI
Future child rural resident at modeled concentrations (RGA groundwater drawn at property boundary)	NA	NA	NA	NA	NA	8.40E-01	Arsenic Selenium 1,1-DCE TCE Manganese	89.2 2.1 0.1 8.5 0.1	Ingestion Dermal contact Inhalation while showering Inhalation household use	94.2 1.1 0 4.6
Future adult rural resident at modeled concentrations (RGA groundwater drawn at property boundary)	6.85E-05	Arsenic 1,1-DCE TCE Technetium-99	90.6 0.2 3.5 5.7	Ingestion Dermal contact Inhalation while showering Inhalation household use	96.7 0.7 0.3 2.3	2.76E-01	Arsenic Selenium 1,1-DCE TCE Manganese	77.9 1.8 0.3 19.9 0.1	Ingestion Dermal contact Inhalation while showering Inhalation household use	82 2 1.8 14.2
Future child rural resident at modeled concentrations (RGA groundwater drawn at Little Bayou Seeps)	NA	NA	NA	NA	NA	3.02E-02	Selenium 1,1-DCE TCE	20 0.6 79.3	Ingestion Dermal contact Inhalation while showering Inhalation household use	47.5 8.6 0.4 43.5
Future adult rural resident at modeled concentrations (RGA groundwater drawn at Little Bayou Seeps)	2.45E-06	1,1-DCE TCE ⁹⁹ Tc	1.8 32.9 65.3	Ingestion Dermal contact Inhalation while showering Inhalation household use	72.1 3.7 2.7 21.4	9.17E-03	Selenium 1,1-DCE TCE	18.9 2.3 78.8	Ingestion Dermal contact Dermal contact Inhalation household use	44.7 17 18.3 20
Future child rural resident at modeled concentrations (RGA groundwater drawn at Ohio River)		Not a POE for groundwater from this SWMU								
Future adult rural resident at modeled concentrations (RGA groundwater drawn at Ohio River)		Not a POE for groundwater from this SWMU								
Future child recreational user at current concentrations (from WAG 22 RI ^b)	NA	NA	NA	NA	NA	4.2E-02	*No COCs		*No COCs	
Future teen recreational user at current concentrations (from WAG 22 RI ^b)	NA	NA	NA	NA	NA	3.8E-02	*No COCs		*No COCs	

Table F.78. Summary of Risk Characterization for SWMU 30 (Continued)

Receptor	Total ELCR ^a	COCs	% Total ELCR	POCs	% Total ELCR	Total HI ^a	COCs	% Total HI	POCs	% Total HI
Future adult recreational user at current concentrations (from WAG 22 RI ^b)	1.5E-05	Arochlor-1260	48.2	Ingestion of deer	8.7	4.3E-02	*No COCs		*No COCs	
		Benzo(a)pyrene	12.9	Ingestion of rabbit	80.0					
Future excavation worker at current concentrations (soil) (from WAG 22 RI ^b)	1.2E-03	Dibenzo(a,h)anthracene	20.8	Ingestion of quail	11.3	4.5E+00	Aluminum Antimony Arsenic Beryllium Cadmium Chromium Copper Iron Manganese Uranium Vanadium	4.6 6.3 3.3 3.8 3.0 10.2 7.6 19.8 14.3 12.2 12.7	Ingestion Dermal External exposure	26.4 73.5
		Arsenic	1.9	Ingestion	6.3					
		Beryllium	93.7	Dermal	91.7					
		Arochlor-1248	0.1	External exposure	3.3					
		Benzo(a)anthracene	0.1							
		Benzo(a)pyrene	0.8							
		Benzo(b)fluoranthene	0.1							
		Dibenzo(a,h)anthracene	0.4							
		Indeno(1,2,3-cd)pyrene	0.1							
		²³⁷ Np	0.3							
²³⁹ Pu	0.2									
²³⁴ U	0.8									
²³⁵ U	0.1									
^{235/238} U	0.8									
²³⁸ U	0.6									

ELCR = excess lifetime cancer risk; HI = hazard index; POC = point of contact; COC = contaminant of concern

Note: NA = ELCR not applicable to child and teen cohorts. ELCR for adult is for lifetime exposure and takes into account exposure as child and teen.

^a Total ELCR and total HI represent total risk or hazard summed across all POCs for all COCs.

^b Remedial Investigation Report for Solid Waste Management Units 7 and 30 of Waste Area Group 22 at the Paducah Gaseous Diffusion Plant, Paducah, Kentucky; DOE/OR/07-1604-V1&D2, January 1998 (DOE 1998a), Tables 1.59 through 1.68, excluding lead as a COC.

Table F.79. Summary of Risk Characterization for SWMU 145

Receptor	Total ELCRA	COCs	% Total ELCR	POCs	% Total ELCR	Total HIa	COCs	% Total HI	POCs	% Total HI
Future child rural resident at current concentrations (RGA groundwater only)	NA	NA	NA	NA	NA	4.17E+01	Antimony Arsenic Manganese	48.0 47.7 4.3	Ingestion Dermal contact	97.8 2.2
Future adult rural resident at current concentrations (RGA groundwater only)	3.27E-02	Arsenic ⁹⁹ Tc Aroclor-1260	5.1 1.7 93.2	Ingestion Dermal contact	6.9 93.1	1.22E+01	Antimony Arsenic Manganese	49.0 46.7 4.3	Ingestion Dermal contact	95.5 4.5
Future child rural resident at modeled concentrations (RGA groundwater drawn at property boundary)	NA	NA	NA	NA	NA	5.16E-01	Arsenic	99.9	Ingestion	99.8
Future adult rural resident at modeled concentrations (RGA groundwater drawn at property boundary)	1.44E-04	Arsenic ⁹⁹ Tc	29.7 70.3	Ingestion	99.9	1.48E-01	Arsenic	99.9	Ingestion	99.6
Future child rural resident at modeled concentrations (RGA groundwater drawn at Ohio River)	NA	NA	NA	NA	NA		*No COCs		*No COCs	
Future adult rural resident at modeled concentrations (RGA groundwater drawn at Ohio River)	5.29E-05	⁹⁹ Tc	100.0	Ingestion	100		*No COCs		*No COCs	
Future child recreational user at current concentrations (soil)	NE	NE	NE	NE	NE	NE	NE	NE	NE	NE

Table F.79. Summary of Risk Characterization for SWMU 145 (Continued)

Receptor	Total ELCR ^a	COCs	% Total ELCR	POCs	% Total ELCR	Total HI ^a	COCs	% Total HI	POCs	% Total HI
Future teen recreational user at current concentrations (soil)	NE	NE	NE	NE	NE	NE	NE	NE	NE	NE
Future adult recreational user at current concentrations (soil)	NE	NE	NE	NE	NE	NE	NE	NE	NE	NE
Future excavation worker at current concentrations	NE	NE	NE	NE	NE	NE	NE	NE	NE	NE

ELCR = excess lifetime cancer risk; HI = hazard index; POC = point of contact; COC = contaminant of concern

Note: NA = ELCR not applicable to child and teen cohorts. ELCR for adult is for lifetime exposure and takes into account exposure as child and teen.

NE = not evaluated; land use scenario was not assessed because surface soil was not assessed for this SWMU.

^a Total ELCR and total HI represent total risk or hazard summed across all POCs for all COCs.

F.6. UNCERTAINTY IN THE RISK ASSESSMENT

Uncertainties are associated with each step of the risk assessment process. The potential effect of the uncertainties on the final risk characterization must be considered when interpreting the results of the risk characterization because a number of assumptions are made during the risk assessment. Types of uncertainties to consider are divided into four broad categories: those associated with data, exposure assessment, toxicity assessment, and risk characterization. For this BHHRA, only groundwater was evaluated, and the evaluation used maximum concentrations modeled at the SWMU unit over a 1,000 year time frame. Some of the uncertainties, therefore, involve the screening of soil analytes to be modeled to groundwater and the model used. These uncertainties are discussed in Section 5 and Appendix E. Other uncertainties arise from the use of the results of previously conducted BHHRAs for soil. Those uncertainties are discussed here, as are uncertainties related to the baseline risk calculations done for groundwater.

Specific uncertainties in each of these categories are discussed in the following sections. Magnitude of the effect of the uncertainty on the risk characterization is categorized as small, moderate, or large. Uncertainties categorized as small are assumed to not affect the risk estimates by more than one order of magnitude; those categorized as moderate are assumed to affect the risk estimates by between one and two orders of magnitude, and uncertainties categorized as large are assumed to affect the risk estimate by more than two orders of magnitude.

In evaluating these uncertainties and their estimated effect on the risk estimates, it should be remembered that the following uncertainties are neither independent nor mutually exclusive; therefore, the total effect of all uncertainties on the risk estimates (i.e., total ELCRs and HIs) is not necessarily the sum of the estimated effects.

F.6.1 UNCERTAINTIES ASSOCIATED WITH DATA AND DATA EVALUATION

F.6.1.1 Determination Of Epcs—Future Conditions

One uncertainty is the potential risk that may develop as COPCs in media at the BGOU sources migrate to groundwater below the SWMU and are transported off-site. To address this uncertainty, results from a fate and transport model were used to estimate potential contributions from each SWMU to POEs for groundwater exposure away from the source area. (See Appendix E.) While the modeling estimated contaminant transport through groundwater based on contaminant concentrations in the surrounding soil, uncertainty still exists in the POE at which exposure may occur in the future and the contaminant mass that is present in the source areas contributing to the future groundwater concentrations of contaminants. This is particularly true for wastes that originally were containerized. For these wastes, the impact on estimation of future contaminant concentrations in soil depends on whether the wastes already have been released from the containers (in which case, the surrounding soil concentrations may reflect the future contaminant mass) or whether the material may escape the containers in the future (which could result in an increase in the source term/contaminant mass). These uncertainties are discussed in Appendix E. Generally, the estimated effect for most of the modeling uncertainties is moderate to small, indicating that the ELCR and HI estimates generated using the modeled concentrations can be expected to vary by less than an order of magnitude. The potential effect of the status of the containerized wastes could have a significant effect on the risk estimates if drum failure has not yet occurred.

F.6.1.2 Determination of EPCs—Data Collected after Previous Risk Assessments

For most SWMUs, no new surface soil samples have been collected and analyzed after the previous BHHRA were completed. At SWMUs 3 and 7, however, additional surface soil samples have been collected. Attachment F1 contains a summary of maximum detected concentrations and a comparison to the NALs from DOE (2001) for these new samples.

At SWMU 7, for all metals and most radionuclides, the maximum detected concentration associated with the new samples is well below the maximum detected concentration used in the previous risk assessment for SWMU 7 and, therefore, would not alter the results of that assessment. The maximum detected concentration of total PCBs (14.8 mg/L) in the new surface samples exceeds the value used in the risk assessment, but PCBs already were retained as a COC for SWMU 7. Uranium -234, uranium-235, and uranium-236 also were detected at higher maximum concentrations in the new surface soil samples at SWMU 7. These radionuclides already were retained as COCs for SWMU 7; the new data therefore do not substantially impact the results of the assessment already conducted and summarized in this appendix.

At SWMU 3, the previous risk assessment included only the industrial worker. The comparison of maximum detected concentrations to NALs in Attachment F1 includes both the industrial worker and the resident child. For the industrial worker, four metals (arsenic, antimony, iron, and uranium metal), TCE, and three radionuclides (cesium-137, thorium-228, and uranium-238) all exceeded the worker NAL in the new samples. Arsenic and uranium-238 already were COCs for the industrial worker at SWMU 3, but the other analytes exceeding the industrial worker NAL would increase the estimated risk and hazard at this SWMU and should be considered in future actions regarding surface soil.

The analyses conducted in this risk assessment for potential migration of soil contaminants to groundwater used the all soil data collected for surface and subsurface soil, including the data collected after approval of the work plan. Those analyses, therefore, included the new sample results collected after the previous BHHRA discussed above for SWMUs 3 and 7.

F.6.2 UNCERTAINTIES ASSOCIATED WITH EXPOSURE ASSESSMENT

Uncertainties associated with the exposure assessment are from three sources. These are uncertainties in biota fate and transport modeling, in use of the RME scenario, and in the development of the CSM and selection of pathways. Each of these uncertainties is discussed in the following material.

F.6.2.1 Uncertainties in Use of RME Scenarios

For each groundwater exposure pathway modeled, assumptions were made about the number of times a year an activity could occur, routes of exposure, and rate of intake of contaminated media. Because site-specific data were not available for many parameters, defaults from the Risk Methods Document were used. Because most of these defaults are conservative to prevent the underestimation of risk estimates, the risk estimates tend to be conservative and may overestimate risk.

F.6.2.2 Uncertainties Related to Development of the Conceptual Site Model

Generally, the level of uncertainty in the development of the CSM is small.

Exposure pathways for some land use scenarios is expected to be small. All pathways that could contribute significantly to exposures to workers, residents, and recreational users at these SWMUs were included in the previous BHHRA for direct soil exposure and in the current BHHRA for residential

exposure to groundwater, including a separate evaluation of vapor intrusion from groundwater into residential buildings. Exposure pathways related to fishing at ponds filled using potentially contaminated groundwater were not evaluated in this or previous BHHRA.

F.6.3 UNCERTAINTIES ASSOCIATED WITH TOXICITY ASSESSMENT

Uncertainties related to the toxicity assessment are from the following three sources: uncertainty because of lack of toxicity values for some COPCs, uncertainty in the calculation of toxicity values by EPA, and uncertainty in the calculation of absorbed dose toxicity values from administered dose toxicity values. Each of these is discussed in the following paragraphs.

F.6.3.1 Uncertainties Because of Lack of Toxicity Values for Some Chemicals

Because virtually all COPCs had a toxicity value for either HI or ELCR, the only uncertainty to consider here is the use of provisional or withdrawn values in the BHHRA. The uncertainty from the use of provisional or withdrawn values is important to the results of the BHHRA. Some COPCs did not have approved toxicity values, so a provisional or withdrawn value was used. The most notable of these COPCs was TCE, which was evaluated using the current KDEP oral slope factor of $0.322 \text{ (mg/kg} \times \text{day)}^{-1}$. This factor is similar to the EPA provisional oral slope factor of $0.4 \text{ (mg/kg} \times \text{day)}^{-1}$, but both these values are very different from the values for TCE from the 2001 Risk Methods Document that were used in previous assessments. If the residential ELCR for TCE in groundwater at SWMU 2 is calculated with the previously used toxicity values, the ELCR would be $1.03\text{E-}03$ instead of the $4.6\text{E-}02$ ELCR derived with the KDEP value. The risk estimated using the new toxicity values is 45 times the risk calculated using the old toxicity factors and indicates the magnitude of the uncertainty associated with the choice of slope factor for this chemical. The uncertainty associated with use of the new oral slope factor for TCE is considered moderate.

F.6.3.2 Exclusion of Lead from Soil Risks

The results of the previous BHHRAs reported here are the risk and hazard calculations done without including lead as one of the COPCs. In the previous assessments, lead had exceedingly high HIs and was the overwhelming risk driver. This finding may be attributed to the use of a very conservative ($1.0\text{E-}07 \text{ mg/kg-day}$) RfD value provided by the Kentucky Department for Environmental Protection (KDEP). That RfD is no longer in use by KDEP. The current EPA screening levels for lead in soil for residential use is 400 mg/kg . As shown in the screening tables (Tables E3.1 through E3.8) show the maximum detected concentration in lead detected in soil at SWMUs 2, 3, 4, 7, 30, and 145. These maximum values are all less than 100 mg/kg . The maximum detected concentrations of lead in soil at SWMUs 5 and 6 both were at or under 200 mg/kg . These maximum detected values all are less than half the EPA screening level for residential soil, indicating that lead does not need to be considered as a COC at any of the BGOU SWMUs based on comparison with the EPA screening value. The uncertainty of excluding lead from soil risks and hazard calculations is considered small.

F.6.3.3 Uncertainties in Deriving Toxicity Values

Standard EPA RfDs and slope factors were used to estimate potential noncarcinogenic and carcinogenic health effects from exposure to chemicals. Considerable uncertainty is associated with the method applied to derive slope factors and RfDs. The EPA has working groups that review all relevant human and animal studies for each compound and select the studies pertinent to the derivation of the specific RfD and slope factor. These studies often involve data from experimental studies in animals, high exposure levels, and exposures under acute or occupational conditions. Extrapolation of these data to humans under low-dose,

chronic conditions introduces uncertainties. The magnitude of these uncertainties is addressed by applying uncertainty factors to the dose response data for each applicable uncertainty. These factors are incorporated to provide a margin of safety for use in human health risk assessments. The effect of uncertainties in calculation of chemical toxicity values is moderate.

Unlike the uncertainty associated with chemical toxicity values, the uncertainty associated with radionuclide toxicity values is small. The dose-response relationship between cancer and ionizing radiation has been evaluated in many reports, some describing exposed human populations, and is well established.

RfDs and slope factors for some constituents used in the soil BHHRA have been updated since the original soil risk assessment was performed. The adjusted values could impact the cancer risk and HIs for SWMUs whose COCs have new values. COCs impacted significantly by updated toxicity factors include these:

- *Beryllium-RfD*. The current oral RfD of 2.00E-03 mg/kg x day and inhalation RfD of 5.71E-06 mg/kg x day are less than the old oral RfD of 5.00E-03 mg/kg x day and inhalation RfD of 5.71E-03 mg/kg x day and would increase the HI for beryllium, especially through the inhalation route of exposure.
- *Beryllium-cancer slope factor*. At the time the WAG 22 RI and the WAG 3 RI were developed, beryllium was still evaluated as a carcinogen. The cancer slope factor for beryllium has been withdrawn from IRIS, and there has been an agreement not to use this withdrawn value for risk assessments. At several SWMUs, beryllium was a significant contributor to the total cancer risk from soil exposure, generally beryllium accounted for greater than 90% of the risk to the industrial worker and greater than 65% of the risk to the resident. When beryllium is removed from consideration as a carcinogen, the total ELCR becomes much lower at those SWMUs for which it is a COC:
 - SWMU 4: 1.6E-05 for future industrial worker and 1.2E-03 for the resident
 - SWMU 5: 2.1E-04 for future industrial worker; no significant change for the resident value
 - SWMU 6: 2.4E-05 for future industrial worker and 1.1E-03 for the resident
 - SWMU 7: 1.6E-04 for future industrial worker and 1.2E-02 for the resident
 - SWMU 30: 1.4E-04 for future industrial worker and 1.1E-02 for the resident

For SWMUs 4 and 6, removal of the contribution of beryllium to the ELCR reduces the total ELCR to within the EPA risk range for the industrial worker scenario.

- *Iron*. The current oral RfD of 7.00E-01 mg/kg x day is higher than the old oral RfD of 3.00E-01, which would result in a decreased hazard index.
- *Total Uranium*. The current oral RfD of 6.00E-04 mg/kg x day is less than the old oral RfD of 3.00E-03 mg/kg x day and would increase the HI.

The remainder of the COCs with updated toxicity factors changed marginally. For example, the oral RfD for copper changed slightly from 3.70E-02 mg/kg x day to 4.00E-02 mg/kg x day. The inhalation slope factor for nickel increased slightly from 8.40E-01 (mg/kg x day)⁻¹ to 9.10E-01 (mg/kg x day)⁻¹. The majority of the toxicity value changes would have minimal impact on the HIs or cancer risk. The uncertainty therefore is considered small for COCs with marginal changes to toxicity values.

Those COCs with significant toxicity value changes would not substantially impact the soil risk assessment. To illustrate, the SWMU 5 excavation worker soil exposure scenario had the lowest total HI for all soil exposure scenarios of 2.16E+00. Beryllium accounted for 3% of the HI and iron accounted for

38%. Although the change in RfDs for beryllium and iron would alter the hazard slightly, the total HI still would be greater than the 0.1. This is also true for total uranium at SWMU 30. Total uranium contributes 9% of the total hazard for the industrial worker soil exposure scenario. With the total HI at 4.40E+00 using the older toxicity value, changing the RfD would not significantly impact the final results of the risk assessment. The effect of changed toxicity values is considered low.

F.6.3.4 Summary of Chemical Contaminants and Historically Available Toxicity Values

Oral RfD values existed for all of the inorganic contaminants included in historical risk assessments, except phosphate, silica, sulfate, and thallium. Oral RfDs exist for all of the organic analytes included in the historical assessments except 1,2-dichloroethane, 2-hexanone, 2-methylnaphthalene, 3-methylcholanthrene, acenaphthylene, adipate, adipate ester, alkylsiloxane, Aroclor 1248, Aroclor 1260, benzene, benz(a)anthracene, benzo(a)pyrene, benzo(b)fluoranthene, benzo(g,h,i)perylene, benzo(k)fluoranthene, chrysene, dibenzo(a,h)anthracene, dibenzothiophene, hexadecanoic acid, indeno(1,2,3-cd)pyrene, octachlorodibenzo-p-dioxin, phenanthrene, phosphoric acid, tris(methylphenyl)ether, phthalate ester, 2-methyl-l-propanoic acid, and vinyl chloride.

All the inorganic contaminants, except barium, cadmium, lead, manganese, and mercury, lack inhalation RfD values. In addition, of the organic contaminants, only 1,1,1-trichloroethane, 1,2-dichloroethane, 2-butanone, 4-methyl-2-pentanone, benzene, carbon tetrachloride, chloroethane, and toluene have inhalation RfC values in the historic risk assessments.

Absorbed dose RfD values were used for all of the inorganic contaminants included in the assessment except phosphate, silica, sulfate, and thallium. Absorbed dose RfDs exist for all of the organic contaminants included in historic assessments, except 1,2-dichloroethane, 2-hexanone, 2-methylnaphthalene, 3-methylcholanthrene, acenaphthylene, adipate, adipate ester, alkylsiloxane, Aroclor 1248, Aroclor 1260, benzene, benz(a)anthracene, benzo(a)pyrene, benzo(b)fluoranthene, benzo(g,h,i)perylene, benzo(k)fluoranthene, chloromethane, chrysene, dibenzo(a,h)anthracene, dibenzothiophene, hexadecanoic acid, indeno(1,2,3-cd)pyrene, octachlorodibenzo-p-dioxin, phenanthrene, tris(methylphenyl) phosphoric acid, phthalate ester, 2-methyl-l-propanoic acid, and vinyl chloride.

Oral slope factors for inorganic compounds used in the historic assessments included arsenic and beryllium. Oral slope factors were not available for 28 of the 30 inorganic compounds included in the historic assessments.

Inhalation slope factors were available for only a few of the contaminants. Inorganic contaminants with inhalation slope factors included arsenic, beryllium, cadmium, and chromium. Organic contaminants with approved inhalation slope factors are 1,1,2-TCE, 1,1-DCE, 1,2-DCE, Aroclor 1016, Aroclor 1248, Aroclor 1254, Aroclor 1260, benzene, benz(a)anthracene, benzo(a)pyrene, benzo(b)fluoranthene, benzo(k)fluoranthene, carbon tetrachloride, chloroform, chloromethane, chrysene, dibenzo(ai)anthracene, indeno(1,2,3-cd)pyrene, octachlorodibenzo-p-dioxin, tetrachloroethene, TCE, and vinyl chloride.

Changes in Toxicity Data Since 1996

A search of changes to toxicity values from 1996 to present was performed to determine the number of changed and/or new toxicity values for chemicals and radionuclides to determine the potential uncertainties introduced by new or changed toxicity data. The following listing is not complete, but shows the changes in toxicity values and the new additions for chemicals that may be associated with the PGDP. Changes have been made to toxicity values of elements, compounds, and radionuclides. The overall effect of the additions or revised values appears to be minimal, except with the noted change of a withdrawn

oral slope factor for beryllium. A summary of the changes is shown in the next two bulleted items and the entire listing, as described above, is shown.

- RfDs for 4-nitrophenol, barium, boron, iron, acetone, uranium (soluble salts), vinyl chloride, 1,1,1-TCA, TCE, naphthalene, chromium (III) and methyl methacrylate. RfCs also were changed for several of the contaminants including acrolein.
- The inhalation slope factor was changed from a HEAST value to a conversion from the IRIS unit risk value for many chemicals, but included beryllium and vinyl chloride, and a change to the slope factor was made for vinyl chloride.

February 2009

Toxicity values, metadata, and the RAGS Part A profile for nitrobenzene were updated based on a new IRIS summary.

September 2005

An oral chronic RfD was added for 4-nitrophenol from NCEA as reported in the EPA Region 6 PRG table.

July 2005

An oral RfD was added for barium from IRIS. A dermal RfD also was calculated.

June 2005

An oral RfD was added for iron. A dermal RfD also was calculated.

August 2004

The toxicity summary for boron and compounds has been revised, and a toxicological review for boron and compounds has been added to the IRIS Web site. The following toxicity values are to be used for boron and compounds:

- The dermal RfD of 1.8E-01 (mg/kg-day) was calculated from the oral RfD.

July 2004

The toxicity summary for 1,2-dibromoethane has been revised, and a toxicological review for 1,2-dibromoethane has been added to the IRIS Web site. The following toxicity values are to be used for 1,2-dibromoethane.

- The new chronic inhalation RfC of 9.0E-03 (mg/m³) has been added. The associated toxicity metadata also have been updated.
- The chronic inhalation RfD of 2.57E-03 (mg/kg-day) was calculated from the RfC.
- The chronic oral RfD is now 9.0E-03 (mg/kg-day) and associated metadata have been updated.
- The dermal RfD of 7.2E-03 (mg/kg-day) was calculated from the oral RfD.

- The oral slope factor is now 2.00E+00 (mg/kg-day)⁻¹ and associated metadata have been added.
- The dermal slope factor is now 2.50E+00 (mg/kg-day)⁻¹ and was calculated from the oral slope factor.
- The oral unit risk is now 6.00E-02 (mg/L)⁻¹ and associated metadata have been added.
- The inhalation unit risk is now 6.0E-01 (mg/m³)⁻¹ associated metadata have been added.
- The inhalation slope factor is now 2.10E-01 (mg/kg-day)⁻¹ and was calculated from the inhalation unit risk.

August 2003

- The toxicity summary for acetone has been revised, and a new toxicological review for acetone has been added to the IRIS Web site. The following toxicity values are to be used for acetone:
 - The chronic oral RfD is now 9.0E-01 (mg/kg-day) based on the critical effect of nephropathy.
 - The dermal RfD is now 7.47E-01 (mg/kg-day).

July 2003

- The ethylbenzene cancer toxicity values have been withdrawn by NCEA. This chemical is now being reassessed for IRIS, which automatically flags further use of any provisional cancer or non-cancer assessments; however, the IRIS database has retained these values and added appropriate footnotes.

June 2003

- The toxicity summary for acrolein has been revised, and a new toxicological review for a has been added to the IRIS Web site. The following toxicity values are to be used for acrolein:
 - The chronic inhalation RfC of 2.0E-05 (mg/m³) has not changed, but the associated toxicity metadata has been updated.
 - The chronic oral RfD is now 5.0E-04 (mg/kg-day) based on critical effects of decreased survival.

May 2001

- For 47 chemicals, the inhalation slope factor was changed from a HEAST value to a conversion from the IRIS unit risk value, which is more current. This change occurred for the following chemicals: acrylamide; acrylonitrile; aldrin; aramite; azobenzene; benzene; beryllium; bis(2-chloroethyl)ether; bis(chloromethyl)ether; bromoform; butadiene, 1,3-; carbon tetrachloride; chlordane; chloroform; chromium VI (chromic acid mists and particulates); coke oven emissions; DDT; dibromoethane, 1,2-; dichloroethylene, 1,1-; dichloropropene, 1,3-; deldrin; diphenylhydrazine, 1,2-; epichlorohydrin; formaldehyde; heptachlor; heptachlor epoxide; hexachlorobenzene; hexachlorobutadiene; hexachlorocyclohexane, alpha-; hexachlorocyclohexane, beta-; hexachlorocyclohexane, technical; hexachloroethane; hydrazine; hydrazine sulfate; nickel refinery dust; nickel subsulfide; nitrosodiethylamine, N-; nitrosodimethylamine, N-; nitroso-di-N-butylamine, N-; nitrosopyrrolidine, N-; propylene oxide; tetrachloroethane, 1,1,2,2-; toxaphene; trichloroethane, 1,1,2; trichlorophenol, 2,4,6; and vinyl chloride.

February 2001

- Updated the oral RfD for uranium (soluble salts). The new value is 6.0E-4 mg/kg/day, and the new dermal RfD is 5.1E-4 mg/kg/day.
- Updated files for the Federal Regulatory Standards. The following is a list of the changes/additions.

January 2001

- There was a new release of toxicity values for radionuclides as presented in Federal Guidance Report No. 13. Essentially, every slope factor was changed.

August 2000

- The IRIS summary for vinyl chloride has been revised, and a new toxicological review for vinyl chloride has been added to the IRIS Web site. The following changes to vinyl chloride have been made on the IRIS database:
 - The oral RfD is now 3E-3 mg/kg/day.
 - The dermal RfD has been added to reflect the addition of the oral RfD.
 - The inhalation RfC is now 1E-1 mg/m³.
 - The inhalation RfD had been added to reflect the addition of the inhalation RfC.
 - The oral slope factor has been changed to 1.4 (mg/kg/day)⁻¹.
 - The dermal slope factor has been changed to reflect the change in the oral slope factor.
 - The oral unit risk has been changed to 4.2E-2 (mg/L)⁻¹.
 - The inhalation unit risk has been changed to 8.8E-3 (mg/m³)⁻¹.
 - The inhalation slope factor has been changed to reflect the change in the inhalation unit risk.

April 2000

- The following chemicals have new **provisional** toxicity values:
 - 1,1- dimethyl hydrazine, Inhalation UR=4.9E0 Oral SF=3E0
 - Ethylbenzene
 - Inhalation UR=1.1E-3
 - Methyl Hydrazine
 - Inhalation UR=4.9E0, Oral SF=3E0
 - 1,1,1-TCA, Chronic RfC=2.2E0

February 2000

- The oral reference dose for methyl methacrylate was updated.

June 1999

- The inhalation RfC for beryllium and compounds has been changed from 2E-2 ug/m³, to 2E-5 mg/m³ and the inhalation RfD for beryllium, and compounds has been changed from 5.71E-3 ug/kg/day to 5.71E-6 mg/kg/day.

October 1998

- Naphthalene (91-20-3) - The chronic oral RfD (RFD_{OC}) changed to 2.0E-2, and the chronic inhalation reference concentration (RFC_{IC}) changed to 3.0E-3. The cancer class changed to 'C'.
- Benzene (71-43-2) - The inhalation unit risk (IUR) changed to 7.8E-3.

September 1998

- Inhalation chronic RfD and RfC added for naphthalene (91-20-3) [RFD_{IC} = 3.71E-04, RFC_{IC} = 1.3E-4] and tetrachloroethylene (127-18-4) [RFD_{IC} = 1.71E-01, RFC_{IC} = 6.0E-1].
- Inhalation slope factor for PCBs and all Aroclors has been changed from 2.0 to 0.4.
- Ingestion and dermal chronic reference dose have been changed for the following:

<u>Chemical</u>	<u>RfD</u>	<u>RfD (dermal)</u>
Trichloroethane, 1,1,1	2.0E-1	1.8E-1
Trichloroethene	6.0E-3	9.0E-4
Naphthalene	4.0E-2	3.2E-2
Chromium (III)	1.5E+0	7.5E-3

July 1998

Inhalation RfDs are defined to be RfC. A database check identified three inhalation RfD that needed to be updated. The following changes resulted:

Chronic:

101-68-8 Methylenediphenyl Diisocyanate

OLD Number = 5.71E-6

NEW Number = 1.71E-4

108-88-3 Toluene

OLD Number = 3.20E-1

NEW Number = 1.1E-1

June 1998

Beryllium. The oral slope factor is withdrawn pending further investigation.

May 1998

- Methyl methacrylate (80-62-6) the chronic oral RfD is changed from 8.00E-02 to 1.40E+00. The chronic inhalation RfC of 7.00E-01 with a footnote of "a" is added. This is multiplied by 20/70 to get a chronic inhalation RfD of 2.00E-1.

February 1998

- Arsenic, inorganic (7440-38-2). The inhalation slope factor of 5.00E+01 has been withdrawn.
- 1, 2- dichloroethane, (107-06-2). The inhalation slope factor of 9.10E-02 has been withdrawn.

Changes from 1996 to 1998 were minimal.

F.6.3.5 Uncertainties Because of Calculation of Absorbed Dose Toxicity Values from Administered Dose Toxicity Values

Uncertainty exists in the validity of the calculations used to convert an administered dose toxicity value to an absorbed dose. Of greatest importance is the lack of consideration of point-of-contact effects in this calculation. For example, some organic analytes can cause a toxic or cancer response in skin. This effect is not considered in the calculation of absorbed dose toxicity values from administered dose toxicity values using EPA protocols. Similarly, the administered dose response for many chemicals relies on the delivery of a high concentration of contaminants to the liver via the portal system after ingestion; this effect is not seen if a contaminant is absorbed through the skin because of the larger distribution space for the contaminant absorbed through the skin. However, even with these uncertainties, the effect of the uncertainty in calculation of absorbed dose toxicity values from administered dose toxicity values upon the risk estimates is estimated to be small because the overall contribution of dermal exposure to total risk and hazard is much smaller due to the use of the new, lower dermal absorption factors.

F.6.4 UNCERTAINTIES ASSOCIATED WITH RISK CHARACTERIZATION

Three uncertainties are related to risk characterization. The first is the method used to combine HQs over pathways and combine pathway HIs to calculate total HI. This method also is used to combine chemical- and pathway-specific ELCRs to derive total ELCRs. The second is the uncertainty added to the assessment by combining risks from chemicals and radionuclides. These uncertainties are discussed in the following subsections.

F.6.4.1 Combining Chemical-specific Risk Values and Pathway Risk Values

The primary uncertainty in risk characterization is the method used to combine HQs and chemical-specific ELCRs over pathways and combine pathway HIs and ELCRs to calculate total HI and ELCR. The uncertainties in this method are discussed in the following text.

The method used to calculate pathway HIs and ELCRs in the BHHRA followed EPA protocols (Risk Methods Document). This guidance calls for the simple summation of HQs and chemical-specific ELCRs to calculate pathway HIs and ELCRs, respectively. This method assumes that all effects between chemicals are additive. EPA makes this assumption because information concerning the effect of chemical mixtures is lacking. Specific limitations of this approach for systemic toxicity effects (HI) have been reported by EPA.

Little is known about the effects of chemical mixtures; although additivity is assumed, the interaction of multiple chemicals possibly could be synergistic or antagonistic.

The RfDs and reference concentrations do not have equal accuracy or precision and are not based on the same severity of effects.

Dose additivity is most properly applied to compounds that induce the same effect by the same mechanism of action. While the approach recommended by EPA is a useful screening-level approach, the potential for at least noncarcinogenic effects to occur can be overestimated for chemicals that act by different mechanisms and on different target organs.

The effect of this uncertainty on the estimate of HI depends on how many contaminants drive HI and if the contaminants have different endpoints. In this BHHRA, several contaminants do affect HI, and these contaminants do have differing endpoints and target organs (see Tables F.40 and F.41). Because only a few “priority COCs” drive HI, as shown in Section 5, and because the HI from each of these “priority COCs” alone is great enough that a systemic toxic effect may be reasonably expected, the effect of this uncertainty on HIs is small.

Specific limitations for this approach in regard to chemical carcinogenesis also have been reported by EPA in RAGS:

- Cancer risks (i.e., ELCRs) are based on slope factors that represent an upper 95th percentile estimate of potency; the upper 95th percentiles of probability distributions are not strictly additive. Summing these risks can result in an overly conservative estimate of lifetime ELCR (EPA 1991).
- Combined cancer risks for chemical carcinogens and radionuclides are presented, but may not be additive because the slope factors used to characterize the risk from chemicals are derived differently from the slope factors used to characterize risk from radionuclides.
- Not all slope factors contain the same weight-of-evidence for human carcinogenicity. As explained in Section 4, EPA recognizes this by placing weight-of-evidence classifications on all slope factors. Those contaminants with an A weight-of-evidence should probably receive more attention in the selection of a remedial design than contaminants with a B or C classification. Similarly, a contaminant with a B classification should probably receive greater attention than one with a C classification. The simple combination of ELCRs does not take this hierarchy into account.

The uncertainties involved in combining chemical-specific ELCRs and pathway ELCRs are considerable. The effect of these uncertainties on the total ELCRs presented in the BHHRA is small because as noted above, only a few “priority COCs” dominate the pathway ELCR for most pathways; therefore, the potential effect of mixtures is reduced.

F.6.4.2 Nature and Extent of Contamination

Uncertainty related to the distribution and concentration of contaminants in the soils may be different than presented by the dataset. That is, the nature and extent of contamination may be over- or under-estimated by the available data. The BGOU is a series of landfills and the large variability of contaminant soil concentrations in landfills is well established.

The results of the BHHRA identified contaminants of concern for soil exposure and groundwater exposure. The priority COCs identified are listed below by SWMU. Table 2.1 of the Draft Risk Methods Documents is a listing of “Significant Chemicals of Potential Concern at the PGDP.” When the COCs of this BHHRA are compared to the list of “Significant Chemicals of Potential Concern at the PGDP,” the COCs identified includes 16 of 23 inorganic compounds, 6 of 12 radionuclides, and 13 of 46 VOCs and SVOCs (this total number of significant VOCs and SVOCs excludes the congeners of dioxins and furans, but includes total dioxins and furans). In addition, all of the constituents listed as “Significant Chemicals of Potential Concern at the PGDP,” were included in the analysis for the BHHRA with exception of

chromium VI and the congeners of dioxins and furans. The uncertainty associated with the nature of the contamination is low, based on this review of the COCs and data analysis.

The extent of contamination within the BGOU is determined by geophysical surveys, maps, and coordinates that determine the physical boundaries of the individual SWMUs. The extent of contamination outside of the BGOU is, in part, based on groundwater concentrations and groundwater modeling. The uncertainty associated with the extent of contamination in the groundwater is low. The extent of the contamination that has migrated from the BGOU to nearby soils is less certain.

The data set for the BGOU RI is comparable to the data available for many landfills. There is a possibility that drummed waste could be intact and contain liquids, uranium, and other unexpected wastes since inventories of the various landfill contents are not complete. Drummed waste may remain intact or partially intact because of the uncertainty regarding past studies (the studies were based on metal drums or containers and it is plausible that some of the containers were lined with plastic or were composed of plastic and may not have degraded as quickly).

F.6.4.3 Uncertainties Regarding Detection Sample Quantitation Limits in Soils

The sample quantitation limits (SQLs) for all chemicals and radionuclides were reviewed by querying the ORIES database containing all the analysis. The SQLs by analyte were extracted from the ORIES database and a comparison of the SQLs to the child residential soil/sediment no action levels (NALs) for the PGDP was done. When compared to the minimum SQL, 56 constituents were greater than the NAL, and 126 constituents had SQLs less than the NAL. Many constituents do not have NALs that were included in the chemical and radionuclide analysis.

For the same chemical, the SQL in one sample maybe higher than, lower than, or equal to SQL values for other samples. In addition, preparation or analytical adjustments, such as dilution of the sample for quantitation of an extremely high level of one chemical, could result in non-detects for other chemicals included in the analysis, even though these chemicals may have been present at trace quantities in the undiluted sample. The SQL is the MDL adjusted to reflect sample-specific action such as dilution or use of a smaller sample aliquot for analysis due to matrix effects or the high concentration of some analytes. (Guidance for Data Useability in Risk Assessment (Part A) EPA/540/R-92/003 December 1991).

SQLs are specific to each sample collected. The database was queried for the minimum SQL for each analyte for each SWMU. For each analyte, there were eight results. Simply examining how many constituent SQLs were below the NAL, as shown above, provides perspective into the SQLs as they compare to the NALs by constituent. When each of the queried results for each of the SWMUs is examined on a percentage basis, it provides better insight into the SQLs for the BGOU. Eighty percent of SQLs were less than the NALs and twenty percent were greater than the constituent specific NALs. The percentage of SQLs, which were lower than the NAL, demonstrates the data set does not introduce significant uncertainty based on SQLs.

The SQLs were sufficient to meet applicable screening criteria. SQLs in soils are variable because of the bulk density differences, physical properties, particle size, soil moisture, and other factors. It is not uncommon to find detection limits SQLs for soil above screening levels, especially for COCs with high slope factors or low Rfds (i.e., low screening levels). The effect of these uncertainties on the total ELCRs and HIs presented in the BHHRA is small.

F.6.4.4 Chemical-specific Risk Values Exceeding 1E-02

At SWMU 2, the risk calculated for TCE exposure through groundwater is 3E-02. Risk estimates greater than 1E-02 are not accurate when calculated using EPA slope factors, which are based on a linearized multistage model. The one-hit model may provide a better estimate of risk in these cases (EPA 1989). One hit values were not calculated for this BHHRA because the purpose of the risk estimates is only to determine which COPCs have risks greater than 1E-04 which indicates the COPC should be a COC for the feasibility study (FS).

F.6.5 SUMMARY OF UNCERTAINTIES

As is shown in the previous subsections, the risk estimates could vary if different assumptions were used in deriving the risk estimates or if better information was available for some parameters. The following text summarizes the estimated effects of each uncertainty mentioned previously.

No uncertainties were estimated to have a large effect on the risk characterization, and only three were estimated to have a moderate effect.

Following is a list of uncertainties with effects estimated to be moderate:

- Exclusion of some potential biota pathways (fish from ponds) for future receptors,
- Migration of groundwater to off-site receptors, and
- Calculation of toxicity values for chemicals (particularly TCE).

Following is a list of uncertainties with effects estimated to be small:

- Determination of exposure points for future concentrations,
- Use of RME default exposure values instead of central tendency exposure values,
- Use of provisional and withdrawn toxicity values,
- Determination of radionuclide toxicity values, and values.

The following is summary information of the historical investigations and risk assessments that were performed for SWMUs that are included in the BGOU. The original uncertainty sections are quite complete, but some of the information is dated. The wording of the uncertainty assessment has not been changed, but some text and tables have been removed to be succinct. Every effort was made to accurately report the uncertainty sections in summary form.

Evaluation of Uncertainty in SWMUs 2 and 3 (Summary)

This section discusses the key assumptions and uncertainties that affect the level of confidence placed on the quantitative risk estimates derived for the SWMUs 2 and 3 risk assessments. Because uncertainties are inherent in any risk assessment, a qualitative discussion of these uncertainties puts into perspective the risks calculated for the site.

Data Evaluation. Of the variables used in performing the risk assessment, the error terms related to the laboratory analyses are probably the best defined and provide less uncertainty than other factors in the assessment. Individual errors or biases in the data are possible, but the size of the database minimizes uncertainties in the overall concentration estimates.

The primary data limitations and uncertainties associated with concentration estimates and data at SWMUs 2 and 3 include the following observations:

- Sampling strategies at SWMUs 2 and 3 were designed to detect migration to off-site areas, not for current or future exposures to surface soil. In some samples, data may reflect "hot spots" and overestimate risks; in other samples, data may reflect contamination adjacent to the site and may underestimate risks.
- Risks from direct contact exposures to surface soils were evaluated using the results from soil samples from zero to 6 ft bgs. Thus, this evaluation closely approximates conditions that might occur during shallow excavations around the SWMUs. Current direct contact exposures to soils 6 inches to 1 ft bgs were not evaluated since only two samples were available at these depths.
- No direct sampling was conducted of the waste itself. No quantification was made, therefore, of the potential risk if excavation into the waste were to occur.
- There is considerable potential variability associated with VOC concentration results because of losses from the soil matrix even with good sampling technique. In addition, with typical laboratory holding times of 14 days at 4° C, a loss in concentration typically occurs (from the time of collection) of 40 to 90 percent of the original concentration, depending on the specific chemical. These uncertainties can lead to underestimates of risks associated with VOCs.
- Disposal records have been shown to be inaccurate; therefore, the low reliability of the buried waste materials inventory introduces uncertainties that may result in under or over estimates of risks.

The discrepancy between maximum detected beta activity levels and maximum detected Tc-99 activity levels is a source of data uncertainty and may result in underestimation of radiological risks.

Exposure Assessment. Worker exposures to contaminated surface soils at SWMUs 2 and 3 are considered conservative; however, the surface water pathway was not quantitatively evaluated in this assessment. SWMUs 2 and 3 are not considered to contribute to the surface water exposure pathway. A reasonable deviation resulting from erosion of sediments in runoff from the site will be evaluated for the surface water integrator OU.

The 250, 8-hr days per year, assumption for workers is excessive for current on-site worker exposures at a single waste management unit (WMU). This exposure level would be appropriate for exposures in areas where continuous activities were required outside the domain of OSHA regulations. Further, it is unreasonable to assume that a worker would remain in the vicinity of a single WMU for a 25-year exposure period.

Current, PGDP worker exposure to SWMUs 2 and 3 is better estimated using the worker/intruder scenario, which reflects 10 percent of a worker's time spent at a single WMU. This scenario also conservatively addresses potential intruder exposures at PGDP. The assumption of biweekly 8-hr exposure periods at a single WMU over a 25-year period overestimates risks to visitors/intruders, even if fences and security measures were eliminated.

The assumption that adult workers ingest 50 mg of soil per day is likely conservative. In addition, the assumption that 100 percent of soil ingested per day comes from the contaminated source is conservative. Thus, both soil ingestion rates and the fraction from the contaminated source tend to overestimate risks.

The assumptions for dermal absorption are also conservative for the amount of soil adhering to skin, skin surface area available for contact, and the amount of a chemical absorbed from soil. These three factors tend to overestimate the amount of chemical absorbed from soil by the dermal route.

Toxicity Assessment. Uncertainty is associated with the use of the method to determine carcinogenic risks in humans. In discussing uncertainty, the EPA expressed the following:

It should be emphasized that the linearized multistage procedure leads to a plausible upper limit to the risk that is consistent with some proposed mechanisms of carcinogenesis. Such an estimate, however, does not necessarily give a realistic prediction of the risk. The true value of risk is unknown, and may be as low as zero. The range of risks, defined by the upper limit given by the chosen model and the lower limit which may be stated as low as zero, should be explicitly stated. (*FR* 51:34013, September 24, 1986).

To assess the overall potential for cancer and noncancer effects posed by multiple chemicals, cancer risks or HIs are summed. This method may be conservative because it does not account for potential differences in toxic end points.

Uncertainty in toxicity assessment can arise from the use of models or test systems that do not accurately describe the exposed population or the relevant exposure environment. This type of uncertainty can be found in the toxicity values derived from animal experiments and in assumptions made about dose-response models, which may or may not be valid.

Several of the constituents reported at the site do not have a current oral, inhalation, and/or dermal slope factor or RfD. Because no dermal toxicity values are available, oral toxicity values were used. No adjustments were made on the basis of absorbance, which tends to underestimate risks via dermal absorption.

Risk Characterization. Standard ground surface conversion factors were used to determine doses and risks associated with external exposures to radiation from contaminated surface soil at SWMUs 2 and 3. The ground surface dose and risk factors are based on assumptions of uniform contamination over a large surface area. Use of generic surface risk factors will result in overestimates of risks from external gamma radiation at SWMUs 2 and 3.

The risk factors used in this report are based on EPA guidance in HEAST and are greater than the risk factors shown in the BEIR III Report, but slightly less than the factors shown in the BEIR V Report. Thus, they represent an estimate of risk that falls within the range of risk estimates from the most recent data. The EPA regards these risk estimates as “reasonable,” but not “conservative.” Consequently, use of the EPA risk factors should not tend to greatly overestimate the risk of low-level radiation exposure. Although several uncertainties produce both over- and underestimated risk calculations in this assessment, factors that tend to overestimate risks outweigh those that underestimate risks; therefore, risks calculated in this assessment are considered conservative.

Some portion of the risks estimated for SWMUs 2 and 3 may be attributed to naturally occurring background concentrations of inorganics and radionuclides in soil and groundwater. For example, arsenic, beryllium, and manganese contribute to risks exceeding 1×10^{-6} and an HI of 1 in reference groundwater and soil samples. This background risk, while not subtracted from site-related risk, presents additional uncertainty in the risk characterization.

SWMUs 7 and 30 Uncertainties from Historical Investigations

In evaluating these uncertainties and their estimated effect on the risk estimates, it must be remembered that the following uncertainties are neither independent nor mutually exclusive. The total effect of all uncertainties discussed in the following subsections on the risk estimates (i.e., total ELCRs and HIs) is not the sum of the estimated effects.

Uncertainties Associated with Data and Data Evaluation

Several uncertainties are associated with the data set and the selection of COPCs. Specific uncertainties that will be discussed in the following subsections are selection of COPCs; determination of exposure point concentrations under current and future conditions; and use of concentrations from total versus filtered samples for inorganic compounds in groundwater.

Selection of COPCs

Some uncertainty is involved with the selection of COPCs. This uncertainty is derived from several sources. The first uncertainty related to the selection of COPCs is the retention of infrequently detected chemicals in the list of COPCs. Several of the chemicals retained in the list of COPCs were detected in less than 10 percent of the samples taken. Of greatest concern is that some of these COPCs are retained as COCs. The difference in total ELCR estimates and total HI when the chemicals detected in less than 10 percent of the samples are retained as COPCs. The infrequently detected COPCs had virtually no effect on the risk or hazard estimates; therefore, the estimated effect of the uncertainty on the risk estimates is small.

The second uncertainty related to selection of COPCs in the BHHRA is that temporal patterns in detection of analytes were not considered when selecting COPCs. If temporal patterns were considered, the final risk results in this BHHRA may be quite different; however, in this BHHRA, the assumed effect of this uncertainty on the risk estimates is small.

The third uncertainty related to selection of COPCs in the BHHRA concerns the quantitation limits used for some analytes. For many organic analytes, the quantitation limit used exceeds a concentration that may result in a significant health effect. For example, at SWMU 7 for groundwater, a total of 47 organic compounds has quantitation limits used for at least one sample that exceed the residential use ELCR preliminary remediation goal (PRG). Similarly, in subsurface soil at this location, 31 organic compounds have quantitation limits used for at least one sample that exceed the residential use ELCR PRG. Because the quantitation limits exceed the PRGs, it is possible that these chemicals are present at concentrations that pose considerable risk, but may not be retained as COPCs and be quantitatively evaluated. Because many of the organic compounds with quantitation limits that exceed PRGs tend to be unrelated to processes at PGDP, the estimated effect of this uncertainty on this risk assessment is small.

A fourth uncertainty related to selection of COPCs is the use of historical data in addition to data collected during the RI. As noted earlier, these data were added to the data set to augment the information collected during the RI. Use of these data is consistent with current EPA guidelines (EPA 1989a). No statistical determination was performed if historical data and data collected during the RI were comparable. However, the estimated effect of this uncertainty on this risk assessment is assumed to be small because of the short period of time between the data collection efforts.

A fifth uncertainty related to the selection of COPCs is the inclusion of common laboratory contaminants in the COPC list. The infrequently detected COPCs had virtually no effect on the risk or hazard estimates; therefore, the estimated effect of the uncertainty on the risk estimates is small.

A sixth uncertainty related to the selection of COPCs is the effect that removal of analytes based on a comparison to blanks may have had on the COPCs list. In this risk assessment, analytes that are common laboratory contaminants had their detected concentrations replaced with the sample quantitation limit when the detected concentration was less than 10 times the concentration detected in an associated blank. Analytes that are not common laboratory contaminants had their detected concentrations replaced with the sample quantitation limit when the detected concentration was less than five times the concentration detected in an associated blank. This process is consistent with current EPA guidance (EPA 1989a) but may have modified the COPCs list if all results for a specific analyte at a SWMU were associated with blanks in which that analyte was detected. For the data used in the BHHRA, there were no analytes for which every sample at a SWMU was related to a contaminated blank; therefore, the effect of this uncertainty on the selection of COPCs is small.

A seventh uncertainty related to the selection of the COPCs is the use of a toxicity screen to determine the final COPCs list. In this BHHRA, the maximum detected concentrations of analytes within each medium at each SWMU were compared to residential human health risk-based screening criteria. The residential risk-based screening levels were used per regulatory agreement (DOE 1996b). Analytes with maximum concentrations less than these screening criteria were removed from the list of COPCs.

To examine the effect the toxicity screen may have had on the COPCs list and on the resulting risk estimates developed in the BHHRA, marginal hazard and risk contributions were calculated for analytes removed on the basis of this screen. Marginal hazard and risk contributions can be defined as the estimated increase in the final hazard and risk estimates under the residential scenario that would have been seen if the analytes removed from the list of COPCs had been left on the list. The marginal contribution of the analytes removed from the COPCs list would have been minimal; therefore, the estimated effect of this uncertainty on the final risk estimates is small.

An eighth uncertainty related to the selection of the COPCs is the use of a background screen to determine the final COPCs list. In this BHHRA, the maximum detected concentrations of analytes within soil and groundwater at each SWMU were compared to background concentrations determined in other related projects at PGDP.

The background concentrations used for this screen were compared to their respective medium-specific human health risk-based screening criteria. Several of the background concentrations used in the BHHRA are greater than their respective risk-based screening criteria. The results indicate that if analytes had not been removed from the COPCs list on the basis of the background screen, the final risk estimates would have been larger. Because this screen relied on the comparison of the maximum detected concentration of each analyte in a medium, it is highly unlikely that analytes would have been removed from the COPCs list in error. The estimated effect of the background screen on the COPCs list is small, and the resulting effect on the final risk estimates is small.

Finally, assuming the ditches contained soil rather than sediment may have affected the selection of COPCs. Generally, this decision may have led to overestimates of exposure by the industrial worker and the rural resident because the default exposure times for these scenarios assuming soil contact are greater than the exposure times assuming contact with sediment. Alternatively, this decision may have led to underestimates of exposure by the recreational user because the default exposure times for this scenario assuming sediment contact is greater than the exposure times assuming contact with sediment. In total, the estimated effect of the decision to consider the medium in the ditches as soil rather than sediment upon the final risk estimates is believed to be small.

Determination of Exposure Point Concentrations—Current Conditions

Little uncertainty is involved in characterizing exposure point concentrations under current conditions in this BHHRA. Sampling data came from sources of known quality. Data sets were generated from samples collected using EPA-approved protocols, and samples were analyzed using EPA-approved methods.

Determination of Exposure Point Concentrations—Future Conditions

Uncertainty is involved in characterizing exposure point concentrations under future conditions in this BHHRA. In calculating the exposure point concentrations at the SWMUs, the concentrations of COPCs are kept constant throughout the exposure period. That is, the risk assessment does not consider that concentrations of some COCs may be lower at some time in the future due to processes such as degradation and attenuation. Because the COCs driving risk at the SWMUs at all times are inorganic compounds that are persistent in the environment, the effect of this uncertainty is estimated to be small.

A second uncertainty is the potential risk that may develop, as COPCs in media at SWMUs 7 and 30 migrate to RGA groundwater below the SWMUs and to surface water bodies near the SWMU. To address this uncertainty, the SESOIL Model was used to estimate potential concentrations of selected COPCs in RGA groundwater below the unit, and a combination of SESOIL and AT123D modeling was used to estimate the potential concentrations of selected COPCs in groundwater at a downgradient point of exposure. Additionally, the SWMM model was used to estimate concentrations migrating to surface water. For SWMUs 7 and 30 and associated ditches, the exposure points to these media were assumed to be at the DOE property boundary for groundwater and at the confluence of the ditches for surface water.

The risk results of the RGA groundwater modeling show that systemic toxicity for the adult and child rural resident is not a land use scenario of concern; however, ELCR is a land use scenario of concern as technetium-99 drives ELCR for the rural resident at all time periods.

The results from the SESOIL, AT123D, and SWMM modeling do indicate that future risks calculated under current concentrations were overestimated; therefore, the effect of this uncertainty is moderate.

Use of Concentrations from Total Versus Filtered Samples

In this BHHRA, all analyte concentrations in water came from the analyses of unfiltered or total samples. The use of data from analyses of total samples is consistent with current EPA guidance (EPA 1989a), but introduces an additional uncertainty to the BHHRA for some water use pathways. It is important to note that the magnitude of the effect of this uncertainty upon the risk estimates is difficult to estimate because it is not known to what extent the quality of water (in terms of total solids) from a residential well would vary from the quality of water taken from a monitoring well. Because the samples used in this study came from wells that were properly developed and were collected using appropriate methods, the effect of this uncertainty is assumed to be small.

For some fate and transport pathways, it may be more appropriate to use dissolved concentrations. The best illustration of this in the BHHRA is the fish consumption pathway. For this pathway, the surface water concentration that should be used is from dissolved samples because fish are normally exposed only to the dissolved contaminants in water and not particulates. The effect of the uncertainty of using data from analyses of total samples on the risk estimates is small.

Uncertainties Associated with Exposure Assessment

Uncertainties associated with the exposure assessment are from five sources. These are uncertainties in use of the RME scenario, in the development of the conceptual site model and selection of pathways, use of default values when estimating dermal absorbed dose, and use of conservative exposure values for the excavation worker. Each of these uncertainties is discussed in the following material.

Uncertainties in Use of Reasonable Maximum Exposure Scenarios

For each exposure pathway modeled, assumptions were made about the number of times a year an activity could occur, the routes of exposure, and the rate of intake of contaminated media. Because site-specific data were not available for most of these parameters, suggested EPA and Commonwealth of Kentucky defaults were used. Because most of these defaults are conservative to prevent the underestimation of risk estimates, the risk estimates tend to be conservative. Generally, when several upper-bound values are combined, the resulting value tends to exceed the level of exposure that may be reasonable at a site. In consideration of this problem, attention should be focused not on the fact that any individual risk model is overly conservative, because most are not, but on the fact that every risk model is conservative. It is the combination of several conservative risk estimates from different exposure routes that lead to undue conservatism and overestimation of risks.

To examine the potential effect of this uncertainty, risks and systemic toxicities for the residential scenario also were estimated using average values for all exposure parameters, (All exposure parameters used in this assessment were taken from the preliminary review draft of EPA's *Superfund's Standard Default Exposure Factors for the Central Tendency and Reasonable Maximum Exposure, Review Draft*. In this assessment, all exposure pathways evaluated were identical to those used in the RME scenario. Similarly, all exposure equations, chemical concentrations, radionuclide activities, and toxicity values were identical to those used for the RME scenario. The results of this assessment shows that the effect of this uncertainty was small for estimates of systemic toxicity and ELCR.

Uncertainties Related to Development of the Site Conceptual Models

Generally, the level of uncertainty in the development of the site conceptual models is small. Data were collected from several previous studies and from local experts to develop these models; however, there are some uncertainties related to specific scenarios that deserve additional explanation. These uncertainties are the consideration or lack of consideration of specific pathways for some scenarios, the lack of consideration of a separate intruder/infrequent recreational user scenario, and the summation of risks across areas and across scenarios.

An uncertainty related to the consideration of specific pathways for some scenarios is the consideration of groundwater ingestion by the future industrial worker and future rural resident. For these scenarios, use of groundwater as drinking water and for showering was assumed for SWMUs 7 and 30. These exposure routes were included to provide risk managers with additional information about the potential risk posed by groundwater at the SWMUs. These pathways were included, although PGDP does not currently use groundwater and does not plan to use groundwater in the future. The effect of including groundwater use in the assessment of the future industrial worker and future rural resident land use scenarios on the total risk estimates was assessed. The effect of this uncertainty was small for both total HI and total ELCR at SWMUs 7 and 30. It also should be noted the groundwater pathways were not included for the ditches. As each ditch spanned both SWMUs 7 and 30, assigning specific groundwater wells was not appropriate or useful, as the ditches will be investigated and remediated as a whole.

The lack of summation of risks across SWMUs and across scenarios in reporting the risk estimates does not impact the risk characterization for each of the scenarios within the SWMUs and does not impact the selection of land use scenarios of concern for each of the SWMUs, as the ECLR and systemic hazard at these units are already very high.

Uncertainties Related to Use of Default Values When Estimating Dermal Absorbed Dose

In this assessment, the default dermal absorption factors for soil provided by the Commonwealth of Kentucky in its *Risk Assessment Guidance* (Commonwealth of Kentucky 1995) were used because chemical-specific absorption values were not available. In this guidance the absorption factors, which estimate the percentage of contaminant in soil or sediment crossing the skin and entering the body are 5 percent for inorganic compounds, 10 percent for semivolatile organic compounds, and 25 percent for volatile organic compounds. These factors are much higher than those recommended by EPA Region 4. These factors are 0.1 percent for inorganic chemicals and 1 percent for organic compounds. The effect of the uncertainty in the dermal absorption factor is moderate for the pathway estimates of systemic toxicity and ELCR and small for the total estimates of systemic toxicity and ELCR.

Uncertainties Due to Lack of Toxicity Values for Some Chemicals

Uncertainties due to lack of toxicity values for some chemicals result from two sources in this BHHRA. These are the uncertainty from the use of provisional or withdrawn values and the uncertainty from extrapolating a toxicity value for an administered dose (oral) to an inhalation dose.

The uncertainty from the extrapolation of toxicity values based on administered doses to inhalation doses did not have a significant effect on the BHHRA. Generally, EPA guidance recommends against extrapolating between oral and inhalation toxicity values (EPA 1989a) because of the differing path a chemical entering through the lungs must follow versus entry through the gut. None of the analytes for which extrapolation from oral toxicity to inhalation toxicity was considered in recent risk assessments presented risk; therefore, the effect of this uncertainty on risk results was small.

Uncertainties Associated With Risk Characterization

Two uncertainties are related to risk characterization. The first is the method used to combine HQs and chemical-specific ELCRs over pathways and combine pathway HIs and ELCRs to calculate total HI and ELCR. The second is the uncertainty added to the assessment by combining risks from chemicals and radionuclides. These uncertainties are discussed in the following subsections.

Combining Chemical-Specific Risk Values and Pathway Risk Values

The primary uncertainty in risk characterization is the method used to combine HQs and chemical-specific ELCRs over pathways and combine pathway HIs and ELCRs to calculate total HI and ELCR. The uncertainties involved in combining chemical-specific ELCRs and pathway ELCRs are considerable. The effect of these uncertainties on the total ELCRs presented in the BHHRA is small because a single chemical dominates the pathway ELCR for most pathways; therefore, the potential effect of mixtures is reduced.

Combining Risks from Chemicals With Those from Radionuclides

Some uncertainty is associated with adding risks from chemical exposure to those from exposure to radionuclides. This uncertainty arises from two sources. The slope factors used to characterize the risk from chemicals are derived differently from the slope factors used to characterize risk from radionuclides.

This difference may result in estimates of chemical exposure risks that may be considered to be upper-bound risk estimates and estimates of radionuclide exposure risks that may be considered to be central tendency (i.e., “best,”) estimates. Combining chemical exposure and radionuclide exposure risk estimates to estimate total risk for a land use scenario may place too much emphasis on chemical exposure risk. Second, the mechanism by which chemicals may cause cancer may vary from the mechanism by which radionuclides may cause cancer. Overall, the effect of this uncertainty on the total risk value for each land use scenario is small because generally one COC drives the risks at the SWMUs assessed. At sites where there are multiple chemicals and radionuclides driving risk, the effect of this uncertainty could be moderate.

Summary of Uncertainties for SWMUs 7 and 30

As is shown in the previous subsections concerning uncertainty, the risk estimates could vary if different assumptions were used in deriving the risk estimates or if better information is available for some parameters. The following text summarizes the estimated effects of each uncertainty mentioned previously.

The only uncertainty with an effect estimated to be large is the use of the provisional toxicity values for lead systemic toxicity.

Uncertainties with effects estimated to be moderate are as follows:

- Migration of groundwater to off-site receptors may underestimate risk,
- Use of KDEP dermal absorption values instead of EPA values on the dermal pathway,
- Use of site-specific exposure values on ELCR for the excavation worker,
- Use of site-specific exposure values on ELCR for the current industrial worker,
- Calculation of toxicity values for chemicals, and
- Combination of chemical with radiological ELCRs.

Uncertainties with effects estimated to be small are as follows:

- Inclusion of infrequently detected COPCs,
- Determination of temporal patterns in data,
- Use of quantitation limits that exceed human health PRGs,
- Use of historical data with data collected as part of the RI,
- Inclusion of common laboratory contaminants in the data,
- Removal of analytes based on comparison to blanks,
- Contribution of analytes removed based on comparison to PRGs,
- Removal of analytes based on comparison to background values,
- Assuming that the ditches contained soil and not sediment,
- Determination of exposure points for current concentrations,
- Determination of exposure points for future concentrations,
- Use of total water samples versus filtered,
- Inclusion of biota exposure pathways,
- Use of RME default exposure values instead of central tendency exposure values,
- Inclusion of groundwater in future land use scenarios,
- Omission of livestock in future rural resident land use scenario,
- Omission of an intruder/infrequent recreator land use scenario,
- Lack of summation across land use scenarios and SWMUs on risk characterization,

- Use of KDEP dermal absorption values instead of EPA values on the total risk,
- Use of site-specific exposure values on systemic toxicity for the excavation worker,
- Use of site-specific exposure values on systemic toxicity for the current industrial worker,
- Use of chronic toxicity values for the excavation worker land use scenario,
- Use of provisional and withdrawn toxicity values, except for lead, on ELCR and HI,
- Selection of toxicity values for PCBs,
- Use of inhalation toxicity values extrapolated from oral toxicity values,
- Determination of radionuclide toxicity values,
- Use of absorbed toxicity values calculated from administered toxicity values,
- Combination of risk from chemicals and radionuclides in pathways, and
- Combination of pathway risks to determine land use scenario risk.

UNCERTAINTY IN THE RISK ASSESSMENT for WAG 3 (SWMUs 4, 5 and 6)

Uncertainties Associated with Data and Data Evaluation

Several uncertainties are associated with the data set and the selection of COPCs. Specific uncertainties discussed in the following sections are

- Selection of COPCs,
- Determination of exposure point concentrations under current and future conditions,
- Use of concentrations from total versus filtered samples for inorganic compounds in groundwater, and
- The effect of combining soil and waste for the future excavator land use scenario.

Selection of COPCs

Uncertainty in the selection of COPCs is derived from several sources. The first uncertainty relating to the selection of COPCs is the retention of infrequently detected or infrequently analyzed chemicals in the list of COPCs. Several of the chemicals retained in the list of COPCs were detected in less than 10 percent of the samples. Others were analyzed only in spot samples. The information most likely was obtained from historic data sets containing chemicals that are not now included in standard target compound lists. Inclusion of these infrequently detected or infrequently analyzed compounds in the risk assessment can contribute to uncertainty in different ways. For example, the use of a single detect as a representative concentration of an analyte that was analyzed in only one sample may overestimate or underestimate hazard or risk to an unknown extent since the nature and extent of this contaminant has not been delineated. By contrast, the incorporation of infrequently detected analytes into the risk determinations at least gives the assurance that the derived representative concentration probably will result in conservative risk estimates provided that all potential “hot-spots” were sampled at the site. For both infrequently detected and infrequently analyzed chemicals, the greatest concern is that some of these COPCs are retained as COCs. The data indicate, in most cases (37/48 comparisons), that the infrequently detected COPCs had no effect on the risk or hazard estimates. Even where a difference was noted, for example, for a future industrial worker exposed to RGA groundwater at SWMU 5, eliminating the ELCR for the infrequently detected carcinogenic contaminant, radium-226, resulted in a reduction in the total ELCR from 5.39E-04 to 1.93E-04, a decrease that falls within the range applicable to a small contribution to uncertainty (approximately 2.5-fold). The estimated effect of this uncertainty on the risk estimates as a whole is small (less than one order of magnitude).

The second uncertainty relating to selection of COPCs is that temporal patterns in detection of analytes were not considered when selecting COPCs. If temporal patterns were considered, the final risk results in this BHHRA may have been quite different depending on the times at which risks were estimated; however, in the time frame considered in this BHHRA (40 years), the assumed effect of this uncertainty on the risk estimates is small.

The third uncertainty relating to selection of COPCs concerns the quantitation limits used for some analytes. For many organic analytes and some radionuclides, the quantitation limit exceeds a concentration or activity that may result in a significant health effect. For example, for RGA groundwater at SWMU 4, 27 organic compounds have quantitation limits for at least one sample that exceed residential use ELCR RBCs. Similarly, in subsurface soil at SWMU 6, 15 organic compounds have quantitation limits for at least one sample that exceed the residential use ELCR RBC. Because the quantitation limits exceed the RBCs, it is possible that these contaminants could be present in amounts that contribute to risk, but they are not retained as COPCs for quantitative evaluation. Because many of these components tend to be unrelated to past practices at WAG 3, the effect of this uncertainty on the risk estimates for this land use scenario is likely to be small.

The fourth uncertainty relating to the selection of COPCs is the inclusion of common laboratory contaminants in the COPC list. Common laboratory contaminants retained as COPCs had almost no effect on hazard and risk estimates; therefore, the estimated effect of the uncertainty on risk estimates is small.

The fifth uncertainty relating to the selection of COPCs is based on the fact that, in this risk assessment, contaminant concentrations were not compared to concentrations found in blank samples. Typically, common laboratory contaminants and other analytes may be deleted from the COPC list if they also are detected in blank samples at appropriate concentrations (RAGS). While not performing this test ensures that all detects for compounds that may be potential laboratory contaminants are treated as site contaminants, it is actually impossible to distinguish unequivocally between the environmental medium and the laboratory-based extraction procedures and processes as the potential source of the observed contamination. The effect of this uncertainty is estimated to be small because, in general, common laboratory contaminants such as acetone, methylene chloride, and the phthalates appear to be sporadically present, or at low concentrations only, thereby not contributing significantly to risk.

The sixth uncertainty relating to the selection of COPCs is the use of a toxicity screen to determine the final COPC list. In this BHHRA, the maximum detected concentrations of analytes within each -medium ~at-each SWMU were compared to residential human health risk-based screening criteria. The residential risk-based screening levels were used per regulatory agreement (Methods Document). Analytes with maximum concentrations less than these screening criteria were removed from the list of COPCs. The marginal contribution of the analytes removed from the COPC list is negligible; therefore, the estimated effect of this uncertainty on the final risk estimates is small.

The seventh uncertainty relating to the selection of COPCs is the use of a soil background screen to determine the final COPC list. The maximum detected concentrations of naturally occurring analytes in surface and subsurface soil were screened against prevailing background concentrations as part of the COPC selection process. While the final hazard and risk estimates are greater in the absence of background screening, overall differences are small to medium. Because this screen relies on a comparison of the maximum detected concentration of each analyte in a medium to the selected soil background concentration, erroneous removal of analytes from the list of COPCs is unlikely.

The eighth uncertainty relating to the selection of COPCs is the omission of a groundwater background screen to determine the final COPC list. Unlike soil, a background screen for groundwater was not used

when developing the list of COPCs, a decision resulting from uncertainty surrounding the suitability of existing background groundwater data for screening purposes. Background values presented in Table A.13 of the Risk Methods Document (DOE 2001) were not available at the time to compare the overall hazards and risks for the future industrial and residential land use scenarios with and without the contribution of these screened COPCs. This gives a measure of the degree of uncertainty bounding the hazard and risk determinations when a background groundwater screen is omitted. The overwhelming majority of comparisons return the same value for the HI or ELCR whether or not the background screen is in place. Where a difference is noted, the difference is far less than an order of magnitude; therefore, the effect of this uncertainty is small.

The final uncertainty associated with COPC selection is related to the manner in which long-lived radionuclides with short-lived decay products are assessed. The method that the risk assessment uses to aggregate data on the radioactivity of isotopic parents and decay products not only impacts COPC selection directly but also, less predictably, the overall contribution of the detected radionuclides to the site-specific ELCR. The impact on quantifiable risk, while uncertain, is likely to be small if the assumption of secular equilibrium is valid.

Use of Concentrations from Total Versus Filtered Groundwater Samples

In this BHHRA, all analyte concentrations in water are from the analyses of unfiltered or total samples. The use of data from analyses of total samples is consistent with current EPA guidance (RAGS), but introduces an additional uncertainty to the BHHRA for some water-use pathways. The magnitude of the effect of this uncertainty upon the risk estimates is difficult to determine because the extent to which the quality of water (in terms of total solids) from a residential well could differ from the quality of water collected during the recent sampling effort is unknown. Because the groundwater samples used in this BHHRA were from boreholes, some samples had high solid content.

The HI estimates calculated using unfiltered water from RGA and McNairy groundwater at WAG 3 differed from those HIs calculated using *only* filtered samples by more than one order of magnitude in almost every case. By contrast, the available ELCRs for filtered and unfiltered groundwater (RGA at SWMU 4) suggest only a small contribution to uncertainty through filtering. Taken together, these results are consistent with the concept that the bulk of the turbid material removed from groundwater during filtering will be those inorganic components contributing most to the calculated systemic toxicity. In summary, the effect of this uncertainty on the ELCR determination is small, but medium-to-large for the HI determinations.

Uncertainties Associated with Toxicity Assessment

Uncertainties related to the toxicity assessment are from the following three sources: 1) uncertainty because toxicity values are lacking for some chemicals, 2) uncertainty in the calculation of toxicity values by EPA, and 3) uncertainty in the calculation of absorbed dose toxicity values from administered dose toxicity values. Each of these is discussed in the following paragraphs.

Uncertainties Because Toxicity Values are Lacking for Some Chemicals

Uncertainties due to lack of toxicity values for some chemicals result from three sources in this BHHRA; these are the uncertainty from the use of provisional or withdrawn values, the uncertainty from extrapolating a toxicity value for an administered dose (oral) to an inhalation dose, and the uncertainty associated with a lack of site-specific information on chemical speciation for such elements as mercury and chromium.

Uncertainty associated with the use of provisional or withdrawn toxicity values has a significant effect on the results of the BHHRA. Some COPCs do not have approved toxicity values, so a provisional or withdrawn value was used. The most notable of these COPCs is lead. This provisional RfD toxicity value was provided by KDEP in a comment package on the WAG 17 RI/BHHRA. As discussed in Sect. 1.5, the systemic toxicity posed by lead dominates all land use land use scenarios at SWMUs where lead was detected. For better interpretation of the HIs for the rest of the COPCs in the BHHRA, results with and without contributions from lead are provided throughout this BHHRA. A further example of the uncertainty of using provisional or withdrawn toxicity values to compute HIs and/or cancer risk estimates is provided by the HQs that can be derived for iron. This issue is important in this risk assessment because iron appears as a priority COC and risk driver for many operative exposure pathways at WAG 3 SWMUs. A number of lines of argument point to the use of an overly conservative provisional RfD of 3.0E-01 mg/kg-day, resulting in a series of unrealistically high contaminant-specific HQs. These have, consequently, in the absence of lead, served as the major components of pathway-specific HIs at WAG 3 SWMUs.

The provisional RfD for iron also has formed the basis for an RBC of 310 mg/kg, a screening value above which most detected concentrations of the element have fallen, as might be expected for locations where an average concentration of 28,000 mg/kg has been established for sitewide background levels of this element. A risk-based concentration approximately two orders of magnitude lower than background ensures that an unrealistically high HQ will be derived for any contaminants whose prevailing concentration maxima (as with iron) fail the background screen, thereby forcing the element's inclusion in the risk/hazard computations.

Because iron is an essential nutrient as well as a possible environmental contaminant, a measure of the overly stringent nature of the provisional RfD for the element is provided by consideration of the RDA for iron of 10 mg/day. For a 14.5 kg child, this is a daily dose of 6.9E-01 mg/kg-day, a level below which a child's intake of iron should not fall if satisfactory health and development is to be maintained. Based on the provisional RfD of 3.0E-01 mg/kg-day for the element, the RDA dose-equivalent is also more than twice that above which a child's intake of iron must not vary. Such a discrepancy points strongly to the uncertainties inherent in the use of the 3.0E-01 mg/kg-day toxicity value for iron in risk/hazard computations.

As a further contribution to the understanding of this uncertainty. ;In general, most risk and hazard estimates fluctuate within relatively narrow limits, the risk to some receptors at certain SWMUs either being identical or differing by factors ranging from close-to-unity to approximately 10. Accordingly, the overall impact on risk of using provisional and withdrawn toxicity values in this risk assessment is small to medium.

Another source of uncertainty associated with choice of toxicity values arises from metals such as chromium that can exist in more than one oxidation state. In this and earlier risk assessments at PGDP, the simplifying assumption was made that all of the element was in the Cr VI form. This is conservative in light of the lower (and therefore more stringent) chronic oral RfD for Cr VI compared to Cr III (3.0E-03 and 1.5E+00 mg/kg-day, respectively), and the fact that, unlike Cr VI, there is no carcinogenic benchmark for Cr III. Because the exact proportions of Cr VI and Cr III in Paducah environmental media are unknown, to make the assumption that all of the element is present as Cr III sets the lower extreme of the range within which the true but unknown contribution to total hazard or risk from this element will fall. The overall impact of such a change can be seen most clearly where, as for the current industrial worker exposed to soil at SWMU 4 (total scenario/exposure route HI of 3.62), chromium is one of the driving COCs (contributing 45 percent to the total HI). Employing the chronic oral and dermal RfDs for Cr HI (there is no RfC for Cr HI) to calculate the HQ applicable to this receptor reduces the contribution from chromium and results in a total HI of 2.02, a 44 percent decrease compared to the default value of

3.62. This change points to the likelihood that the overall impact of this uncertainty is small according to established criteria (less than one order of magnitude).

In the past, there was uncertainty in the selection of the appropriate toxicity value for PCBs (e.g., Aroclor-1254, -1260, etc.) because of 1) difficulty in identifying specific Aroclors in a mixture, 2) different rates of decay among the Aroclors in environmental media, and (3) the effects of weathering processes on the congener-specific “fingerprint” over time, a process making the Aroclors appear to be more chlorinated than they are actually. To address these concerns and to ensure that the risk numbers for Aroclors are suitably conservative, KDEP requires that all PCBs be evaluated as Aroclor-1260. This assessment conforms to KDEP guidance because oral slope factors for all Aroclors were assumed to be equal to 2.0 (mg/kg-day),¹ consistent with recent EPA guidance (EPA 1996a) and identical to the current upper-bound slope factor for total PCBs found on IRIS (EPA 2000). Results for exposures to multiple Aroclors are summed to generate a total PCB-specific value. Unlike earlier assessments performed at PGDP, in which the effect of uncertainty in the selection of toxicity values for PCBs on the final risk values may have been moderate, the effect of this uncertainty on ELCR determinations is likely to be small.

Although the impact of addressing the ELCR due to multiple Aroclors as total PCBs appears to be small, the potential exists for this approach to result in lower overall total HIs, because there are no verified or provisional chronic oral RfDs or chronic inhalation RfCs for either Total PCBs or Aroclor-1260. The potential contribution of these compounds to systemic toxicity is not represented in this risk assessment; however, this does not mean that the Aroclors are likely to be without non-carcinogenic toxicological impacts, because verified chronic oral RfDs are available for Aroclors-1016 and -1254 (7.0E-05 and 2.0E-05 mg/kg-day, respectively). To address the semiquantitative impact of this possible underestimation of the noncarcinogenic hazard due to the presence of total PCBs at WAG 3, the conservative assumption was made that the chronic oral RfD for Aroclor-1254 (2.0E-05 mg/kg-day) would be applicable to total PCBs, and the HI for a future on-site child resident at SWMU 4 was recalculated with the potential toxicity of these compounds taken into account. Driven by the ingestion of vegetables exposure route (HQ of 22.3), a compound-specific HI of 23 was obtained by this strategy resulting in an increase in the scenario-specific HI from 98.2 (lead was not present in the surface soil at SWMU 4) to 121.2. The consequence of the possible underestimation of the systemic toxicological hazard due to non-consideration of total PCBs is probably small.

Including inhalation toxicity values extrapolated from administered doses in the risk characterization would not have significantly affected the results of the BHHRA. EPA guidance (RAGS) recommends against extrapolating between oral and inhalation toxicity values because of the differing path a chemical entering through the lungs must follow before exerting its effect compared to that of a chemical entering via the gut. Examination of this form of extrapolation as an uncertainty in assessments for PGDP was requested by the regulatory community. Previous work at PGDP, in which this effect was examined quantitatively, determined that including extrapolated inhalation toxicity values in the risk characterization resulted in insignificant changes in the final risk estimates. Therefore, the estimated effect of this uncertainty on the risk results is small.

Uncertainties in Deriving Toxicity Values

Standard EPA RfDs and slope factors were used to estimate potential noncarcinogenic and carcinogenic health effects from exposure to chemicals. Considerable uncertainty is associated with the method applied to derive slope factors and RfDs, even though EPA has working groups that review all relevant human and animal studies for each compound and select studies pertinent to the derivation of the specific RfD and slope factor. For example, the primary information often involves data from experimental studies in animals, high exposure levels, and exposures under acute or occupational conditions.

Extrapolation of these data to humans under low-dose and chronic conditions introduces uncertainties, the magnitude of which is addressed by applying uncertainty factors to the dose response data for each applicable uncertainty. These factors are incorporated to provide a margin of safety for use in human health risk assessments. The effect of uncertainties in calculating chemical toxicity values is moderate.

Unlike that associated with chemical toxicity, quantitative uncertainty associated with radionuclide toxicity values is probably small. Thus, the dose-response relationship between cancer and ionizing radiation has been evaluated in many reports and is well established. In addition, unlike toxicity values for chemicals, risk factors for radionuclides are extrapolated from the cancer risk established using the Japanese Atomic Bomb Survivors database and a relative risk projection model. Accordingly, carcinogenic slope factors for radionuclides are based on human data and are likely to be more accurate, thereby contributing only minimally to uncertainty.

Uncertainties Due to Calculation of Absorbed Dose Toxicity Values from Administered Dose Toxicity Values

Uncertainty exists in the validity of the calculations used to convert an administered dose toxicity value to an absorbed dose. Of particular importance is the lack of consideration of point-of-contact effects in this calculation. For example, some organic analytes (e.g., PAHs) can cause a toxic or carcinogenic response in skin, an effect that is not considered in the calculation of absorbed dose toxicity values from administered dose toxicity values using EPA protocols. Similarly, the administered dose-response for many chemicals relies on the delivery of a high concentration of contaminants to the liver via the portal system after ingestion. This effect is not seen if a contaminant is absorbed through the skin because of the larger distribution space for the contaminant absorbed through the skin. Even with these uncertainties, the effect of the uncertainty in calculation of absorbed dose toxicity values from administered dose toxicity values upon the risk estimates is likely to be small.

Uncertainties Associated with Risk Characterization

Two uncertainties are related to risk characterization. The first is the validity of combining HQs and chemical-specific ELCRs over pathways and of combining pathway HIs and ELCRs to derive a total HI and ELCR. The second is the justification for combining risks from chemicals and radionuclides. These uncertainties are discussed in the following sections.

Combining Chemical-Specific Risk Values and Pathway Risk Values

The method used to calculate pathway HIs and ELCRs in the BHHRA follows EPA protocols (RAGS, Methods Document) and involves the simple addition of chemical-specific HQs and ELCRs to obtain pathway HIs and ELCRs, respectively. The method assumes that all effects among chemicals are additive, an assumption made by the EPA in the absence of any evidence to the contrary. Certain limitations of the approach have been noted by EPA in RAGS, as follows:

- Little is known about the effects of chemical mixtures. Although additivity is assumed, the interaction of multiple chemicals possibly could be synergistic or antagonistic.
- The RfDs and RfCs do not have equal accuracy or precision and are not based on the same severity of effects.
- Dose additivity is most properly applied to compounds that induce the same effect by the same mechanism of action. While the approach recommended by EPA is a useful screening-level approach,

the cumulative systemic toxicity could be overestimated for chemicals that act by different mechanisms and/or on different target organs.

The effect of this uncertainty on the estimate of systemic toxicity depends on how many contaminants drive systemic toxicity and whether the contaminants have different endpoints. In this BHHRA, comparatively few contaminants drive systemic toxicity for most land use scenarios, with individual, contaminants typically contributing significant levels of risk for some exposure scenarios; therefore, the overall effect of this uncertainty on HIs is small.

EPA has reported specific limitations for this approach in regard to chemical carcinogenesis (RAGS):

- Cancer risks (i.e., ELCRs) are based on slope factors that represent an upper 95th percentile estimate of potency, the probability distributions of which are unlikely to be strictly additive; therefore, summing these risks can result in an overly conservative estimate of lifetime ELCR.
- Cancer risks may not be additive. By analogy to systemic toxicity effects, the endpoints may differ, and mechanisms of effect may vary.
- EPA assigns qualitative weight-of-evidence descriptors to its designation of chemicals as carcinogens or non-carcinogens [A, B, C, D or E, as specified in the Agency's current guidelines (EPA 1986), or "known/likely," "cannot be determined," or "not likely," as suggested in their recent proposed guidelines (EPA 1996b)]. Where the quantitative data are sufficient, the agency develops slope factors for compounds assigned to the A, B, C or "known/likely" categories, for which the level of uncertainty might be expected to be less for those contaminants at the high-confidence end of the range of available qualitative descriptors (i.e., "A" or "known"). However, the simple combination of ELCRs for potential carcinogens "across-the board" in this BHHRA does not take this hierarchy into account.

Thus, the uncertainties involved in combining chemical-specific ELCRs and pathway ELCRs are considerable. The effect of these uncertainties on the total ELCRs presented in the BHHRA is small because, for the most part, a single (or very few) chemical dominates the pathway ELCR for most pathways at some SWMUs. In such circumstances, the uncertainty associated with the potential effect of mixtures is reduced.

Summary of Uncertainties for SWMUs 4, 5 and 6

As shown in the previous sections, risk estimates may vary if different assumptions are used in deriving risk estimates or if better information is available for some parameters. The following text summarizes the estimated effects of each uncertainty discussed previously.

One uncertainty with an effect that is estimated to be large is the use of a provisional toxicity value for the systemic toxicity of lead. Because the uncertainty regarding the RfD for lead was identified as being large and easy to quantify, the summary discussions in this BHHRA are more detailed than for some of the other uncertainties. This discussion is not meant to imply that the authors believe the provisional toxicity value for lead provided by KDEP is incorrect.

Another factor in the risk assessment that makes a large contribution to uncertainty is the use of KDEP defaults versus site-specific estimates for the exposure duration and frequency at which a current industrial worker will be exposed to contamination at the subject SWMUs.

In addition to the above, the use of total water samples versus filtered can contribute greatly to systemic toxicity for certain scenario/exposure route combinations.

Following are uncertainties with effects estimated to be moderate:

- Use “of site-specific exposure” values on systemic toxicity and cancer risks “for the excavation worker”
- Underestimation of risk due to migration of groundwater to off-site receptors
- Calculation of toxicity values for chemicals
- Evaluation of groundwater separately from soil in future land use scenarios
- Use of total water samples versus filtered
- Removal of analytes as COPCs based on a comparison to soil background values
- Use of site-specific exposure values on systemic toxicity for the current industrial worker
- Use of provisional and withdrawn toxicity values on the total systemic toxicity and cancer risks
- Use of KDEP dermal absorption values instead of EPA values on the total systemic toxicity and cancer risks

Following is a list of uncertainties with effects estimated to be small:

- Use of total water samples versus filtered
- Use of KDEP versus EPA dermal absorption values in systemic toxicity and risk calculations
- Applying the toxicity values for Cr VI to “total” chromium
- Use of quantification limits that exceed human health RBCs
- Inclusion of biota exposure pathways
- Inclusion of infrequently detected/infrequently analyzed COPCs
- Lack of determination of temporal patterns in data
- Inclusion of common laboratory contaminants
- Lack of analyte comparison to blanks
- Contribution of analytes removed based on a toxicity screen
- Removal of analytes based on comparison to background values
- Lack of approved groundwater background concentrations
- Determination of exposure points for current concentrations
- Determination of exposure points for future concentrations
- Use of RME default exposure values instead of central tendency exposure values
- Evaluation of groundwater separately from soil in future land use scenarios
- Omission of livestock in future rural resident land use scenario
- Omission of an intruder/infrequent recreator land use scenario
- Use of KDEP dermal absorption values versus EPA values on total systemic toxicity and cancer risks
- Use of site-specific exposure values on systemic toxicity and cancer risks for the excavation worker
- Use of site-specific exposure values on systemic toxicity for the current industrial worker

- Use of chronic toxicity values for the excavation worker land use scenario
- Use of provisional and withdrawn toxicity values on the total systemic toxicity and cancer risks
- Selection of toxicity values for PCBs
- Use of inhalation toxicity values extrapolated from oral toxicity values
- Determination of radionuclide toxicity values
- Use of absorbed toxicity values calculated from administered toxicity values
- Combination of risk from chemicals and radionuclides in pathways
- Combination of pathway risks to determine land use scenario risk
- Omission of the chemical properties of uranium from the toxicity evaluation
- Exclusion of VOC data rejected because of missed holding times
- Heterogeneity of pit contents and soil as sources for SWMU-specific risks and hazards
- Use of “+D” slope factors to derive carcinogenic risk for all members of an isotopic decay series

F.7. CONCLUSIONS

This section summarizes the results of this and previous BHHRA and draws conclusions from the results. The primary purpose of this section is to provide a concise summary of each of the BHHRA steps without the use of tables, extensive explanations, or justifications. This section also includes a series of observations in which the results of the BHHRA are combined with the uncertainties in the risk assessment.

F.7.1 CHEMICALS OF POTENTIAL CONCERN

Through a series of screening steps, which follow the Risk Methods Document (DOE 2001) and other regulatory agency approved procedures, the data sets were reduced to lists of COPCs for the entire BGOU. Conclusions are compiled from the previous BHHRA [i.e., WAG 22 RI Addendum for SWMUs 2 and 3 (DOE 1994); WAG 3 RI for SWMUs 3, 4, and 6 (DOE 2000)]; and WAG 22 RI for SWMUs 7 and 30 (DOE 1998a)] and Tables F.3 through F.10.

F.7.2 EXPOSURE ASSESSMENT

Historical information and newly collected data were used to develop a CSM. After consideration of the available data and scope of the SI, the potential receptor population under current conditions at the source units is industrial workers, and the potential receptor populations under future conditions are industrial workers, excavation workers, recreational users, and on-site rural residents. The potential receptor populations under future conditions in BGOU areas are recreational and residential. Within these broad categories, the recreational users and rural residents contain age cohorts. For the recreational users, the cohorts include the child (aged 1 to 7), teen (aged 8 to 20), and the adult (older than 21). For rural residents, the cohorts include children (aged 1 to 7) and older individuals (termed adults in this and previous BHHRA). The recreational user and the rural resident population may also contain sensitive subpopulations such as pregnant women, young children (aged 0 to 1), the elderly, and the infirm. In this and earlier BHHRA, exposure by these subpopulations is not quantified because much of the information that is needed is not available. Finally, this and earlier assessments assume that the recreational user is a rural resident who has repeated access to the study area. Recreational users not residing in the study area are not considered separately because nearby residents were determined to be the individuals most likely to take part in recreational activities at PGDP on a continual basis. In addition, the exposure assessment determined that little information useful in remedy selection would be obtained by including a separate visiting recreational user in the assessment. The groundwater exposure scenarios are hypothetical (i.e., at the plant boundary, property boundary, and near the Ohio River) because the areas containing the POEs currently are used for recreational and industrial purposes and do not contain residences. Table F.15 in the CSM section shows the scenarios and media evaluated in this risk assessment. The exposure routes considered are listed below.

Industrial Worker

- Ingestion of surface soil and sediments (considered as one media type)
- Dermal contact with surface soil
- Inhalation of vapors and particulates emitted by surface soil
- External exposure to ionizing radiation

Groundwater exposure was evaluated in previous risk assessments for the industrial worker scenario, but this exposure was not included in the BHHRA. Groundwater exposures are not commonly included in the industrial worker scenario because institutional controls are assumed to be in place.

Future Excavation Worker

Ingestion of surface and subsurface soil
Dermal contact with surface and subsurface soil
Inhalation of vapors and particulates emitted by surface and subsurface soil
External exposure to ionizing radiation

Future Recreational User

Ingestion of surface soil
Dermal contact with surface soil
Inhalation of vapors and particulates emitted by surface soil
Ingestion of game
External exposure to ionizing radiation

The listing above includes all exposure routes included in each previous risk assessment. SWMUs 2, 3, and 145 did not have a recreational user scenario; however, when it was performed at other SWMUs the ELCR from ingestion of game was the driver for this scenario.

Future On-Site Rural Resident

Ingestion of surface soil
Dermal contact with surface soil
Inhalation of vapors and particulates emitted by surface soil
External exposure to ionizing radiation
Ingestion of groundwater
Vapor intrusion
Dermal contact with groundwater while showering
Inhalation of vapors emitted by groundwater during household use and
Inhalation of vapors emitted by groundwater while showering
Ingestion of produce

Future Off-Site Rural Resident

Ingestion of surface soil
Dermal contact with surface soil
Inhalation of vapors and particulates emitted by surface soil
External exposure to ionizing radiation
Ingestion of groundwater
Dermal contact with groundwater while showering
Inhalation of vapors emitted by groundwater during household use and
Inhalation of vapors emitted by groundwater while showering
Vapor intrusion

F.7.3 TOXICITY ASSESSMENT

The majority of toxicity values used in the risk assessment were taken from the RAIS and the TCE slope factor was taken from KDEP. After compiling toxicity information, the determination was made that the majority of the COPCs had a toxicity value available for one or more routes of exposure (see Section F.4.6).

F.7.4 RISK CHARACTERIZATION

Quantitative risks were computed by integrating the CDIs tabulated from the exposure assessment and toxicity values calculated from the toxicity assessment. For soil, results from previous risk assessments for SWMUs 2, 3, 4, 5, 6, 7, and 30 (DOE 1994; DOE 1998a; DOE 2000) were used. For most of the SWMUs, no new surface data have been collected since the previous risk assessments were performed. In addition, the soils at these units are outside the scope of this BGOU BHHRA as noted in the approved work plan; therefore, a new quantitative risk assessment was not performed for soils. Significant findings are summarized below (see Table F.80).

F.7.4.1 Land Use Scenarios of Concern

The following are land uses of concern for BGOU:

Industrial: SWMUs 2, 3, 4, 5, 6, 7, and 30

Excavation: SWMUs 4, 5, 6, 7, and 30

Recreational: SWMUs 5, 7, and 30

On-Site Residential: SWMUs 2, 3, 4, 5, 6, 7, 30, and 145

Off-Site Residential: SWMUs 2, 3, 4, 5, 7, 30, and 145

F.7.4.2 Contaminants of Concern for Soil

To make a determination about whether contaminants are of concern in soil, quantitative risk and hazard results over all pathways from the previous risk assessments for SWMUs 2, 3, 4, 5, 6, 7, and 30 (DOE 1994; DOE 1998a; DOE 2000) were compared to risk and hazard benchmarks for land use scenarios of concern. The benchmarks used for this comparison were a) 0.1 for HI and b) 1×10^{-6} for ELCR. Lead results in the previous risk assessments were derived with toxicity values that are no longer used.

Contaminants with chemical-specific HIs or ELCRs exceeding these benchmarks are deemed COCs. Priority COCs are contaminants whose chemical-specific HI is greater than 1 or whose ELCR is greater than 1×10^{-4} for one or more scenarios. The following are priority COCs found in soil at individual SWMUs.

SWMU 2—none

SWMU 3—none

SWMU 4—barium, beryllium, cadmium, chromium, iron, nickel, uranium, vanadium

SWMU 5—aluminum, arsenic, beryllium, chromium, nickel, and Total PAHs

SWMU 6—Total PAHs, beryllium, chromium, nickel

SWMU 7–aluminum, antimony, arsenic, beryllium, cadmium, chromium, iron, manganese, nickel, uranium, vanadium, benzo(a)pyrene, dibenzo(a,h)anthracene, Aroclor-1254, Aroclor-1260, ²³⁹Pu, ²³⁴U, ²³⁵U, ^{235/236}U, and ²³⁸U

SWMU 30–aluminum, antimony, arsenic, barium, beryllium, cadmium, chromium, copper, iron, manganese, mercury, nickel, uranium, vanadium, benzo(a)anthracene, benzo(a)pyrene, benzo(b)fluoranthene, dibenzo(a,h)anthracene, indeno(1,2,3-cd)pyrene, Aroclor-1254, Aroclor-1260, ²³⁴U, ²³⁵U, ^{235/236}U, and ²³⁸U

Table F.80. Scenarios for Which Human Health Risk Exceeds *De Minimis* Levels^a

Scenario	Location							
	SWMU 2	SWMU 3	SWMU 4	SWMU 5	SWMU 6	SWMU 7	SWMU 30	SWMU 145
Results for excess lifetime cancer risk:								
Current On-site Industrial Worker	X	X	X	X	X	X	X	NA
Exposure to Soil								
Future On-site Industrial Worker	X	X	X	X	X	X	X	NA
Exposure to Soil	NA	NA	NA	NA	NA	NA	NA	NA
Exposure to Surface Water								
Future On-site Excavation Worker	NA	NA	X	X	X	X	X	NA
Exposure to Soil/Soil and Waste								
Future On-site Recreational User	NA	NA	---	X	X	X	X	NA
Exposure to Game	NA	NA	NA	NA	NA	NA	NA	NA
Exposure to Soil								
Future On-site Rural Resident	NA	NA	X	X	X	X	X	NA
Exposure to Soil	X	X	X	X	---	X	X	X
Exposure to Groundwater ^b	X	X	X	---	---	X	X	---
Vapor Intrusion ^c								
Future Off-site Rural Resident	X	X	X	X	---	X	X	X
Exposure to Groundwater ^b	X	---	X	---	---	---	X	---
Vapor Intrusion ^c								
Result for Systematic Toxicity^b								
Current On-site Industrial Worker	---	---	X	---	---	X	X	NA
Exposure to Soil								
Future On-site Industrial Worker	---	---	X	---	---	X	X	NA
Exposure to Soil								
Future On-site Excavation Worker	NA	NA	X	X	X	X	X	NA
Exposure to Soil/Soil and Waste								
Future On-site Recreational User	NA	NA	---	---	---	---	---	NA
Exposure to Game	NA	NA	NA	NA	NA	NA	NA	NA
Exposure to Soil	NA	NA	NA	NA	NA	NA	NA	NA
Exposure to Surface Water								

Table F.80. Scenarios for Which Human Health Risk Exceeds *De Minimis* Levels^a (Continued)

Scenario	Location							
	SWMU 2	SWMU 3	SWMU 4	SWMU 5	SWMU 6	SWMU 7	SWMU 30	SWMU 145
Future On-site Rural Resident	NA	NA	X	X	X	X	X	NA
Exposure to Soil	X	X	X	X	---	X	X	X
Exposure to Groundwater ^b	X	X	X	---	---	X	X	X
Vapor Intrusion ^c								
Future Off-site Rural Resident	X	---	X	---	---	X	X	---
Exposure to Groundwater ^b	---	---	---	---	---	---	---	---
Vapor Intrusion ^c								

Notes: Scenarios where risk exceeds *de minimis* levels are marked with an X. Scenarios where risk did not exceed *de minimis* levels are marked with a ---. NA indicates that the scenario/land use combination was not assessed because the scenario is not applicable, or the medium is not present.

^a Consistent with the PGDP Risk Methods Document (DOE 2001), the *de minimis* levels used are a cumulative ELCR of 1×10^{-6} and a cumulative HI of 1.

^b Systemic toxicity results summarized here for the resident and recreational user are for the child. The off-site POE considered is the property boundary.

^c Based on results of preliminary deterministic contaminant transport modeling. The POE is the property boundary. X indicates that the location contains a source of unacceptable off-site contamination, and --- indicates that the location is not a source of off-site contamination

F.7.4.3 Contaminants of Concern for Groundwater

Similarly for groundwater, to determine whether modeled concentrations of contaminants are of concern, quantitative risk and hazard results over all pathways were compared to risk and hazard benchmarks for land use scenarios of concern. The benchmarks used for this comparison were a) 0.1 for HI and b) 1×10^{-4} for ELCR.

Contaminants with chemical-specific HIs or ELCRs exceeding these benchmarks were deemed COCs. Priority COCs are contaminants whose chemical-specific HI is greater than 1 or whose ELCR is greater than 1×10^{-4} for one or more scenarios. The following presents priority COCs found in groundwater at individual SWMUs.

SWMU 2—arsenic; manganese; uranium; *cis*-1,2-DCE; TCE; and Aroclor-1248

SWMU 3—arsenic; manganese; uranium; and ⁹⁹Tc

SWMU 4—arsenic, manganese; *cis*-1,2-DCE; TCE; vinyl chloride; and ⁹⁹Tc

SWMU 5—arsenic; manganese, uranium and naphthalene

SWMU 6—none

SWMU 7—arsenic, 1,1-DCE; *cis*-1,2-DCE; Aroclor-1254, TCE, vinyl chloride

SWMU 30—arsenic

SWMU 145—antimony; arsenic; manganese; Aroclor-1260; and ⁹⁹Tc

“Priority COCs”² are identified in this section as an aid to risk managers during decision making. The priority COCs identified above in this risk assessment are based on the modeled groundwater concentrations at all POEs.

F.7.4.4 Pathways of Concern

Each of the pathways included in the BHHRA is a POC.

² “Priority COCs” are identified as an aid to risk managers during decision making; however, all COCs will be addressed through remediation, removal, management, or other enforceable control.

F.7.4.5 Media of Concern

Media of concern are those media that appear in at least one POC. Because they contribute to at least one POC, soil and RGA groundwater are media of concern at all eight SWMUs.

F.7.5 OBSERVATIONS

Consistent with regulatory guidance and agreements contained in the PGDP Risk Methods Document, this and previous BHHRA presents risks for land use scenarios representing current use, as well as several reasonable future uses. Risk evaluation of surface soil was conducted for all SWMUs as part of the evaluation of the scenarios specified in the work plan. The oral slope factor for beryllium was withdrawn in 1998. The inhalation slope factor for beryllium is $8.4 \text{ (mg/kg-day)}^{-1}$. The Inhalation Slope Factor was calculated from inhalation unit risk as described in Supplemental Guidance from *RAGS: Region 4 Bulletins, Human Health Risk Assessment* (Interim Guidance) as described in the RAIS online toxicity values. A review of the pathways for exposure reveals inhalation of particulates is a minor contributor to cumulative ELCR. Inhalation of particulates is consistently less than 1% of the cumulative ELCR; therefore, cumulative ELCRs are shown with and without the contribution from beryllium in this section. In addition, total PAH and individual PAH compounds (depending on how the risks were calculated in previous risk assessments) contribute to risk in many scenarios. PAH compounds are ubiquitous and may be products of anthropogenic activities other than the PGDP. However, there is not a definitive study that describes the concentrations of PAHs in nearby areas that are not influenced by the PGDP. Until a definitive study is prepared, the risks of PAH compounds are included in this document. The scenarios described in the BHHRA are as follows:

- Future on-site industrial use—direct contact with surface soil (soil found 0 to 1 ft bgs). Groundwater was assessed in some the previous risk assessments as part of the industrial use scenario. These results for groundwater were not included in the on-site industrial scenario for this BHHRA. Future on-site excavation worker—direct contact with surface and subsurface soil (soil 0 to 10 ft bgs).
- Future recreational user—direct contact with surface soils and consumption of game exposed to surface soils.
- Future on-site rural resident—direct contact with surface soil at and use of groundwater drawn from the RGA at source areas and vapor intrusion into basements.
- Future off-site rural resident—use in the home of groundwater drawn from the RGA at the DOE plant boundary, property boundary, and the Ohio River.

Specific observations for this BHHRA are presented below.

F.7.5.1 Observations—Future Industrial Worker

Cumulative HIs for the industrial worker were greater than 1 at SWMUs 4, 7 and 30 based on soil exposure. At SWMU 4, chromium, iron, and vanadium were the primary drivers contributing 45%, 24%, and 24% to the HI, respectively. At SWMU 7, beryllium, chromium, iron, manganese, uranium, and vanadium were the major drivers (> 5%) contributing 9.6%, 13.6%, 20.6%, 10.7%, 13.7%, 13.7%, and 17.7% to the HI, respectively. At SWMU 30, aluminum, beryllium, chromium, iron, manganese, uranium, and vanadium were the major drivers contributing 5.1%, 10.8%, 13.5%, 19.8%, 11.3%, 9.0%, and 17.6% to the HI, respectively.

Cumulative ELCRs exceeded 1E-04 for all SWMUs and were greater than 1E-03 at SWMU 7 and SWMU 30 for exposure to soil (SWMU 145 was not evaluated for this scenario). The following summarizes the cumulative risk estimates and major contributors (> 5%) to the ELCR for these SWMUs.

- SWMU 2 cumulative ELCR 1.20E-04; drivers are ^{235}U +daughters at 83.9%, ^{238}U +daughters at 10.7%.
- SWMU 3 cumulative ELCR 1.20E-04; drivers are ^{235}U +daughters at 83.9% and ^{238}U +daughters at 10.7%.
- SWMU 4 cumulative ELCR 5.40E-04; the primary driver is beryllium at 97%. If beryllium risk is not included the cumulative ELCR is 1.62E-05.
- SWMU 5 cumulative ELCR 4.10E-04; drivers are arsenic at 6%, beryllium at 49%, and Total PAH at 45%. If the contribution from beryllium is discounted, the cumulative ELCR is 2.13E-04.
- SWMU 6 cumulative ELCR 2.40E-04; drivers are beryllium at 90% and Total PAH at 10%. Excluding beryllium, the cumulative ELCR is 2.4E-05.
- SWMU 7 cumulative ELCR 3.90E-03 the primary driver is beryllium at 96%. If beryllium is not included in the cumulative ELCR, the risk is 1.5E-04 with ^{238}U and arsenic contributing to the risk.
- SWMU 30 cumulative ELCR 3.80E-03; the primary driver is beryllium at 96.2%. If beryllium is not incorporated in the total, the cumulative ELCR for soils is 1.42E-04, with arsenic contributing 13% and various PAHs contributing 35%.

Cumulative HIs for the current industrial/maintenance worker were the same as the cumulative HIs for the future industrial worker (i.e., same HI values and drivers including their percent contributions) with the exception of SWMUs 2 and 3. SWMUs 2 and 3 calculated risk and systemic toxicity by using exposure frequencies of 25 days per year for the current industrial worker and 250 days per year for future industrial workers.

The current industrial/maintenance worker had ELCRs exceeding 1E-04 for all SWMUs. Specifically, the cumulative ELCRs at SWMUs 7 and 30 were 3.80E-03 and 3.70E-03, respectively. Approximately 96% of the risk at SWMUs 7 and 30 under this scenario was due to the risk from beryllium. As previously noted, the oral slope factor for beryllium has been withdrawn. If the risks from exposure to beryllium are discounted, the cumulative ELCR is reduced, as described in the summary for future industrial workers above.

F.7.5.2 Observations—Future Excavation Worker

Cumulative HIs for the future excavation worker were greater than 1 for SWMUs 4, 5, 6, 7, and 30 based on soil exposure. This exposure scenario was not evaluated for SWMU 2, SWMU 3, or SWMU 145. The following summarizes the cumulative HIs and major contributors to elevated hazards at these SWMUs.

- SWMU 4 cumulative HI 2.61; drivers are aluminum at 8%, chromium at 24%, iron at 24%, manganese 14%, and vanadium 20%.
- SWMU 5 cumulative HI 2.16; drivers are aluminum at 9%, arsenic at 7%, chromium at 18%, iron at 38%, and manganese at 22%.

- SWMU 6 cumulative HI 2.44; drivers are aluminum at 8%, chromium at 15%, iron at 32%, manganese at 15%, and vanadium at 26%.
- SWMU 7 cumulative HI 5.40; drivers are antimony at 11.3%, chromium at 17.6%, iron at 21.3%, manganese at 11%, uranium at 7.5%, and vanadium at 10.9%.
- SWMU 30 cumulative HI 4.50; drivers are antimony at 6.3%, chromium at 10.2%, copper at 7.6%, iron at 19.8%, manganese at 14.3%, uranium at 12.2%, and vanadium at 12.7%.

Cumulative ELCRs exceeded 1E-04 for all SWMUs (except SWMUs 2, 3, and 145) and were greater than 1E-03 at SWMU 4, SWMU 7, and SWMU 30 for exposure to soil. The following summarizes the cumulative risk estimates and major contributors (> 5%) to the ELCR for these SWMUs.

- SWMU 4 cumulative ELCR 2.70E-03; drivers are beryllium at 7% and total uranium (the total uranium risk was calculated using ²³⁸U slope factors) at 83%. If the contribution from beryllium were discounted, the cumulative is ELCR 2.51E-03.
- SWMU 5 cumulative ELCR 2.9E-04 ; drivers are arsenic at 8%, beryllium at 62%, and Total PAH at 28%. If the risk from beryllium is not included, the cumulative ELCR would be 1.1E-04.
- SWMU 6 cumulative ELCR 2.30E-04; drivers are beryllium at 90% and Total PAH at 9%. If the risk from beryllium is removed, the cumulative ELCR would be 2.3E-05.
- SWMU 7 cumulative ELCR 1.60E-03; drivers are beryllium at 42.2%, ²³⁵U at 9.1%, and ²³⁸U at 41.3%. If the risk from beryllium were discounted, the cumulative ELCR would be 9.2E-04.
- SWMU 30 cumulative ELCR 1.20E-03; driver is beryllium at 93.7%. If the risk from beryllium were discounted, the cumulative ELCR would be 7.6E-05. PAHs, arsenic and radionuclides contribute the remaining risk.

F.7.5.3 Observations—Future Recreational Users

Cumulative HIs for the child, teen, and adult recreational users were less than 1 for SWMUs 4, 5, 6, 7, and 30 based on soil exposure. Cumulative ELCRs exceeded 1E-06 for future adult recreational users only at SWMUs 5, 7, and 30 based on consumption of game. This exposure scenario was not evaluated at SWMUs 2, 3, and 145. The contributors to cumulative ELCR include beryllium and PAH compounds. As stated previously, inclusion of these contaminants, especially beryllium, may artificially increase risk. The following summarizes the cumulative risk estimates and major contributors to the ELCR for these SWMUs.

- SWMU 5 cumulative ELCR 1.0E-05; driver is Total PAH at 96%.
- SWMU 7 cumulative ELCR 1.1E-05; drivers are Aroclor-1260 at 18.6%, benzo(a)pyrene at 9.5%, dibenzo(a,h)anthracene at 42.5%, and ²³⁸U at 15.7%.
- SWMU 30 cumulative ELCR 1.5E-05; drivers are Aroclor-1260 at 48.2%, benzo(a)pyrene at 12.9%, and dibenzo(a,h)anthracene at 20.8%.

F.7.5.4 Observations—Future On-Site Rural Residents

Because of the nature of residential use, risk and hazard contributions were noted for both soil and groundwater exposure. The following summarizes the cumulative HIs and ELCRs observed for each resident.

Hazards—Future Child Residential Exposure to Soil. Cumulative HIs based on direct contact with soil for the child rural resident were greater than 1 for SWMUs 4, 5, 6, 7, and 30. This exposure scenario was not evaluated for SWMUs 2, 3, and 145. The major contributors to elevated hazards are as follows:

- SWMU 4 cumulative HI 98.2: chromium at 24%, iron 60%, and vanadium at 9%
- SWMU 5 cumulative HI 46.2: aluminum at 24%, arsenic at 53%, and chromium at 17%
- SWMU 6 cumulative HI 9.38: beryllium at 8%, chromium at 72%, and nickel at 15%
- SWMU 7 cumulative HI 370: arsenic at 6.2%, iron at 19.7%, and uranium at 58.4%
- SWMU 30 cumulative HI 260: arsenic at 7.5%, iron at 22.6%, and uranium at 46.8%

Hazard—Future Adult Resident Exposure to Soil. Cumulative HIs for the future on-site adult resident were greater than 1 for SWMUs 4, 5, 6, 7, and 30. SWMU 145 was evaluated for excavation worker scenario only. This exposure scenario was not evaluated for SWMUs 2, 3, and 145. The major contributors to elevated hazards are as follows:

- SWMU 4 cumulative HI 28.4: chromium at 22%, iron at 63%, and vanadium at 8%
- SWMU 5 cumulative HI 13.9: aluminum at 24%, arsenic at 55%, and chromium at 15%
- SWMU 6 cumulative HI 2.57: beryllium at 7%, zinc at 6%, nickel at 17% and chromium at 70%
- SWMU 7 cumulative HI 110: arsenic at 6.5%, iron at 19.8%, and uranium at 59.5%
- SWMU 30 cumulative HI 79: arsenic at 7.9%, iron at 22.8%, and uranium at 47.5%

Risks—Future Adult Residential Exposure to Soil. Cumulative ELCRs exceeding 1E-03 from direct contact with soil was observed for SWMUs 4, 5, 6, 7, and 30. This exposure scenario was not evaluated for SWMUs 2, 3, and 145. The contributors to cumulative ELCR include beryllium and PAH compounds. As stated previously, inclusion of these contaminants, especially beryllium, may artificially increase risk. Cumulative ELCRs greater than 1E-02 were identified for SWMUs 3, 5, 7, and 30. The major contributors to elevated risks are as follows:

- SWMU 4 cumulative ELCR 4.3E-03: beryllium at 72%, total PCBs at 5%, ²³⁴U at 6%, and ²³⁸U at 17%. The cumulative ELCR, without the contribution from beryllium, is 1.24E-03.
- SWMU 5 cumulative ELCR 1.0E-02: arsenic at 21%, beryllium at 9%, and Total PAH at 68%. The cumulative ELCR, without the contribution from beryllium, is 9.1E-03.
- SWMU 6 cumulative ELCR 2.4E-03: beryllium at 54% and Total PAH at 46%. The cumulative ELCR, without the contribution from beryllium, is 1.1E-03.
- SWMU 7 cumulative ELCR 3.4E-02: arsenic at 7.3%, beryllium at 65.4%, and ²³⁸U at 17.6%. The cumulative ELCR, without the contribution from beryllium, is 1.19E-02.
- SWMU 30 cumulative ELCR 3.2E-02: arsenic at 6.8%, various PAH compounds totaling 7%, beryllium at 66.7%, and ²³⁸U at 11.5%. The cumulative ELCR, without the contribution from beryllium, is 1.06E-02.

Hazards—Future Resident Exposure to Groundwater. Cumulative HIs based on exposure to groundwater for the future on-site rural resident were greater than 1 for all of the SWMUs, except SWMU 6, which was evaluated for manganese, but did not have an HI greater than 1. The following lists those constituents that contributed to the elevated HIs. The major contaminants driving the hazard were ingestion of arsenic and uranium metal, and ingestion and inhalation of TCE and *cis*-1,2 DCE.

The following lists those constituents that contributed to elevated HIs by SWMU for the Child Resident:

- SWMU 2: TCE at 52% and *cis*-1,2-DCE at 47%
- SWMU 3: arsenic at 52%, uranium at 39%, and manganese at 10%
- SWMU 4: TCE at 93% and *cis*-1,2-DCE at 6%
- SWMU 5: uranium at 90%, arsenic at 3.6%, naphthalene at 3.4%, and manganese at 2.6 %
- SWMU 7: arsenic at 30%, TCE at 26%, Aroclor-1254 at 22%, and *cis*-1,2-DCE at 6.6%
- SWMU 30: arsenic at 64%, manganese at 8.8%, uranium at 15%, 1,1-DCE at 5%, and TCE at 5%
- SWMU 145: antimony at 48% and arsenic at 48%

Risks—Future Residential Exposure to Groundwater. Cumulative ELCRs exceeding both 1E-06 and 1E-04 from direct exposure to groundwater was observed for all of the SWMUs except SWMU 6, which did not have carcinogenic COCs. The major contaminants driving risk were ingestion of arsenic and TCE.

The following lists those constituents that contributed to elevated risks by SWMU:

- SWMU 2: TCE at 98%
- SWMU 3: arsenic at 72% and ⁹⁹Tc at 25%
- SWMU 4: TCE at 68% and vinyl chloride at 31%
- SWMU 5: arsenic at 97%
- SWMU 7: 1,1-DCE at 66%, arsenic at 15%, TCE at 4.1% and vinyl chloride at 12%
- SWMU 30: arsenic at 89% and TCE at 5.2%
- SWMU 145: arsenic at 5.1% and Aroclor-1260 at 93%

F.7.5.5 Observations—Future Off-Site Rural Residents

Risk and hazard estimates for future off-site residential use are based on peak modeled groundwater concentrations. The following summarizes the results of the quantitative assessment at the plant boundary, property boundary, and at the Ohio River (or seeps).

Future Residential Exposure to Groundwater—Plant Boundary. SWMU 6 was not evaluated for groundwater exposure, and SWMU 145 lies outside the plant boundary. Cumulative HIs based on exposure to groundwater at the DOE plant boundary were greater than one for SWMU 2, SWMU 4, SWMU 5, SWMU 7, and SWMU 30. The major contaminants contributing to hazard were TCE, *cis*-1,2-DCE, arsenic, manganese, and PCBs. The cumulative ELCR was greater than 1E-06 for SWMU 2, SWMU 3, SWMU 4, SWMU 5, SWMU 7, and SWMU 30. The cumulative ELCR was greater than 1E-04 for SWMU 2, SWMU 3, SWMU 4, SWMU 7, and SWMU 30. The major contaminants contributing to risk were TCE, 1,1-DCE, vinyl chloride, ⁹⁹Tc, and arsenic.

Below are descriptions of the risk at the plant boundary from exposure to groundwater over time, and the contaminants that contribute to the risk during various time periods. The information below provides a summary of the Tables in Section F.5.5 which describe risk over time. Section 5 also provides additional detail of groundwater modeling results.

SWMU 2—At the plant boundary, the risks from SWMU 2 peak during 15 to 40 years due to the concentration of TCE. By 200 years, the concentration of TCE has decreased and the risk has decreased to approximately 1E-06. The risk continues this downward trend until the concentration of arsenic begins to increase at approximately 700 years. The risk increases moderately to the end of the simulation (1000 years) where it is approximately 7.7E-05.

SWMU 3—At the plant boundary, the risks from SWMU 3 peak at approximately 65 years, primarily due to ⁹⁹Tc. At year 125, the risk from SWMU 3 at the plant boundary is at 1E-06 and continues to decline to the end of the simulation.

SWMU 4—At the plant boundary, the risks from SWMU 4 peak at approximately 15 years, primarily due to TCE and vinyl chloride concentrations. At approximately 50 years, ⁹⁹Tc begins to contribute to the total risk. The risk decreases to 1E-06 at approximately 240 years and decreases until approximately 400 years into the simulation. At approximately 400 years, the risk from arsenic concentrations begins to increase until 1000 years when the risk is approximately 7E-05.

SWMU 5—At the plant boundary, the risks from SWMU 5 peak briefly in the first 50 years just above 1E-06 due to TCE and to some degree ⁹⁹Tc. The risk declines until about 300 years when arsenic concentration begins to increase risk back to 1E-06 and then increases to approximately 5E-5 at 1,000 years.

SWMU 7—At the plant boundary, the risks from SWMU 7 peak in the first 5 years because of TCE, 1,1 DCE, and vinyl chloride concentrations. The risk decreases in 170 years from these contaminants to a risk near 1E-06. The risk begins to increase primarily due to increased concentrations of arsenic after 170, with the risk remaining a approximately 3 E-04 until the end of the simulation (1,000 years).

SWMU 30—At the plant boundary, the risks from SWMU 30 peak in the initial 10 to 20 years due to concentrations of TCE, 1,1 DCE, and ⁹⁹Tc. At approximately 100 years, the concentrations of these contaminants decrease sufficiently that arsenic concentrations begin to be the primary contributors to risk. The risk from the organic compounds is at approximately 4E-04 at 100 years. The risk due primarily to arsenic after 100 years remains near 1E-04 until the end of the simulation (1,000 years).

Future Residential Exposure to Groundwater—Property Boundary. Cumulative HIs based on exposure to groundwater at the DOE property boundary were greater than 1 for SWMU 2, SWMU 4, SWMU 7, and SWMU 30. The major contaminants driving hazard were ingestion of arsenic, TCE, *cis*-1,2-DCE, and PCBs.

Cumulative ELCR exceeded 1E-06 for groundwater exposure for all of the SWMUs, except SWMU 6. Cumulative ELCRs greater than 1E-04 from groundwater use were identified for SWMUs 2, 4, 7, 30, and 145. The major contaminants driving risk were ingestion of arsenic, TCE, 1,1-DCE, vinyl chloride, and ⁹⁹Tc.

Below are descriptions of the risk at the property boundary from exposure to groundwater over time, and the contaminants that contribute to the risk during various time periods. The information below provides a summary of the Tables in Section F.5.5, which describe risk over time. Section 5 also provides additional detail of groundwater modeling results. The descriptions below address the SWMUs that produced a cumulative ELCR greater than 1E-04.

SWMU 2—At the property boundary, the risks from SWMU 2 peak at approximately 40 years due to the concentrations of TCE. The peak risk is greater than 3E-03, but, after 40 years, the concentrations decrease and at approximately 190 years the risk is near 1E-06. The risk decreases until approximately

600 years when arsenic concentrations contribute to risk and begin to increase the risk for the next 400 years to an approximate average of 5E-05.

SWMU 4—At the property boundary, the risks from SWMU 4 peak at approximately 15 years due to TCE and vinyl chloride. The risk from these compounds diminishes to 1E-06 in approximately 220 years. The risk remains below 1E-06 until the end of the simulation (1,000 years).

SWMU 7—At the property boundary, the risks from SWMU 7 peak at approximately 15 years due to TCE, 1,1-DCE, and vinyl chloride. The risks decrease from these contaminants and reach 1E-06 at approximately year 120. After year 120, the influence of arsenic begins to increase risk until the end of the simulation. The maximum risk primarily from arsenic is 7E-05 at 1,000 years.

SWMU 30—At the property boundary, the risks from SWMU 30 peak at about 15 years primarily due to TCE and 1,1-DCE. Technetium-99 contributes to the risk, but the concentration of ⁹⁹Tc does not peak until approximately 50 years. The risk from TCE, ⁹⁹Tc, and 1,1-DCE decreases to 1E-06 at approximately 120 years. At approximately 500 years, risk primarily due to arsenic begins to increase from 1 E-06 to 6E-05 at 1,000 years.

Future Residential Exposure to Groundwater—Ohio River or Seeps. Cumulative HIs based on exposure to groundwater for the future off-site rural resident at the Ohio River were greater than 1 for SWMUs 2, 4, and 30. The major contaminants driving hazard were ingestion of TCE and *cis*-1,2-DCE.

Cumulative ELCRs of 1E-06 from groundwater exposure were observed for SWMUs 2, 3, 4, 7, 30, and 145. Cumulative ELCRs greater than 1E-04 from groundwater use were identified for SWMUs 2, 4, 7, and 30. The contaminants driving risk were ingestion of TCE, 1,1-DCE, vinyl chloride, and ⁹⁹Tc.

Below are descriptions of the risk at the Ohio River or the Little Bayou seeps from exposure to groundwater over time, and the contaminants which contribute to the risk during various time periods. The information below provides a summary of the Tables in Section F.5.5, which describes risk over time. Section 5 also provides additional detail of groundwater modeling results. The descriptions below address the SWMUs that produced a cumulative ELCR greater than 1E-04.

SWMU 2—At the Ohio River, the risks from SWMU 2 peaked at approximately 90 years due primarily to TCE. After 90 years, the risk continued to decrease during the 1,000 year simulation and reached a risk of 1E-6 at approximately 200 years.

SWMU 4—At the Ohio River, the risks from SWMU 4 peaked at approximately 90 years due primarily to TCE and vinyl chloride. After 90 years, the risk continued to decrease during the 1,000 year simulation and reached a risk of 1E-6 at approximately 240 years.

SWMU 7—At the Little Bayou Seeps, the risks from SWMU 7 peaked at approximately 35 years due primarily to 1,1-DCE, TCE, and vinyl chloride. After the peak, the risk decreased to 1E-06 at approximately 120 years.

F.7.5.6 Summary of Observations

The following summarize the observations noted for this and previous BHHRA for BGOU. The discussion focuses on the individual exposure scenarios examined for the assessment.

F.7.5.6.1 Future Industrial Worker

SWMUs 4, 7, and 30 hazard levels exceed 1 for industrial worker exposure to soil, with chromium, iron, and vanadium serving as the primary hazard drivers for elevated HIs. All SWMUs (SWMU 145 was not evaluated for this scenario) exceed risk levels of 1E-04 for industrial worker exposure to soil, with ²³⁵U+daughters, ²³⁸U+daughters, and beryllium serving as the primary risk drivers. Other COCs contributing to elevated risks include Total PAH and arsenic. SWMUs 2, 3, 5, 7 and 30 exceed risk levels of 1E-04 for industrial worker exposure to soil, if beryllium is not include in the cumulative ELCR for each of the SWMUs.

F.7.5.6.2 Future Excavation Worker

SWMUs 4, 5, 6, 7, and 30 exceed a hazard level of 1 for excavation worker exposure to soil (SWMUs 4, 5, and 6 were evaluated for exposure to soil and waste), with aluminum, antimony, chromium, iron, manganese, uranium, and vanadium serving as the primary hazard driver for elevated HIs. Other COCs contributing to hazards include arsenic and copper. SWMUs 4, 5, 6, 7, and 30 exceed the risk level of 1E-04 for excavation worker exposure to soil, with beryllium, uranium, Total PAH, and ²³⁸U serving as the primary risk drivers. Other COCs contributing to elevated risks include arsenic and ²³⁵U. SWMUs 4, 5, and 7 exceed the risk level of 1E-04 for excavation worker exposure to soil when beryllium is not included in the cumulative ELCR. SWMUs 4, 5, and 6 included exposure to soil and waste, which was included in this BHHRA but referred to as one media type, soil.

F.7.5.6.3 Future On-Site Residents (Groundwater)

For residential groundwater use at the SWMU boundary, ELCR was greater than 1E-04 and HI was greater than 1 for all SWMUs except SWMU 6. The primary risk drivers are TCE, arsenic, vinyl chloride, 1,1-DCE, and ⁹⁹Tc.

F.7.5.6.4 Future Off-Site Residents (Groundwater)

SWMUs 2, 4, 5, 7, and 30 exceed a hazard level of 1 for off-site residential exposure to groundwater at the PGDP plant boundary. SWMUs 2, 4, 7, and 30 exceed a hazard level of 1 at the property boundary. SWMUs 2, 4, and 30 exceed a hazard level of 1 at the Ohio River (or seeps). The primary drivers for hazard are arsenic, TCE, *cis*-1,2-DCE, and 1,1-DCE. SWMUs 2, 3, 4, 7, and 30 at the plant boundary, SWMUs 2, 4, 7, 30, and 145 at the property boundary, and SWMUs 2, 4, 7, and 30 at the Ohio River (or seeps) exceed a risk level of 1E-04 for off-site residential exposure to groundwater. The primary risk drivers are TCE, 1,1-DCE, and ⁹⁹Tc.

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F.8. REMEDIAL GOAL OPTIONS

This section presents RGOs for the COCs identified in Section 5 and the methods used to calculate the RGOs. These RGOs should not be interpreted as being cleanup goals, but as risk-based values that may be used to guide the development of cleanup goals by risk managers. Cleanup goals will be determined in later decision documents.

RGOs were calculated for each groundwater COC from the modeled groundwater concentrations considering use of groundwater at each source. When calculating the HI-based RGOs, the more conservative child-based values are reported. In addition, for comparison to the RGOs, the maximum contaminant level (MCL) for each COC is presented. Note, MCLs are not clean-up criteria. The National Contingency Plan notes that clean-up criteria different from MCLs may be required if multiple contaminants are present or if contaminants may reach a receptor through exposure routes different from those considered in the development of MCLs. Risks for use of contaminated groundwater must be presented in addition to a simple screen against MCLs so that risk managers can make appropriate decisions.

F.8.1 CALCULATION OF GROUNDWATER RGOs

EPA guidance (EPA 1991) directs that RGOs are to be calculated for all COCs identified in a BHHRA. The COCs identified in this risk assessment and their RGOs are presented in Table F.81. These RGOs were calculated using the following equation.

$$\frac{\text{Concentration}}{\text{Risk}} = \frac{\text{RGO}}{\text{Target Risk}}$$

Where:

- Concentration is the exposure concentration for the medium.
- Risk is the risk posed by exposure to the contaminated medium.
- RGO is the remedial goal option.
- Target Risk is one of the values listed in Table F.80.

F.8.2 PRESENTATION OF RGOs

RGOs for soil COCs determined in the previous risk assessments are the 2008 draft NALs (DOE 2008) for the listed analyte and receptor. The previous risk assessments for soil calculated RGOs for soil COCs, but both toxicity factors and exposure factors for some of the COCs have been updated since that time. Therefore, the current NALs for the same soil exposure scenarios (residential, industrial, and excavation) are presented as the RGOs in Table F.81 for soil.

The equation developed in the previous subsection was applied for each groundwater COC. The RGOs developed for all COCs using this equation are presented in Table F.82. In addition, these tables present the groundwater EPCs used in the BHHRA.

Table F.81. RGOs for Soil COCs of the BGOU SWMUs

COC ^A	Cancer NAL	Noncancer NAL	RGO ^B at		RGO at HI=3	RGO at ELCR= 1 x 10 ⁻⁶	RGO at ELCR= 1 x 10 ⁻⁵	RGO at ELCR= 1 x 10 ⁻⁴	Units
			HI=0.1	HI=1					
Residential User Soil Exposure									
Aluminum		9.69E+02	9.69E+01	9.69E+02	2.91E+03				mg/kg
Antimony		8.69E-02	8.69E-03	8.69E-02	2.61E-01				mg/kg
Arsenic	1.44E-01	1.16E+00	1.16E-01	1.16E+00	3.48E+00	1.44E-01	1.44E+00	1.44E+01	mg/kg
Barium		1.40E+02	1.40E+01	1.40E+02	4.20E+02				mg/kg
Beryllium and compounds	1.19E-03	2.20E-01	2.20E-02	2.20E-01	6.60E-01	1.19E-03	1.19E-02	1.19E-01	mg/kg
Cadmium	2.00E+00	3.26E+00	3.26E-01	3.26E+00	9.78E+00	2.00E+00	2.00E+01	2.00E+02	mg/kg
Chromium	1.10E+02	8.32E+01	8.32E+00	8.32E+01	2.50E+02	1.10E+02	1.10E+03	1.10E+04	mg/kg
Cobalt	4.69E+02	6.95E+01	6.95E+00	6.95E+01	2.09E+02	4.69E+02	4.69E+03	4.69E+04	mg/kg
Copper		9.39E+01	9.39E+00	9.39E+01	2.82E+02				mg/kg
Iron		4.14E+02	4.14E+01	4.14E+02	1.24E+03				mg/kg
Manganese		5.60E+01	5.60E+00	5.60E+01	1.68E+02				mg/kg
Nickel	5.06E+03	4.35E+01	4.35E+00	4.35E+01	1.31E+02	5.06E+03	5.06E+04	5.06E+05	mg/kg
Uranium		2.57E+00	2.57E-01	2.57E+00	7.71E+00				mg/kg
Vanadium		7.71E-01	7.71E-02	7.71E-01	2.31E+00				mg/kg
Zinc		5.21E+02	5.21E+01	5.21E+02	1.56E+03				mg/kg
Aroclor 1260	6.08E-02					6.08E-02	6.08E-01	6.08E+00	mg/kg
Benzo[a]anthracene	7.48E-02					7.48E-02	7.48E-01	7.48E+00	mg/kg
Benzo[a]pyrene	7.48E-03					7.48E-03	7.48E-02	7.48E-01	mg/kg
Benzo[b]fluoranthene	7.48E-02					7.48E-02	7.48E-01	7.48E+00	mg/kg
Dibenz[a,h]anthracene	7.48E-03					7.48E-03	7.48E-02	7.48E-01	mg/kg
Total Dioxins/Furans	6.78E-07					6.78E-07	6.78E-06	6.78E-05	mg/kg
Indeno[1,2,3-cd]pyrene	7.48E-02					7.48E-02	7.48E-01	7.48E+00	mg/kg
Total PCBs	5.78E-02					5.78E-02	5.78E-01	5.78E+00	mg/kg
Total PAHs	7.48E-03					7.48E-03	7.48E-02	7.48E-01	mg/kg
Neptunium-237+D	8.39E-02					8.39E-02	8.39E-01	8.39E+00	pCi/g
Plutonium-239*	3.15E+00					3.15E+00	3.15E+01	3.15E+02	pCi/g
Radium-226+D	7.94E-03					7.94E-03	7.94E-02	7.94E-01	pCi/g
Uranium-234	5.47E+00					5.47E+00	5.47E+01	5.47E+02	pCi/g
Uranium-235+D	1.22E-01					1.22E-01	1.22E+00	1.22E+01	pCi/g
Uranium-238+D	5.17E-01					5.17E-01	5.17E+00	5.17E+01	pCi/g

Table F.81. RGOs for Soil COCs of the BGOU SWMUs (Continued)

COC	Cancer NAL	Noncancer NAL	RGO at			RGO at HI=3	RGO at ELCR= 1 x 10 ⁻⁶	RGO at ELCR= 1 x 10 ⁻⁵	RGO at ELCR= 1 x 10 ⁻⁴	Units
			HI=0.1	HI=1	HI=3					
Industrial Worker Soil Exposure										
Aluminum		4.22E+03	4.22E+02	4.22E+03	1.27E+04					mg/kg
Antimony		3.46E-01	3.46E-02	3.46E-01	1.04E+00					mg/kg
Arsenic	4.84E-01	7.78E+00	7.78E-01	7.78E+00	2.33E+01	4.84E-01	4.84E+00	4.84E+01	4.84E+01	mg/kg
Barium		5.92E+02	5.92E+01	5.92E+02	1.78E+03					mg/kg
Beryllium and compounds	2.83E-03	8.68E-01	8.68E-02	8.68E-01	2.60E+00	2.83E-03	2.83E-02	2.83E-01	2.83E-01	mg/kg
Cadmium	1.49E+01	1.97E+01	1.97E+00	1.97E+01	5.91E+01	1.49E+01	1.49E+02	1.49E+03	1.49E+03	mg/kg
Chromium	2.11E+02	3.26E+02	3.26E+01	3.26E+02	9.78E+02	2.11E+02	2.11E+03	2.11E+04	2.11E+04	mg/kg
Cobalt	9.05E+02	4.48E+02	4.48E+01	4.48E+02	1.34E+03	9.05E+02	9.05E+03	9.05E+04	9.05E+04	mg/kg
Copper		4.91E+02	4.91E+01	4.91E+02	1.47E+03					mg/kg
Iron		1.90E+03	1.90E+02	1.90E+03	5.70E+03					mg/kg
Manganese		2.29E+02	2.29E+01	2.29E+02	6.87E+02					mg/kg
Nickel	9.75E+03	2.22E+02	2.22E+01	2.22E+02	6.66E+02	9.75E+03	9.75E+04	9.75E+05	9.75E+05	mg/kg
Uranium		1.88E+01	1.88E+00	1.88E+01	5.64E+01					mg/kg
Vanadium		3.04E+00	3.04E-01	3.04E+00	9.12E+00					mg/kg
Zinc		2.50E+03	2.50E+02	2.50E+03	7.50E+03					mg/kg
Aroclor 1260	1.75E-01					1.75E-01	1.75E+00	1.75E+01	1.75E+01	mg/kg
Benz[a]anthracene	1.94E-01					1.94E-01	1.94E+00	1.94E+01	1.94E+01	mg/kg
Benzo[a]pyrene	1.94E-02					1.94E-02	1.94E-01	1.94E+00	1.94E+00	mg/kg
Benzo[b]fluoranthene	1.94E-01					1.94E-01	1.94E+00	1.94E+01	1.94E+01	mg/kg
Dibenz[a,h]anthracene	1.94E-02					1.94E-02	1.94E-01	1.94E+00	1.94E+00	mg/kg
Total Dioxins/Furans	1.89E-06					1.89E-06	1.89E-05	1.89E-04	1.89E-04	mg/kg
Indeno[1,2,3-cd]pyrene	1.94E-01					1.94E-01	1.94E+00	1.94E+01	1.94E+01	mg/kg
Total PCBs	1.63E-01					1.63E-01	1.63E+00	1.63E+01	1.63E+01	mg/kg
Total PAHs	1.94E-02					1.94E-02	1.94E-01	1.94E+00	1.94E+00	mg/kg
Neptunium-237+D	2.71E-01					2.71E-01	2.71E+00	2.71E+01	2.71E+01	pCi/g
Plutonium-239*	1.07E+01					1.07E+01	1.07E+02	1.07E+03	1.07E+03	pCi/g
Radium-226+D	2.56E-02					2.56E-02	2.56E-01	2.56E+00	2.56E+00	pCi/g
Uranium-234	1.89E+01					1.89E+01	1.89E+02	1.89E+03	1.89E+03	pCi/g
Uranium-235+D	3.95E-01					3.95E-01	3.95E+00	3.95E+01	3.95E+01	pCi/g
Uranium-238+D	1.70E+00					1.70E+00	1.70E+01	1.70E+02	1.70E+02	pCi/g

Table F.81. RGOs for Soil COCs of the BGO SWMUs (Continued)

COC	Cancer NAL	Noncancer NAL	RGO at			RGO at HI=3	RGO at ELCR=1 x 10 ⁻⁶	RGO at ELCR=1 x 10 ⁻⁵	RGO at ELCR=1 x 10 ⁻⁴	Units
			HI=0.1	HI=1	HI=3					
Excavation Worker Soil Exposure										
Aluminum		4.84E+03	4.84E+02	4.84E+03	1.45E+04					mg/kg
Antimony		4.52E-01	4.52E-02	4.52E-01	1.36E+00					mg/kg
Arsenic	3.13E-01	5.03E+00	5.03E-01	5.03E+00	1.51E+01	3.13E-01	3.13E+00	3.13E+01	3.13E+01	mg/kg
Barium		7.11E+02	7.11E+01	7.11E+02	2.13E+03					mg/kg
Beryllium and compounds	3.83E-03	1.15E+00	1.15E-01	1.15E+00	3.45E+00	3.83E-03	3.83E-02	3.83E-01	3.83E-01	mg/kg
Cadmium	2.12E+00	1.45E+01	1.45E+00	1.45E+01	4.35E+01	2.12E+00	2.12E+01	2.12E+02	2.12E+02	mg/kg
Chromium	2.85E+02	4.36E+02	4.36E+01	4.36E+02	1.31E+03	2.85E+02	2.85E+03	2.85E+04	2.85E+04	mg/kg
Cobalt	1.22E+03	3.11E+02	3.11E+01	3.11E+02	9.33E+02	1.22E+03	1.22E+04	1.22E+05	1.22E+05	mg/kg
Copper		4.37E+02	4.37E+01	4.37E+02	1.31E+03					mg/kg
Iron		2.02E+03	2.02E+02	2.02E+03	6.06E+03					mg/kg
Manganese		2.90E+02	2.90E+01	2.90E+02	8.70E+02					mg/kg
Nickel	1.32E+04	2.05E+02	2.05E+01	2.05E+02	6.15E+02	1.32E+04	1.32E+05	1.32E+06	1.32E+06	mg/kg
Uranium		1.10E+01	1.10E+00	1.10E+01	3.30E+01					mg/kg
Vanadium		4.03E+00	4.03E-01	4.03E+00	1.21E+01					mg/kg
Zinc		2.50E+03	2.50E+02	2.50E+03	7.50E+03					mg/kg
Atroclor 1260	1.55E-01					1.55E-01	1.55E+00	1.55E+01	1.55E+01	mg/kg
Benz[a]anthracene	2.16E-01					2.16E-01	2.16E+00	2.16E+01	2.16E+01	mg/kg
Benzo[a]pyrene	2.16E-02					2.16E-02	2.16E-01	2.16E+00	2.16E+00	mg/kg
Benzo[b]fluoranthene	2.16E-01					2.16E-01	2.16E+00	2.16E+01	2.16E+01	mg/kg
Dibenz[a,h]anthracene	2.16E-02					2.16E-02	2.16E-01	2.16E+00	2.16E+00	mg/kg
Total Dioxins/Furans	1.79E-06					1.79E-06	1.79E-05	1.79E-04	1.79E-04	mg/kg
Indeno[1,2,3-cd]pyrene	2.16E-01					2.16E-01	2.16E+00	2.16E+01	2.16E+01	mg/kg
Total PCBs	1.48E-01					1.48E-01	1.48E+00	1.48E+01	1.48E+01	mg/kg
Total PAHs	2.16E-02					2.16E-02	2.16E-01	2.16E+00	2.16E+00	mg/kg
Neptunium-237+D	3.27E-01					3.27E-01	3.27E+00	3.27E+01	3.27E+01	pCi/g
Plutonium-239*	1.62E+00					1.62E+00	1.62E+01	1.62E+02	1.62E+02	pCi/g
Radium-226+D	3.30E-02					3.30E-02	3.30E-01	3.30E+00	3.30E+00	pCi/g
Uranium-234	2.83E+00					2.83E+00	2.83E+01	2.83E+02	2.83E+02	pCi/g
Uranium-235+D	4.55E-01					4.55E-01	4.55E+00	4.55E+01	4.55E+01	pCi/g
Uranium-238+D	1.17E+00					1.17E+00	1.17E+01	1.17E+02	1.17E+02	pCi/g

* COC = contaminant of concern

^b RGO = remedial goal option. RGOs for soil for both HI and ELCR are calculated from the 2008 draft NALs (DOE 2008).

Table F.82. RGOs for Groundwater COCs of the BGOU SWMUs

Residential User Groundwater Exposure											
COC ^A	EPC ^B	SWMU ^C	ELCR at EPC	HI at EPC	RGO ^D at HI=0.1	RGO at HI=1	RGO at HI=3	RGO at ELCR=1 x 10 ⁻⁶	RGO at ELCR=1 x 10 ⁻⁵	RGO at ELCR=1 x 10 ⁻⁴	mg/L
MCL											
Antimony	7.99E-02	145		5.97E+00	1.34E-03	1.34E-02	1.34E-01	3.76E-05	3.76E-04	3.76E-03	0.006
Arsenic	6.21E-02	145	1.65E-03	1.99E+01	3.12E-04	3.12E-03	9.36E-03	3.76E-05	3.76E-04	3.76E-03	0.010
Manganese	1.01E+00	5		2.15E+00	4.70E-02	4.70E-01	1.41E+00				---
Selenium	1.51E-02	30		2.90E-01	5.21E-03	5.21E-02	1.56E-01				0.05
Uranium	4.89E-02	3		7.82E+00	604	603	1.88E-02				0.03
Total PCBs	5.23E-05	7	7.09E-06	4.20E+00	1.25E-06	1.25E-05	3.74E-05	7.38E-06	7.38E-05	7.38E-04	0.0005
1,1-DCE	8.98E-02	7	2.08E-03	8.51E-01	1.06E-02	1.06E-01	3.17E-01	4.32E-05	4.32E-04	4.32E-03	---
<i>cis</i> -1,2-DCE	1.15E+01	2		6.07E+02	1.89E-03	1.89E-02	5.68E-02				0.07
Naphthalene	5.55E-03	5		2.80E+00	1.98E-04	1.98E-03	5.95E-03				---
TCE	1.18E+00	4	3.67E-02	5.39E+02	2.19E-04	2.19E-03	6.57E-03	3.22E-05	3.22E-04	3.22E-03	0.005
Vinyl Chloride	2.61E-02	4	1.65E-02	1.21E+00	2.16E-03	2.16E-02	6.47E-02	1.58E-06	1.58E-05	1.58E-04	0.002
Technetium-99	1.01E+04	145	5.54E-04					1.82E+0	1.82E+02	1.82E+03	900 ^E
Uranium-234	7.94E+00	7	1.11E-05					1	7.12E+00	7.12E+01	20 ^F
Uranium-238	1.59E+01	3	2.76E-05					5.76E-01	5.76E+00	5.76E+01	20 ^F

^A COC = contaminant of concern

^B EPC = exposure point concentration; represents maximum EPC value for all SWMUs where constituent was a COC for the applicable scenario

^C SWMU = the SWMU associated with the maximum EPC value

^D RGO = remedial goal option

^E converted from MCL of 4 mrem/yr dose (DOE 2001)

^F converted from MCL for total uranium of 0.03 mg/L (DOE 2001)

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ATTACHMENT F1

SUMMARY TABLES FOR NEW SURFACE SOIL DATA

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Table F1.1 Summary of New Maximum Detected Concentrations in SWMU 3 Surface Soil

Analyte	New Max Value for SWMU 3	Residential	Exceeds Residential	Industrial	Exceeds Industrial	Background	Background	COPC	COPC
		Child No Action Limit*	Child NAL	Worker No Action Limit*	Worker NAL	Concentration	Exceeded?	for Child Resident?	for Industrial Worker?
Metals (mg/kg)									
Aluminum	8,570	732	Yes	4,640	Yes	13,000	No	No	No
Antimony	15.7	0.0635	Yes	0.379	Yes	0.21	Yes	Yes	Yes
Arsenic	7.62	0.132	Yes	0.523	Yes	12	No	Yes	Yes
Barium	87.7	37	Yes	229	No	200	No	No	No
Calcium	3,5200	NA	NA	NA	NA	200,000	No	No	No
Chromium	12.5	60.5	No	356	No	16	Yes	No	No
Cobalt	6.41	209	No	1920	No	14	No	No	No
Copper	10.6	68.1	No	493	No	19	No	No	No
Iron	15,700	314	Yes	2070	Yes	28,000	No	Yes	Yes
Lead	12.7	50	No	50	No	36	No	No	No
Magnesium	2,500	NA	NA	NA	NA	7,700	No	No	No
Manganese	558	7.46	Yes	45.2	Yes	1,500	No	No	No
Mercury	0.024	100,000 or 0.158	No	100,000 or 0.982	No	0.2	No	No	No
Molybdenum	6.21	10.9	Yes	83	No	NA	NA	Yes	No
Nickel	10.7	0.34	Yes	242	No	21	Yes	Yes	No
Sodium	112	NA	NA	NA	NA	320	No	No	No
Tin	15	439	No	2790	No	NA	NA	No	No
Uranium	43	2.16	Yes	20.2	Yes	4.9	Yes	Yes	Yes
Vanadium	33.7	0.562	Yes	3.32	Yes	38	No	No	No
Zinc	31.8	401	No	2,730	No	65	Yes	No	No
PCBs (mg/kg)									
None Detected									
Organics (mg/Kg)									
Acetone	0.00811	53.4	No	358	No	NA	NA	No	No
Trichloroethene	0.00633	0.741	Yes	2.51	Yes	NA	NA	Yes	Yes

Table F1.1 Summary of New Maximum Detected Concentrations in SWMU 3 Surface Soil (Continued)

Analyte Radionuclides (pCi/g)	New Max Value for SWMU 3	Residential		Exceeds		Industrial Worker		Background Concent- ration	Background Exceeded?	COPC for Child Resident?	COPC for Industrial Worker?
		Child No Action Limit*	Child NAL	Residential Child NAL	Exceeds Residential Child NAL	Industrial Worker No Action Limit*	Exceeds Industrial Worker NAL				
Cesium-137	0.344	0.0128	0.0128	Yes	Yes	0.0858	Yes	0.49	No	Yes	Yes
Plutonium-239	0.0562	2.22	2.22	Yes	Yes	11.5	No	0.025	Yes	Yes	No
Technetium-99	21.6	67.4	67.4	No	No	362	No	2.5	Yes	No	No
Thorium-228	0.442	0.00418	0.00418	Yes	Yes	0.028	Yes	NA	NA	Yes	Yes
Thorium-230	0.46	2.85	2.85	No	No	14.9	No	1.5	Yes	No	No
Thorium-232	0.519	2.61	2.61	Yes	Yes	13.5	No	NA	NA	Yes	No
Uranium	7.02	NA	NA	NA	NA	NA	NA	NA	NA	No	No
Uranium-234	0.958	3.81	3.81	Yes	Yes	19.8	No	2.5	Yes	Yes	No
Uranium-235/236	0.079	0.0591	0.0591	Yes	Yes	0.395	No	0.14	Yes	Yes	No
Uranium-238	5.99	0.261	0.261	Yes	Yes	1.71	Yes	1.2	Yes	Yes	Yes

* NAL values are from DOE 2001. *Methods for Conducting Human Health Risk Assessment and Risk Evaluations at the Paducah Gaseous Diffusion Plant*, DOE/OR/07-1506&D1, U. S. Department of Energy, Paducah, KY, December.

Note: -- means that there was no value recorded in the applicable table for that analyte

Table F1.2 Summary of New Maximum Detected Concentrations in SWMU 7 Surface Soil

Analyte	New Max Value for SWMU 7	No Action Limit*	Exceeds Residential Child NAL	Background Concentration	Background Exceeded?	COPC?	SWMU 7 COPC
Metals (mg/kg)							
Aluminum	8,910	732	Yes	13,000	No	No	Yes
Arsenic	3.14	0.132	Yes	12	No	No	Yes
Barium	61.2	37	Yes	200	No	No	No
Calcium	2,100	--	NA	200,000	No	No	No
Chromium	17.9	60.5	No	16	Yes	No	Yes
Cobalt	6.75	209	No	14	No	No	Yes
Copper	53.9	68.1	No	19	Yes	No	No
Iron	13,100	314	Yes	28,000	No	No	Yes
Lead	11.9	50	No	36	No	No	Yes
Magnesium	1,040	--	NA	7,700	No	No	No
Manganese	122	7.46	Yes	1,500	No	No	No
		100,000					
		or					
Mercury	0.043	0.158	No	0.2	No	No	No
Nickel	29.8	0.34	Yes	21	Yes	Yes	Yes
Uranium	1,270	2.16	Yes	4.9	Yes	Yes	Yes
Vanadium	31	0.562	Yes	38	No	No	Yes
Zinc	37.8	401	No	65	Yes	No	Yes
PCBs (ug/kg)							
PCB-1248	14,800	57.4	Yes	--	NA	Yes	--
PCB-1254	130	38.8	Yes	--	NA	Yes	Yes
		1,640					
PCB Total	14,800	or 57.4	Yes	--	NA	Yes	--

Table F1.2 Summary of New Maximum Detected Concentrations in SWMU 7 Surface Soil (Continued)

Analyte	New Max Value for SWMU 7	No Action Limit*	Exceeds Residential Child NAL	Background Concentration	Background Exceeded?	COPC?	SWMU 7 COPC
Organics (ug/Kg)							
Acetone	83.7	5,340	No	--	NA	No	--
Di-n-butylphthalate	940	264,000	No	--	NA	No	--
Methylene chloride	5.7	3,920	No	--	NA	No	No
Radionuclides (pCi/g)							
Alpha activity	65.1	--	NA	--	NA	No	Yes
Beta activity	139	--	NA	--	NA	No	Yes
Neptunium-237	0.0775	0.0405	Yes	0.1	No	No	Yes
Plutonium-239	0.141	2.22	No	0.025	Yes	No	Yes
Technetium-99	11.1	67.4	No	2.5	Yes	No	No
Thorium-228	3.36	0.00418	Yes	1.6	Yes	Yes	--
Thorium-230	1.94	2.85	No	1.5	Yes	No	No
Thorium-232	0.459	2.61	No	1.5	No	No	--
Thorium-234	2,650	--	NA	--	NA	No	--
Uranium	2,750	--	NA	--	NA	No	--
Uranium-234	318	3.81	Yes	2.5	Yes	Yes	Yes
Uranium-235/236	42.1	0.0591	Yes	0.14	Yes	Yes	Yes
Uranium-238	2,390	0.261	Yes	1.2	Yes	Yes	Yes

Note: -- means that there was no value recorded in the applicable table for that analyte

* NAL values are from DOE 2001. Methods for Conducting Human Health Risk Assessment and Risk Evaluations at the Paducah Gaseous Diffusion Plant, DOE/OR/07-1506&D1, U. S. Department of Energy, Paducah, KY, December.

ATTACHMENT F2
HISTORICAL RISK ASSESSMENT TABLES

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Table F2.1. Cancer Risk Estimates for Direct Contact to Soil

Future Industrial Exposure

Scenario: unrestricted worker (250 days/year)

Chemical	Slope Factor (mg/kg-day)⁻¹	Soil Conc. mg/kg	Chronic Daily Intake mg/kg-day	Excess Lifetime Cancer Risk
INGESTION				
Pentachlorophenol	0.12	0.1000	1.7E-08	2.1E-09
OCDD (total)	150	0.0033	5.8E-10	8.6E-08
PCB-1248	7.7	0.2100	3.7E-08	2.8E-07
PCB-1260	7.7	0.1300	2.3E-08	1.7E-07
Arsenic	1.75	10.1700	1.8E-06	3.1E-06
Beryllium	4.3	0.7400	1.3E-07	5.6E-07
pathway sum=				4E-06
DERMAL ABSORPTION				
Pentachlorophenol				
OCDD (total)	0.12	0.1000	1.1E-08	1.3E-09
PCB-1248	150	0.0033	3.6E-10	5.4E-08
PCB-1260	7.7	0.2100	2.3E-08	1.8E-07
Arsenic	7.7	0.1300	1.4E-08	1.1E-07
Beryllium	1.75	10.1700	1.1E-07	1.9E-07
	4.3	0.7400	8.1E-09	3.5E-08
pathway sum=				6E-07
	Unit Risk (ug/m³)-1			
INHALATION				
OCDD (total)	0.000000033	0.0033	5.0E-14	5.8E-18
Arsenic	0.0043	10.1700	1.5E-10	2.3E-09
Beryllium	0.0024	0.7400	1.1E-11	9.4E-11
Chromium VI	0.012	19.0000	2.9E-10	1.2E-08
Nickel (soluble salt)	0.00024	25.1000	3.8E-10	3.2E-10
pathway sum=				1E-08
sum of pathways=				5E-06
Ingestion:	Intake (mg/kg-d)=(conc. in soil*IngR*CF*FI*EF*ED)/(BW*AT)			
Dermal Absorption:	Abs dose(mg/kg-d)=(soil conc.*CF*SA*AF*ABS*EF*ED)/(BW*AT)			
Inhalation:	Inh dose (mg/kg-d)=(soil conc.*EF*ED*InhR*(1/PEF))/(BW*AT)			
	SFI=Unit Risk*(BW/InhR)*1000			
exposure parameters				
IngR=Ingestion rate (mg soil/day)			50	
CF=Conversion factor (10E-6)			1E-06	
FI=Fraction ingested			1	
EF=Exposure frequency (days/year)			250	
ED=Exposure duration (year)			25	
BW=Body weight (kg)			70	
AT=Averaging time (days)			25,550	
SA=Skin surface area (cm ²)			3,120	
AF=Soil to skin adherence (mg/cm ²)			1	
ABS=Absorption (0.1 % metals; 1 % organics)			0.001	0.01
PEF=Particulate emission (m ³ /kg)			4.63E+09	
InhR=Inhalation rate (m ³ /day)			20	

Table taken from Attachment 2-1 Remedial Investigation Addendum for Waste Area Grouping 22, Burial Grounds, Solid Waste Management Units 2 and 3, at the Paducah Gaseous Diffusion Plant, Paducah, Kentucky DOE/OR/07-1141&D2 (KY/ER-32 & D2) September 1994 Revision 2.

Table F2.2. Chronic Hazard Index Estimates for Direct Contact to Soil

Future Industrial Exposure

Scenario: unrestricted worker (250 days/year)

Chemical	Reference Dose mg/kg-day	Soil Conc. mg/kg	Chronic Daily Intake mg/kg-day	Chronic Hazard
INGESTION				
Pentachlorophenol	0.03	0.10	4.9E-08	2E-06
Arsenic	0.0003	10.17	5.0506	2E-02
Barium	0.07	132.68	6.5E-05	9E-04
Beryllium	0.005	0.74	3.6E-07	7E-05
Chromium VI	0.005	19.00	9.3E-06	2E-03
Copper	0.037	24.71	1.2E-05	3E-04
Manganese	0.14	2,541.05	1.2E-03	9E-03
Mercury	0.0003	0.15	7.3E-08	2E-04
Nickel (soluble salt)	0.02	25.10	1.2E-05	6E-04
Selenium	0.005	0.40	2.0E-07	4E-05
Silver	0.005	5.38	2.6E-06	5E-04
Vanadium	0.007	31.80	1.6E-05	2E-03
Zinc	0.3	67.05	3.3E-05	1 E-04
Uranium (soluble salt)	0.003	83.58	4.1 E-05	1 E-02
pathway sum=				0.05
DERMAL ABSORPTION				
Pentachlorophenol	0.03	0.10	3.1E-08	1E-06
Arsenic	0.0003	10.17	3.1E-07	1E-03
Barium	0.07	132.68	4.1E-06	6E-05
Beryllium	0.005	0.74	2.3E-08	5E-06
Chromium VI	0.005	19.00	5.8E-07	1E-04
Copper	0.037	24.71	7.5E-07	2E-05
Manganese	0.005	2,541.05	7.8E-05	2E-02
Mercury	0.0003	0.15	4.6E-09	2E-05
Nickel (soluble salt)	0.02	25.10	7.7E-07	4E-05
Selenium	0.005	0.40	1.2E-08	2E-06
Silver	0.005	5.38	1.6E-07	3E-05
Vanadium	0.007	31.80	9.7E-07	1E-04
Zinc	0.3	67.05	2.0E-06	7E-06
Uranium (soluble salt)	0.003	83.58	2.6E-06	9E-04
pathway sum=				0.02
	Reference Conc. mg/m³			
INHALATION				
Barium	0.0005	132.68	5.6E-09	4E-05
Chromium VI	0.000002	19.00	8.0E-10	1 E-03
Manganese	0.0004	2,541.05	1.1 E-07	9E-04
Mercury	0.0003	0.15	6.3E-12	7E-08
pathway sum=				0.002
sum of pathways=				0.07
Ingestion:	Intake (mg/kg-d)=(conc. in soil*IngR*CF*FI*EF*ED)/(BW*AT)			
Dermal Absorption:	Abs dose(mg/kg-d)=(soil conc.*CF*SA*AF*ABS*EF*ED)/(BW*AT)			
Inhalation:	Inh dose (mg/kg-d)=(soil conc.*EF*ED*InhR*(1/PEF))/(BW*AT)			
	SFI=Unit Risk*(BW/InhR)*1000			

Table F2.2. Chronic Hazard Index Estimates for Direct Contact to Soil (Continued)

exposure parameters

IngR=Ingestion rate (mg soil/day)	50	
CF=Conversion factor (10E-6)	1E-06	
FI=Fraction ingested	1	
EF=Exposure frequency (days/year)	250	
ED=Exposure duration (year)	25	
BW=Body weight (kg)	70	
AT=Averaging time (days)	25,550	
SA=Skin surface area (cm ²)	3,120	
AF=Soil to skin adherence (mg/cm ²)	1	
ABS=Absorption (0.1 % metals; 1 % organics)	0.001	0.01
PEF=Particulate emission (m ³ /kg)	4.63E+09	
InhR=Inhalation rate (m ³ /day)	20	

Table taken from Attachment 2-2 Remedial Investigation Addendum for Waste Area Grouping 22, Burial Grounds, Solid Waste Management Units 2 and 3, at the Paducah Gaseous Diffusion Plant, Paducah, Kentucky DOE/OR/07-1141&D2 (KY/ER-32 & D2) September 1994 Revision 2.

Table F2.3. Excess Lifetime Risk of Cancer Incidence for Direct Contact to Soil

Future Industrial Exposure Scenario: unrestricted worker (250 days/year)								
Radionuclide ^a	Soil Concentration (pCi/g) (SC)	Annual Intake (pCi)	Total Intake (pCi)	Dose Conversion Factor ^b (mrem/pCi) or (mrem*g/pCi/h)	Committed Effective Dose	Total Committed Effective Dose	Cancer Incidence Risk	Risk of Cancer Incidence
					Equivalent 1 Yr Intake (mrem/yr) ^c	Equivalent Dose (mrem)	Factor (pCi)-1 ^d or (g/pCi-1)	
INGESTION								
Neptunium-237+D	0.32	4.00	100.0	4.4E-03	1.8E-02	4.4E-01	2.2E-10	2.2E-08
Plutonium-239	7.90	98.75	2,468.8	3.7E-04	3.6E-02	9.1E-01	2.3E-10	5.7E-07
Thorium-230	14.00	175.00	4,375.0	5.5E-04	9.6E-02	2.4E+00	1.3E-11	5.7E-08
Uranium-234	18.00	225.00	5,625.0	2.8E-04	6.4E-02	1.6E+00	1.6E-11	9.0E-08
Ursnium-235+D	1.70	21.25	531.3	2.7E-04	5.7E-03	1.4E-01	1.6E-11	8.5E-09
Uranium-238+D	69.00	862.50	21,562.5	2.6E-04	2.2E-01	5.5E+00	2.8E-11	6.0E-07
Technetium-99	58.00	725.00	18,125.0	1.5E-06	1.1 E-03	2.6E-02	1.3E-12	2.4E-08
Pathway sum=					4.4E-01	1.1E+01		1 E-06
INHALATION								
Neptunium-237+D	0.32	3.5E-04	8.6E-03	5.4E-01	1.9E-04	4.7E-03	2.9E-08	2.5E-10
Plutonium-239	7.90	8.5E-03	2.1E-01	3.1E-01	2.6E-03	6.6E-02	3.8E-08	8.1E-09
Technetium-99	14.00	1.5E-02	3.8E-01	8.3E-06	1.3E-07	3.1 E-06	8.3E-12	3.1E-12
Thorium-230	18.00	1.9E-02	4.9E-01	2.6E-01	5.1E-03	1.3E-01	2.9E-08	1.4E-08
Uranium-234	1.70	1.8E-03	4.6E-02	1.3E-01	2.4E-04	6.1E-03	2.6E-08	1.2E-09
Ursnium-235+D	69.00	7.5E-02	1.9E+00	1.2E-01	9.2E-03	2.3E-01	2.5E-08	4.7E-08
Uranium-238+D	58.00	6.3E-02	1.6E+00	1.2E-01	7.4E-03	1.8E-01	5.2E-08	8.1E-08
Pathway sum=					2.5E-02	6.2E-01		2 E-07
EXPOSURE TO EXTERNAL RADIATION								
Neptunium-237+D	0.32			1.0E-04	8.4E-02	1.6E+00	4.3E-07	8.2E-07
Plutonium-239	7.90			4.2E-08	6.6E-04	1.7E-02	1.7E-11	8.0E-10
Technetium-99	14.00			1.5E-08	4.2E-04	1.0E-02	6.0E-13	5.0E-11
Thorium-230	18.00			1.2E-07	4.3E-03	1.1E-01	5.4E-11	5.8E-09
Uranium-234	1.70			5.7E-08	1.9E-04	4.8E-03	3.0E-11	3.0E-10
Ursnium-235+D	69.00			3.8E-05	5.2E+00	1.3E+02	2.4E-07	9.9E-05
Uranium-238+D	58.00			7.5E-06	8.7E-01	2.2E+01	3.6E-08	1.2E-05
Pathway sum=					6.2E+00	1.5E+02		1E-04
Sum of the Pathways=					6.6E+00	1.7E+02		1E-04

exposure assumptions

Ingestion Rate IR (g/day)	0.05	Ingestion Risk= SCx IRxEFx EDx RF
Exposure Frequency (EF) (day/yr)	250	Inhalation Risk = SC + IR x EF x ED x CF x 1/PEF x RF
Exposure Duration (ED) (years)	25	External Radiation Risk = SC x ED x Te x (1-SF) x RF
Particulate emission factor (m ³ /kg):	4.63E+09	
Worker inhalation rate (m ³ /day):	20	
Conversion factor(1000 g/kg):	1000	
Exposure Time (ET) (hr/day)	8	
Shielding factor (SF):	0	
Fraction of year exposed (Te):	0.24	Te = (ET x EF)/(8400 HR/YR)

NOTES:

- (a) Radionuclides shown with +D include short lived daughter products in risk calculations.
- (b) Ingestion and inhalation dose factors were taken from Federal Guidance Report 11. "Limiting Values of Radionuclide Intake and Air Concentration and Dose Factors for Inhalation, Submersion, and Ingestion"(EPA-520/1-88-020). Dose after intake of parent radionuclide. External Radiation dose factors were taken from NUREG/CR-5512 "Residual Radioactive Contamination from Decommissioning, Technical Basis for Translating Contamination Levels to Annual Dose."
- (c) Committed effective dose equivalent expressed as committed (50 yr) dose (mrem) due to one year of exposure(mrem/yr).
- (d) Cancer risk factors taken from January 1992 HEAST tables.

Table taken from Attachment 2-3 Remedial Investigation Addendum for Waste Area Grouping 22, Burial Grounds, Solid Waste Management Units 2 and 3, at the Paducah Gaseous Diffusion Plant, Paducah, Kentucky DOE/OR/07-1141&D2 (KY/ER-32 & D2) September 1994 Revision 2.

Table F2.4. Cancer Risk Estimates for Direct Contact to Soil

Current Industrial Exposure Scenario: worker/intruder (25 days/year)				
Chemical	Slope Factor (mg/kg-day) ⁻¹	Soil Conc. (mg/kg)	Chronic Daily Intake (mg/kg-day)	Excess Lifetime Cancer Risk
INGESTION				
Pentachlorophenol	0.12	0.10	1.7E-09	2.1E-10
OCDD (total)	150	0.0033	5.8E-11	8.6E-09
PCB-1248	7.7	0.21	3.7E-09	2.8E-08
PCB-1260	7.7	0.130	2.3E-09	1.7E-08
Arsenic	1.75	10.17	1.8E-07	3.1E-07
Beryllium	4.3	0.74	1.3E-08	5.6E-08
Pathway sum=				4E-07
DERMAL ABSORPTION				
Pentachlorophenol	0.12	0.10	1.1 E-09	1.3E-10
OCDD (total)	150	0.0033	3.6E-11	5.4E-09
PCB-1248	7.7	0.21	2.3E-09	1.8E-08
PCB-1260	7.7	0.130	1.4E-09	1.1 E-08
Arsenic	1.75	10.17	1.1E-08	1.9E-08
Beryllium	4.3	0.74	8.1E-10	3.5E-09
Pathway sum=				6E-08
Unit Risk (ug/m³)-1				
INHALATION				
OCDD (total)	0.000000033	0.0033	5.0E-15	5.8E-19
Arsenic	0.0043	10.17	1.5E-11	2.3E-10
Beryllium	0.0024	0.74	1.1E-12	9.4E-12
Chromium VI	0.01 2	19.00	2.9E-11	1.2E-09
Nickel (soluable salts)	0.00024	25.10	3.8E-11	3.2E-11
Pathway sum=				1E-09
Sum of pathways=				5E-07
Ingestion:	Intake (mg/kg-d)=(conc. in soil*IngR*CF*FI*ED)/(BW*AT) Abs dose(mg/kg-d)=(soil conc. *CF*SA*AF*ABS*EF*ED)/(BW*AT)			
Dermal Absorption:				
Inhalation:	Inh dose (mg/kg-d)=(soil conc. *EF*ED*InhR*(I/PEF))/(BW*AT) SFi=Unit Risk*(BW/InhR)*1000			
exposure parameters				
IngR=Ingestion rate (mg soil/day)	50			
CF=Conversion factor (10E-6)	1E-06			
FI=Fraction ingested	1			
EF=Exposure frequency (days/year)	25			
ED=Exposure duration (year)	70			
BW=Body weight (kg)	25			
AT=Averaging time (days)	70			
SA=Skin surface area (cm ²)	25,550			
AF=Soil to skin adherence (mg/cm ²)	3,120			
ABS=Absorption (0.1 % metals; 1 % organics)	1			
PEF=Particulate emission (m ³ /kg)	0.001			
InhR=Inhalation rate (m ³ /day)	4.63E+09			
	20			

Table taken from Attachment 2-4 Remedial Investigation Addendum for Waste Area Grouping 22, Burial Grounds, Solid Waste Management Units 2 and 3, at the Paducah Gaseous Diffusion Plant, Paducah, Kentucky DOE/OR/07-1141&D2 (KY/ER-32 & D2) September 1994 Revision 2.

Table F2.5. Chronic Hazard Index Estimates for Direct Contact to Soil

Current Industrial Exposure

Scenario: worker/intruder (25 days/year)

Chemical	Reference Dose mg/kg-day	Soil Conc. mg/kg	Chronic Daily Intake mg/kg-day	Hazard Quotient
INGESTION				
Pentachlorophenol	0.03	0.10	4.9E-09	1.6E-07
Arsenic	0.0003	10.17	5.0E-07	1.7E-03
Barium	0.07	132.68	6.5E-06	9.3E-05
Beryllium	0.005	0.74	3.6E-08	7.2E-06
Chromium VI	0.005	19.00	9.3E-07	1.9E-04
Copper	0.037	24.71	1.2E-06	3.3E-05
Manganese	0.14	2,541.05	1.2E-04	8.9E-04
Mercury	0.0003	0.15	7.3E-09	2.4E-05
Nickel (soluble salt)	0.02	25.10	1.2E-06	6.1E-05
Selenium	0.005	0.40	2.0E-08	3.9E-06
Silver	0.003	5.38	2.6E-07	8.8E-05
Vanadium	0.007	31.80	1.6E-06	2.2E-04
Zinc	0.30	67.05	3.3E-06	1.1E-05
Uranium (soluble salt)	0.003	83.58	4.1 E-06	1.4E-05
pathway sum=				0.005
DERMAL ABSORPTION				
Pentachlorophenol	0.03	0.10	3.1E-09	1.0E-07
Arsenic	0.0003	10.17	3.1E-08	1.0E-04
Barium	0.07	132.68	4.1E-07	5.8E-06
Beryllium	0.005	0.74	2.3E-09	4.5E-07
Chromium VI	0.005	19.00	5.8E-08	1.2E-05
Copper	0.037	24.71	7.5E-08	2.0E-06
Manganese	0.005	2,541.05	7.8E-06	1.6E-03
Mercury	0.0003	0.15	4.6E-10	1.5E-06
Nickel (soluble salt)	0.02	25.10	7.7E-08	3.8E-06
Selenium	0.005	0.40	1.2E-09	2.4E-07
Silver	0.003	5.38	1.6E-08	5.5E-06
Vanadium	0.007	31.80	9.7E-08	1.4E-05
Zinc	0.3	67.05	2.0E-07	6.8E-07
Uranium (soluble salt)	0.003	83.58	2.6E-07	8.5E-05
pathway sum=				0.002
	Reference Conc. mg/m³			
INHALATION				
Barium	0.0005	132.68	5.6E-10	3.9E-06
Chromium VI	0.000002	19.00	8.0E-11	1.4E-04
Manganese	0.0004	2,541.05	1.1E-08	9.4E-05
Mercury	0.0003	0.15	6.3E-13	7.4E-09
pathway sum=				0.0002
sum of pathways=				0.007
Ingestion:	Intake (mg/kg-d)=(conc. in soil*IngR*CF*FI*EF*ED)/(BW*AT)			
Dermal Absorption:	Abs dose(mg/kg-d)=(soil conc.*CF*SA*AF*ABS*EF*ED)/(BW*AT)			
Inhalation:	Inh dose (mg/kg-d)=(soil conc.*EF*ED*InhR*(1/PEF)/(BW*AT)			
	SFI=Unit Risk*(BW/InhR)*1000			

Table F2.5. Chronic Hazard Index Estimates for Direct Contact to Soil (Continued)

exposure parameters

IngR=Ingestion rate (mg soil/day)	50	
CF=Conversion factor (10E-6)	1E-06	
FI=Fraction ingested	1	
EF=Exposure frequency (days/year)	25	
ED=Exposure duration (year)	25	
BW=Body weight (kg)	70	
AT=Averaging time (days)	25,550	
SA=Skin surface area (cm ²)	3,120	
AF=Soil to skin adherence (mg/cm ²)	1	
ABS=Absorption (0.1 % metals; 1 % organics)	0.001	0.01
PEF=Particulate emission (m ³ /kg)	4.63E+09	
InhR=Inhalation rate (m ³ /day)	20	

Table taken from Attachment 2-5 Remedial Investigation Addendum for Waste Area Grouping 22, Burial Grounds, Solid Waste Management Units 2 and 3, at the Paducah Gaseous Diffusion Plant, Paducah, Kentucky DOE/OR/07-1141&D2 (KY/ER-32 & D2) September 1994 Revision 2.

Table F2.6. Excess Lifetime Risk of Cancer Incidence for Direct Contact to Soil

Current Industrial Exposure Scenario: worker/intruder (25 days/year)								
Radionuclide ^a	Soil Concentration (pCi/g) (SC)	Annual Intake (pCi)	Total Intake (pCi)	Dose Conversion Factor ^b (mrem/pCi) or (mrem*g/pCi/h)	Committed Effective Dose Equivalent 1	Total Committed Effective Dose Equivalent	Cancer Risk Factor	Risk of Cancer Incidence
					(mrem/yr) ^c	(mrem)	(pCi)-1 ^d or (g/pCi-1)	
INGESTION								
Neptunium-237+D	0.36	0.45	11.3	4.4E-03	2.0E-03	5.0E-02	2.2E-10	2.5E-09
Plutonium-239	7.90	9.88	246.9	3.7E-04	3.6E-03	9.1E-01	2.3E-10	5.7E-08
Thorium-230	14.00	17.50	437.5	5.5E-04	9.6E-03	2.4E-01	1.3E-11	5.7E-09
Uranium-234	18.00	22.50	562.5	2.8E-04	6.4E-03	1.6E-01	1.6E-11	9.0E-09
Ursnium-235+D	1.70	2.13	53.1	2.7E-04	5.7E-04	1.4E-02	1.6E-11	8.5E-10
Uranium-238+D	69.00	86.25	2156.3	2.6E-04	2.2E-02	5.5E-01	2.8E-11	6.0E-08
Technetium-99	58.00	72.50	1812.5	1.5E-06	1.1 E-04	2.6E-03	1.3E-12	2.4E-09
Pathway sum=					4.4E-02	1.1E+00		1 E-07
INHALATION								
Neptunium-237+D	0.36	3.9E-05	9.7E-04	5.4E-01	2.1E-05	5.2E-04	2.9E-08	2.8E-11
Plutonium-239	7.90	8.5E-04	2.1E-02	3.1E-01	2.6E-04	6.6E-03	3.8E-08	8.1E-10
Technetium-99	14.00	1.5E-03	3.8E-02	8.3E-06	1.3E-08	3.1 E-07	8.3E-12	3.1E-13
Thorium-230	18.00	1.9E-03	4.9E-02	2.6E-01	5.1E-04	1.3E-02	2.9E-08	1.4E-09
Uranium-234	1.70	1.8E-04	4.6E-03	1.3E-01	2.4E-05	6.1E-04	2.6E-08	1.2E-10
Ursnium-235+D	69.00	7.5E-03	1.9E-01	1.2E-01	9.2E-04	2.3E-02	2.5E-08	4.7E-09
Uranium-238+D	58.00	6.3E-03	1.6E-01	1.2E-01	7.4E-04	1.8E-02	5.2E-08	8.1E-09
Pathway sum=	7.90				2.5E-03	6.2E-02		2 E-08
EXPOSURE TO EXTERNAL RADIATION								
Neptunium-237+D	0.36			1.0E-04	7.2E-03	1.8E-01	4.3E-07	9.2E-08
Plutonium-239	7.90			4.2E-08	6.6E-05	1.7E-03	1.7E-11	8.0E-11
Technetium-99	14.00			1.5E-08	4.2E-05	1.0E-03	6.0E-13	5.0E-12
Thorium-230	18.00			1.2E-07	4.3E-04	1.1E-02	5.4E-11	5.8E-10
Uranium-234	1.70			5.7E-08	1.9E-05	4.8E-04	3.0E-11	3.0E-11
Ursnium-235+D	69.00			3.8E-05	5.2E-01	1.3E+01	2.4E-07	9.9E-06
Uranium-238+D	58.00			7.5E-06	8.7E-02	2.2E+00	3.6E-08	1.2E-06
Pathway sum=	7.90				6.2E-01	1.5E+01		1E-05
Sum of the Pathways=					6.7E-01	1.7E+01		1E-05

exposure assumptions

Ingestion Rate IR (g/day)	0.05	Ingestion Risk= SCx IRxEFx EDx RF
Exposure Frequency (EF) (day/yr)	25	Inhalation Risk = SC + IR x EF x ED x CF x I/PEF x RF
Exposure Duration (ED) (years)	25	External Radiation Risk = SC x ED x Te x (1-SF) x RF
Particulate emission factor (m ³ /kg):	4.63E+09	
Worker inhalation rate (m ³ /day):	20	
Conversion factor(1000 g/kg):	1,000	
Exposure Time (ET) (hr/day)	8	
Shielding factor (SF):	0	
Fraction of year exposed (Te):	0.2	Te = (ET x EF)/(8,400 HR/YR)

NOTES:

- (a) Radionuclides shown with +D include short lived daughter products in risk calculations.
- (b) Ingestion and inhalation dose factors were taken from Federal Guidance Report 11. "Limiting Values of Radionuclide Intake and Air Concentration and Dose Factors for Inhalation, Submersion, and Ingestion"(EPA-520/1-88-020). Dose after intake of parent radionuclide. External Radiation dose factors were taken from NUREG/CR-5512 "Residual Radioactive Contamination from Decommissioning, Technical Basis for Translating Contamination Levels to Annual Dose."
- (c) Committed effective dose equivalent expressed as committed (50 yr) dose (mrem) due to one year of exposure(mrem/yr).
- (d) Cancer risk factors taken from January1992 HEAST tables.

Table taken from Attachment 2-6 Remedial Investigation Addendum for Waste Area Grouping 22, Burial Grounds, Solid Waste Management Units 2 and 3, at the Paducah Gaseous Diffusion Plant, Paducah, Kentucky DOE/OR/07-1141&D2 (KY/ER-32 & D2) September 1994 Revision 2.

Table F2.7. Cancer Risks Estimated for Domestic Use of Groundwater

Scenario: Future Potable Use of Groundwater
 MW093 (RGA)

Chemical	Oral Slope Factor (mg/kg-day)⁻¹	Concentration MW093 ug/L	Chronic Daily Intake mg/kg-day	Excess Lifetime Cancer Risk	Total Pathway Risk
INGESTION OF GROUNDWATER					
2,4-Dinitrotoluene	0.68	18.50	2.2E-04	1.5E-04	
N-Nitroso-di-npropylamine	7.0	22.00	2.6E-04	1.8E-03	
Pentachlorophenol	0.12	57.00	6.7E-04	8.0E-05	
Arsenic	1.75	3.35	3.9E-05	6.9E-05	
Sum=					2E-03

Ingestion: Intake (mg/kg-d)=(conc. in gw*IngR*CF* EF*ED)/(BW*AT)

exposure parameters

IngR=Ingestion rate (L/day) 2
 CF=Conversion factor (mg/ug) 0.001
 EF=Exposure frequency (days/year) 350
 ED=Exposure duration (year) 30
 BW=Body weight (kg) 70
 AT=Averaging time (days) 25,550

Table taken from Attachment 2-7 Remedial Investigation Addendum for Waste Area Grouping 22, Burial Grounds, Solid Waste Management Units 2 and 3, at the Paducah Gaseous Diffusion Plant, Paducah, Kentucky DOE/OR/07-1141&D2 (KY/ER-32 & D2) September 1994 Revision 2.

Table F2.8. Cancer Risks Estimated for Domestic Use of Groundwater

Scenario: Future Potable Use of Groundwater

MW074 (UCRS)

Chemical	Oral Slope Factor (mg/kg-day)⁻¹	Concentration MW074 ug/L	Chronic Daily Intake mg/kg-day	Excess Lifetime Cancer Risk	Total Pathway Risk
INGESTION OF GROUNDWATER					
Beryllium	4.3	15.8	1.9E-04	8.0E-04	8.0E-04
INHALATION OF VOLATILE COMPOUNDS DURING DOMESTIC USE OF GROUNDWATER					0.0E+00
Sum of Pathways=					8E-04

Ingestion: Intake (mg/kg-d)=(conc. in gw*IngR*CF* EF*ED)/(BW*AT)
 Inhalation: Inh dose (mg/kg-d)=(conc. in gw*VF*InhR*EF*ED)/(BW*AT)
 SFi=Unit Risk*(BW/InhR)*1000

exposure parameters

IngR=Ingestion rate (L/day) 2
 CF=Conversion factor (mg/ug) 0.001
 EF=Exposure frequency (days/year) 350
 ED=Exposure duration (year) 30
 BW=Body weight (kg) 70
 AT=Averaging time (days) 25,550
 InhR=Indoor Inhalation Rate (m³/day) 15
 VF=Volatilization Factor (L/m³) 0.5

Table taken from Attachment 2-8 Remedial Investigation Addendum for Waste Area Grouping 22, Burial Grounds, Solid Waste Management Units 2 and 3, at the Paducah Gaseous Diffusion Plant, Paducah, Kentucky DOE/OR/07-1141&D2 (KY/ER-32 & D2) September 1994 Revision 2.

Table F2.9. Hazard Index Estimates for Domestic Use of Groundwater

Scenario: Future Potable Use of Groundwater
 MW074 (UCRS)

Chemical	Reference Dose (mg/kg-day)	Concentration MW074 ug/L	Chronic Daily Intake mg/kg-day	Hazard Quotient	Pathway Hazard Index
INGESTION OF GROUNDWATER					
Nickel	0.02	125.4	3.4E-03	0.172	
Barium	0.07	634	1.7E-02	0.248	
Zinc	0.3	343.3	9.4E-03	0.031	
Vanadium	0.007	410.1	1.1E-02	1.605	
Chromium	0.005	139.8	3.8E-03	0.766	
Cadmium	0.0005	4.6	1.3E-04	0.252	
Silver	0.003	42.1	1.2E-03	0.384	
Manganese	0.005	1,535.3	4.2E-02	8.413	
Copper	0.037	95.9	2.6E-03	0.000	
Beryllium	0.005	15.8	4.3E-04	0.087	
Uranium (soluble salts)	0.003	10.68	2.9E-04	0.098	

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Equations:

Ingestion: $\text{Intake (mg/kg-d)} = (\text{conc. in gw} * \text{IngR} * \text{CF} * \text{EF} * \text{ED}) / (\text{BW} * \text{AT})$

exposure parameters

IngR=Ingestion rate (L/day) 2
 CF=Conversion factor (mg/ug) 0.001
 EF=Exposure frequency (days/year) 350
 ED=Exposure duration (year) 30
 BW=Body weight (kg) 70
 AT=Averaging time (days) 10,950

Table taken from Attachment 2-9 Remedial Investigation Addendum for Waste Area Grouping 22, Burial Grounds, Solid Waste Management Units 2 and 3, at the Paducah Gaseous Diffusion Plant, Paducah, Kentucky DOE/OR/07-1141&D2 (KY/ER-32 & D2) September 1994 Revision 2.

Table F2.10. Hazard Index Estimates for Domestic Use of Groundwater

Scenario: Future Potable Use of Groundwater
 MW089 (RGA)

Chemical	Reference Dose (mg/kg-day)	Concentration MW074 ug/L	Chronic Daily Intake mg/kg-day	Hazard Quotient	Pathway Hazard Index
INGESTION OF GROUNDWATER					
Nickel	0.02	14.6	4.0E-04	0.020	
Barium	0.07	253	6.9E-03	0.099	
Zinc	0.3	34.3	9.4E-04	0.003	
Vanadium	0.007	7.6	2.1E-04	0.030	
Chromium	0.005	7.8	2.1E-04	0.043	
Cyanide	0.02	3	8.2E-05	0.004	
Thallium (Carbonate)	0.00008	0.9	2.5E-05	0.308	
Manganese	0.005	3,630	9.9E-02	19.890	
Copper	0.037	9	2.5E-04	0.000	
Arsenic	0.0003	3.9	1.1E-04	0.356	
					20.75
INHALATION OF VOLATILE COMPOUNDS DURING DOMESTIC USE OF GROUNDWATER					
Sum of Pathways=					0.0
					20.75

Equations:

Ingestion: $\text{Intake (mg/kg-d)} = (\text{conc. in gw} * \text{IngR} * \text{CF} * \text{EF} * \text{ED}) / (\text{BW} * \text{AT})$
 Inhalation: $\text{Inh dose (mg/kg-d)} = (\text{conc. in gw} * \text{VF} * \text{InhR} * \text{EF} * \text{ED}) / (\text{BW} * \text{AT})$

exposure parameters

IngR=Ingestion rate (L/day) 2
 CF=Conversion factor (mg/ug) 0.001
 EF=Exposure frequency (days/year) 350
 ED=Exposure duration (year) 30
 BW=Body weight (kg) 70
 AT=Averaging time (days) 10,950
 InhR=Indoor Inhalation Rate (m³/day) 15
 VF=Volatilization Factor (L/m³) 0.5

Table taken from Attachment 2-10 Remedial Investigation Addendum for Waste Area Grouping 22, Burial Grounds, Solid Waste Management Units 2 and 3, at the Paducah Gaseous Diffusion Plant, Paducah, Kentucky DOE/OR/07-1141&D2 (KY/ER-32 & D2) September 1994 Revision 2.

Table F2.11. Risk of Cancer Incidence for Domestic Use of Groundwater

Scenario: Future Potable Use of Groundwater
 MW154 (UCRS)

Radionuclide ^a	Groundwater Concentration (pCi/L) ^b	Annual Intake (pCi/yr)	Total Intake (pCi)	Ingestion Dose Conversion Factor ^c (mrem/pCi)	Committed Effective Dose Equivalent 1 Yr Intake (mrem/yr) ^d	Total Committed Effective Dose Equivalent (mrem)	Cancer Incidence Risk Factor (pCi)-1 ^e	Risk of Cancer Incidence
Np-237	0.32	224.0	6720.0	4.4E-03	9.9E-01	3.0E+01	2.2E-10	1.5E-06
Pu-239	0.18	126.0	3780.0	3.7E-04	4.6E-02	1.4E+00	2.3E-10	6.7E-07
	1000	7.0E+05	2.1E+07	1.5E-06	1.0E+00	3.1E+01	1.3E-12	2.7E-05
Tc-99	3.6	2,520.0	75,600.0	2.8E-04	7.1E-01	2.1E+01	1.6E-11	1.2E-06
	0.14	98.0	2,940.0	2.7E-04	2.6E-02	7.8E-01	1.6E-11	4.7E-08
U-234	27	18,900.0	567,000.0	2.6E-04	4.8E+00	1.4E+02	2.8E-11	1.6E-05
U-235+D								
U-238+D								
Pathway totals= exposure assumptions					7.6E+00	2.3E+02		5E-05

Ingestion Rate IR (g/day) 2 Ingestion Risk= WC x IR x EF x ED x RF
 Exposure Frequency (EF) (day/yr) 350
 Exposure Duration (ED) (years) 30

NOTES:

^aRadionuclides shown with +D include short lived daughter products in risk calculations.

^bSample concentrations are actual values. Results are shown as calculated by the lab, even if they are less than the detection limit for this analysis. ND is shown if the actual value was negative.

^cIngestion and inhalation dose factors were taken from Federal Guidance Report 11. "Limiting Values of Radionuclide Intake and Air Concentration and Dose Factors for Inhalation, Submersion, and Ingestion"(EPA-520/1-88-020). Dose factors include the contribution to dose from ingrowth of decay products after intake of parent radionuclide.

^dCommitted effective dose equivalent expressed as committed (50 yr) dose (mrem) due to one year of exposure (mrem/yr).

^eCancer risk factors taken from January 1992 HEAST tables.

Table taken from Attachment 2-11 Remedial Investigation Addendum for Waste Area Grouping 22, Burial Grounds, Solid Waste Management Units 2 and 3, at the Paducah Gaseous Diffusion Plant, Paducah, Kentucky DOE/OR/07-1141&D2 (KY/ER-32 & D2) September 1994 Revision 2.

Table F2.12. Risk of Cancer Incidence for Domestic Use of Groundwater

Scenario: Future Potable Use of Groundwater
 MW84 (RGA)

Radionuclide ^a	Groundwater Concentration (pCi/L) ^b	Annual Intake (pCi/yr)	Total Intake (pCi)	Ingestion Dose Conversion Factor ^c (mrem/pCi)	Committed Effective Dose Equivalent 1 Yr Intake (mrem/yr) ^d	Total Committed Effective Dose Equivalent (mrem)	Cancer Incidence Risk Factor (pCi)-1 ^e	Risk of Cancer Incidence
Np-237	ND	0.0	0.0	4.4E-03	0.0E+00	0.0E+00	2.2E-10	0.0E+00
Pu-239	0.03	21.0	630.0	3.7E-04	7.7E-03	2.3E-01	2.3E-10	1.4E-07
	466	3.3E+05	9.8E+06	1.5E-06	4.8E-01	1.4E+01	1.3E-12	1.3E-05
Tc-99	0.14	98.0	2940.0	2.8E-04	2.8E-02	8.3E-01	1.6E-11	4.7E-08
	0.01	7.0	210.0	2.7E-04	1.9E-03	5.6E-02	1.6E-11	3.4E-09
U-234	0.23	161.0	4,830.0	2.6E-04	4.1E-02	1.2E+00	2.8E-11	1.4E-07
U-235+D								
U-238+D								
Totals					5.5E-01	1.7E+01		1E-05

exposure assumptions

Ingestion Rate IR (g/day) 2
 Exposure Frequency (EF) (day/yr) 350
 Exposure Duration (ED) (years) 30
 Ingestion Risk= WC x IR x EF x ED x RF

NOTES:

^aRadionuclides shown with +D include short lived daughter products in risk calculations.

^bSample concentrations are actual values. Results are shown as calculated by the lab, even if they are less than the detection limit for this analysis. ND is shown if the actual value was negative.

^c Ingestion and inhalation dose factors were taken from Federal Guidance Report 11. "Limiting Values of Radionuclide Intake and Air Concentration and Dose Factors for Inhalation, Submersion, and Ingestion"(EPA-520/1-88-020). Dose factors include the contribution to dose from ingrowth of decay products after intake of parent radionuclide.

^d Committed effective dose equivalent expressed as committed (50 yr) dose (mrem) due to one year of exposure (mrem/yr).

^e Cancer risk factors taken from January 1992 HEAST tables.

Table taken from Attachment 2-12 Remedial Investigation Addendum for Waste Area Grouping 22, Burial Grounds, Solid Waste Management Units 2 and 3, at the Paducah Gaseous Diffusion Plant, Paducah, Kentucky DOE/OR/07-1141&D2 (KY/ER-32 & D2) September 1994 Revision 2.

Table F2.13. Summary of Human Health Risk Characterization for SWMU 4 Without Lead as a COPC

Receptor	Total ELCR	COCs	% Total ELCR	POCs	% Total ELCR	Total HI	COCs	% Total HI	POCs	% Total HI
Current industrial worker at current concentrations (soil)	5.4E-04	Beryllium Uranium-238	97 2	Dermal contact External exposure	97 2	3.62	Beryllium Chromium Iron Vanadium Barium	5 45 24 24 2	Dermal contact	99
Future industrial worker at current concentrations (soil)	5.4E-04	Beryllium Uranium-238	97 2	Dermal contact External exposure	97 2	3.62	Beryllium Chromium Iron Vanadium Barium	5 45 24 24 2	Dermal contact	99
Future industrial worker at current concentrations (RGA groundwater)	4.7E-04	Arsenic Beryllium 1,1-DCE Carbon tetrachloride Chloroform TCE Vinyl chloride	15 48 8 7 2 20 2	Incidental ingestion Dermal contact Inhalation while showering	72 18 10	32.6	Aluminum Arsenic Cadmium Chromium Iron Manganese Vanadium Carbon tetrachloride TCE	4 1 1 1 66 5 2 4 14	Ingestion Dermal contact Inhalation while showering	88 6 6
Future industrial worker at current concentrations (McNairy groundwater)	3.1E-03	Arsenic Beryllium	18 82	Ingestion Dermal contact	78 22	75.9	Aluminum Arsenic Barium Beryllium Cadmium Chromium Iron Manganese Vanadium	4 5 1 1 1 3 63 8 14	Ingestion Dermal contact	93 7
Future child rural resident at current concentrations (soil)	NA	NA	NA	NA	NA	98.2	Barium Beryllium Cadmium Chromium Iron Nickel Vanadium	2 2 2 24 60 2 9	Ingestion Dermal contact Ingestion of vegetables	1 21 78

Table F2.13. Summary of Human Health Risk Characterization for SWMU 4 Without Lead as a COPC (Continued)

Receptor	Total ELCR	COCs	% Total ELCR	POCs	% Total ELCR	Total HI	COCs	% Total HI	POCs	% Total HI
Future child rural resident at current concentrations (RGA groundwater)	NA	NA	NA	NA	NA	487	Aluminum Arsenic Boron Chromium Iron Manganese Vanadium Carbon tetrachloride Chloroform TCE <i>cis</i> -1,2-DCE	3 1 1 1 49 3 1 10 1 29 1	Ingestion Dermal contact Inhalation while showering/household Ingestion of vegetables	40 1 30 29
Future child rural resident at current concentrations (McNairy groundwater)	NA	NA	NA	NA	NA	798	Aluminum Arsenic Barium Beryllium Cadmium Chromium Iron Manganese Mercury Vanadium Zinc	4 5 1 1 1 3 66 6 1 12 1	Ingestion Dermal contact Ingestion of vegetables	60 2 39
Future adult rural resident at current concentrations (soil)	4.3E-03	Beryllium Total PCBs Uranium-234 Uranium-238	72 5 6 17	Dermal contact External exposure Ingestion of vegetables	6 2 1	28.4	Barium Beryllium Cadmium Chromium Iron Nickel Vanadium	2 2 2 22 63 2 8	Dermal contact Ingestion of vegetables	14 85
Future adult rural resident at current concentrations (RGA groundwater)	7.0E-03	Arsenic Beryllium 1,1-DCE Carbon tetrachloride Chloroform TCE Vinyl chloride Technetium-99	8 22 15 7 5 20 2 21	Ingestion Dermal contact Inhalation while showering/household Ingestion of vegetables	26 3 30 41	158	Aluminum Arsenic Boron Chromium Iron Manganese Vanadium Carbon tetrachloride TCE	3 1 1 1 57 4 1 7 22	Ingestion Dermal contact Inhalation of vapors/particles Ingestion of vegetables	51 2 19 28

Table F2.13. Summary of Human Health Risk Characterization for SWMU 4 Without Lead as a COPC (Continued)

Receptor	Total ELCR	COCs	% Total ELCR	POCs	% Total ELCR	Total HI	COCs	% Total HI	POCs	% Total HI
Future adult rural resident at current concentrations (McNairy groundwater)	> 1.0E-02*	Arsenic Beryllium Technetium-99	21 77 2	Ingestion Dermal contact Ingestion of vegetables	58 8 35	303	Aluminum Arsenic Barium Beryllium Cadmium Chromium Iron Manganese Vanadium Zinc	4 5 1 1 3 66 6 12 1	Ingestion Dermal contact Ingestion of vegetables	65 2 32
Future child recreational user at current concentrations (soil)	NA	NA	NA	NA	NA	<1	-	-	-	-
Future teen recreational user at current concentrations (soil)	NA	NA	NA	NA	NA	<1	-	-	-	-
Future adult recreational user at current concentrations (soil)	< 1.0E-06	-	-	-	-	<1	-	-	-	-
Future excavation worker at current concentrations (soil and waste)	2.7E-03	Arsenic Beryllium Total dioxins/furans Total PCBs Radium-226 Total uranium Uranium-238	1 7 4 2 2 83 1	Ingestion Dermal contact External exposure	37 10 54	2.61	Aluminum Arsenic Barium Beryllium Cadmium Chromium Iron Manganese Vanadium	8 4 2 2 1 24 24 14 20	Ingestion Dermal contact	13 87

Table taken from Remedial Investigation Report for Waste Area Grouping 3 at the Paducah Gaseous Diffusion Plant, Paducah, Kentucky, DOE/OR-07-1895&D1, July 2000.

NA = ELCR not applicable to child and teen cohorts. Values for adult include exposure as child and teen.

- = There are no COCs or POCs.

* = The ELCR is approximate because the linearized multistage model returns imprecise values at risks > 1.0E-02.

Table F2.14. Summary of Human Health Risk Characterization for SWMU 5 Without Lead as a COPC

Receptor	Total ELCR	COCs	% Total ELCR	POCs	% Total ELCR	Total HI	COCs	% Total HI	POCs	% Total HI
Current industrial worker at current concentrations (soil)	4.1E-04	Arsenic Beryllium Total PAHs	6 49 45	Ingestion Dermal contact	2 98	<1	-	-	-	-
Future industrial worker at current concentrations (soil)	4.1E-04	Arsenic Beryllium Total PAHs	6 49 45	Ingestion Dermal contact	2 98	<1	-	-	-	-
Future industrial worker at current concentrations (RGA groundwater)	5.4E-04	Beryllium 1,1-DCE Radium-226	35 1 64	Ingestion Dermal contact	90 9	26.8	Aluminum Barium Cadmium Chromium Iron Manganese Vanadium	4 1 1 2 73 16 2	Ingestion Dermal contact	96 4
Future industrial worker at current concentrations (McNairy groundwater)	1.2E-03	Beryllium Radium-226	42 58	Ingestion Dermal contact	89 11	63	Aluminum Cadmium Chromium Iron Manganese Vanadium	4 1 7 79 3 5	Ingestion Dermal contact	95 5
Future child rural resident at current concentrations (soil)	NA	NA	NA	NA	NA	46.2	Aluminum Arsenic Beryllium Chromium Nickel Zinc	24 53 1 17 3 1	Ingestion Dermal contact Ingestion of vegetables	1 12 87
Future child rural resident at current concentrations (RGA groundwater)	NA	NA	NA	NA	NA	283	Aluminum Barium Cadmium Chromium Iron Manganese Vanadium	4 1 1 2 77 12 1	Ingestion Dermal contact Ingestion of vegetables	61 1 37
Future child rural resident at current concentrations (McNairy groundwater)	NA	NA	NA	NA	NA	680	Aluminum Cadmium Chromium Iron Manganese Vanadium	4 1 6 81 3 4	Ingestion Dermal contact Ingestion of vegetables	60 1 39
Future adult rural resident at current concentrations (soil)	> 1.0E-02*	Arsenic Beryllium Total PAHs Total PCBs	21 9 68 2	Dermal contact Ingestion of vegetables	9 90	13.9	Aluminum Arsenic Beryllium Chromium Nickel Zinc	24 55 1 15 3 1	Dermal contact Ingestion of vegetables	8 92

Table F2.14. Summary of Human Health Risk Characterization for SWMU 5 Without Lead as a COPC (Continued)

Receptor	Total ELCR	COCs	% Total ELCR	POCs	% Total ELCR	Total HI	COCs	% Total HI	POCs	% Total HI
Future adult rural resident at current concentrations (RGA groundwater)	3.9E-03	Beryllium I,1-DCE Radium-226 Technetium-99	33 4 57 5	Ingestion Dermal contact Inhalation while showering/household Ingestion of vegetables	56 3 4 37	107	Aluminum Barium Cadmium Chromium Iron Manganese Vanadium	4 1 1 2 76 13 1	Ingestion Dermal contact Ingestion of vegetables	67 2 31
Future adult rural resident at current concentrations (McNairy groundwater)	8.2E-03	Beryllium Radium-226	43 57	Ingestion Dermal contact Ingestion of vegetables	61 4 34	257	Aluminum Cadmium Chromium Iron Manganese Vanadium	4 1 6 81 3 4	Ingestion Dermal contact Ingestion of vegetables	65 2 33
Future child recreational user at current concentrations (soil)	NA	NA	NA	NA	NA	<1	-	-	-	-
Future teen recreational user at current concentrations (soil)	NA	NA	NA	NA	NA	<1	-	-	-	-
Future adult recreational user at current concentrations (soil)	1.0E-05	Arsenic Total PAHs Total PCBs	2 96 2	Ingestion of venison Ingestion of rabbit Ingestion of quail	16 63 21	<1	-	-	-	-
Future excavation worker at current concentrations (soil and waste)	2.9E-04	Arsenic Beryllium Total PAHs Total PCBs	8 62 28 1	Ingestion Dermal contact	13 87	2.16	Aluminum Arsenic Barium Beryllium Chromium Iron Manganese	9 7 2 3 18 38 22	Ingestion Dermal contact	18 82

Table taken from Remedial Investigation Report for Waste Area Grouping 3 at the Paducah Gaseous Diffusion Plant, Paducah, Kentucky, DOE/OR-07-1895&D1, July 2000.

NA = ELCR not applicable to child and teen cohorts. Values for adult include exposure as child and teen.

- = There are no COCs or POCs.

* = The ELCR is approximate because the linearized multistage model returns imprecise values at risks > 1.0E-02.

Table F2.15. Summary of Human Health Risk Characterization for SWMU 6 Without Lead as a COPC

Receptor	Total ELCR	COCs	% Total ELCR	POCs	% Total ELCR	Total HI	COCs	% Total HI	POCs	% Total HI
Current industrial worker at current concentrations (soil)	2.4E-04	Beryllium Total PAHs	90 10	Dermal contact	99	< 1	-	-	-	-
Future industrial worker at current concentrations (soil)	2.4E-04	Beryllium Total PAHs	90 10	Dermal contact	99	< 1	-	-	-	-
Future industrial worker at current concentrations (RGA groundwater)	2.3E-04	Arsenic Beryllium TCE	15 74 11	Ingestion Dermal contact Inhalation while showering	76 22 2	19.1	Aluminum Arsenic Barium Cadmium Chromium Iron Manganese Vanadium TCE	3 1 1 2 2 61 20 3 6	Ingestion Dermal contact Inhalation while showering	92 6 2
Future industrial worker at current concentrations (McNairy groundwater)	7.8E-04	Arsenic Beryllium	24 76	Ingestion Dermal contact	79 21	41.7	Aluminum Arsenic Barium Cadmium Chromium Iron Manganese Vanadium	5 3 1 1 6 74 3 5	Ingestion Dermal contact	95 5
Future child rural resident at current concentrations (soil)	NA	NA	NA	NA	NA	9.38	Beryllium Chromium Nickel Zinc	8 72 15 5	Dermal contact Ingestion of vegetables	34 65
Future child rural resident at current concentrations (RGA groundwater)	NA	NA	NA	NA	NA	223	Aluminum Arsenic Barium Cadmium Chromium Iron Manganese Vanadium TCE	3 1 1 1 2 58 14 2 17	Ingestion Dermal contact Inhalation while showering/household Ingestion of vegetables	54 1 12 33
Future child rural resident at current concentrations (McNairy groundwater)	NA	NA	NA	NA	NA	451	Aluminum Arsenic Barium Cadmium Chromium Iron Manganese Vanadium	5 3 1 1 6 76 2 5	Ingestion Dermal contact Ingestion of vegetables	59 1 39

Table F2.15. Summary of Human Health Risk Characterization for SWMU 6 Without Lead as a COPC (Continued)

Receptor	Total ELCR	COCs	% Total ELCR	POCs	% Total ELCR	Total HI	COCs	% Total HI	POCs	% Total HI
Future adult rural resident at current concentrations (soil)	2.4E-03	Beryllium Total PAHs	54 46	Dermal contact Ingestion of vegetables	30 69	2.57	Beryllium Chromium Nickel Zinc	7 70 17 6	Dermal contact Ingestion of vegetables	24 75
Future adult rural resident at current concentrations (RGA groundwater)	2.3E-03	Arsenic Beryllium TCE Technetium-99	12 51 16 21	Ingestion Dermal contact Inhalation while showering/household Ingestion of vegetables	41 6 8 46	79.9	Aluminum Arsenic Barium Cadmium Chromium Iron Manganese Vanadium TCE	3 1 1 1 2 61 15 2 12	Ingestion Dermal contact Inhalation while showering/household Ingestion of vegetables	62 2 7 29
Future adult rural resident at current concentrations (McNairy groundwater)	5.7E-03	Arsenic Beryllium	28 72	Ingestion Dermal contact Ingestion of vegetables	59 7 34	170	Aluminum Arsenic Barium Cadmium Chromium Iron Manganese Vanadium	5 3 1 1 6 76 2 5	Ingestion Dermal contact Ingestion of vegetables	65 2 33
Future child recreational user at current concentrations (soil)	NA	NA	NA	NA	NA	<1	-	-	-	-
Future teen recreational user at current concentrations (soil)	NA	NA	NA	NA	NA	<1	-	-	-	-
Future adult recreational user at current concentrations (soil)	< 1.0E-06	-	-	-	-	<1	-	-	-	-
Future excavation worker at current concentrations (soil and waste)	2.3E-04	Beryllium Total PAHs	90 9	Ingestion Dermal contact	5 95	2.44	Aluminum Barium Beryllium Chromium Iron Manganese Vanadium	8 2 3 15 32 15 26	Ingestion Dermal contact	12 88

Table taken from Remedial Investigation Report for Waste Area Grouping 3 at the Paducah Gaseous Diffusion Plant, Paducah, Kentucky, DOE/OR-07-1895&D1, July 2000.
 NA = ELCR not applicable to child and teen cohorts. Values for adult include exposure as child and teen.
 - = There are no COCs or POCs.

Table F2.16. Summary of Human Health Risk Characterization for SWMU 7 Without Lead as a COPC

Receptor	Total ELCR	ELCR COCs	% Total ELCR	ELCR POCs	% Total ELCR	Total HI ^a	Systematic Toxicity COCs	% Total HI	Systematic Toxicity POCs	% Total HI
Current industrial worker at current concentrations	4 x 10 ⁻³	Arsenic Beryllium Benz(a)anthracene Benzo(a)pyrene Benzo(b)fluoranthene Indeno(1,2,3-cd)pyrene ²³⁷ Np ²³⁸ U ²³⁵ U ²³⁸ U	<1 97.4 <1 <1 <1 <1 <1 <1 <1 <1	Ingestion of soil Dermal contact with soil External exposure to soil	<1 97.4 2.5	5	Aluminum Antimony Arsenic Beryllium Chromium Iron Manganese Uranium Vanadium	4.1 4.4 2.6 9.6 13.6 20.7 10.7 13.7 17.7	Ingestion of soil Dermal contact with soil	3.6 96.4
Future industrial worker at current concentrations	6 x 10 ⁻³	Arsenic Beryllium 1,1-DCE PCB-1248 PCB-1254 Benz(a)anthracene Benzo(a)pyrene Benzo(b)fluoranthene Carbon tetrachloride Dibenzo(a,h)anthracene Indeno(1,2,3-cd)pyrene TCE Vinyl chloride ²³⁷ Np ²³⁸ Pu ²²⁷ Rn ⁹⁹ Tc ²³⁰ Th ²³⁴ U ²³⁵ U ²³⁸ U	12.0 70.9 <1 <1 1.3 <1 <1 <1 <1 <1 <1 2.4 4.9 1.7 <1 2.5 <1 <1 <1 <1 <1 2.4	Ingestion of groundwater Dermal contact with groundwater Inhalation while showering Ingestion of soil Dermal contact with soil External exposure to soil	25.5 2.9 3.5 <1 67.3 1.7	62	Aluminum Antimony Arsenic Barium Beryllium Cadmium Chromium Iron Manganese Nickel Uranium Vanadium 1,2- <i>cis</i> -DCE 1,2- DCE (total) 2,4-Dimethylphenol 4-Methylphenol PCB-1254 Carbon tetrachloride TCE	2.3 1.0 6.7 <1 <1 <1 2.9 8.4 8.8 <1 1.2 3.2 <1 13.8 <1 <1 40.0 <1 7.0	Ingestion of groundwater Dermal contact with groundwater Ingestion of soil Dermal contact with soil	63.7 27.2 <1 7.8
Future child recreational user at current concentrations	NA	NA	NA	NA	NA	0.07	NE	NE	NE	NE
Future teen recreational user at current concentrations	NA	NA	NA	NA	NA	0.06	NE	NE	NE	NE
Future adult recreational user at current concentrations	1 x 10 ⁻⁵	PCB-1260 Benzo(a)pyrene Dibenzo(a,h)anthracene ²³⁸ U	18.2 9.1 41.8 16.4	Ingestion of deer Ingestion of rabbit Ingestion of quail	10.0 70.9 21.8	0.07	NE	NE	NE	NE

Table F2.16. Summary of Human Health Risk Characterization for SWMU 7 Without Lead as a COPC (Continued)

Receptor	Total ELCR	ELCR COCs	% Total ELCR	ELCR POCs	% Total ELCR	Total HI ^a	Systematic Toxicity COCs	% Total HI	Systematic Toxicity POCs	% Total HI
Future child rural resident at current concentrations	NA	NA	NA	NA	NA	1320	Aluminum Antimony Arsenic Barium Beryllium Cadmium Chromium Cobalt Copper Fluoride Iron Manganese Mercury Molybdenum Nickel Nitrate-nitrite Selenium Silver Tin Uranium Vanadium Zinc 1,2- <i>cis</i> -DCE 1,2- DCE (total) 2,4-Dimethylphenol 2-Methylphenol 4-Methylphenol Acetone PCB-1254 Benzene Carbon tetrachloride TCE	1.8 <1 5.2 <1 <1 <1 <1 1.6 <1 <1 <1 8.9 3.9 <1 <1 <1 <1 <1 <1 <1 16.4 1.4 <1 <1 40.8 <1 <1 <1 <1 12.4 <1 <1 4.0	Ingestion of groundwater Dermal contact with groundwater Inhalation while showering Inhalation from household use Ingestion of vegetables from groundwater Ingestion of soil Dermal contact with soil Ingestion of vegetables from soil	20.7 3.5 <1 <1 <1 47.6 <1 2.2 25.5

Table F2.16. Summary of Human Health Risk Characterization for SWMU 7 Without Lead as a COPC (Continued)

Receptor	Total ELCR	ELCR COCs	% Total ELCR	ELCR POCs	% Total ELCR	Total HI ^a	Systematic Toxicity COCs	% Total HI	Systematic Toxicity POCs	% Total HI		
Future adult rural resident at current concentrations	5 x 10 ⁻²	Arsenic	14.6	Ingestion of groundwater	13.1	446	Aluminum	1.8	Ingestion of groundwater	25.1		
		Beryllium	42.6	Dermal contact with groundwater	<1		Antimony	<1	Dermal contact with groundwater	5.3		
		1,1,2-TCA	<1	Inhalation while showering	<1		Arsenic	5.4	Barium	<1	Inhalation from household use	<1
		1,1-DCE	<1	Inhalation from household use	3.5		Beryllium	<1	Cadmium	<1	Ingestion of vegetables from groundwater	44.2
		1,2-DCE	<1	Ingestion of soil	20.3		Chromium	1.6	Cobalt	<1	Ingestion of soil	<1
		PCB-1248	1.5	Dermal contact with soil	<1		Copper	<1	Fluoride	<1	Dermal contact with soil	1.2
		PCB-1254	<1	External exposure to soil	20.3		Iron	8.8	Iron	8.8	Ingestion of vegetables from soil	23.8
		PCB-1260	<1	Ingestion of vegetables from soil	38.9		Manganese	4.3	Manganese	<1		
		Benzene	<1	Carbon tetrachloride	<1		Mercury	<1	Molybdenum	<1		
		Benz(a)anthracene	<1	Chloroform	<1		Nickel	<1	Nickel	<1		
		Benzo(a)pyrene	1.0	Chloromethane	<1		Nitrate-nitrite	<1	Nitrate-nitrite	<1		
		Benzo(b)fluoranthene	<1	Dibenz(o,a,h)anthracene	1.2		Uranium	15.0	Uranium	15.0		
		Bis(2ethylhexyl)phthalate	<1	Indeno(1,2,3-cd)pyrene	<1		Vanadium	<1	Vanadium	<1		
		Carbon tetrachloride	<1	Tetrachloroethene	<1		Zinc	<1	Zinc	<1		
		Chloroform	<1	TCE	3.3		1,2-cis-DCE	<1	1,2-cis-DCE	<1		
		Chloromethane	<1	Vinyl chloride	10.2		1,2-DCE (total)	39.0	1,2-DCE (total)	39.0		
		Dibenz(o,a,h)anthracene	<1	24 ^{Am}	<1		2,4-Dimethylphenol	<1	2,4-Dimethylphenol	<1		
		Indeno(1,2,3-cd)pyrene	1.2	237 ^{Np}	1.2		2-Methylphenol	<1	2-Methylphenol	<1		
		Tetrachloroethene	<1	238 ^{Pu}	<1		4-Methylphenol	<1	4-Methylphenol	<1		
		TCE	<1	238 ^{Rn}	<1		Acetone	<1	Acetone	<1		
Vinyl chloride	7.4	99 ^{Tc}	7.4	PCB-1254	14.8	PCB-1254	14.8					
24 ^{Am}	<1	230 Th	<1	Carbon tetrachloride	<1	Carbon tetrachloride	<1					
237 ^{Np}	2.2	233 ^U	2.2	TCE	4.3	TCE	4.3					
238 ^{Pu}	<1	235 ^U	<1									
238 ^{Rn}	<1	238 ^U	<1									
99 ^{Tc}	11.5	238 ^U	11.5									
230 Th	1.8	Arsenic	1.8	Ingestion of soil	25.6	5	Aluminum	5.0	Ingestion of soil	18.4		
233 ^U	42.5	Beryllium	42.5	Dermal contact with soil	43.8		Antimony	11.3	Dermal contact with soil	81.5		
235 ^U	<1	Benzo(a)pyrene	<1	External exposure to soil	32.5		Arsenic	3.4				
238 ^U	<1	Dibenzo(a,h)anthracene	1.7				Chromium	17.6				
238 ^U	<1	237 ^{Np}	<1				Copper	2.9				
238 ^U	3.4	238 ^{Pu}	<1				Iron	21.3				
238 ^U	9.4	238 ^U	9.4				Manganese	11.0				
238 ^U	<1	238 ^U	<1				Nickel	3.9				
238 ^U	41.3	238 ^U	41.3				Uranium	7.5				
238 ^U	<1	238 ^U	<1				Vanadium	10.9				
Future excavation worker at current concentrations	2 x 10 ⁻³	Arsenic	1.8	Ingestion of soil	25.6		5	Aluminum	5.0	Ingestion of soil	18.4	
		Beryllium	42.5	Dermal contact with soil	43.8			Antimony	11.3	Dermal contact with soil	81.5	
		Benzo(a)pyrene	<1	External exposure to soil	32.5			Arsenic	3.4			
		Dibenzo(a,h)anthracene	1.7					Chromium	17.6			
		237 ^{Np}	<1					Copper	2.9			
		238 ^{Pu}	<1					Iron	21.3			
		238 ^U	3.4					Manganese	11.0			
		238 ^U	9.4					Nickel	3.9			
		238 ^U	<1					Uranium	7.5			
		238 ^U	41.3					Vanadium	10.9			

Table taken from Executive Summary Remedial Investigation Report for Solid Waste Management Units 7 and 30 of Waste Area Grouping 22 at the Paducah Gaseous Diffusion Plant, Paducah, Kentucky DOE/OR-007-1604V2&D1 July 1997.
 NA = ELCR not applicable to child and teen cohorts. Values for adult include exposure as child and teen.
 NE = Land use scenario not evaluated or not of concern. Total ELCR and total HI columns reflect values from Tables 1.59 to 1.70 of DOE 1997 without lead included.
^a %Total HI for COCs and PCBs values do not include lead as a COC. Tables 1.59 to 1.70 of DOE 1997 present the systemic toxicity with lead included.

Table F2.17. Summary of Human Health Risk Characterization for SWMU 30 Without Lead as a COPC

Receptor	Total ELCR	ELCR COCs	% Total ELCR	ELCR POCs	% Total ELCR	Total HI ^a	Systematic COCs	% Total HI	Systematic Toxicity POCs	% Total HI
Current industrial worker at current concentrations	4 x 10 ⁻³	Arsenic Beryllium PCB-1260 Benz(a)anthracene Benzo(a)pyrene Benzo(b)fluoranthene Indeno(1,2,3-cd)pyrene 237Np 234U 235U 235/236U 238U	<1 97.3 <1 <1 <1 <1 <1 <1 <1 <1 <1 1.4	Ingestion of soil Dermal contact with soil External exposure to soil	<1 97.3 1.7	4	Aluminum Antimony Arsenic Beryllium Cadmium Chromium Iron Manganese Uranium Vanadium	5.1 3.7 2.7 10.7 3.5 13.5 19.8 11.3 9.0 17.6	Ingestion of soil Dermal contact with soil	2.9 97.0
Future industrial worker at current concentrations	4 x 10 ⁻³	Arsenic Beryllium 1,1-DCE PCB-1260 Benz(a)anthracene Benzo(a)pyrene Benzo(b)fluoranthene Bis(2ethylhexyl)phthalate Carbon tetrachloride Dibenzo(a,h)anthracene Indeno(1,2,3-cd)pyrene Tetrachloroethene TCE Vinyl chloride 241Am 237Np 239Pu 222Rn 99Tc 234U 235U 235/236U 238U	1.3 90.0 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 3.0 <1 <1 <1 <1 <1 2.5 <1 <1 <1 <1 1.4	Ingestion of groundwater Dermal contact with groundwater Inhalation while showering Ingestion of soil Dermal contact with soil External exposure to soil	4.0 <1 3.0 <1 90.0 1.6	12	Aluminum Antimony Arsenic Beryllium Cadmium Chromium Iron Manganese Nitrate Uranium Vanadium Carbon tetrachloride Di-n-octylphthalate TCE	2.0 1.4 2.7 4.0 1.3 6.5 7.7 4.3 1.9 3.7 7.1 1.3 19.0 34.2	Ingestion of groundwater Dermal contact with groundwater Ingestion of soil Dermal contact with soil	33.1 28.8 1.1 36.3
Future child recreational user at current concentrations	NA	NA	NA	NA	NA	0.04	NE	NE	NE	NE
Future teen recreational user at current concentrations	NA	NA	NA	NA	NA	0.04	NE	NE	NE	NE

Table F2.17. Summary of Human Health Risk Characterization for SWMU 30 Without Lead as a COPC (Continued)

Receptor	Total ELCR	ELCR COCs	% Total ELCR	ELCR POCs	% Total ELCR	Total HI ^a	Systematic Toxicity COCs	% Total HI	Systematic Toxicity POCs	% Total HI
Future adult recreational user at current concentrations	2 x 10 ⁻⁵	PCB-1260 Benzo(a)pyrene Dibenzo(a,h)anthracene	48.0 12.7 20.7	Ingestion of deer Ingestion of rabbit Ingestion of quail	8.7 80.0 11.3	0.04	NE	NE	NE	NE
Future child rural resident at current concentrations	NA	NA	NA	NA	NA	334	Aluminum Antimony Arsenic Barium Beryllium Cadmium Chromium Copper Fluoride Chromium VI Iron Manganese Mercury Nickel Nitrate Nitrate as nitrogen Selenium Uranium Vanadium Zinc 1,2-DCE (total) PCB-1254 Bis(2ethylhexyl)phthalate Carbon tetrachloride Di-n-octylphthalate TCE	3.3 <1 6.6 <1 1.4 1.7 3.0 <1 <1 <1 18.0 2.0 <1 <1 <1 70.7	Ingestion of groundwater Dermal contact with groundwater Inhalation while showering Inhalation from household use Ingestion of vegetables from groundwater Ingestion of soil Dermal contact with soil Ingestion of vegetables from soil	8.0 2.7 <1 1.3 8.7 1.0 7.4 70.7

Table F2.17. Summary of Human Health Risk Characterization for SWMU 30 Without Lead as a COPC (Continued)

Receptor	Total ELCR	ELCR COCs	% Total ELCR	ELCR POCs	% Total ELCR	Total HF ^a	Systematic Toxicity COCs	% Total HI	Systematic Toxicity POCs	% Total HI
Future adult rural resident at current concentrations	4 x 10 ⁻²	Arsenic Beryllium 1,1,2-TCA 1,1-DCE PCB-1254 PCB-1260 Benz(a)anthracene Benzo(a)pyrene Benzo(b)fluoranthene Benzo(k)fluoranthene Bis(2ethylhexyl)phthalate Carbon tetrachloride Chloroform Chrysene Dibenzo(a,h)anthracene Indeno(1,2,3-cd)pyrene Tetrachloroethene TCE ²⁴¹ Am ²³⁷ Np ²³⁹ Pu ²²⁶ Ra ²²² Rn ⁹⁹ Tc ²³⁰ Th ²³⁴ U ²³⁵ U ^{235/236} U ²³⁸ U	5.5 47.7 <1 <1 <1 1.3 <1 3.2 <1 <1 <1 <1 <1 <1 <1 <1 1.3 <1 3.9 <1 <1 <1 <1 21.4 3.2 <1 <1 8.4	Ingestion of groundwater Dermal contact with groundwater Inhalation while showering Inhalation from household use Ingestion of vegetables from groundwater Ingestion of soil Dermal contact with soil External exposure to soil Ingestion of vegetables from soil	1.8 <1 <1 2.1 22.7 <1 25.0 <1 45.5	105	Aluminum Antimony Arsenic Barium Beryllium Cadmium Chromium Copper Iron Manganese Mercury Nickel Nitrate Uranium Vanadium Zinc 1,2-DCE (total) PCB-1254 Carbon tetrachloride Di-n-octylphthalate TCE	3.1 <1 6.7 <1 1.1 1.7 2.8 <1 17.3 1.6 <1 <1 <1 35.9 2.0 <1 <1 2.0 1.3 3.0 16.9	Ingestion of groundwater Dermal contact with groundwater Inhalation from household use Ingestion of vegetables from groundwater Ingestion of soil Dermal contact with soil Ingestion of vegetables from soil	10.4 4.6 <1 8.8 <1 4.6 70.4

Table F2.17. Summary of Human Health Risk Characterization for SWMU 30 Without Lead as a COPC (Continued)

Receptor	Total ELCR	ELCR COCs	% Total ELCR	ELCR POCs	% Total ELCR	Total HI ^a	Systematic TOXCOCs	% Total HI	Systematic Toxicity POCs	% Total HI	
Future excavation worker at current concentrations	1 x 10 ⁻³	Arsenic	2.0	Ingestion of soil	6.3	4	Aluminum	5.0	Ingestion of soil	20.0	
		Beryllium	91.7	Dermal contact with soil	91.7		Antimony	6.8	Dermal contact with soil		80.0
		PCB-1248	<1	External exposure to soil	3.3		Arsenic	3.6			
		Benz(a)anthracene	<1				Beryllium	3.4			
		Benzo(a)pyrene	<1				Cadmium	3.2			
		Benzo(b)fluoranthene	<1				Chromium	11.1			
		Dibenzo(a,h)anthracene	<1				Iron	21.5			
		Indeno(1,2,3-cd)pyrene	<1				Manganese	15.6			
		²³⁷ Np	<1				Uranium	13.3			
		²³⁹ Pu	<1				Vanadium	13.8			
		²³⁴ U	<1								
		²³⁵ U	<1								
		^{235/236} U	<1								
²³⁸ U	4.1										

Table taken from Executive Summary Remedial Investigation Report for Solid Waste Management Units 7 and 30 of Waste Area Grouping 22 at the Paducah Gaseous Diffusion Plant, Paducah, Kentucky DOE/OR/07-1604/V2&DI July 1997

NA = ELCR not applicable to child and teen cohorts. Values for adult include exposure as child and teen.

NE = Land use scenario no evaluated or not of concern. Total ELCR and total HI columns reflect values from Tables 1.59 to 1.70 of DOE 1997 without lead included.

^a %Total HI for COCs and PCBs values do not include lead as a COC. Tables 1.59 to 1.70 of DOE 1997 present the systemic toxicity with lead included.

Table F2.18. Summary of Human Health Risk Characterization for the North Ditch Without Lead as a COPC

Receptor	Total ELCR	ELCR COCs	% Total ELCR	ELCR POCs	% Total ELCR	Total HI ^a	Systematic COCs	% Total HI	Systematic Toxicity POCs	% Total HI
Current industrial worker at current concentrations	4 x 10 ⁻⁴	Arsenic Beryllium PCB-1260 235/236U 238U	7.6 84.2 <1 4.5	Ingestion of soil Dermal contact with soil External exposure to soil	1.0 92.1 5.0	5	Aluminum Antimony Arsenic Chromium Iron Manganese Uranium Vanadium	6.6 5.7 3.7 14.4 24.9 15.3 3.2 22.5	Dermal contact with soil	98.8
Future industrial worker at current concentrations	4 x 10 ⁻⁴	Arsenic Beryllium PCB-1260 235/236U 238U	7.6 84.2 1.2 8.2 4.5	Ingestion of soil Dermal contact with soil External exposure to soil	1.0 92.1 5.0	5	Aluminum Antimony Arsenic Chromium Iron Manganese Uranium Vanadium	6.6 5.7 3.7 14.4 24.9 15.3 3.2 22.5	Dermal contact with soil	98.8
Future child recreational user at current concentrations	NA	NA	NA	NA	NA	0.004	NE	NE	NE	NE
Future teen recreational user at current concentrations	NA	NA	NA	NA	NA	0.003	NE	NE	NE	NE
Future adult recreational user at current concentrations	1 x 10 ⁻⁶	NE	NE	NE	NE	0.004	NE	NE	NE	NE
Future child rural resident at current concentrations	NA	NA	NA	NA	NA	229	Aluminum Antimony Arsenic Barium Beryllium Cadmium Chromium Copper Iron Manganese Nickel Uranium Vanadium Zinc	7.0 1.9 14.1 1.1 <1 <1 4.4 <1 36.2 4.3 <1 23.4 4.7 <1	Ingestion of soil Dermal contact with soil Ingestion of vegetables from soil	1.3 12.3 86.4

Table F2.18. Summary of Human Health Risk Characterization for the North Ditch Without Lead as a COPC (Continued)

Receptor	Total ELCR	ELCR COCs	% Total ELCR	ELCR POCs	% Total ELCR	Total HI ^a	Systematic COCs	% Total HI	Systematic Toxicity POCs	% Total HI
Future adult rural resident at current concentrations	9 x 10 ⁻³	Arsenic Beryllium PCB-1260 ²³⁷ Np ²³⁴ U ^{235/236} U ²³⁸ U	37.8 21.1 9.7 <1 5.0 1.1 24.4	Ingestion of soil Dermal contact with soil External exposure to soil Ingestion of vegetables from soil	<1 11.1 2.6 84.4	68	Aluminum Antimony Arsenic Barium Beryllium Cadmium Chromium Copper Iron Manganese Nickel Uranium Vanadium Zinc	7.0 1.6 14.6 1.1 <1 <1 4.0 <1 37.3 3.8 1.0 23.8 3.8 <1	Ingestion of soil Dermal contact with soil Ingestion of vegetables from soil	<1 7.9 91.6
Future excavation worker at current concentrations	NE	NE	NE	NE	NE	NE	NE	NE	NE	NE

Table taken from Executive Summary Remedial Investigation Report for Solid Waste Management Units 7 and 30 of Waste Area Grouping 22 at the Paducah Gaseous Diffusion Plant, Paducah, Kentucky DOE/OR/07-1604/V2&DI July 1997

NA = ELCR not applicable to child and teen cohorts. Values for adult include exposure as child and teen.

NE = Land use scenario no evaluated or not of concern. Total ELCR and total HI columns reflect values from Tables 1.59 to 1.70 of DOE 1997 without lead included.

^a %Total HI for COCs and PCBs values do not include lead as a COC. Tables 1.59 to 1.70 of DOE 1997 present the systemic toxicity with lead included.

Table F2.19. Summary of Human Health Risk Characterization for the South Ditch Without Lead as a COPC

Receptor	Total ELCR	ELCR COCs	% Total ELCR	ELCR POCs	% Total ELCR	Total HI ^a	Systematic COCs	% Total HI	Systematic Toxicity POCs	% Total HI
Current industrial worker at current concentrations	4 x 10 ⁻⁴	Arsenic Beryllium PCB-1260 Benz(a)anthracene Benzo(a)pyrene Benzo(b)fluoranthene Dibenzo(a,h)anthracene e Indeno(1,2,3-cd)pyrene 237Np 235/236U 238U	4.7 58.3 1.9 1.3 16.9 3.1 2.3 1.0 <1 2.2 6.9	Ingestion of soil Dermal contact with soil External exposure to soil	1.7 88.9 9.2	5	Aluminum Antimony Arsenic Cadmium Chromium Iron Manganese Nickel Uranium Vanadium	6.9 16.0 2.1 2.4 10.7 17.3 9.8 4.6 9.2 17.6	Dermal contact with soil	98.2
Future industrial worker at current concentrations	4 x 10 ⁻⁴	Arsenic Beryllium PCB-1260 Benz(a)anthracene Benzo(a)pyrene Benzo(b)fluoranthene Dibenzo(a,h)anthracene e Indeno(1,2,3-cd)pyrene 237Np 235/236U 238U	4.7 58.3 1.9 1.3 16.9 3.1 2.3 1.0 <1 2.2 6.9	Ingestion of soil Dermal contact with soil External exposure to soil	1.7 88.9 9.2	5	Aluminum Antimony Arsenic Cadmium Chromium Iron Manganese Nickel Uranium Vanadium	6.9 16.0 2.1 2.4 10.7 17.3 9.8 4.6 9.2 17.6	Dermal contact with soil	98.2
Future child recreational user at current concentrations	NA	NA	NA	NA	NA	0.005	NE	NE	NE	NE
Future teen recreational user at current concentrations	NA	NA	NA	NA	NA	0.005	NE	NE	NE	NE
Future adult recreational user at current concentrations	2 x 10 ⁻⁶	PCB-1260	66.7	Ingestion of rabbit	83.3	0.006	NE	NE	NE	NE

Table F2.19. Summary of Human Health Risk Characterization for the South Ditch Without Lead as a COPC (Continued)

Receptor	Total ELCR	ELCR COCs	% Total ELCR	ELCR POCs	% Total ELCR	Total HI ^a	Systematic Toxicity COCs	% Total HI	Systematic Toxicity POCs	% Total HI	
Future child rural resident at current concentrations	NA	NA	NA	NA	NA	334	Aluminum Antimony Arsenic Barium Beryllium Cadmium Chromium Copper Iron Manganese Nickel Uranium Vanadium Zinc PCB-1260	4.9 3.5 5.6 <1 <1 1.4 2.3 1.6 17.9 1.9 <1 9.5 46.4 2.7 <1	Ingestion of soil Dermal contact with soil Ingestion of vegetables from soil	1.3 8.5 90.1	
Future adult rural resident at current concentrations	1 x 10 ⁻²	Arsenic Beryllium PCB-1016 PCB-1260 Benz(a)anthracene Benzo(a)pyrene Benzo(b)fluoranthene Benzo(k)fluoranthene Bis(2-ethylhexyl)-phthalate Chrysene Dibenzo(a,h)anthracene Indeno(1,2,3-cd)pyrene ²³⁷ Np ²³⁴ U ^{235/236} U ²³⁸ U	14.2 8.6 <1 10.0 1.6 20.7 3.8 <1 <1 <1 2.8 1.2 1.1 8.57 1.8 23.6	Ingestion of soil Dermal contact with soil External exposure to soil Ingestion of vegetables from soil	<1 6.7 2.9 85.7	101	Aluminum Antimony Arsenic Barium Cadmium Chromium Copper Iron Manganese Mercury Nickel Uranium Vanadium Zinc PCB-1016	5.0 3.1 5.8 <1 1.3 2.0 1.6 17.8 1.6 <1 9.8 47.2 2.2 1.4 <1	Ingestion of soil Dermal contact with soil Ingestion of vegetables from soil	<1 5.5 94.2	
Future excavation worker at current concentrations	NE	NE	NE	NE	NE	NE	NE	NE	NE	NE	NE

Table taken from Executive Summary Remedial Investigation Report for Solid Waste Management Units 7 and 30 of Waste Area Grouping 22 at the Paducah Gaseous Diffusion Plant, Paducah, Kentucky DOE/OR/07-1604/V2&D1 July 1997

NA = ELCR not applicable to child and teen cohorts. Values for adult include exposure as child and teen.

NE = Land use scenario no evaluated or not of concern. Total ELCR and total HI columns reflect values from Tables 1.59 to 1.70 of DOE 1997 without lead included.

^a % Total HI for COCs and PCBs values do not include lead as a COC. Tables 1.59 to 1.70 of DOE 1997 present the systemic toxicity with lead included.

Table F2.20. Summary Human Health Risk Characterization for Future Risk at Future Modeled Concentrations for SWMUs 7 and 30 and Associated Ditches

Receptor	Total ELCR	ELCR COCs	% Total ELCR	ELCR POCs	% Total ELCR	Total HI	Systematic Toxicity COCs	% Total HI	Systematic Toxicity POCs	% Total HI
Future child rural resident at future modeled concentrations – 30 years	NA	NA	NA	NA	NA	0.08	NE	NE	NE	NE
Future child rural resident at future modeled concentrations – 100 years	NA	NA	NA	NA	NA	0.3	NE	NE	NE	NE
Future adult rural resident at future modeled concentrations – 30 years	5×10^{-5}	Vinyl chloride ⁹⁹ Tc	7.5 91.7	Ingestion of groundwater Inhalation from household use	95.8 3.75	0.03	NE	NE	NE	NE
Future adult rural resident at future modeled concentrations – 100 years	2×10^{-4}	Vinyl chloride ⁹⁹ Tc	7.0 90.0	Ingestion of groundwater Inhalation from household use	95.0 2.75	0.1	NE	NE	NE	NE

Table taken from Executive Summary Remedial Investigation Report for Solid Waste Management Units 7 and 30 of Waste Area Grouping 22 at the Paducah Gaseous Diffusion Plant, Paducah, Kentucky DOE/OR/07-1604/V2&D1 July 1997

NA = ELCR not applicable to child and teen cohorts. Values for adult include exposure as child and teen.

NE = Land use scenario no evaluated or not of concern. Total ELCR and total HI columns reflect values from Tables 1.59 to 1.70 of DOE 1997 without lead included.

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APPENDIX G
REVIEW OF ECOLOGICAL
RISK ASSESSMENTS

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ACRONYMS

BGOU	Burial Grounds Operable Unit
bgs	below ground surface
BRA	baseline risk assessment
COPC	contaminant of potential concern
CSM	conceptual site model
DNAPL	dense nonaqueous phase liquid
DOE	U.S. Department of Energy
EPC	exposure point concentration
ERA	ecological risk assessment
HQ	hazard quotient
HI	hazard index
NFA	no further action
NOAEL	No Observed Adverse Effects Level
PAH	polyaromatic hydrocarbon
PCB	polychlorinated biphenyl
PGDP	Paducah Gaseous Diffusion Plant
PRG	preliminary remediation goal
RCRA	Resource Conservation and Recovery Act
RGA	Regional Gravel Aquifer
RI	remedial investigation
RGO	remedial goal option
SFF	site foraging factor
SI	site investigation
SVOC	semivolatile organic compound
SWOU	surface water operable unit
SWMU	solid waste management unit
⁹⁹ Tc	technetium-99
TCE	trichloroethene
UCL	upper confidence limit
VOC	volatile organic compound
WAG	waste area group
WKWMA	West Kentucky Wildlife Management Area

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G.1. INTRODUCTION

G.1.1 SITE LOCATION

This appendix provides summaries of the results of ecological risk assessments (ERAs) previously completed for Solid Waste Management Units (SWMUs) 2, 4, 5, 6, 7, and 30 within the Burial Grounds Operable Unit (BGOU) of Paducah Gaseous Diffusion Plant (PGDP) (Figure G.1). SWMUs 3 and 145 are not included because SWMU 3 is covered by a RCRA cap and SWMU 145 is sited on 44 acres that now lie beneath the C-746-S&T Landfills. SWMUs 2, 3, 4, 5, 6, 7, and 30 are located within the developed area of the PGDP facility. SWMUs 2, 3, and 4 are in the west-central area of the plant inside the security fence-lined area. SWMU 4 is bounded on all sides by plant roads and an active railroad spur (see Section 1.3 of the main text). SWMUs 5, 6, 7, and 30 are in the northwestern section of the PGDP secured area. Some of the area surrounding the PGDP facility is a recreational wildlife area, the West Kentucky Wildlife Management Area (WKWMA), with residential areas lying beyond the WKWMA. Private land in rural residential and agricultural areas also borders the PGDP facility.

G.1.2 SITE HISTORY

All the SWMUs considered in the ERAs originally were burial pits or landfills for process wastes from PGDP. The individual waste streams, burial practices, and operating time frames for each SWMU are described in-depth in Section 1.3 of this Remedial Investigation (RI) Report, and that material is briefly summarized here.

SWMU 2 was used primarily for the disposal of uranium metal pieces, uranium oxides, oils [that may have contained polychlorinated biphenyls (PCBs)], trichloroethene (TCE), and uranyl fluoride. Some technetium-99 (⁹⁹Tc) also may be present at this SWMU associated with the uranium wastes. SWMU 3 was an aboveground earth and clay surface impoundment with an overflow weir that subsequently was converted for disposal of solid uranium-contaminated wastes (uranium metal, uranium oxides, smelter furnace liners, and radioactively contaminated trash). SWMU 3 also includes an adjacent ditch that carried leachate from the surface impoundment. The landfill ceased operation in 1986 and was covered with a Resource Conservation and Recovery Act (RCRA) multilayered cap. SWMU 4 is an open vegetated field that was used to bury wastes in designated burial cells. This SWMU may have received uranium- and ⁹⁹Tc-contaminated sludge as well as TCE. SWMU 4 was covered in 1982 with 2 to 3 ft of soil material and a 6-inch clay cap. SWMU 5 contained disposal cells laid out in a grid pattern. Slag from nickel and aluminum smelters was disposed of here along with radioisotopes. Waste cells in SWMU 5 were covered with 2 to 3 ft of soil after they were filled. SWMU 6 also was divided into discrete burial cells: two for magnesium scrap, one for exhaust fan hoods contaminated with perchloric acid (see Section 2.5.4 of the main text for further discussion of the instability of perchloric acid), one for contaminated aluminum scrap, and one area for a single contaminated modine trap. SWMU 7 contains burial pits that were used for disposal of noncombustible, uranium-contaminated and uncontaminated trash, material, and equipment. SWMU 30 contained an area used for burning combustible trash that may have contained uranium contamination. The ash and debris from the incineration were buried below ground in a pit at SWMU 30. SWMU 145 contains landfills used to discard scrap and waste materials, including construction debris.

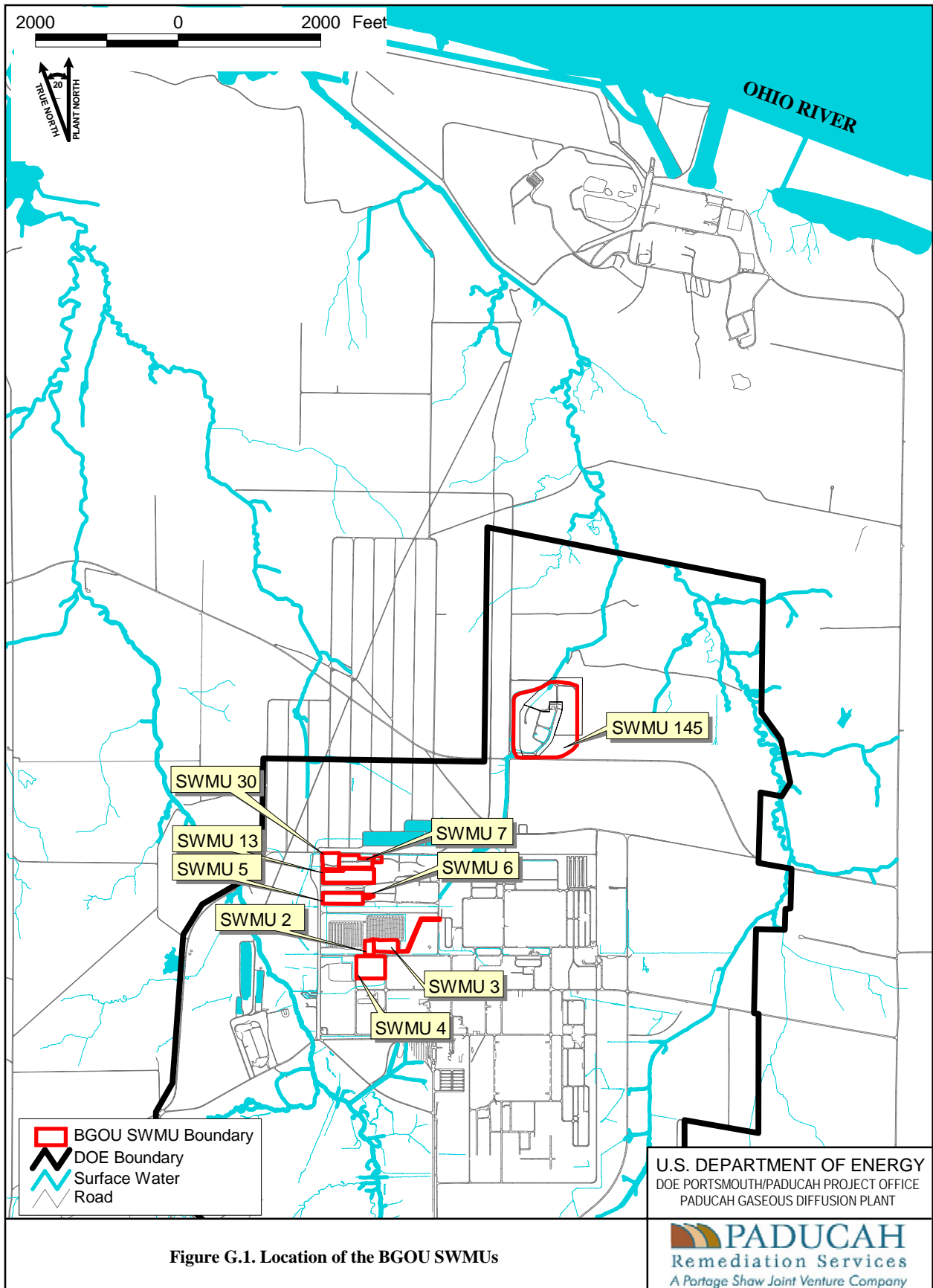


Figure G.1. Location of the BGOU SWMUs

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Trends in TCE concentration in soil and groundwater suggest dense nonaqueous-phase liquid (DNAPL) is present at SWMUs 4 and the area joining SWMUs 7 and 30. Some of the burial ground SWMUs ceased operation in the 1950s, but others were in use as late as the 1980s. After each unit ceased to be used for waste burial, it was covered with a soil cap. SWMU 3 was closed with the addition of a RCRA cap in 1987. Previous sampling identified a groundwater plume of dissolved-phase TCE in the subsurface under most of these SWMUs.

G.2. PROBLEM FORMULATION

The first step in an ERA includes the problem formulation. This step encompasses development of the preliminary conceptual site model (CSM), determination of potentially complete exposure pathways and potentially contaminated media, selection of exposure endpoints, and selection of screening levels protective of the endpoints and potentially exposed receptors at the site. The problem formulation presented here is consistent with those presented in the original ERAs that are summarized in this appendix.

G.2.1 PRELIMINARY CONCEPTUAL SITE MODEL

The preliminary CSM includes a description of the environmental setting, known site contaminants, and a figure representing the potential exposure pathways. This preliminary CSM is used as the basis for selection of benchmark values used to screen the site for potential ecological risk.

G.2.1.1 Site Environmental Setting and Habitat Descriptions

The SWMUs included in this ERA are generally similar in topography and process history. All the SWMUs, except SWMU, 3 originally served as burial grounds or landfills for process wastes from PGDP operations. Once the pits within the burial grounds no longer were being used, they were topped with soil covers and most sites were revegetated. SWMU 3, however, was covered with a RCRA cap. Although there is potential for contamination below the surface to migrate laterally toward surface water, the direction of shallow groundwater flow is primarily downward and represents limited risks to terrestrial receptors near these sites. Figures G.2 to G.9 show the surface conditions at the SWMUs considered in this report.

The terrestrial ecosystems occurring in the area of these SWMUs are described more fully in the work plan for the BGOU (DOE 2006). This section presents a brief summary of the ecosystem relevant to defining the CSM and exposure pathways. The primary ecosystem in the area outside the industrial area around the SWMUs is upland grassland interspersed with developed industrial areas. The vegetation over these SWMUs is maintained with routine mowing (see section 3.1) approximately eight times per year. Most of the SWMUs also are surrounded by fencing and/or roads. The buffer area and areas bordering the PGDP facility include forest, thickets, and agricultural land. Much of the PGDP facility is surrounded by the WKWMA, which includes managed native prairie and deciduous forest. Species documented to occur in the area include numerous small mammals, particularly shrews, mice, and voles. Numerous bird species, including doves, turkey, quail, bluebirds and other songbirds, as well as hawks and owls, are found in this area. There also are amphibians, reptiles (primarily lizards and turtles), and bats. Table G.1 lists species observed in the nonindustrial areas of the PGDP and at the adjacent WKWMA.



Figure G.2. Surface of SWMU 2



Figure G.3. Surface of SWMU 3



Figure G.4. Surface of SWMU 4 (Area behind Fence)



Figure G.5. Surface of SWMU 5 (Area behind Fence)



Figure G.6. Surface of SWMU 6



Figure G.7. Surface of SWMU 7 (Field with Tree Surrounded by Road)



Figure G.8. SWMU 30 (Foreground, SWMU 7 is in Upper Left)

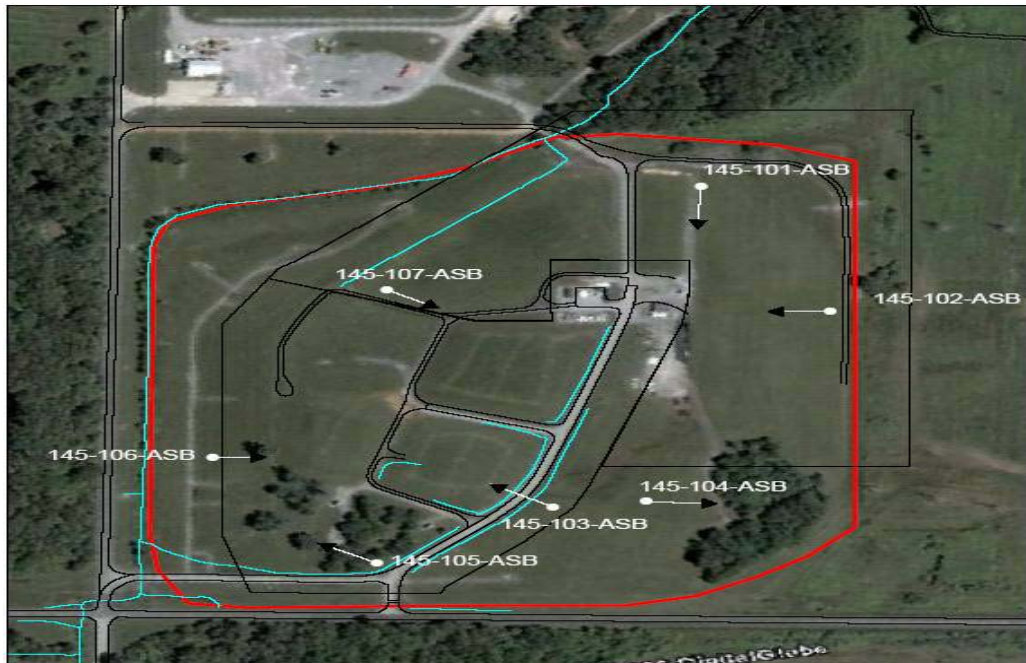


Figure G.9. Aerial View of Landfills at SWMU 145

Table G.1. Wildlife Species Present or Potentially Present at the PGDP Site^a

Common Name	Scientific Name
<i>Fish</i>	
Black Buffalo	<i>Ictiobus niger</i>
Blackspotted Topminnow	<i>Fundulus olivaceus</i>
Creek Chub	<i>Semotilus atromaculatus</i>
Bluegill sunfish	<i>Lepomis macrochirus</i>
Green sunfish	<i>Lepomis cyanellus</i>
Redspotted Sunfish	<i>Lepomis miniatus</i>
Largemouth bass	<i>Micropterus salmoides</i>
Longear sunfish	<i>Lepomis megalotis</i>
Stoneroller	<i>Campostoma sp.</i>
<i>Reptiles and Amphibians</i>	
American Toad	<i>Bufo americanus</i>
Bull frog	<i>Rana catesbeiana</i>
Eastern box turtle	<i>Terrapene carolina</i>
Leopard frog	<i>Rana sphenocephala</i>
Salamanders	Various species
Snakes	Various species
Green Treefrog	<i>Hyla cinerea</i>
Woodhouse toad	<i>Bufo woodhousei</i>
Northern crawfish frog	<i>Rana areolata circulosa</i>
Green frog	<i>Rana clamitans melanota</i>
	<i>Pseudacris triseriata</i>
Upland chorus frog	<i>ferriarum</i>
<i>Birds</i>	
American robin	<i>Turdus migratorius</i>
American woodcock	<i>Scolopax minor</i>
Bald eagle	<i>Haliaeetus leucocephalus</i>
Barred owl	<i>Strix varia</i>
Belted kingfisher	<i>Ceryle alcyon</i>
Blue jay	<i>Cyanocitta cristata</i>
Blue-winged teal	<i>Anas discors</i>
Canada goose	<i>Branta canadensis</i>
Coot	<i>Fulica americana</i>
American Crow	<i>Corvus brachyrhynchos</i>
Downy woodpecker	<i>Picoides pubescens</i>
Eastern bluebird	<i>Sialis sialis</i>
Eastern kingbird	<i>Tyrannus tyrannus</i>
Eastern meadowlark	<i>Sturnella magna</i>
Eastern phoebe	<i>Sayornis phoebe</i>

Table G.1. Wildlife Species Present or Potentially Present at the PGDP Site^a (Continued)

Common Name	Scientific Name
<i>Bird (Continued)</i>	
Eastern wood pewee	<i>Contopus virens</i>
Gadwall duck	<i>Anas strepera</i>
Great Blue Heron	<i>Ardea herodias</i>
Great Crested Flycatcher	<i>Myiarchus crinitus</i>
Great-horned owl	<i>Bubo virginianus</i>
Hairy woodpecker	<i>Picoides villosus</i>
Hawks	Various species
Hérons and egrets	Various species
Killdeer	<i>Charadrius vociferus</i>
Loggerhead shrike	<i>Lanius ludovicianus</i>
Mallard duck	<i>Anas platyrhynchos</i>
Mourning dove	<i>Zenaida macroura</i>
Northern bobwhite (aka bobwhite quail)	<i>Colinus virginianus</i>
Northern cardinal	<i>Cardinalis cardinalis</i>
Northern flicker	<i>Colaptes auratus</i>
Pileated woodpecker	<i>Dryocopus pileatus</i>
Red-bellied woodpecker	<i>Melanerpes erythrocephalus</i>
Red-shouldered hawk	<i>Buteo lineatus</i>
Red-tailed hawk	<i>Buteo jamaicensis</i>
Red-winged blackbird	<i>Agelaius phoeniceus</i>
Ruby-throated hummingbird	<i>Archilochus colubris</i>
Screech owl	<i>Megascops asio</i>
Song sparrow	<i>Melospiza melodia</i>
Swallows	Various species
vireos	Various vireo sp.
Tufted titmouse	<i>Baeolophus bicolor</i>
Turkey Vulture	<i>Cathartes aura</i>
Warblers	Various species
Chuck-will's widow	<i>Caprimulgus carolinensis</i>
White-breasted nuthatch	<i>Sitta carolinensis</i>
Whip-poor-will	<i>Caprimulgus vociferous</i>
Wild turkey	<i>Meleagris gallopavo</i>
Wood cock	<i>Scolopax minor</i>
Wood duck	<i>Aix sponsa</i>
Wrens	Various species
Yellow-billed cuckoo	<i>Coccyzus americanus</i>

Table G.1. Wildlife Species Present or Potentially Present at the PGDP Site (Continued)

Common Name	Scientific Name
<i>Mammals</i>	
American beaver	<i>Castor canadensis</i>
American mink (aka mink)	<i>Mustela vison</i>
Bobcat	<i>Lynx rufus</i>
Common muskrat	<i>Ondatra zibethicus</i>
Coyote	<i>Canis latrans</i>
Eastern cottontail	<i>Sylvilagus floridanus</i>
Eastern grey squirrel and fox squirrel	<i>Sciurus carolinensis</i>
Evening bat	<i>Nycticeius humeralis</i>
Groundhog	<i>Marmota monax</i>
Indiana bat	<i>Myotis sodalists</i>
Mice	Various species
Moles	Various species
Opposum	<i>Didelphis virginiana</i>
Raccoon	<i>Procyon lotor</i>
Red fox	<i>Vulpes vulpes</i>
Grey fox	<i>Urocyon cinereoargenteus</i>
Shrews	Various species
Skunk	<i>Mephitis mephitis</i>
Southeastern myotis bat	<i>Myotis sodalis</i>
Voles	Various species
White-tailed deer	<i>Odocoileus virginianus</i>

^aThe listed species are from the Surface Water Operable Unit Report (DOE 2008) and the WKWMA species information website (<http://fw.ky.gov/kfwis/arcims/WmaSpecies.asp?strID=137>)

A number of state and federal listed, threatened, and endangered species may be present on the buffer areas within PGDP and the surrounding WKWMA land, though they are unlikely to be found on the maintained surface within the SWMUs. These species are listed in Table G.2 of this document. As noted in the footnote to Table G.2, none of the species listed in the table have been reported as sighted on the U.S. Department of Energy (DOE) Reservation.

Table G.2. Federally Listed, Proposed, and Candidate Species Potentially Occurring within the Paducah Site Study Area^a

Common Name	Scientific Name	Animal Type	Endangered Species Act Status
Indiana bat ^b	<i>Myotis sodalists</i>	Mammal	Listed endangered
Interior least tern	<i>Sterna antillarum athalassos</i>	Bird	Listed endangered
Pink mucket	<i>Lampsilis abrupta</i>	Mussel	Listed endangered
Ring pink	<i>Obovaria retusa</i>	Mussel	Listed endangered
Orangefoot pimpleback	<i>Plethobasus cooperianus</i>	Mussel	Listed endangered
Fat pocketbook	<i>Potamilus capax</i>	Mussel	Listed endangered

^a All of the listed species are discussed in *Environmental Investigations at the Paducah Gaseous Diffusion Plant and Surrounding Area, McCracken County, Kentucky, Volume III*, COE Nashville District, May 1994. Note that the area evaluated in the referenced report encompasses 11,719 acres and extends to include the Ohio River, which is over three miles north of the DOE Reservation. None of these species have been reported as sighted on the DOE Reservation, although potential summer habitat exists there for the Indiana bat. No critical habitat for any of these species has been designated anywhere in the area.

^b Specimens of the Indiana bat were collected from WKWMA property in 1991 and 1999.

G.2.1.2 Existing Data

The dataset for surface soils used in the ERAs summarized in this appendix is described in the RI documents containing the original assessments (DOE 1994; DOE 1997; DOE 1998; and DOE 2000). No new surface soil samples have been collected since the original assessments were done, except at SWMUs 3 and 7. The uncertainty section discusses the results and potential impact of new surface soil samples.

G.2.1.3 Site Contaminants

Only surface soil contaminants at the SWMUs were considered in the ERAs summarized in this appendix. Site contaminants at all SWMUs included inorganic chemicals, organic chemicals, and radionuclides.

G.2.1.4 Fate and Transport Mechanisms

Potential migration pathways for contaminants from waste and soil at the BGOU include transport of contaminated surface soil off-site by surface water, migration of contaminants to the subsurface soil, migration to groundwater, and uptake of soil contaminants through the on-site food chain. In addition, subsurface contaminants may be brought to the surface through bioturbation by burrowing animals or uptake by vegetation on the site. The surface soils at most of the BGOU SWMUs considered here are held in place by vegetation or, for SWMU 3, by the presence of a RCRA cap with a vegetative cover. Transport of surface soil off-site is likely to be minimal. Migration of contaminants to subsurface soil and through subsurface soil to groundwater is likely to occur. Contaminants in groundwater may be discharged to surface water at areas away from the BGOU SWMUs. Contaminants in surface soil are likely to be taken up into plants and soil invertebrates at these sites and would enter higher trophic level organisms through the food chain.

G.2.2 POTENTIALLY COMPLETE EXPOSURE PATHWAYS

Only surface samples (0–1 ft) were included in the assessments summarized in this appendix. Subsurface contamination may be accessible to burrowing ecological receptors. The subsurface interval of interest for this type of exposure would be in the 1–5 ft bgs depth. Insufficient analytical data are available in this depth range at these SWMUs to conduct a quantitative screening of potential exposure to burrowing animals. Burrowing animals also could encounter buried waste, but insufficient characterization is available for the waste to conduct a screening for exposure to materials in waste. Potential ecological risk from exposure to subsurface soil and waste to burrowing animals therefore is addressed qualitatively in the uncertainty section of this appendix. Significant contaminant transport through runoff directly to surface water is unlikely because most of the sites have vegetated surfaces. The pathways through which receptors could contact contaminants in surface soil include direct ingestion of soil, ingestion of plant or animals from the site as food, external exposure to ionizing radiation, and dermal contact with soil or surface water. A CSM representative of the CSMs from the past assessments and reflective of current site conditions is shown in Figure G.10.

G.2.3 POTENTIALLY CONTAMINATED MEDIA

Potential sources within the pits in the burial grounds may have contaminated both surface and subsurface soil. Only surface samples (0-1 ft) were included in the assessments described here. Subsurface contamination and buried waste are addressed qualitatively in the uncertainty section.

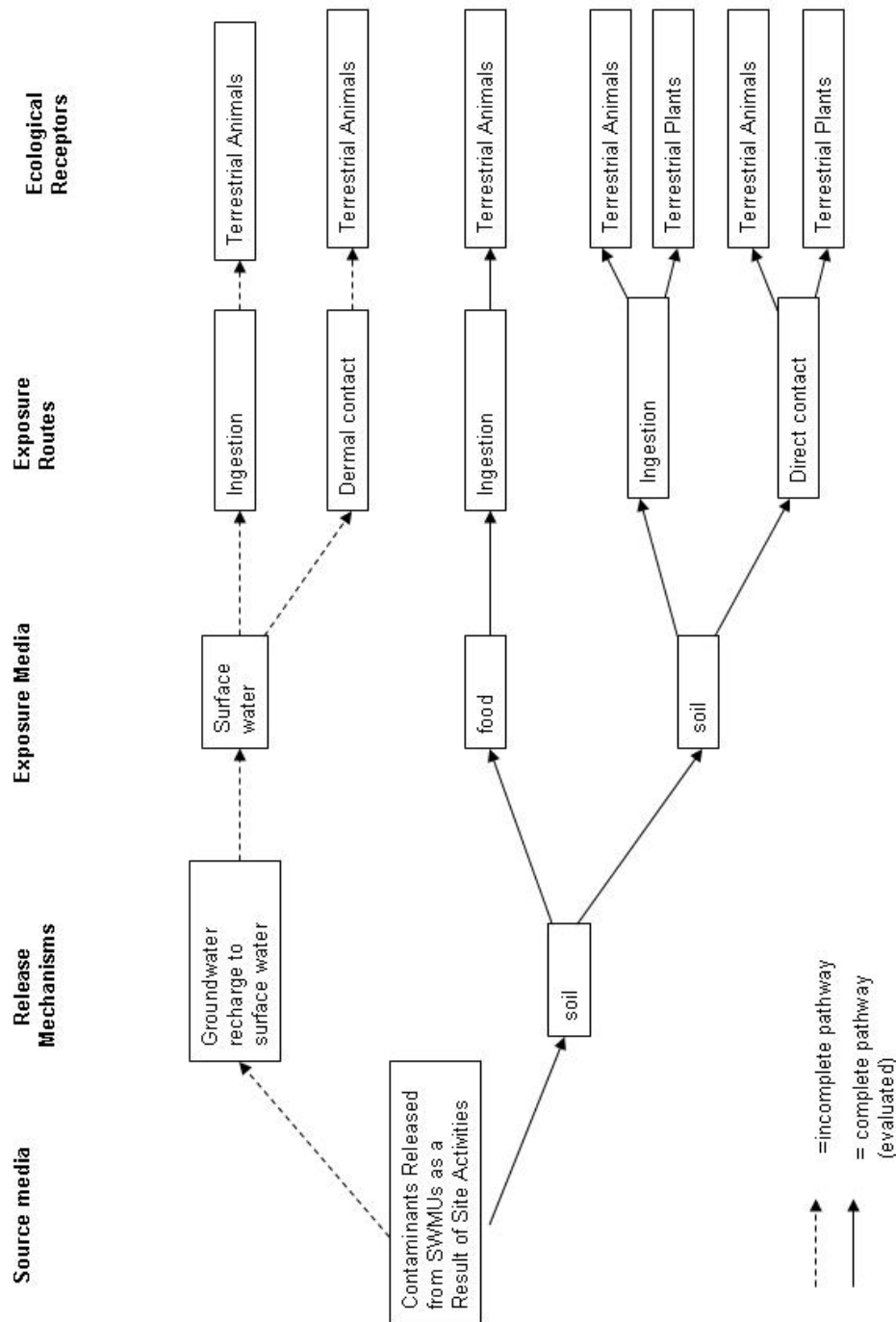


Figure G.10. Preliminary Conceptual Site Model for BGOU SWMUs

G.3. SCREENING-LEVEL EFFECTS EVALUATION

The ERAs summarized in this appendix all were completed prior to the development of the no further action (NFA) levels provided in Appendix A of the PGDP Ecological Risk Assessment methods document (DOE 2001). For SWMU 2, the maximum detected concentration of each potential contaminant was used as the exposure point concentration (EPC) and compared to a single ecological screening level selected from the literature. For SWMUs 4, 5, 6, 7, and 30, maximum detected concentrations of potential contaminants were used as the EPC and compared to screening levels developed for plant and soil receptors. For wildlife receptors, which are more mobile, the upper confidence limit (UCL) concentrations of potential contaminants were used as the EPC and compared to screening levels for each of those receptors unless the UCL exceeded the maximum detected concentration; in this case, the maximum detected concentration was used.

G.4. SUMMARY OF PREVIOUSLY CONDUCTED ERAS

For the ecological risk characterization for soil, the results of the previous ERAs are summarized here. For most of the SWMUs, no new surface data have been collected since the previous risk assessments were performed. In addition, the soils at some or all of the units are outside the scope of the BGOU RI as defined in the approved work plan (DOE 2006); therefore, a new quantitative risk assessment was not performed for soils. Previous ERAs were conducted for SWMUs 2, 4, 5, 6, 7, and 30 at PGDP. SWMU 3 is covered with a RCRA cap, and SWMU 145 lies beneath a landfill closed under RCRA; therefore, surface soil risks at these two units do not require further evaluation for ecological risk.

Screening levels for some classes of compounds [PCBs and polyaromatic hydrocarbons (PAHs)] were also used in the previous risk assessments. SWMU 5 was the only SWMU with any COPCs that were PAHs. Because screening levels were available for these two PAHs (fluorine and phenanthrene), they were evaluated using their individual screening levels instead of the class screening level. The maximum detected concentration for each detected PCB was summed to develop a value for the PCB class. If no PCBs were detected at a SWMU, there was no evaluation in the ERA of the PCB class for that SWMU.

A summary of the results of the comparison in previous assessments of the site data to the ecological screening levels is provided in Table G.3. This table lists the number of COPCs in each suite retained for each site and the medium for further consideration. As shown, all BGOU SWMUs with an ERA had one or more COPCs, and the majority of COPCs retained at the BGOU SWMUs are metals.

Table G.3. Summary of Suite of COPCs Retained in Surface Soil

Area	Media	Metal	Rad	PCB	SVOC	VOC
SWMU 2	Soil	6	----	---	----	----
SWMU 3	Soil	NE	NE	NE	NE	NE
SWMU 4	Soil	5	----	1	----	----
SWMU 5	Soil	5	----	1	3	----
SWMU 6	Soil	2	----	----	1	----
SWMU 7	Soil	19	Total*	1	----	----
SWMU 30	Soil	17	Total*	1	----	----
SWMU 145	Soil	NE	NE	NE	NE	NE

----: no COPCs

NE: SWMU did not undergo an ecological evaluation

*Radionuclide risk was assessed based on a total dose benchmark for all radionuclides

SVOC=semivolatile organic compound

VOC=volatile organic compound

G.4.1 SWMU 2

The waste area grouping (WAG) 22 RI addendum for SWMUs 2 and 3 contained only a preliminary identification of COPCs for ecological risk (DOE 1994). The subsequent data summary and interpretation report (DOE 1997) was completed for SWMU 2 and contains a comparison of surface soil concentrations to ecological preliminary remediation goals (PRGs). That comparison was conducted for sediment in the ditch, surface soil (from 0 to 1 ft bgs), and the deeper Regional Gravel Aquifer (RGA) and McNairy soil. The ecological PRGs used for the comparison were the Oak Ridge values from Efroymsen *et al.* (1996). Comparisons for deeper soil may be relevant for burrowing animals and are presented in Section 5 of the data summary and interpretation report. Only the results from the data summary and interpretation report for surface soil are summarized in Table G.4; further evaluation of ditch sediment in this area of the facility is provided in the *Surface Water Operable Unit (On-Site) Site Investigation and Baseline Risk Assessment Report at the Paducah Gaseous Diffusion Plant, Paducah, Kentucky* [Surface Water Operable Unit (SWOU) Site Investigation (SI) and Baseline Risk Assessment (BRA)] (DOE 2008). Only the surface soil COPCs exceeding their screening criteria based on the maximum detected concentration are provided in Table G.4; the comparison of all COPCs can be found in Table 5.15 of the data summary and interpretation report.

Table G.4. Summary of COPCs Retained in Surface Soil for SWMU 2

Analyte	Frequency of Detection	Max Detected Concentration (mg/kg)	EcoPRG (mg/kg)	Hazard Quotient (HQ)
Arsenic	3/3	24.6	2.66	9.3
Chromium	3/3	11.3	0.4	28
Manganese	3/3	655	100	6.5
Nickel	3/3	28.6	24.0	1.2
Silver	3/3	6.3	2.0	3.1
Vanadium	3/3	18.9	2.0	9.5

Based on the data summary and interpretation report (DOE 1997)

G.4.2 SWMU 4

The WAG 3 RI report (DOE 2000) included an ecological risk assessment for surface soil at SWMU 4. This assessment included a comparison to both No Observed Adverse Effects Level (NOAEL)-based benchmark screening levels for the receptors listed in the table and to the background values for surface soil. Tables G.5 and G.6 include only chemicals exceeding both benchmark and background values. Table G.5 presents the frequency of detection and the concentrations used as EPCs for each analyte. The hazard quotient (HQ) for each receptor using those EPCs is provided in Table G.6. Blank cells in Table G.6 indicate the HQ was one or less; a dash indicates that no screening level was available for that receptor/COPC combination. All HQs listed in Table G.6 are compared to NOAEL-based HQs. NOAEL-based benchmarks for plants and soil invertebrates are from Efroymsen *et al.* (1997a; 1997b). NOAEL-based benchmarks for wildlife are derived from the literature sources shown in Exhibits 2.1 and 2.14 of the original document and following the methods in Sample, Opresko, and Suter II (1996).

Table G.5. COPCs Retained in Surface Soil for SWMU 4

Analyte	Frequency of Detection	Max detected concentration (EPC for soil receptors) mg/kg	UCL (EPC for wildlife) mg/kg
Chromium	13/13	296	10,539.31
Copper	13/13	30.1	14.7
Nickel	11/13	153	32.12
Vanadium	13/13	47.8	28.63
Zinc	12/13	72.5	48.43
PCBs (total)	4/11	0.494	0.49

Based on the WAG 3 RI report (DOE 2000)

Table G.6. HQs for COPCs Retained in Surface Soil for SWMU 4

Analyte	Plant HQ	Soil Invertebrates HQ	Meadow vole HQ	Long-tailed weasel HQ	White-footed mouse HQ	Short-tailed shrew HQ	American Woodcock HQ
Chromium	296	740	1.63			4.41	
Nickel	5.1						
Vanadium	23.9	----	2.99			9.15	
Zinc	1.45						4.22
PCBs (total)		----				2.83	1.27

Blank cells indicate the HQ was one or less; a dash indicates that no screening level was available for that receptor/COPC combination. Based on the WAG 3 RI report (DOE 2000).

G.4.3. SWMU 5

The WAG 3 RI report (DOE 2000) included an ecological risk assessment for surface soil at SWMU 5. This assessment included a comparison to both NOAEL-based benchmark screening levels for the receptors listed in the table and to the background values for surface soil. Tables G.7 and G.8 include only chemicals exceeding both benchmark and background values. Table G.7 presents the frequency of detection and the concentrations used as EPCs for each analyte. The HQ for each receptor using those EPCs is provided in Table G.8. Blank cells in Table G.8 indicate the HQ was one or less; a dash indicates

that no screening level was available for that receptor/COPC combination. All HQs listed in Table G.8 are compared to NOAEL-based HQs. NOAEL-based benchmarks for plants and soil invertebrates are from Efroymsen *et al.* (1997a; 1997b). NOAEL-based benchmarks for wildlife are derived from the literature sources shown in Exhibits 2.13 and 2.14 of the original document and following the methods in Sample, Opresko, and Suter II (1996).

Table G.7. COPCs Retained in Surface Soil for SWMU 5

Analyte	Frequency of Detection	Max detected concentration (EPC for soil receptors) mg/kg	UCL (EPC for wildlife) mg/kg
Aluminum	13/13	13,800	10,761.77
arsenic	5/13	12.2	7.55
Chromium	13/13	20.5	15.01
Nickel	12/13	119	24.11
Zinc	13/13	163	84.34
Fluoranthene	7/13	53.3	3.85
Phenanthrene	7/13	34.6	2.63
di-n-butyl phthalate	3/13	1.7	0.91
PCBs (total)	5/13	0.618	0.55

Based the WAG 3 RI report (DOE 2000)

Table G.8. HQs for COPCs Retained in Surface Soil for SWMU 5

Analyte	Plant HQ	Soil Invertebrates HQ	Meadow vole HQ	Long-tailed weasel HQ	White-footed mouse HQ	Short-tailed shrew HQ	American Woodcock HQ
Aluminum	276	-----	159	42.5	24.3	49.4	10.6
Arsenic	1.22		3.08			7.39	
Chromium	20.5	51.25					
Nickel	3.97						
Zinc	3.26	1.63					6.8
Fluoranthene	-----	1.57					
Phenanthrene	-----	1.02					
di-n-butyl phthalate							1.8
PCBs (total)		-----				3.64	1.64

Blank cells indicate the HQ was one or less; a dash indicates that no screening level was available for that receptor/COPC combination. Based on the WAG 3 RI report (DOE 2000).

G.4.4 SWMU 6

The WAG 3 RI report (DOE 2000) included an ecological risk assessment for surface soil at SWMU 6. This assessment included a comparison to both NOAEL-based benchmark screening levels for the receptors listed in the table and to the background values for surface soil. Tables G.9 and G.10 include only chemicals exceeding both benchmark and background values. Table G.9 presents the frequency of detection and the concentrations used as EPCs for each analyte. The HQ for each receptor using those EPCs is provided in Table G.10. Blank cells in Table G.10 indicate the HQ was one or less; a dash indicates that no screening level was available for that receptor/COPC combination. All HQs listed in Table G.10 are compared to NOAEL-based HQs. NOAEL-based benchmarks for plants and soil invertebrates are from Efroymsen *et al.* (1997a; 1997b). NOAEL-based benchmarks for wildlife are derived from the literature sources shown in Exhibits 2.13 and 2.14 of the original document and following the methods in Sample, Opresko, and Suter II (1996).

Table G.9. COPCs Retained in Surface Soil for SWMU 6

Analyte	Frequency of Detection	Max detected concentration (EPC for soil receptors) mg/kg	UCL (EPC for wildlife) mg/kg
Nickel	5/7	43.2	24.84
Zinc	7/7	128	78.44
di-n-butyl phthalate	2/7	1.7	0.99

Based on the WAG 3 RI report (DOE 2000)

Table G.10. HQs for COPCs Retained in Surface Soil for SWMU 6

Analyte	Plant HQ	Soil Invertebrates HQ	Meadow vole HQ	Long-tailed weasel HQ	White-footed mouse HQ	Short-tailed shrew HQ	American Woodcock HQ
Nickel	1.44						
Zinc	2.56	1.28					5.58
di-n-butyl phthalate							1.80

Blank cells indicate the HQ was one or less; a dash indicates that no screening level was available for that receptor/COPC combination. Based on the WAG 3 RI report (DOE 2000).

G.4.5 SWMU 7

The WAG 22 RI report (DOE 1998), which contains the ecological risk assessment for SWMU 7, was completed prior to development of the NFA levels in the PGDP Risk Methods Document in 2001; therefore, the document developed NOAEL-based benchmarks for each of six ecological receptors. Three of the receptors (plants, soil invertebrates, and soil microbes) were relatively nonmobile and were screened by comparison of the benchmarks to maximum detected values. Three of the receptors (white-tailed deer, white-footed mouse, and short-tailed shrew) were considered relatively mobile and were screened by comparison of the benchmarks to the UCL (if the UCL exceeded the maximum detected concentration, then the maximum detected concentration was used). Table 1.11 in Appendix A of the WAG 22 RI report presents the summary of COPCs for SWMU 7. Table G.11 summarizes the EPCs for COPCs for the two classes of receptors for SWMU 7. Only analytes exceeding their screening levels are summarized in Table G.11. The full comparison of all analytes to screening levels is shown in Appendix A of the WAG 22 RI report.

Table G.11. COPCs Retained in Surface Soil for SWMU 7

Analyte	Frequency of Detection	Max detected concentration (EPC for soil receptors) mg/kg	UCL (EPC for wildlife) mg/kg
Aluminum	13/13	14,800	9,670
Arsenic	13/13	16.0	7.21
Barium	13/13	120	75.5
Beryllium	12/13	24.0	11.4
Cadmium	9/13	3.0	1.87
Chromium	13/13	44.0	32.2
Cobalt	13/13	29.0	12.0
Copper	13/13	99.0	34.1
Fluoride	6/6	32.0	24.7
Iron	13/13	30,000	21,900
Lead	13/13	120	33.0
Manganese	13/13	1,160	472
Mercury	11/13	0.1	.0961
Nickel	13/13	55.0	26.4
Selenium	9/13	0.88	0.524
Thallium	12/13	2.0	1.28
Uranium	7/7	1,400	701
Vanadium	13/13	52.0	29.5
Zinc	13/13	200	97.5
Aroclor-1260	6/13	1.8	0.295

Based on the WAG 22 RI report (DOE 1998)

Table 2.4 in Appendix A of the WAG 22 RI report presents the screening levels developed for three receptors: plants, soil microbes, and earthworms. Table 2.5 in Appendix A of the WAG 22 RI presents the screening levels developed for three wildlife receptors: white-tailed deer, white-footed mouse, and the short-tailed shrew. The wildlife benchmarks are all NOAEL-based levels.

Hazard quotients for analytes with HQs exceeding 1 for any of the receptors as well as the total hazard index (HI) (the sum of all the HQs for a receptor) for SWMU 7 are listed in Table G.12. The HQs for all analytes are provided in Tables 2.6 and 2.7 of Appendix A of the WAG 22 RI.

A separate dose-based analysis was done to assess risk to ecological receptors from radionuclides and is presented in Table 2.9 of Appendix A of the WAG 22 RI. The dose contributions were less than the benchmark dose of 100 mrad per day for all the receptors listed in Table G.12, except that the HQ for plants was 1.09. The isotopes U-234 and U-235 were the main contributors to the total dose for plants.

Table G.12. HQs for COPCs Retained in Surface Soil for SWMU 7

Analyte	Plants HQ	Soil microbes HQ	Earthworm HQ	White-tailed deer HQ	White-footed mouse HQ	Short-tailed shrew HQ
Aluminum	296	24.7	-----	40.6	59.8	620
Arsenic	1.6				3.54	27.1
Barium			-----			1.11
Beryllium	2.4	-----	-----			6.17
Cadmium						3.45
Chromium	44	4.4	110		3.19	22.6
Cobalt	1.45		-----			
Copper			1.98			
Fluoride		1.07	-----	-----	-----	-----
Iron	-----	150	-----			
Lead	2.4					
Manganese	2.32	11.6	-----			
Mercury						3.76
Nickel	1.83					1.06
Selenium						1.09
Thallium	2.0	-----	-----			6.09
Uranium	280	-----	-----		1.41	22.6
Vanadium	26.0	2.6	-----	1.56	1.12	9.02
Zinc	4.0	2.0				1.10
Aroclor-1260	-----	-----	-----		4.66	33.2

Blank cells indicate the HQ was one or less; a dash indicates that no screening level was available for that receptor/COPC combination. Based on the WAG 22 RI report (DOE 1998)

G.4.6 SWMU 30

The WAG 22 RI report (DOE 1998), which also contains the ecological risk assessment for SWMU 30, was completed prior to development of the NFA levels in the PGDP risk methods document in 2001; therefore, the document developed NOAEL-based benchmarks for each of six ecological receptors. Three of the receptors (plants, soil invertebrates, and soil microbes) were relatively nonmobile and were screened by comparison to maximum detected values. Three of the receptors (white-tailed deer, white-footed mouse, and short-tailed shrew) were considered relatively mobile and were screened by comparison to the UCL (if the UCL exceeded the maximum detected concentration, then the maximum detected concentration was used). Table 1.11 in Appendix A of the WAG 22 RI report presents the summary of COPCs for SWMU 30. Table G.13 summarizes the EPCs for COPCs for the two classes of receptors for SWMU 30. Only analytes exceeding their NFA level are summarized in the table. The full comparison of all analytes to NFA levels is shown in Appendix A of the WAG 22 RI report.

Table G.13. COPCs Retained in Surface Soil for SWMU 30

Analyte	Frequency of Detection	Max detected concentration (EPC for soil receptors) mg/kg	UCL (EPC for wildlife) mg/kg
Aluminum	14/14	15,000	10,500
Arsenic	14/14	9.0	6.19
Barium	14/14	161	104
Beryllium	14/14	24.0	11.1
Cadmium	13/14	9.0	3.58
Chromium	14/14	38.0	27.9
Cobalt	14/14	12.0	8.08
Copper	14/14	89.0	50.0
Iron	14/14	29,600	18,000
Manganese	14/14	624	434
Mercury	14/14	2.0	0.272
Nickel	14/14	66.0	39.6
Silver	9/14	4.0	2.09
Thallium	12/14	1.2	0.578
Uranium	2/4	450	394
Vanadium	14/14	34.0	25.5
Zinc	14/14	155	76.7
Aroclor-1260	7/17	150	1.32

Based on the WAG 22 RI report (DOE 1998)

Table 2.4 in Appendix A of the WAG 22 RI report presents the screening levels developed for three receptors: plants, soil microbes, and earthworms. Table 2.5 in Appendix A of the WAG 22 RI report presents the screening levels developed for three wildlife receptors: white-tailed deer, white-footed mouse, and the short-tailed shrew. The wildlife benchmarks are all NOAEL-based levels.

Analytes with HQs exceeding 1 for any of the receptors as well as the total HI for SWMU 30 are listed in Table G.14. The HQs for all analytes are provided in Tables 2.6 and 2.7 of Appendix A of the WAG 22 RI report.

A separate dose-based analysis was done to assess risk to ecological receptors from radionuclides and is presented in Table 2.9 of Appendix A of the WAG 22 RI. The dose contributions were less than the benchmark dose of 100 mrad per day for all the receptors listed in Table G.14, except that the HQ for plants was 1.32. The isotopes U-234 and U-235 were the main contributors to the total dose for plants.

Table G.14. HQs for COPCs Retained in Surface Soil for SWMU 30

Analyte	Plant HQ	Soil microbe HQ	Earthworm HQ	White-tailed deer HQ	White-footed mouse HQ	Short-tailed shrew HQ
Aluminum	300	25	----	44.0	65.1	678
Arsenic					3.04	23.3
Barium			----			1.54
Beryllium	2.4		----			6.01
Cadmium	3.0				1.08	6.65
Chromium	38.0	3.8	95.0		2.78	19.7
Copper			1.78			
Iron	----	148	----	----	----	----
Manganese	1.25	6.24	----			
Mercury	6.67		20.0		1.50	10.6
Nickel	2.20					1.60
Silver	2.0		----			
Thallium	1.20	----	----			2.75
Uranium	90.0	----	----			12.7
Vanadium	17.0	1.70	----	1.35		7.78
Zinc	3.10	1.55				
Aroclor-1260	----	----	----		20.9	148

Blank cells indicate the HQ was one or less; a dash indicates that no screening level was available for that receptor/COPC combination. Based on the WAG 22 RI report (DOE 1998)

G.5. FOOD WEB MODELING FOR PCBS

The ecological evaluations done as part of the previous RI reports include comparisons to screening levels designed to be protective of adverse effects from direct exposure to the media (direct contact and

ingestion of soil). For contaminants that bioaccumulate through the food chain, such as PCBs, additional risks may be present because of the additional dose ingested through the food chain. The ERAs in the previous RIs did not include a food web model assessment; however, the assessment conducted as part of the screening ERA in the SWOU SI and BRA (DOE 2008) assessed the bioaccumulation potential for several species of mammalian and avian receptors. The receptor species evaluated for soil (short-tailed shrew, meadow vole, American kestrel, American woodcock, and American robin) also are appropriate receptors for the SWMUs within the BGOU.

The approach used for the food web model is described in Section E.3.2 of the SWOU SI and BRA (DOE 2008). Food web modeling for total PCBs involved calculation of the potential dose ingested per kg body weight per day to the toxicity reference benchmarks developed in *Toxicological Benchmarks for Wildlife: 1996 Revision* (Sample, Opresko, and Suter II 1996). The toxicity reference benchmarks are provided in Table E.3 of the SWOU SI and BRA and the HQs for the food web model appear in Section E.5.2 of the same document. This document covers different sites than the BGOU RI, but the modeling results can be used directly because the dose of PCBs (and therefore the HQ) for each receptor is directly proportional to the soil concentration. Table E.3.2 and Table E.3 of the SWOU SI and BRA show the NOAEL-based HQs for total PCBs for each species. The ratio of these soil concentrations to HQs is valid for the BGOU units as well, as long as they are calculated for a site foraging factor (SFF) of 1.0. The SFF is 1.0 for the short-tailed shrew, the meadow vole, the American robin, and the bobwhite quail; therefore, the HQs for these receptors were used to evaluate the total PCB results of the BGOU RI units. The ratios used to convert the total PCB value into an equivalent food web model HQ are shown in Table G.15. The conversion factors are the same for both high and low concentrations (except for rounding differences); the larger value was used if a slight difference existed between the two values.

Table G.15. Factors for Converting Soil Concentrations into Food Web Model HQs

Receptor Species	Soil Concentration in SWOU Appendix E	NOAEL HQ in SWOU Appendix E	Soil-to-HQ Conversion Factor
Short-tailed Shrew	0.4	10.2	25.5
Meadow vole	0.4	0.035	0.0875
American robin	0.4	1.2	3.0
Bobwhite Quail	0.4	0.014	0.035
Short-tailed Shrew	20	511	25.6
Meadow vole	20	1.7	0.085
American robin	20	57.5	2.9
Bobwhite Quail	20	0.69	0.035

Table G.16 shows the HQs associated with the maximum total PCBs detected at each SWMU in the BGOU SWMU soil. These HQs are >1 for the short-tailed shrew and for the American robin at SWMUs 2, 4, 5, and 7. At SWMU 30, HQs are greater than 1 for all mammal species evaluated and the American robin.

Table G.16. HQs Associated with the Maximum Total PCBs at Each SWMU¹

	Conversion Factor	SWMU 2 HQ	SWMU 4 HQ	SWMU 5 HQ	SWMU 6 HQ	SWMU 7 HQ	SWMU 30 HQ
Total PCBs (mg/kg)		0.39	0.494	0.618	NA	1.99	17.1
Short-tailed Shrew	25.5	9.9	13	16	NA	51	436
Meadow vole	0.0875	0.034	0.043	0.054	NA	0.17	1.5
American robin	3.0	1.2	1.5	1.9	NA	6.0	51
Bobwhite Quail	0.035	0.014	0.017	0.022	NA	0.07	0.60

¹ SWMU 3 surface soil was never evaluated except in combination with SWMU 2 surface soil. SWMU 145 was not evaluated because of the presence of a cap on the site.
NA: PCBs not detected.

G.6. UNCERTAINTIES

There are a number of uncertainties that impact the potential usefulness of the results of the previously conducted ERAs. At SWMU 7, additional surface soil samples were collected at two sampling locations after completion of the previous ERA. The maximum detected concentration of metals, organics, and radionuclides for these new SWMU 7 samples appears in Attachment F1 of Appendix F (the human health risk assessment). For all metals and most radionuclides, the maximum detected concentration associated with the new samples is well below the maximum detected concentration used in the previous ecological risk assessment for SWMU 7 and therefore would not alter the results of that assessment. The maximum detected concentration of Total PCBs (14.8 mg/L) in the new surface samples exceeds the value used in the risk assessment, but PCBs already were retained as a COPC for SWMU 7. The new samples contained a detection of di-n-butyl phthalate, but the detected concentration of 0.94 would have an HQ of 1.0 using the di-n-butyl phthalate benchmark from the WAG 3 RI report. Uranium-234, uranium-235, and uranium-236 also were detected at higher maximum concentrations in the new surface soil samples at SWMU 7. Radionuclides already were retained for SWMU 7 based on dose to plants from uranium; the new data therefore do not substantially impact the results of the assessment already conducted and summarized in this appendix. The screening evaluations summarized here included the surface soil interval. Sufficient sampling results were not available for subsurface soil in the depth range of interest (0-5 ft bgs) to which burrowing animals may be exposed to conduct a quantitative screen of the subsurface soil. In addition, characterization of the buried waste was not adequate to quantitatively assess potential exposure for burrowing animals. Because the screening for surface soil indicated that additional evaluation of all the BGOU SWMUs was necessary, the uncertainties relating to exposures of burrowing animals will be addressed as part of that future evaluation, which will resolve the uncertainties related to these potential exposure pathways.

Another uncertainty in these screening assessments is that the ecological screening levels are protective of entire suites of receptors, some of which may not be present at these disturbed sites. The grassy areas of these sites would be attractive to ecological receptors, but the surrounding industrial area may limit the extent to which ecological receptors use these areas.

The exposure parameters used for the wildlife receptors in the previous ERAs were assembled from literature values. These general values are representative of the population as a whole or may come from a population in a different area of the United States than the facility. Exposure parameter values for

populations of these receptors in western Kentucky may differ from those used in the assessment, which could result in an over- or underestimate of dose and the resulting risk or hazard.

The food web modeling for PCBs is based on bioaccumulation and transfer of contaminants through the food chain, sometimes for several levels of the food chain. Uncertainty in the values for bioaccumulation factors used to estimate the concentration in food items for the food web model generally leads to an overestimation of potential risk, because of conservatism in the bioaccumulation factors available in the literature.

These uncertainties, combined with the results of the ERAs, indicate the need for further evaluation of these sites. Risk managers may determine that sites do not need further evaluation (if exposure pathways are not complete or planned actions will eliminate the exposure pathway) or may recommend additional evaluation of the sites to better define the potential ecological risk indicated by the results of the previous ERAs. Alternatively, the benchmarks used in the screenings presented here and in the NFA levels in the PGDP risk ecological risk methods documents (DOE 2001 and the 2008 draft revision of the same document) may be used as the ecologically based remedial goal options (RGOs).

G.7. CONCLUSIONS

Each of the sites evaluated in the ERAs summarized in this section retained a number of COPCs. Some metals at concentrations above background were retained as contaminants of concern for ecological risk at each SWMU. Total PCBs were retained as COPCs for all SWMUs, except SWMU 6. This is based on direct risk from soil as well as risks to some wildlife receptors from bioaccumulation through the food chain. The only other COPCs retained are three SVOCs (fluorene, phenanthrene, and di-n-butyl phthalate) at SWMU 5 and di-n-butyl phthalate at SWMU 6.

The current plan is to conduct additional ecological risk assessment in future RI activities. In the absence of these future activities, the benchmarks used in the screenings presented here and in the NFA levels in the PGDP risk ecological Risk Methods Documents (DOE 2001 and the 2008 draft revision of the same document) will be used to develop ecologically based RGOs.

G.8. REFERENCES

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APPENDIX H

APPLICABLE OR RELEVANT AND APPROPRIATE REQUIREMENTS AND TO BE CONSIDERED GUIDANCE

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ACRONYMS

ALARA	as low as reasonably achievable
ARAR	applicable or relevant and appropriate requirement
BGOU	Burial Grounds Operable Unit
CERCLA	Comprehensive Environmental Response, Compensation, and Liability Act
<i>CFR</i>	<i>Code of Federal Regulations</i>
COE	U.S. Army Corps of Engineers
DOE	U.S. Department of Energy
EPA	U.S. Environmental Protection Agency
<i>FR</i>	<i>Federal Register</i>
FS	feasibility study
NCP	National Oil and Hazardous Substances Pollution Contingency Plan
NWP	Nationwide Permit
OSHA	Occupational Safety and Health Association
PGDP	Paducah Gaseous Diffusion Plant
RI	remedial investigation
SWMU	solid waste management unit
TBC	To Be Considered
<i>USC</i>	<i>United States Code</i>

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H.1 INTRODUCTION

Congress specified in the Comprehensive Environmental Response, Compensation, and Liability Act (CERCLA) § 121 (42 USC 9621) that remedial actions for the cleanup of hazardous substances must require a level or standard of control that attains those requirements, criteria, standards, or limitations under federal or more stringent state environmental laws that are legally applicable or relevant and appropriate (ARAR) to the hazardous substances or circumstances at a site (unless an ARAR is waived).

This appendix supplies a preliminary list of available federal and state ARARs that may be associated with potential remedial actions at the Burial Grounds Operable Unit (BGOU) at the Paducah Gaseous Diffusion Plant (PGDP). The process of ARAR identification is an iterative one that is continually changing as the remedial investigation/feasibility study (RI/FS) progresses; therefore, the ARARs that are identified represent a compilation of potential ARARs that are subject to change as site-specific contamination at the BGOU is further characterized and alternatives are further evaluated. Site-specific ARARs will be identified further during the remedial action selection for the FS. The U.S. Environmental Protection Agency (EPA) differentiates ARARs as either “applicable” or “relevant and appropriate” to a site. The terms and conditions of these categories are as follows:

- *Applicable requirements* are “those cleanup standards, standards of control, and other substantive requirements, criteria, or limitations promulgated under federal environmental or state environmental or facility siting laws that specifically address a hazardous substance, pollutant, contaminant, remedial action, location, or other circumstance found at a CERCLA site” (40 CFR § 300.5); and
- *Relevant and appropriate requirements* are “those cleanup standards, standards of control, and other substantive requirements, criteria, or limitations promulgated under federal environmental or state environmental or facility siting laws that, while not applicable to a hazardous substance, pollutant, contaminant, remedial action, location, or other circumstance at a CERCLA site, address problems or situations sufficiently similar to those encountered at the CERCLA site that their use is well suited to the particular site” (40 CFR § 300.5).

The EPA also categorizes ARARs based on whether they are specific to the chemical(s) present at the site (chemical-specific), the remedial action being evaluated (action-specific), or the location of the site (location-specific). The EPA designated these categories to assist in the identification of ARARs; however, they are not necessarily precise [53 FR 51437 (1988)]. Some ARARs may fit into more than one category, while others may not definitively fit into any one category. Terms and conditions relevant to this categorization are included in the list that follows:

- *Chemical-specific ARARs* usually are “health- or risk-based numerical values or methodologies that, when applied to site-specific conditions, result in the establishment of numerical values” [53 FR 51437 (1988)]. These values establish the acceptable amount or concentration of a chemical that may remain in, or be discharged to, the ambient environment.
- *Action-specific ARARs* usually are “technology- or activity-based requirements or limitations placed on actions taken with respect to hazardous wastes, or requirements to conduct certain actions to address particular circumstances at a site” [53 FR 51437 (1988)]. Selection of a particular remedial action at a site will trigger action-specific ARARs that specify appropriate technologies and performance standards.
- *Location-specific ARARs* “generally are restrictions placed upon the concentration of hazardous substances or the conduct of activities solely because they are in special locations” [53 FR 51437

(1988)]. Some examples of special locations include floodplains, wetlands, historic places, and sensitive ecosystems or habitats.

Pursuant to CERCLA § 121(e) [42 *USC* 9621(e)(1)], response actions, or portions of response actions conducted entirely on-site, as defined in 40 *CFR* 300.5, must comply with the substantive portions of ARARs, but not the procedural or administrative requirements. Additionally, CERCLA § 121(d)(4) [42 *USC* 9621(d)(4)] provides six ARAR waiver options that may be invoked, provided that human health and the environment are protected.

Published, unpromulgated information that does not necessarily meet the definition of an ARAR may be necessary, under certain circumstances, to determine what is protective of human health and the environment. This type of information is known as To Be Considered (TBC) guidance and also may be useful in developing CERCLA remedies. Because ARARs do not exist for every chemical or circumstance that may be found at a CERCLA site, the EPA believes that it may be necessary, when determining cleanup requirements or designing a remedy, to consult reliable information that otherwise would not be considered a potential ARAR. Criteria or guidance developed by the EPA, other federal agencies, or states may assist in determining, for example, health-based levels for a particular contaminant or the appropriate method for conducting an action for which there are no ARARs. The TBC guidance generally falls within four categories: (1) health effects information; (2) technical information on how to perform or evaluate investigations or response actions; (3) policy; and (4) proposed regulations, if the proposed regulation is noncontroversial and likely to be promulgated as drafted.

The EPA requires compliance with Occupational Safety and Health Association (OSHA) standards through § 300.150 of the National Oil and Hazardous Substances Pollution Contingency Plan (NCP), not through the ARARs process. Worker health and safety requirements typically are not addressed as ARARs. The regulations at 29 *CFR* 1910.120 are designed to protect workers involved in cleanup operations at uncontrolled hazardous waste sites and to provide for worker protection during initial site characterization and analysis, monitoring activities, materials handling activities, training, and emergency response.

As mentioned above, ARARs identification is an iterative process that continually changes as the RI/FS progresses. There are no chemical-specific ARARs for this action. The action-specific ARARs will be identified as part of the FS, based upon the remedial alternatives under consideration; therefore, the ARARs discussed are focused on location-specific ARARs. The final set of ARARs will be included as part of the Record of Decision based on the selected remedy.

H.2 LOCATION-SPECIFIC APPLICABLE OR RELEVANT AND APPROPRIATE REQUIREMENTS

No threatened or endangered species or their potential habitats, critical habitats, 100-year floodplains, wetlands, prime farmland, or cultural resources have been identified in the boundaries of the BGOU solid waste management units (SWMUs); however, a 100-year floodplain has been identified near SWMU 4 and wetlands have been identified in the ditches south of SWMU 4 and across a roadway to the north. Wetlands also have been identified south of SWMUs 5 and 6 on the other side of a roadway (CDM 1994; COE 1994; LMES 1996). ARARs discussed in this section will be met by avoidance of the resource to the extent practicable. If impacts become apparent, measures to mitigate adverse effects will be taken.

Construction activities must avoid or minimize adverse impacts on wetlands and act to preserve and enhance their natural and beneficial values (10 *CFR* 1022). If the action involves the discharge of dredged

or fill material into waters of the United States, the response action will comply with the substantive requirements of Nationwide Permit (NWP) 38 (Cleanup of Hazardous and Toxic Waste), however, the specific requirement of notification is not required for CERCLA actions under this NWP. Consequently, although wetlands should be delineated and avoided to the extent possible, the delineation does not have to be sent to the U.S. Army Corps of Engineers (COE), and the COE does not have to be notified for this action [61 *FR* 65905-65906 (1996)].

No federally listed or candidate species or their habitats are known to occur in the vicinity of the project area. The Commonwealth of Kentucky has no threatened and endangered species regulations promulgated at this time.

H.3 ACTION-SPECIFIC APPLICABLE OR RELEVANT AND APPROPRIATE REQUIREMENTS

Action-specific ARARs will be developed, as appropriate, in the FS.

H.4 CHEMICAL-SPECIFIC APPLICABLE OR RELEVANT AND APPROPRIATE REQUIREMENTS

There are no chemical-specific ARARs for this action.

H.5 REFERENCES

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