

DEPARTMENT OF ENERGY Environmental Management Los Alamos Field Office (EM-LA) Los Alamos, New Mexico 87544

April 9, 2020

EMLA-2020-1364-02-001

Mr. Kevin Pierard Bureau Chief Hazardous Waste Bureau New Mexico Environment Department 2905 Rodeo Park Drive East, Building 1 Santa Fe, NM 87505-6313

Subject: Submittal of the Addendum to the Phase II Investigation Report for Middle Los Alamos Canyon Aggregate Area, Revision 1

Dear Mr. Pierard:

Enclosed please find two hard copies with electronic files of the "Addendum to the Phase II Investigation Report for Middle Los Alamos Canyon Aggregate Area, Revision 1." Also enclosed is an electronic copy of a redline strikeout version of the report that includes all changes made in response to the New Mexico Environment Department's (NMED's) comments dated July 2, 2019. Responses to NMED's comments were submitted on September 20, 2019, and approved by NMED on September 27, 2019.

If you have any questions, please contact Brenda Bowlby at (505) 551-2957 (brenda.bowlby@emla.doe.gov) or Cheryl Rodriguez at (505) 257-7941 (cheryl.rodriguez@em.doe.gov).

Sincerely,

Arturo Duran Date: 2020.04.02 11:28:21 -06'00'

Arturo Q. Duran Compliance and Permitting Manager Environmental Management Los Alamos Field Office

Enclosures:

- Two hard copies with electronic files (including a redline strikeout version) Addendum to the Phase II Investigation Report for Middle Los Alamos Canyon Aggregate Area, Revision 1 (EM2020-0081)
- 2. Response to Draft New Mexico Environment Department Comments on the Addendum to Middle Los Alamos Canyon Area Phase II Investigation Report, Dated July 2, 2019 (EM2019-0350)

CC (letter with hard-copy enclosure[s]): Brenda Bowlby, N3B Cheryl Rodriguez, EM-LA

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Response to Draft New Mexico Environment Department Comments on the Addendum to Middle Los Alamos Canyon Area Phase II Investigation Report, Dated July 2, 2019

INTRODUCTION

To facilitate review of this response, the New Mexico Environment Department's (NMED's) comments are included verbatim (in italics). The U.S. Department of Energy (DOE) Environmental Management Los Alamos Field Office responses follow each NMED comment.

NMED Comment

1. Section 3.4.4, Nature and Extent of Contamination, page 10:

Permittee Statement: Aroclor-1260 was detected in 242 samples with a maximum concentration of 23.9 mg/kg. Concentrations increased with depth at locations 02-600449, 02-612452, 02-613626, 02-613668, 02-613700, 02-61436, 02-61444, 02-61479, 02-61488, 02-61489, and 02-61553 and did not change substantially with depth (0.001 mg/kg to 0.0426 mg/kg) at locations 02-61435, 02-61451, 02-61482, 02-61486, and 02-61490; only 1 depth was sampled at locations 02-612451, 02-613292, 02-613699, 02-61478, 02-61492, and 02-61539 through 02-61543; and concentrations decreased with depth at all other locations. Concentrations decreased downgradient. Vertical extent is defined at locations where only 1 depth was sampled by decreasing concentrations in deeper samples from nearby locations. The maximum concentration at locations where concentrations increased with depth or did not change substantially with depth (3.3 mg/kg at location 02-613626) was approximately 1.4 times the residential SSL, and the industrial SSL was approximately 3.4 times this concentration. Aroclor-1260 does not pose an unacceptable risk under the industrial and recreational scenarios (Appendix F, Tables F-4.2-1 and F-4.2-3). Lateral extent of Aroclor-1260 is defined and further sampling for vertical extent is not warranted.

NMED Comment: Data presented in Table 3.4-4, PCBs Detected at SWMU 02-014, indicate that concentrations of Aroclor-1260 also increased with depth at location 02-613289 with a maximum concentration of 20.7 mg/kg in the deepest sample collected from this location. Characterization samples were collected from this location in 2010, 2017, and 2018. The three deepest samples, ranging from 6-11.25 ft bgs were collected in January 2018 though it is unclear if the samples were collected before or after excavation of contaminated soil from 0-5 ft bgs. The maximum concentration at location 02-613289 is 1.9 times the industrial SSL and 8.5 times the residential SSL. The results indicate that vertical extent is not defined and additional sampling is warranted. Revise the discussion for accuracy and reevaluate vertical extent of Aroclor-1260 at location 02-613289.

DOE Response

 During 2017, sampling was conducted to characterize the nature and extent of polychlorinated biphenyl (PCB) contamination identified during previous sampling at Area of Concern (AOC) 02-011(a). The results of this sampling led to investigation of the source of the PCBs and the identification of Solid Waste Management Unit (SWMU) 02-014 as a new SWMU. Included in this sampling were 187 samples collected during July and August 2017, which should have been assigned to new SWMU 02-014 when it was formally identified in April 2018. The review of the data to evaluate vertical extent at location 02-613289 revealed that these 187 samples had not been assigned to SWMU 02-014 and as such, had not been included in the addendum to the Phase II investigation report. These samples have now been added to the data set for SWMU 02-014 and will be included in the revised addendum. The updated PCB results for SWMU 02-014 are shown in the attached revision to Plate 2.

NMED's comment notes that the concentration of Aroclor-1254 in the deepest sample at location 02-613289 is 1.9 times the industrial soil screening level (SSL) and 8.5 times the residential SSL and further sampling is warranted. As shown on Plate 2, there are 5 sample locations around location 02-613289 where samples were collected to a depth of 20 ft. These locations are 02-61432 (11 ft from location 02-613289), 02-61440 (9 ft from location 02-613289), 02-61441 (10 ft from location 02-613289), 02-61448 (6 ft from location 02-613289), and 02-61455 (13 ft from location 02-613289). Aroclor-1260 concentrations decreased with depth at all 5 locations. A total of 15 samples were collected from these 5 locations at depths greater than the deepest sample at location 02-613289 (14.0 to 15.0 ft below ground surface [bgs], 16.0 to 17.0 ft bgs, and 19.0 to 20.0 ft bgs at each location). As shown on Plate 2, Aroclor-1260 was detected in 7 of these 15 samples at concentrations ranging from 0.00316 mg/kg to 0.0707 mg/kg, all more than 2 orders of magnitude less than the concentration in the deepest sample at location 02-613289. Aroclor-1260 was not detected in the samples from 19.0 ft to 20.0 ft bgs at any of the 5 locations except location 02-61448, where it was detected at 0.0186 mg/kg. Based on the results of sampling at these 5 locations, further sampling for vertical extent of PCBs at location 02-613289 is not warranted. The text in the addendum will be revised to include these lines of evidence that further sampling is not warranted.

Although the Aroclor-1260 results for location 02-613289 presented in Table 3.4-4 show concentrations increasing with depth, these results do not include those from samples collected from 0.0–5.0 ft below bgs that were removed during excavation. Those results, which are included in Appendix E, show concentrations decrease with depth, with a maximum Aroclor-1260 concentration of 1520 mg/kg detected in the sample from 0.0–1.0 ft bgs. The text will be revised to clarify that Aroclor-1260 concentrations decreased with depth at this location by providing a reference to the excavated surface sample results in Appendix E. Although the concentrations increased with depth in post-excavation sample results presented in Table 3.4-4, concentrations decreased with depth in deeper samples at five adjacent locations as indicated above.

NMED Comment

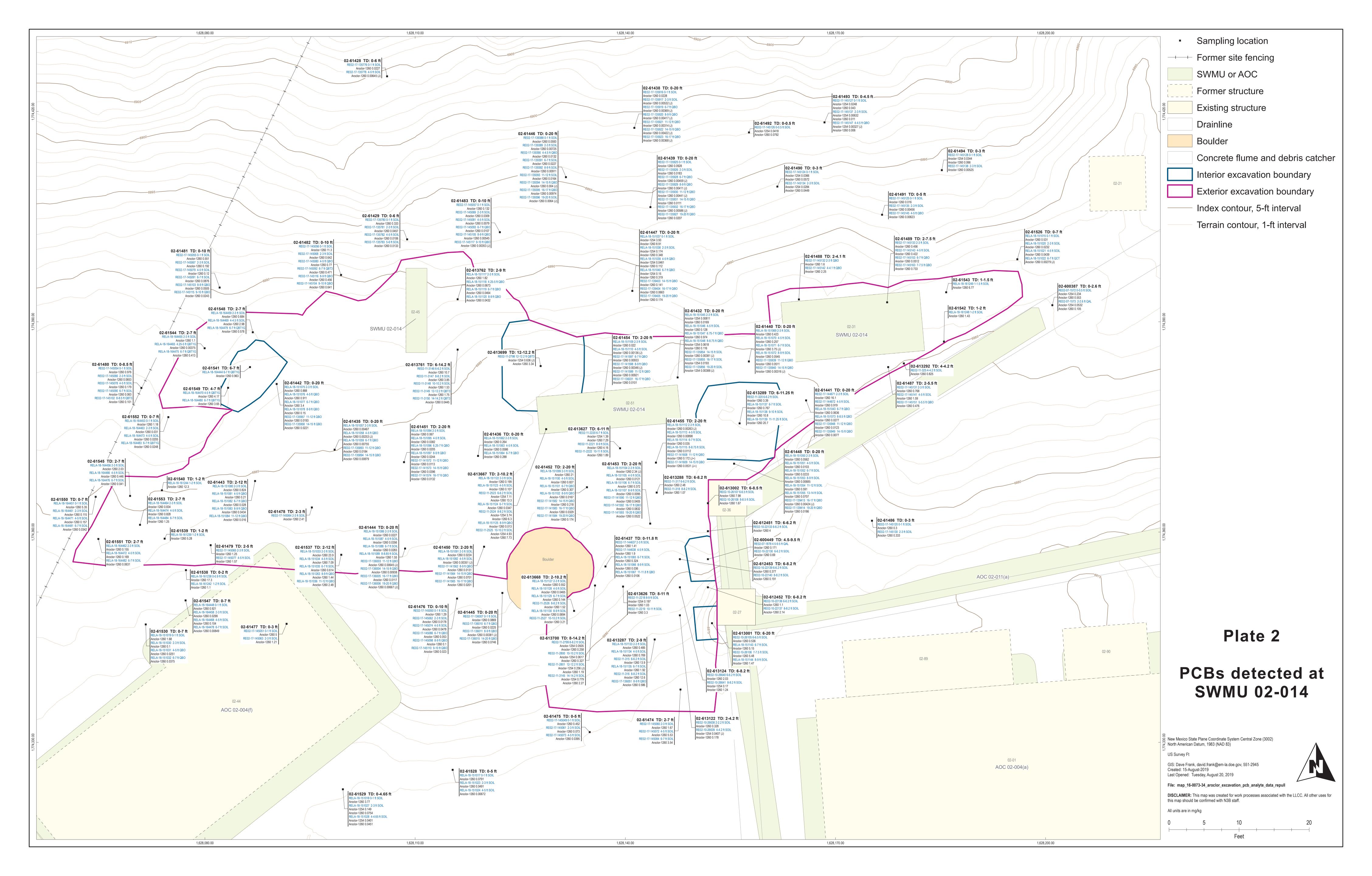
2. Section 3.6 Summary of Ecological Risk Screening, page 11:

Permittee Statement: SWMU 02-014 is within the TA-02 core area. Ecological risk for the TA-02 core area was evaluated in the "Phase II Investigation Report for Middle Los Alamos Canyon Aggregate Area, Revision 2" (N3B 2018, 700091). Based on evaluations of the minimum ESLs, HI analyses, potential effects to populations (individuals for threatened or endangered [T&E] species), lowest observed adverse effect level (LOAEL) analyses, the relationship of detected concentrations and screening levels to background concentrations, and results of site-specific ecological risk studies, the Phase II IR concluded no potential ecological risks exist for the TA-02 core area, which includes SWMU 02-014.

NMED Comment: Due to the ecological risk assessment for SWMU 02-014 being presented in the "Phase II Investigation Report for Middle Los Alamos Canyon Aggregate Area, Revision 2" (report) NMED must complete a review of the report prior to finalizing the review of the "Addendum to Middle Los Alamos Canyon Aggregate Area Phase II Investigation Report".

DOE Response

2. Comment noted.



March 2020 EM2020-0081

Addendum to the Phase II Investigation Report for Middle Los Alamos Canyon Aggregate Area, Revision 1



Newport News Nuclear BWXT-Los Alamos, LLC (N3B), under the U.S. Department of Energy Office of Environmental Management Contract No. 89303318CEM000007 (the Los Alamos Legacy Cleanup Contract), has prepared this document pursuant to the Compliance Order on Consent, signed June 24, 2016. The Compliance Order on Consent contains requirements for the investigation and cleanup, including corrective action, of contamination at Los Alamos National Laboratory. The U.S. government has rights to use, reproduce, and distribute this document. The public may copy and use this document without charge, provided that this notice and any statement of authorship are reproduced on all copies.

Addendum to the Phase II Investigation Report for Middle Los Alamos Canyon Aggregate Area, Revision 1

March 2020

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EXECUTIVE SUMMARY

This addendum to the 2018 "Phase II Investigation Report for Middle Los Alamos Canyon Aggregate Area, Revision 2" (Phase II IR) evaluates the nature and extent of contamination and potential human health and ecological risks for Solid Waste Management Unit (SWMU) 02-014. This SWMU is located within Technical Area 02 (TA-02) at Los Alamos National Laboratory (LANL or the Laboratory). This site was identified as a new SWMU during efforts to discover the source of polychlorinated biphenyl (PCB) contamination from investigation sampling at Area of Concern (AOC) 02-011(a)(ii). SWMU 02-014 consists of three former electrical transformer stations that served buildings in TA-02. The investigations revealed that PCB contamination at this site was more extensive than expected and remediation of soil contamination was required. As a result, investigation and remediation activities could not be completed in time for the results to be included in the Phase II IR. Remediation activities were planned for completion in late 2018/early 2019 with the results to be provided in an addendum to the Phase II IR.

Characterization data for SWMU 02-014 consist of results from samples collected in 2007, 2010, 2017, and 2018. Removal of PCB-contaminated soil was conducted to address potentially unacceptable risk for industrial workers and recreational users in the depth interval 0.0–1.0 ft below ground surface (bgs) and to meet the Toxic Substances Control Act bulk PCB remediation waste cleanup level for low-occupancy areas. Soil was excavated during 2018 and removal areas were expanded both laterally and vertically based on confirmation sampling results. A total of 282 yd³ of PCB-contaminated soil was excavated and packaged for transportation to an off-site disposal facility.

Following completion of investigation sampling and remediation activities, characterization data for SWMU 02-014 were evaluated in the same manner as described in the Phase II IR for the other SWMUs and AOCs within Middle Los Alamos Canyon Aggregate Area to identify chemicals of potential concern, evaluate nature and extent of contamination, and assess risk to human health. SWMU 02-014 was determined to not pose an unacceptable human health risk or dose under the industrial, recreational, residential, and construction worker scenarios. As described in the Phase II IR, ecological risk was evaluated for all SWMUs and AOCs within the TA-02 core area, including SWMU 02-014, rather than by individual SWMU or AOC. The Phase II IR concluded no potential ecological risks exist for the TA-02 core area. Based on the results of data evaluations presented in this addendum, the Department of Energy Environmental Management Los Alamos Field Office and Newport News Nuclear BWXT-Los Alamos, LLC, recommend corrective action complete without controls for SWMU 02-014.

CONTENTS

1.0	INTRO	DUCTIC)N	1
	1.1	General	Site Information	1
	1.2	Purpose	e of Investigation	1
	1.3	Docume	ent Organization	2
2.0	SCOP	E OF AC	TIVITIES	2
	2.1	Premob	ilization Activities	2
	2.2	Summa	ry of Field Activities	2
		2.2.1	Geodetic Survey	3
		2.2.2	Field Screening	3
		2.2.3	Surface and Shallow-Subsurface Soil Investigation	3
		2.2.4	Equipment Decontamination	3
		2.2.5	Excavation	4
		2.2.6	Health and Safety Measures	
		2.2.7	Waste Management	
	2.3	•	Analyses	
	2.4	Deviatio	ons	5
3.0	FIELD	INVEST	IGATION RESULTS FOR SWMU 02-014	5
	3.1	Site Des	scription and Operational History	5
	3.2	Relatior	nship to Other SWMUs and AOCs	6
	3.3	Summa	ry of Previous Investigations	6
	3.4	Site Cor	ntamination	6
		3.4.1	Soil and Rock Sampling	
		3.4.2	Soil and Rock Sample Field-Screening Results	
		3.4.3	Soil and Rock Sample Analytical Results	
		3.4.4	Nature and Extent of Contamination	
	3.5		ry of Human Health Risk Screening	
	3.6		ry of Ecological Risk Screening	
4.0	CONC	LUSION	S	15
	4.1		and Extent of Contamination	
	4.2	Summa	ry of Risk-Screening Assessments	
		4.2.1	Human Health Risk-Screening Assessment	
		4.2.2	Ecological Risk-Screening Assessment	15
5.0	RECO	MMEND	ATIONS	16
6.0	REFE	RENCES	AND MAP DATA SOURCES	16
	6.1		Ces	
	6.2	Map Da	ta Sources	17

Figures

Figure 3.4-1	Organic chemicals other than PCBs detected at SWMU 02-014	19
Figure 3.4-2	Radionuclides detected or detected above BVs/FVs at SWMU 02-014	20

Tables

Table 2.2-1	Surveyed Coordinates of Sample Locations at SWMU 02-014	21
Table 2.2–2	Field–Screening Results for Samples Collected at SWMU 02-014	23
Table 3.4-1	Samples Collected and Analyses Requested at SWMU 02-014	35
Table 3.4-2	Inorganic Chemicals Detected or Detected above BVs at SWMU 02-014	49
Table 3.4-3	Organic Chemicals other than PCBs Detected at SWMU 02-014	54
Table 3.4-4	PCBs Detected at SWMU 02-014	57
Table 3.4-5	Radionuclides Detected or Detected above BV/FV at SWMU 02-014	67

Appendixes

Appendix A	Acronyms and Abbreviations, Metric Conversion Table, and Data Qualifier Definitions
Appendix B	Field Methods
Appendix C	Waste Management
Appendix D	Analytical Program
Appendix E	Analytical Suites and Results and Analytical Reports (on CD included with this document)
Appendix F	Risk Assessments
Appendix G	Box Plots and Statistical Results

Plates

Plate 1	Site map of SWMU 02-014
Plate 2	Inorganic chemicals detected or detected above BVs at SWMU 02-014
Plate 3	PCBs detected at SWMU 02-014

1.0 INTRODUCTION

Los Alamos National Laboratory (LANL or the Laboratory) is a multidisciplinary research facility owned by the U.S. Department of Energy (DOE). The Laboratory is located in north-central New Mexico, approximately 60 mi northeast of Albuquerque and 20 mi northwest of Santa Fe. The Laboratory site covers approximately 36 mi² of the Pajarito Plateau, which consists of a series of fingerlike mesas that are separated by deep canyons containing perennial and intermittent streams running from west to east. Mesa tops range in elevation from approximately 6200 ft to 7800 ft above mean sea level.

The Laboratory has been a participant in a national effort by DOE to clean up sites and facilities formerly involved in weapons research and development. The goal of this effort is to ensure past operations do not threaten human or environmental health and safety in and around Los Alamos County, New Mexico. To achieve this goal, the Laboratory has investigated sites potentially contaminated by past Laboratory operations.

This investigation report addendum addresses one solid waste management unit (SWMU) within the Middle Los Alamos Canyon Aggregate Area at the Laboratory. This site is potentially contaminated with both hazardous and radioactive components. The New Mexico Environment Department (NMED), pursuant to the New Mexico Hazardous Waste Act, regulates cleanup of hazardous wastes and hazardous constituents. DOE regulates cleanup of radioactive contamination, pursuant to DOE Order 5400.5, "Radiation Protection of the Public and the Environment"; DOE Order 435.1, "Radioactive Waste Management"; and DOE Order 458.1, "Administrative Change 3, Radiation Protection of the Public and the Environment." Information on radioactive materials and radionuclides, including the results of sampling and analysis of radioactive constituents, is voluntarily provided to NMED in accordance with DOE policy.

Corrective actions at the Laboratory are subject to a Compliance Order on Consent (the Consent Order). This investigation report addendum describes work activities that were completed in accordance with the Consent Order.

1.1 General Site Information

The Middle Los Alamos Canyon Aggregate Area consists of 80 SWMUs and areas of concern (AOCs), 40 of which did not warrant investigation (LANL 2008, 101669.12). The remaining 40 SWMUs and AOCs underwent sampling activities in 2007 and Phase II investigation sampling activities in 2010. These 40 sites are located at Technical Area 02 (TA-02), TA-21, and TA-26 and include 13 SWMUs and 27 AOCs. Details of previous investigations, including the results of the 2007 sampling activities, are provided in the "Investigation Report for Middle Los Alamos Canyon Aggregate Area, Revision 1" (LANL 2008, 101669.12). Results from sampling activities conducted in 2010 for the 40 sites, as well as supplemental sampling performed in 2017 are provided in the "Phase II Investigation Report for Middle Los Alamos Canyon Aggregate Area, Revision 7" (Phase II IR) (N3B 2018, 700091).

1.2 Purpose of Investigation

During the course of the Phase II investigation activities being conducted to discover a source of polychlorinated biphenyl (PCB) contamination at storm drain AOC 02-011(a)(ii), a new SWMU (SWMU 02-014) was discovered. SWMU 02-014 consists of three former electrical transformer stations in TA-02. The investigations revealed that PCB contamination at this site was more extensive than expected and remediation of soil contamination was required. As a result, investigation and remediation activities could not be completed in time for the results to be included in the Phase II IR. Remediation activities

were planned for completion in late 2018/early 2019 with the results to be provided in an addendum to the Phase II IR.

All analytical data collected during the 2018–2019 remediation and investigation activities are presented and evaluated in this addendum, in conjunction with decision-level data previously associated with AOC 02-011(a)(ii), and are now associated with SWMU 02-014.

1.3 Document Organization

This investigation report addendum is organized in six sections, including this introduction, with multiple supporting appendixes. Section 2 provides an overview of the scope of the activities performed at the site. Section 3 presents an overview of the operational history of the site, summaries of previous investigations, results of the field activities performed during the 2018–2019 investigation, site contamination, evaluation of the nature and extent of contamination, and summaries of human health and ecological risk-screening assessments. Section 4 presents the conclusions of the nature and extent of contaminations based on applicable data and the risk-screening assessments. Section 6 includes a list of references cited and the map data sources used in all the figures and plates.

The appendixes include acronyms, a metric conversion table, and definitions of data qualifiers (Appendix A); field methods (Appendix B); waste management (Appendix C); analytical program descriptions and summaries of data quality (Appendix D); analytical suites and results and analytical reports (Appendix E); risk-screening assessments (Appendix F); and box plots and statistical results (Appendix G).

2.0 SCOPE OF ACTIVITIES

This section presents an overview of activities performed during the implementation of investigation and remediation activities at SWMU 02-014 in 2018–2019. Field activities during previous investigations at AOC 02-011(a) were described in the Phase II IR (N3B 2018, 700091). The field investigation results are presented in detail in section 3 and in the appendixes. The scope of activities for the 2018–2019 investigation included geodetic surveys, surface and shallow-subsurface sampling, soil excavation, health and safety monitoring, waste management activities, and sample analysis.

2.1 Premobilization Activities

Premobilization activities included preparation of work planning documents including a site-specific safety and health plan, site-specific environmental safety and health plan, integrated work document, and quality assurance plan.

2.2 Summary of Field Activities

This section describes the field activities conducted during the 2018–2019 investigation and remediation activities. Additional details regarding the field methods and procedures used to perform these field activities are presented in Appendix B.

2.2.1 Geodetic Survey

Real-time kinematic (RTK) global positioning system (GPS) surveying was conducted to establish the coordinates of sample locations and to lay out planned soil excavation areas. Surveying was performed using Topcon HiPer V Navigation Satellite System antennas coupled with a Topcon FC-5000 Data Collector Controller. This system was used to stake sampling locations, locations to be left unexcavated (i.e., uncontaminated locations), locations excavated, planned excavation boundaries, and pre- and post-excavation topographic elevation data collection, specifically for excavations too deep to safely check via RTK GPS. The surveyed coordinates for all sampling locations not previously reported are presented in Table 2.2-1. All coordinates are expressed as State Plane Coordinate System 83, New Mexico Central, U.S. feet. All surveyed coordinates for sampling locations were submitted for upload to the Environmental Information Management Database.

2.2.2 Field Screening

All samples collected were field screened for radioactivity and some were also screened for volatile organic compounds (VOCs) using a photoionization detector. A radiological control technician (RCT) conducted radiological screening using a ThermoFisher Model SHP-380, with Eberline Model E600 Geiger Counter, for detection of low-energy radiation. All field results for alpha and beta/gamma radioactivity were recorded on the field sample collection log/chain-of-custody (SCL/COC) forms. The SCLs/COC forms are provided on CD in Appendix E. The radiological and VOC screening results not previously reported are presented in Table 2.2-2.

2.2.3 Surface and Shallow-Subsurface Soil Investigation

Surface and shallow-subsurface samples were collected using a stainless-steel hand auger to collect material within the prescribed sampling intervals. For samples collected at depths greater than 3 ft, 4-in. polyvinyl chloride pipe was decontaminated and inserted into the hole to prevent hole collapse and cross-contamination of samples. A stainless-steel bowl and scoop were used to capture the sample from the hand-auger bucket. The sample was then transferred to sterile sample collection jars for transport to the Sample Management Office (SMO).

Quality assurance (QA)/quality control (QC) samples consisted of field duplicates and rinsate blanks. Field duplicate samples were collected at a minimum rate of 1 per 10 investigation samples. Rinsate blanks were also collected at a minimum rate of 1 per 10 investigation samples to confirm decontamination of the sampling equipment.

All sample collection activities were coordinated with the SMO. Upon collection, samples remained at all times in the controlled custody of the field team until delivered to the SMO. Sample custody was then relinquished to the SMO for delivery to a preapproved off-site contract analytical laboratory.

2.2.4 Equipment Decontamination

Between collection of each sample and between sampling locations, all field equipment with the potential to contact sample material (e.g., hand augers, sampling scoops, and bowls) was decontaminated using dry decontamination methods to prevent cross-contamination of samples and locations. Rinsate blanks were used to check the effectiveness of decontamination.

2.2.5 Excavation

Excavation of PCB-contaminated soil was performed during 2018–2019. Results of the previous investigations at SWMU 02-014 were used to define the area requiring excavation. The following cleanup goals were considered in defining the excavation area:

- industrial and recreational soil screening levels (SSLs) (11 mg/kg and 5.53 mg/kg, respectively for Aroclor-1254 and 11.1 mg/kg and 10.3 mg/kg, respectively for Aroclor-1260) for the depth interval 0.0–1.0 ft below ground surface (bgs) and
- the Toxic Substances Control Act (TSCA) bulk PCB remediation waste cleanup level (25 mg/kg for total PCBs) for low-occupancy areas.

Based on consideration of the above goals, an excavation area was defined where all soil within the depth interval 0.0–1.0 ft bgs would be removed to address potential risk to industrial workers and recreational users. This area is shown on Plate 1. Within this area, four additional areas requiring deeper excavation were defined to meet the TSCA cleanup level. Excavation depths in these areas ranged from 4.5 ft bgs to 10.5 ft bgs.

The planned excavation areas were staked in the field and soil was excavated to the required depths using a Yanmar mini-excavator and Volvo excavator. Approximately 255 yd³ of soil was excavated from these areas. Following excavation, confirmatory samples were collected to determine whether additional removal was needed to meet cleanup goals. Based on this evaluation, additional excavation from the interval 0.0–1.0 ft bgs was required to the east of the northeast corner of the original excavation area and to the west at two locations on the west boundary of the original excavation area. One area where additional deeper excavation was required was also identified at the northwest corner of the original excavation from these areas resulted in removal of approximately 27 yd³ of soil.

2.2.6 Health and Safety Measures

Health and safety measures were identified based on the results of previous investigations at SWMU 02-014. Monitoring of airborne particulates was required to monitor for potential exposure to PCBs during soil excavation activities. Airborne particulate measurements were made using a Thermo Andersen MIE Personal Dust Monitor Model 1000. Because heavy equipment was being used for soil excavation, noise monitoring was also conducted using a Quest NoisePro DLX noise level meter.

During excavation of deep interior area I-1, two asbestos-wrapped pipes (assumed to be decommissioned gas and water lines based on conversations with LANL Utilities Management) were exposed within the first 1–2 feet bgs. A "pause work" commenced the afternoon of November 19, 2018, and went to December 03, 2018. During this time, an asbestos abatement plan and integrated work document addendum were developed. The approved asbestos abatement included use of personal protective equipment and collection of airborne asbestos samples. The approved asbestos abatement plan was then implemented and the exposed pipes and wrapping were removed from the excavation and properly packaged and labeled. About 1 yd³ of waste (pipe, asbestos wrapping, contact waste materials) was generated in the process.

2.2.7 Waste Management

The waste streams associated with investigation and remediation activities included excavated soil, contact waste, and asbestos-containing materials. All waste, including excavated soil, was managed in accordance with the project waste characterization strategy form (WCSF). All excavated media at SWMU 02-014 was placed in 5.18-yd³ soft-sided IP-1 bags. IP-1 bags were positioned on pallets for loading and staging. After a bag was sealed, an RCT screened all sides of the bag before releasing the bag for staging at TA-41. Contact waste was stored in labeled 1-gal. plastic bags in the radioactive waste accumulation area on-site until it could be transferred to a 5.18-yd³ soft-sided IP-1 bag. Temporary storage was within a posted radiological waste storage area at TA-41. All waste containers that were staged at TA-41 were covered with tarps for additional protection from the elements.

The management of wastes is described in greater detail in Appendix C. The WCSF is provided in Appendix C (Attachment C-1 on CD).

2.3 Sample Analyses

The SMO shipped all samples to off-site contract analytical laboratories for the requested analyses. All samples collected during the 2018–2019 investigation and remediation activities were submitted for analysis of PCBs.

Analytical methods and summaries of data quality are presented in Appendix D. Analytical results, analytical reports, and SCLs/COCs are included on CD in Appendix E.

2.4 Deviations

The 2018–2019 sampling and remediation activities at SWMU 02-014 were not addressed in the approved Phase II investigation work plan for Middle Los Alamos Canyon Aggregate Area (LANL 2009, 105073; NMED 2009, 105595). The need for these activities was identified as a result of the Phase II sampling performed at AOC 02-011(a)(ii) and the subsequent discovery of SWMU 02-014.

3.0 FIELD INVESTIGATION RESULTS FOR SWMU 02-014

3.1 Site Description and Operational History

SWMU 02-014 consists of three former electrical transformer stations (structures 02-31, 02-45, and 02-51) that served buildings in TA-02 (Plate 1). This site was identified during efforts to discover the source of PCB contamination identified during investigation sampling at storm drain AOC 02-011(a)(ii). Historical records, including engineering drawings and photographs, were reviewed and three potential sources of PCBs were identified. Former structure 02-31 was an electrical transformer station located 40 ft behind building 02-1. The transformer station was built in 1944 and was removed in 1950. Former structure 02-45 was built in 1954 to serve building 02-44. The transformer station consisted of three transformers mounted across two telephone poles approximately 14 ft above the ground. The transformer station was replaced with another transformer station (structure 02-51). Former structure 02-31 and 20 ft southeast of former structure 02-45. Historical records indicated PCB-containing transformer oil had been used at this former transformer station. Structure 02-51 was constructed in 1961 and demolished in 2003.

3.2 Relationship to Other SWMUs and AOCs

SWMU 02-014 is adjacent to and northwest of AOC 02-011(a) segments (i), (ii), and (iii). SWMU 02-014 is the source of the PCB contamination previously detected at AOC 02-011(a).

3.3 Summary of Previous Investigations

No previous investigations have been performed at SWMU 02-014. The area of SWMU 02-014 was previously sampled as part of the investigations performed for AOC 02-011(a).

3.4 Site Contamination

3.4.1 Soil and Rock Sampling

SWMU 02-014 was sampled during 2017 to characterize the areas potentially requiring removal of PCBcontaminated soil. A total of 256 samples were collected at 46 locations around areas where PCB contamination was previously detected during the investigation of AOC 02-011(a). Samples were collected at various depth intervals based on previous results, to a maximum of 20 ft bgs or until refusal. Following evaluation of initial results, additional sampling was performed in 2018 to better characterize extent of potential excavation areas. During January 2018, 102 samples were collected from 21 existing locations and 7 new locations. In September 2018, 6 samples were collected from 5 new locations to refine the area requiring excavation. Based on these results, PCB-contaminated soil was excavated during November and December 2018, and 30 confirmation samples were collected at 10 new locations on the western side of the site.

3.4.2 Soil and Rock Sample Field-Screening Results

No radiological-screening results exceeded twice the daily site background levels and no VOCs were detected.

3.4.3 Soil and Rock Sample Analytical Results

Decision-level data at SWMU 02-014 consist of results from 436 samples collected from 83 locations in 2007, 2010, 2017, and 2018. The 436 samples include 272 soil/Qal, 6 Qbt 3, 7 Qbt 1g, 1 Qct, and 150 Qbo samples. Plate 1 shows the sample locations and Table 3.4-1 presents the samples collected and the analyses requested for SWMU 02-014.

Inorganic Chemicals

A total of 92 samples (70 soil/Qal, 1 Qbt 3, and 21 Qbo) were analyzed for target analyte list (TAL) metals, and 3 soil/Qal samples were analyzed for nitrate, perchlorate, and total cyanide. Table 3.4-2 presents the inorganic chemicals detected or detected above background values (BVs). Plate 2 shows the spatial distribution of inorganic chemicals detected or detected above BVs.

Aluminum was detected above the Qbt 1g, Qct, Qbo BV (3560 mg/kg) in 18 samples with a maximum concentration of 15,000 mg/kg. The Gehan and quantile tests indicated site concentrations of aluminum in tuff are statistically different from background (Figure G-1 and Table G-1). Aluminum is retained as a chemical of potential concern (COPC).

Arsenic was detected above the Qbt 1g, Qct, Qbo BV (0.56 mg/kg) in 18 samples with a maximum concentration of 0.94 mg/kg. The quantile and slippage tests indicated site concentrations of arsenic in tuff are statistically different from background (Figure G-2 and Table G-1). Arsenic is retained as a COPC.

Barium was detected above the Qbt 1g, Qct, Qbo BV (25.7 mg/kg) in 17 samples with a maximum concentration of 96.1 mg/kg. The Gehan and quantile tests indicated site concentrations of barium in tuff are statistically different from background (Figure G-3 and Table G-1). Barium is retained as a COPC.

Cadmium was not detected above the soil and Qbt 1g, Qct, Qbo BVs (0.4 mg/kg for both) but had detection limits (DLs) (0.497 mg/kg to 0.517 mg/kg) above BV in five soil samples and one tuff sample. The DLs were only 0.097 mg/kg and 0.117 mg/kg above the BVs, and the maximum DL is below or equivalent to the three highest concentrations (0.6 mg/kg, 1.4 mg/kg, and 2.6 mg/kg) and the highest DL (2 mg/kg) in the soil background data set. Cadmium is not a COPC.

Calcium was detected above the Qbt 1g, Qct, Qbo BV (1900 mg/kg) in three samples with a maximum concentration of 7260 mg/kg. The quantile and slippage tests indicated site concentrations of calcium in tuff are not statistically different from background (Figure G-4 and Table G-1). Calcium is not a COPC.

Chromium was detected above the Qbt 1g, Qct, Qbo BV (2.6 mg/kg) in 12 samples with a maximum concentration of 14.5 mg/kg. The Gehan and quantile tests indicated site concentrations of chromium in tuff are statistically different from background (Figure G-5 and Table G-1). Chromium is retained as a COPC.

Copper was detected above the soil and Qbt 1g, Qct, Qbo BVs (14.7 mg/kg and 3.96 mg/kg) in one soil sample and two tuff samples with a maximum concentration of 47.5 mg/kg. The Gehan and quantile tests indicated site concentrations of copper in soil are not statistically different from background (Figure G-6 and Table G-2). The Gehan and quantile tests indicated site concentrations of copper in tuff are statistically different from background (Figure G-7 and Table G-1). Copper is retained as a COPC.

Iron was detected above the Qbt 1g, Qct, Qbo BV (3700 mg/kg) in 17 samples with a maximum concentration of 7560 mg/kg. The Gehan and quantile tests indicated site concentrations of iron in tuff are statistically different from background (Figure G-8 and Table G-1). Iron is retained as a COPC.

Magnesium was detected above the Qbt 1g, Qct, Qbo BV (739 mg/kg) in ten samples with a maximum concentration of 1710 mg/kg. The Gehan and quantile tests indicated site concentrations of magnesium in tuff are statistically different from background (Figure G-9 and Table G-1). Magnesium is retained as a COPC.

Manganese was detected above the Qbt 1g, Qct, Qbo BV (189 mg/kg) in eight samples with a maximum concentration of 274 mg/kg. The Gehan and quantile tests indicated site concentrations of manganese in tuff are statistically different from background (Figure G-10 and Table G-1). Manganese is retained as a COPC.

Mercury was detected above the soil BV (0.1 mg/kg) in six samples with a maximum concentration of 0.538 mg/kg. Mercury is retained as a COPC.

Nickel was detected above the Qbt 1g, Qct, Qbo BV (2 mg/kg) in 15 samples with a maximum concentration of 4.37 mg/kg. The quantile test indicated site concentrations of nickel in tuff are statistically different from background (Figure G-11 and Table G-1). Nickel is retained as a COPC.

Nitrate was detected in two samples with a maximum concentration of 1.77 mg/kg. Nitrate is naturally occurring, and the concentrations likely reflect naturally occurring levels. In addition, SWMU 02-014 consists of former electrical transformer stations and is not a source of nitrate. Nitrate is not a COPC.

Perchlorate was detected in one sample at a concentration of 0.000813 mg/kg. Perchlorate is retained as a COPC.

Selenium was detected above the Qbt 3 and Qbt 1g, Qct, Qbo BVs (0.3 mg/kg for both) in 1 Qbt 3 sample and 5 Qbo samples with a maximum concentration of 1.11 mg/kg and was not detected but had DLs (0.403 mg/kg to 2.5 mg/kg) above the soil BV (1.52 mg/kg) and Qbt 1g, Qct, Qbo BV in 15 soil samples and 12 Qbo samples. Selenium is retained as a COPC.

Thallium was not detected above the soil and Qbt 1g, Qct, Qbo BVs (0.73 mg/kg and 1.22 mg/kg) but had a DL (2.5 mg/kg) above BV in three soil samples and two tuff samples. The slippage test indicated site concentrations of thallium in soil are statistically different from background (Figure G-12 and Table G-2). The quantile and slippage tests indicated site concentrations of thallium in tuff are not statistically different from background (Figure G-13 and Table G-1). Thallium is retained as a COPC.

Vanadium was detected above the Qbt 1g, Qct, Qbo BV (4.59 mg/kg) in eight samples with a maximum concentration of 6.43 mg/kg. The Gehan and quantile tests indicated site concentrations of vanadium in tuff are statistically different from background (Figure G-14 and Table G-1). Vanadium is retained as a COPC.

Zinc was detected above the soil BV (48.8 mg/kg) in four samples with a maximum concentration of 105 mg/kg. The quantile and slippage tests indicated site concentrations of zinc in soil are not statistically different from background (Figure G-15 and Table G-2). Zinc is not a COPC.

Organic Chemicals

A total of 353 samples (209 soil/Qal, 5 Qbt 3, 7 Qbt 1g, 1 Qct, and 131 Qbo) were analyzed for PCBs, 3 soil/Qal samples were analyzed for semivolatile organic compounds (SVOCs) and dioxins/furans, and 2 soil/Qal samples were analyzed for VOCs. Table 3.4-3 presents detected organic chemicals other than PCBs, and Table 3.4-4 presents detected PCBs. Figure 3.4-1 shows the spatial distribution of detected organic chemicals other than PCBs, and Plate 3 shows the spatial distribution of PCBs.

Organic chemicals detected at SWMU 02-014 include anthracene; Aroclor-1254; Aroclor-1260; benzo(a)anthracene; benzo(a)pyrene; benzo(b)fluoranthene; benzo(g,h,i)perylene; benzo(k)fluoranthene; chrysene; fluoranthene; 1,2,3,4,6,7,8-heptachlorodibenzodioxin; 1,2,3,4,6,7,8-heptachlorodibenzofuran; 1,2,3,4,7,8,9-heptachlorodibenzofuran; 1,2,3,4,7,8-hexachlorodibenzodioxin; 1,2,3,6,7,8-hexachlorodibenzodioxin; 1,2,3,7,8,9-hexachlorodibenzofuran; 1,2,3,4,7,8-hexachlorodibenzofuran; 1,2,3,6,7,8-hexachlorodibenzofuran; 1,2,3,7,8,9-hexachlorodibenzofuran; 2,3,4,6,7,8-hexachlorodibenzofuran; 1,2,3,4,6,7,8,9-octachlorodibenzofuran; 1,2,3,4,6,7,8,9-octachlorodibenzofuran; 1,2,3,7,8-pentachlorodibenzodioxin; 1,2,3,7,8-pentachlorodibenzofuran; 2,3,4,7,8-pentachlorodibenzofuran; phenanthrene; pyrene; 2,3,7,8-tetrachlorodibenzofuran; and toluene. The detected organic chemicals are retained as COPCs.

Radionuclides

A total of 92 samples (70 soil/Qal, 1 Qbt 3, and 21 Qbo) were analyzed for gamma-emitting radionuclides, isotopic plutonium, isotopic uranium, and strontium-90; 7 soil/Qal samples were analyzed for americium-241; and 3 soil/Qal samples were analyzed for tritium. Table 3.4-5 presents the radionuclides detected or detected above BVs/fallout values (FVs). Figure 3.4-2 shows the spatial distribution of radionuclides detected or detected above BVs/FVs.

Cesium-134 was detected in three soil samples and two tuff samples with a maximum activity of 0.066 pCi/g. Cesium-134 is retained as a COPC.

Cesium-137 was detected in two soil samples below 0.0 ft to 1.0 ft bgs with a maximum activity of 0.759 pCi/g. Cesium-137 is retained as a COPC.

Plutonium-238 was detected in two soil samples below 0.0 ft to 1.0 ft bgs and one tuff sample with a maximum activity of 0.019 pCi/g. Plutonium-238 is retained as a COPC.

Plutonium-239/240 was detected above the soil FV (0.054 pCi/g) in three samples, detected in seven soil samples below 0.0 ft to 1.0 ft bgs, and detected in two tuff samples with a maximum activity of 0.076 pCi/g. Plutonium-239/240 is retained as a COPC.

3.4.4 Nature and Extent of Contamination

The nature and extent of inorganic and organic COPCs at SWMU 02-014 are discussed below.

Inorganic Chemicals

Inorganic COPCs at SWMU 02-014 include aluminum, arsenic, barium, chromium, copper, iron, magnesium, manganese, mercury, nickel, perchlorate, selenium, thallium, and vanadium.

Aluminum was detected above the Qbt 1g, Qct, Qbo BV in 18 samples with a maximum concentration of 15,000 mg/kg. Concentrations increased with depth at locations 02-61435, 02-61439, 02-61442, 02-61450, 02-61451, 02-61452, and 02-61454; did not change substantially with depth (170 mg/kg) at location 02-61445; decreased with depth at all other locations; and decreased downgradient (the concentration in a shallow sample at location 02-61445 was 10,200 mg/kg and below the soil BV [Appendix E, Pivot Tables]). The residential and industrial SSLs were approximately 5 times and 86 times the maximum concentration, respectively (63,000 mg/kg and 1,280,000 mg/kg below the respective SSLs). The lateral extent of aluminum is defined and further sampling for vertical extent is not warranted.

Arsenic was detected above the Qbt 1g, Qct, Qbo BV in 18 samples with a maximum concentration of 0.94 mg/kg. Concentrations decreased with depth and decreased downgradient. The lateral and vertical extent of arsenic are defined. Arsenic concentrations did not vary substantially vertically or horizontally. The range of concentrations detected in soil (0.321 mg/kg to 2.38 mg/kg), Qbt 3 (0.791 mg/kg), and Qbo (0.354 mg/kg to 0.94 mg/kg) were similar and do not indicate any spatial concentration trends. The residential and industrial SSLs were approximately 8 times and 38 times the maximum concentration above BV (0.94 mg/kg). Further sampling for extent of arsenic is not warranted.

Barium was detected above the Qbt 1g, Qct, Qbo BV in 17 samples with a maximum concentration of 96.1 mg/kg. Concentrations increased with depth at locations 02-61435, 02-61442, 02-61446, and 02-61454; decreased with depth at all other locations; and decreased downgradient (the concentrations in shallow samples at locations 02-61443, 02-61445, 02-61450, and 02-61451 were 31.2 mg/kg, 73.9 mg/kg, 29.7 mg/kg, and 69.5 mg/kg, respectively, and below the soil BV [Appendix E, Pivot Tables]).

The residential SSL was approximately 162 times the maximum concentration. The lateral extent of barium is defined and further sampling for vertical extent is not warranted.

Chromium was detected above the Qbt 1g, Qct, Qbo BV in 12 samples with a maximum concentration of 14.5 mg/kg. Concentrations increased with depth at locations 02-61436 and 02-61454, did not change substantially with depth (0.12 mg/kg) at locations 02-61442 and 02-61451, decreased with depth at all other locations, and decreased downgradient (concentrations in shallow samples at locations 02-613763, 02-61435, 02-61439, 02-61442, 02-61446, and 02-61451 were 3.86 mg/kg, 4.92 mg/kg, 3.52 mg/kg, 2.73 mg/kg, 4.16 mg/kg, and 3.15 mg/kg, respectively, and below the soil BV [Appendix E, Pivot Tables]). As described in section 4.2 of the Phase II IR (LANL 2018, 700091), SWMU 02-014 is not a potential source of hexavalent chromium and use of the SSL for trivalent chromium to determine whether additional sampling is warranted is appropriate. The residential SSL for trivalent chromium (117,000 mg/kg) was approximately 8070 times the maximum concentration. The lateral extent of chromium is defined and further sampling for vertical extent is not warranted.

Copper was detected above the soil and Qbt 1g, Qct, Qbo BVs in one soil sample and two tuff samples with a maximum concentration of 47.5 mg/kg. Concentrations increased with depth at location 02-61438, decreased with depth at location 02-61435, and decreased downgradient. The residential SSL was approximately 66 times the maximum concentration. The lateral extent of copper is defined and further sampling for vertical extent is not warranted.

Iron was detected above the Qbt 1g, Qct, Qbo BV in 17 samples with a maximum concentration of 7560 mg/kg. Concentrations increased with depth at locations 02-61442, 02-61451, 02-61452, and 02-61454; did not change substantially with depth (60 mg/kg and 80 mg/kg) at locations 02-61443 and 02-61450; decreased with depth at all other locations; and decreased downgradient (concentrations in shallow samples at locations 02-61435, 02-61436, 02-61438, 02-61439, 02-61441, 02-61443, and 02-61450 were 12,800 mg/kg, 5380 mg/kg, 6140 mg/kg, 4850 mg/kg, 7020 mg/kg, 4060 mg/kg, and 4720 mg/kg, respectively, and below the soil BV [Appendix E, Pivot Tables]). The residential and industrial SSLs were approximately 7 times and 120 times the maximum concentration, respectively (47,200 mg/kg and 972,000 mg/kg below the respective SSLs). The lateral extent of iron is defined and further sampling for vertical extent is not warranted.

Magnesium was detected above the Qbt 1g, Qct, Qbo BV in ten samples with a maximum concentration of 1710 mg/kg. Concentrations increased with depth at locations 02-61435, 02-61438, 02-61442, 02-61452, and 02-61454; did not change substantially with depth (12 mg/kg) at location 02-61451; decreased with depth at all other locations; and decreased downgradient (the concentration in a shallow sample at location 02-61451 was 839 mg/kg and below the soil BV [Appendix E, Pivot Tables]). The residential essential nutrient SSL was approximately 12,200 times the maximum concentration. The lateral extent of magnesium is defined and further sampling for vertical extent is not warranted.

Manganese was detected above the Qbt 1g, Qct, Qbo BV in eight samples with a maximum concentration of 274 mg/kg. Concentrations increased with depth at locations 02-61435, 02-61436, 20-61442, and 02-61445; did not change substantially with depth (11 mg/kg and 29 mg/kg) at locations 02-61451 and 02-61454; decreased with depth at all other locations; and decreased downgradient (concentrations in shallow samples at locations 02-61438, 02-61450, 02-61451, and 02-61454 were 236 mg/kg, 247 mg/kg, 218 mg/kg, and 245 mg/kg, respectively, and below the soil BV [Appendix E, Pivot Tables]). The residential SSL was approximately 38 times the maximum concentration. The lateral extent of manganese is defined and further sampling for vertical extent is not warranted.

Mercury was detected above the soil BV in six samples with a maximum concentration of 0.538 mg/kg. Concentrations decreased with depth at all locations and decreased downgradient. The lateral and vertical extent of mercury are defined.

Nickel was detected above the Qbt 1g, Qct, Qbo BV in 15 samples with a maximum concentration of 4.37 mg/kg. Concentrations increased with depth at locations 02-61435, 02-61442, and 02-61454; did not change substantially with depth (0.05 mg/kg to 0.2 mg/kg) at locations 02-61436, 02-61439, and 02-61451; decreased with depth at all other locations; and decreased downgradient. The residential SSL was approximately 357 times the maximum concentration. The lateral extent of nickel is defined and further sampling for vertical extent is not warranted.

Perchlorate was detected in one sample at a concentration of 0.000813 mg/kg. Concentrations increased with depth and increased laterally at location 02-600387. The detected concentration was below the estimated quantitation limit (EQL). Perchlorate was not detected in samples collected downgradient of location 02-600387 at AOC 02-011(a)(i,ii,iii,iv,v,vi) location 02-600407. The residential SSL was approximately 67,400 times the detected concentration. The lateral extent of perchlorate is defined and further sampling for vertical extent is not warranted.

Selenium was detected above the Qbt 3 and Qbt 1g, Qct, Qbo BVs in 1 Qbt 3 sample and 5 Qbo samples with a maximum concentration of 1.11 mg/kg and was not detected but had DLs (0.403 mg/kg to 2.5 mg/kg) above the soil BV and Qbt 1g, Qct, Qbo BV in 15 soil samples and 12 Qbo samples. Concentrations increased with depth at locations 02-61442, 02-61443, and 02-61446; decreased with depth at all other locations; and decreased downgradient (concentrations in shallow samples at locations 02-61432, 02-61435, and 02-61445 were 0.375 mg/kg, 1.15 mg/kg, and 0.537 mg/kg, respectively, and below the soil BV [Appendix E, Pivot Tables]). The residential SSL was approximately 352 times the maximum concentration and approximately 156 times the maximum DL. The lateral extent of selenium is defined and further sampling for vertical extent is not warranted.

Thallium was not detected above the soil and Qbt 1g, Qct, Qbo BVs but had a DL (2.5 mg/kg) above the soil and Qbt 1g, Qct, Qbo BVs in three soil samples and two tuff samples. The maximum DL was approximately 3.2 times the residential SSL, and the industrial SSL was approximately 5.2 times the maximum DL. Samples with DLs above BV were collected at locations 02-61436, 02-61441, 02-61444, and 02-61454. At locations 02-61436, 02-61441, and 02-61444, thallium was not detected or detected above BVs in any other samples and DLs were all below BV (0.043 mg/kg to 0.14 mg/kg [Appendix E, Pivot Tables]). At location 02-61454, thallium was not detected (with a DL of 2.5 mg/kg) in samples from 2.0 to 3.0 ft bgs and 4.0 to 5.0 ft bgs and was detected at 0.053 mg/kg (below BV) in a sample from 6.0 to 7.0 ft bgs (Appendix E, Pivot Tables). At all other locations, thallium was not detected in 53 samples, with DLs ranging from 0.027 mg/kg to 0.312 mg/kg and was not detected in 53 samples, with DLs ranging from 0.044 mg/kg to 0.316 mg/kg (Appendix E, Pivot Tables). All concentrations and DLs at these locations were below the residential SSL. Further sampling for extent of thallium is not warranted.

Vanadium was detected above the Qbt 1g, Qct, Qbo BV in eight samples with a maximum concentration of 6.43 mg/kg. Concentrations increased with depth at locations 02-61442 and 02-61454, decreased with depth at all other locations, and decreased downgradient. The residential SSL was approximately 61 times the maximum concentration. The lateral extent of vanadium is defined and further sampling for vertical extent is not warranted.

Organic Chemicals

Organic COPCs at SWMU 02-014 include anthracene; Aroclor-1254; Aroclor-1260; benzo(a)anthracene; benzo(a)pyrene; benzo(b)fluoranthene; benzo(g,h,i)perylene; benzo(k)fluoranthene; chrysene; fluoranthene; 1,2,3,4,6,7,8-heptachlorodibenzodioxin; 1,2,3,4,6,7,8-heptachlorodibenzofuran;

- fluorantnene; 1,2,3,4,6,7,8-neptachiorodibenzodioxin; 1,2,3,4,6,7,8-neptachiorodibenzotura
- 1,2,3,4,7,8,9-heptachlorodibenzofuran; 1,2,3,4,7,8-hexachlorodibenzodioxin;
- $1,2,3,6,7,8-hexa chlorodibenzodioxin;\ 1,2,3,7,8,9-hexa chlorodibenzodioxin;$
- 1,2,3,4,7,8-hexachlorodibenzofuran; 1,2,3,6,7,8-hexachlorodibenzofuran;
- 1,2,3,7,8,9-hexachlorodibenzofuran; 2,3,4,6,7,8-hexachlorodibenzofuran; indeno(1,2,3-cd)pyrene;
- 1,2,3,4,6,7,8,9-octachlorodibenzodioxin; 1,2,3,4,6,7,8,9-octachlorodibenzofuran;
- 1,2,3,7,8-pentachlorodibenzodioxin; 1,2,3,7,8-pentachlorodibenzofuran;

2,3,4,7,8-pentachlorodibenzofuran; phenanthrene; pyrene; 2,3,7,8-tetrachlorodibenzofuran; and toluene.

The polycyclic aromatic hydrocarbons (PAHs) anthracene; benzo(a)anthracene; benzo(a)pyrene; benzo(b)fluoranthene; benzo(g,h,i)perylene; benzo(k)fluoranthene; chrysene; fluoranthene; indeno(1,2,3-cd)pyrene; phenanthrene; and pyrene were each detected in 2 samples at location 02-600387 at concentrations ranging from 0.00727 mg/kg to 0.0984 mg/kg. Concentrations of pyrene decreased with depth and concentrations of all other PAHs did not change substantially with depth (0.00176 mg/kg to 0.0455 mg/kg). Concentrations increased laterally at location 02-600387. Of the 22 detected concentrations, 11 were below EQLs. The residential SSLs ranged from approximately 20 times to 1,930,000 times the maximum concentrations, and the residential SSLs of all PAHs except benzo(a)anthracene, benzo(a)pyrene, and indeno(1,2,3-cd)pyrene were more than 100 times the maximum concentration. The residential SSL for benzo(a)anthracene was approximately 30 times the maximum concentration, and the industrial SSL was approximately 626 times the maximum concentration (0.0516 mg/kg). The residential SSL for benzo(a)pyrene was approximately 20 times the maximum concentration, and the industrial SSL was approximately 428 times the maximum concentration (0.0551 mg/kg). The residential SSL for indeno(1,2,3-cd)pyrene was approximately 61 times the maximum concentration, and the industrial SSL was approximately 1280 times the maximum concentration (0.0252 mg/kg). Further sampling for extent of PAHs is not warranted.

Aroclor-1254 was detected in 31 samples with a maximum concentration of 7.11 mg/kg. Concentrations increased with depth at locations 02-613122, 02-613124, and 02-613700 and did not change substantially with depth (0.0082 mg/kg) at location 02-61490; only 1 depth was sampled at locations 02-613699 and 02-61492; concentrations decreased with depth at all other locations; and concentrations decreased downgradient. The residential SSL was approximately 1.5 times and the industrial SSL was approximately 14 times the maximum concentration where vertical extent is not defined (0.779 mg/kg at location 02-613700). Aroclor-1254 does not pose an unacceptable risk under the industrial and recreational scenarios (Appendix F, Tables F-4.2-1 through F-4.2-4). Lateral extent of Aroclor-1254 is defined and further sampling for vertical extent is not warranted.

Aroclor-1260 was detected in 317 samples with a maximum concentration of 23.9 mg/kg. Concentrations increased with depth at locations 02-600449, 02-612452, 02-613626, 02-613668, 02-613700, 02-61479, 02-61488, 02-61489, and 02-61553 and did not change substantially with depth (0.0078 mg/kg to 0.033 mg/kg) at locations 02-61436, 02-61445, 02-61486, and 02-61490; only 1 depth was sampled at locations 02-612451, 02-613292, 02-613699, 02-61478, 02-61492, and 02-61539 through 02-61543; and concentrations decreased with depth at all other locations. Concentrations decreased downgradient. Vertical extent is defined at locations where only 1 depth was sampled by decreasing concentrations in deeper samples from nearby locations. Concentrations in samples collected at 0.0 to 1.0 ft bgs, 2.0 to 3.0 ft bgs, 4.0 ft to 4.2 ft bgs, and 4.0 ft to 5.0 ft bgs at location 02-613289 were 1520 mg/kg, 7.51 mg/kg, 0.68 mg/kg, and 33.1 mg/kg, respectively, and concentrations decreased with depth at this location. These samples were excavated and the results are not included in Table 3.4-4 but are included in Appendix E.

The maximum concentration at locations where concentrations increased with depth or did not change substantially with depth (3.3 mg/kg at location 02-613626) was approximately 1.4 times the residential SSL, and the industrial SSL was approximately 3.4 times this concentration. Although concentrations decreased with depth at location 02-613289, the concentration of Aroclor-1260 in the deepest sample at this location (11.0 ft to 11.25 ft bgs) is 1.9 times the industrial SSL and 8.5 times the residential SSL. Further sampling at this location is not warranted based on the results in deeper samples at adjacent locations 02-61432 (11 ft from location 02-613289), 02-61440 (9 ft from location 02-613289), 02-61441 (10 ft from location 02-613289), 02-61448 (6 ft from location 02-613289), and 02-61455 (13 ft from location 02-613289). Aroclor-1260 concentrations decreased with depth at all 5 locations. A total of 15 samples were collected from these 5 locations at depths greater than the deepest sample at location 02-613289 (14.0 to 15.0 ft bgs, 16.0 to 17.0 ft bgs, and 19.0 to 20.0 ft bgs at each location). As shown on Plate 3, Aroclor-1260 was detected in 7 of these 15 samples at concentrations ranging from 0.00316 mg/kg to 0.0707 mg/kg, all more than 2 orders of magnitude less than the concentration in the deepest sample at location 02-613289. Aroclor-1260 was not detected in the samples from 19.0 ft to 20.0 ft bgs at any of the 5 locations except location 02-61448, where it was detected at 0.0186 mg/kg. Aroclor-1260 does not pose an unacceptable risk under the industrial and recreational scenarios (Appendix F, Tables F-4.2-1 and F-4.2-3). Lateral extent of Aroclor-1260 is defined and further sampling for vertical extent is not warranted.

A total of 16 dioxin/furan congeners were each detected in 2 or 3 samples with maximum concentrations ranging from 0.000000204 mg/kg to 0.00123 mg/kg. Concentrations at location 02-600387 did not change substantially with depth (0.000000028 mg/kg to 0.00003 mg/kg) or decreased with depth, and only 1 depth was sampled at location 02-600449. Concentrations of 2,3,4,7,8-pentachlorodibenzofuran and 2,3,7,8-tetrachlorodibenzofuran increased downgradient at location 02-600449 and concentrations of all other congeners decreased downgradient or did not change substantially downgradient (0.00000001 mg/kg to 0.00000155 mg/kg). The maximum concentrations were converted to toxicity equivalency concentrations (TECs) using congener-specific toxicity equivalency factors (NMED 2017, 602273). The residential SSL for 2.3,7,8-tetrachlorodibenzodioxin (2.3,7,8-TCDD [not detected but the only dioxin with an SSL]) ranged from approximately 13 times to 2080 times the TECs and was more than 100 times the TECs for all congeners except 1,2,3,4,6,7,8-heptachlorodibenzodioxin; 1,2,3,4,7,8-hexachlorodibenzofuran; 2,3,4,7,8-pentachlorodibenzofuran; and 2,3,7,8-tetrachlorodibenzofuran. The residential SSL for 2,3,7,8-TCDD was approximately 30 times the TEC, and the industrial SSL for 2,3,7,8-TCDD was approximately 143 times the TEC for 1,2,3,4,6,7,8-heptachlorodibenzodioxin (0.00000166 mg/kg). The residential SSL for 2,3,7,8-TCDD was approximately 76 times the TEC, and the industrial SSL for 2,3,7,8-TCDD was approximately 371 times the TEC for 1,2,3,4,7,8-hexachlorodibenzofuran (0.000000641 mg/kg). The residential SSL for 2,3,7,8-TCDD was approximately 13 times the TEC, and the industrial SSL for 2,3,7,8-TCDD was approximately 62 times the TEC for 2,3,4,7,8-pentachlorodibenzofuran (0.00000384 mg/kg). The residential SSL for 2,3,7,8-TCDD was approximately 77 times the TEC, and the industrial SSL for 2,3,7,8-TCDD was approximately 372 times the TEC for 2,3,7,8-tetrachlorodibenzofuran (0.000000639 mg/kg). Further sampling for extent of dioxin and furan congeners is not warranted.

Toluene was detected in one sample at a concentration of 0.000465 mg/kg. Only one depth was sampled at location 02-600387 and concentrations decreased downgradient. The residential SSL is approximately 11,200,000 times the maximum concentration. Lateral extent of toluene is defined and further sampling for vertical extent is not warranted.

Radionuclides

Radionuclide COPCs at SWMU 02-014 include cesium-134, cesium-137, plutonium-238, and plutonium-239/240.

Cesium-134 was detected in three soil samples and two tuff samples with a maximum activity of 0.066 pCi/g. Activities increased with depth at location 02-61452; decreased with depth at locations 02-61442, 02-61454, and 02-61455; and decreased downgradient. The residential SAL was approximately 76 times the maximum activity. Lateral extent of cesium-134 is defined and further sampling for vertical extent is not warranted.

Cesium-137 was detected in two soil samples below 0.0 ft to 1.0 ft bgs with a maximum activity of 0.759 pCi/g. Activities decreased with depth at location 02-61451 and decreased downgradient. The lateral and vertical extent of cesium-137 are defined.

Plutonium-238 was detected in two soil samples below 0.0 ft to 1.0 ft bgs and one tuff sample with a maximum activity of 0.019 pCi/g. Activities did not change substantially with depth (0.003 pCi/g) at location 02-61450, decreased with depth at location 02-61428, and decreased downgradient. The residential SAL was approximately 4420 times the maximum activity. Lateral extent of plutonium-238 is defined and further sampling for vertical extent is not warranted.

Plutonium-239/240 was detected above the soil FV (0.054 pCi/g) in three samples, detected in seven soil samples below 0.0 ft to 1.0 ft bgs, and detected in two tuff samples with a maximum activity of 0.076 pCi/g. Activities increased with depth at locations 02-61452 and 02-61453, did not change substantially with depth (0.007 pCi/g) at location 02-61448, decreased with depth at all other locations, and decreased downgradient. The residential SAL was approximately 1040 times the maximum activity. Lateral extent of plutonium-239/240 is defined and further sampling for vertical extent is not warranted.

Summary of Nature and Extent

The lateral and vertical extent of inorganic and organic COPCs is defined or no further sampling for extent is warranted at SWMU 02-014.

3.5 Summary of Human Health Risk Screening

Industrial Scenario

The total excess cancer risk for the industrial scenario is 5×10^{-6} , which is less than the NMED target risk level of 1×10^{-5} (NMED 2017, 602273). The industrial hazard index (HI) is 0.06, which is less than the NMED target HI of 1 (NMED 2017, 602273). The total dose is 0.0009 mrem/yr, which is less than the target dose of 25 mrem/yr as authorized by DOE Order 458.1.

Recreational Scenario

The total excess cancer risk for the recreational scenario is 5×10^{-6} , which is less than the NMED target risk level of 1×10^{-5} (NMED 2017, 602273). The recreational HI is 0.2, which is less than the NMED target HI of 1 (NMED 2017, 602273). The total dose is 0.0008 mrem/yr, which is less than the target dose of 25 mrem/yr as authorized by DOE Order 458.1.

Construction Worker Scenario

The construction worker HI is 0.8, which is less than the NMED target risk level of 1 (NMED 2017, 602273). The residential exposure scenario is protective of construction workers for carcinogenic risk and total dose.

Residential Scenario

The total excess cancer risk for the residential scenario is 1×10^{-5} , which is equivalent to the NMED target risk level of 1×10^{-5} (NMED 2017, 602273). The residential HI is 0.6, which is less than the NMED target HI of 1 (NMED 2017, 602273). The total dose is 2 mrem/yr, which is less than the target dose of 25 mrem/yr as authorized by DOE Order 458.1.

Based on the risk-screening assessment results, no potential unacceptable risks or doses exist for the industrial, recreational, construction worker, and residential scenarios at SWMU 02-014.

3.6 Summary of Ecological Risk Screening

SWMU 02-014 is within the TA-02 core area. Ecological risk for the TA-02 core area was evaluated in the "Phase II Investigation Report for Middle Los Alamos Canyon Aggregate Area, Revision 2" (N3B 2018, 700091). Based on evaluations of the minimum ESLs, HI analyses, potential effects to populations (individuals for threatened or endangered [T&E] species), lowest observed adverse effect level (LOAEL) analyses, the relationship of detected concentrations and screening levels to background concentrations, and results of site-specific ecological risk studies, the Phase II IR concluded no potential ecological risks exist for the TA-02 core area, which includes SWMU 02-014.

4.0 CONCLUSIONS

4.1 Nature and Extent of Contamination

Based on the evaluation of the sampling data, the nature and extent of contamination have been defined and/or no further sampling for extent is warranted for SWMU 02-014.

4.2 Summary of Risk-Screening Assessments

4.2.1 Human Health Risk-Screening Assessment

There were no potential unacceptable risks or doses for SWMU 02-014 under the industrial, recreational, and residential scenarios. There was no potential noncarcinogenic risk to construction workers, and the residential scenario demonstrated protection of construction workers for carcinogenic risk and dose.

SWMU 02-014 is not accessible by the public. Therefore, an as low as reasonably achievable (ALARA) evaluation for radiological exposure to the public is not currently required. An ALARA evaluation will be conducted should DOE plan to release this area.

4.2.2 Ecological Risk-Screening Assessment

Ecological risk was evaluated collectively for sites within the TA-02 core area, including SWMU 02-014, in the "Phase II Investigation Report for Middle Los Alamos Canyon Aggregate Area, Revision 2" (N3B 2018, 700091). Based on evaluations of the minimum ESLs, HI analyses, potential effects to populations (individuals for T&E species), LOAEL analyses, the relationship of detected concentrations and screening levels to background concentrations, and results of site-specific ecological risk studies conducted within the TA-02 core area, no potential ecological risks exist for the TA-02 core area, which includes SWMU 02-014.

5.0 RECOMMENDATIONS

SWMU 02-014 was found to pose no potential unacceptable risks to human health under the industrial, recreational, construction worker, and residential scenarios and to ecological receptors. Residual PCB contamination is below the TSCA cleanup level of 25 mg/kg for low-occupancy areas. This site is appropriate for corrective action complete without controls.

6.0 REFERENCES AND MAP DATA SOURCES

6.1 References

The following reference list includes documents cited in this report. Parenthetical information following each reference provides the author(s), publication date, and ERID, ESHID, or EMID. This information is also included in text citations. ERIDs were assigned by the Laboratory's Associate Directorate for Environmental Management (IDs through 599999); ESHIDs were assigned by the Laboratory's Associate Directorate for Environment, Safety, and Health (IDs 600000 through 699999); and EMIDs are assigned by Newport News Nuclear BWXT-Los Alamos, LLC (N3B) (IDs 700000 and above). IDs are used to locate documents in N3B's Records Management System and in the Master Reference Set. The NMED Hazardous Waste Bureau and N3B maintain copies of the Master Reference Set. The set ensures that NMED has the references to review documents. The set is updated when new references are cited in documents.

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6.2 Map Data Sources

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Drainage: WQH Drainage_arc; Los Alamos National Laboratory, ENV Water Quality and Hydrology Group; 1:24,000 Scale Data; 03 June 2003.

Existing sampling locations: Point Feature Locations of the Environmental Restoration Project Database, er_location_ids_pnt; Los Alamos National Laboratory, Waste and Environmental Services Division, EP2010-0035; 21 January 2010.

Former fences: Phase II Investigation Work Plan for Middle Los Alamos Canyon Aggregate Area, Revision 1; Los Alamos National Laboratory, Environmental Programs; Report LA-UR-09-1206, EP2009-0080; February 2009.

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Primary Landscape Features, ksl_landscape_arc; Los Alamos National Laboratory, KSL Site Support Services, Planning, Locating and Mapping Section; 06 January 2004; as published 28 May 2009.

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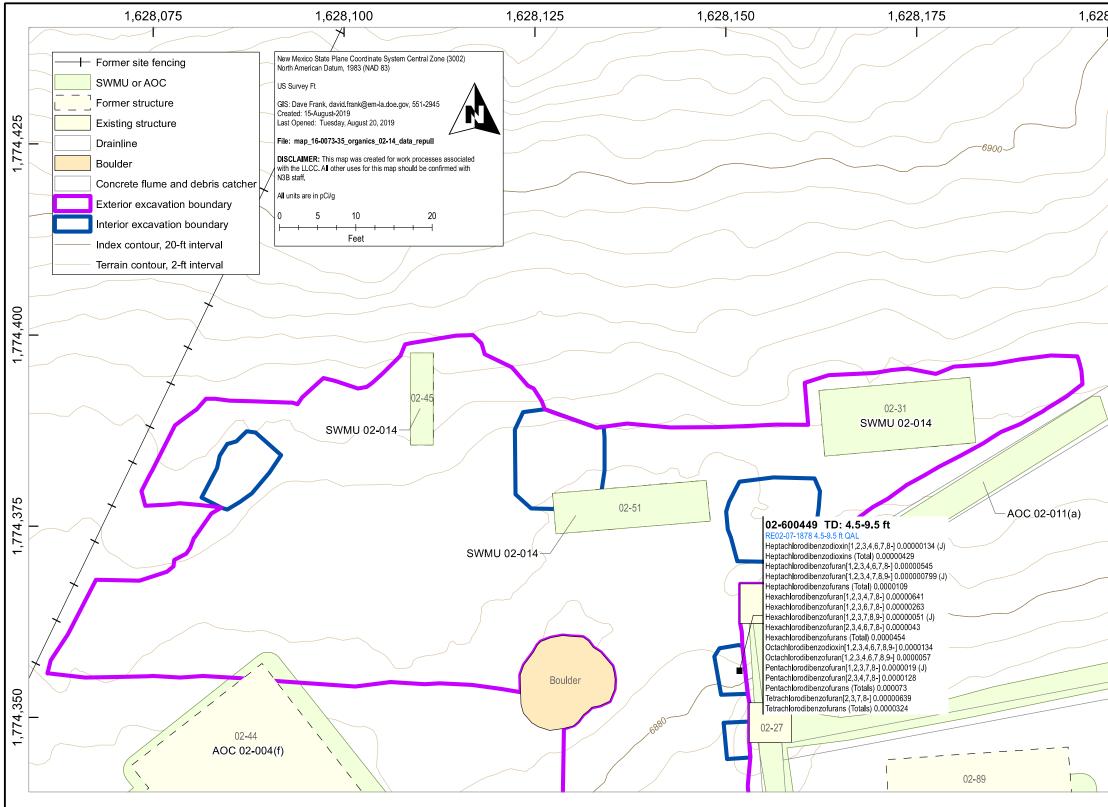


Figure 3.4-1 Organic chemicals other than PCBs detected at SWMU 02-014

8,200	1,628,225	5
\overline{V}		6900
	02-600387 TD: 0-2.6 ft	
	RE02-07-1572 0-0.5 ft SOIL	
	Anthracene 0.00903 (J)	
	Benzo(a)anthracene 0.0516 Benzo(a)pyrene 0.0551	
	Benzo(b)fluoranthene 0.0702	
	Benzo(g,h,i)perylene 0.0271 (J)	
	Benzo(k)fluoranthene 0.041	
	Chrysene 0.0598	
	Fluoranthene 0.0984	\sim
	Heptachlorodibenzodioxin[1,2,3,4,6,7,8-] 0.00016	
	Heptachlorodibenzodioxins (Total) 0.001	
	Heptachlorodibenzofuran[1,2,3,4,6,7,8-] 0.0000475 Heptachlorodibenzofuran[1,2,3,4,7,8,9-] 0.00000235 (J)	
	Heptachlorodibenzofurans (Total) 0.000205	
	Hexachlorodibenzodioxin[1,2,3,4,7,8-] 0.00000071 (J)	
	Hexachlorodibenzodioxin[1,2,3,6,7,8-] 0.00000332	
	Hexachlorodibenzodioxin[1,2,3,7,8,9-] 0.0000013 (J)	
	Hexachlorodibenzodioxins (Total) 0.0000622	
	Hexachlorodibenzofuran[1,2,3,4,7,8-] 0.00000486	
	Hexachlorodibenzofuran[1,2,3,6,7,8-] 0.00000166 (J) Hexachlorodibenzofuran[1,2,3,7,8,9-] 0.0000005 (J)	
	Hexachlorodibenzofuran[2,3,4,6,7,8-] 0.0000003(0)	
	Hexachlorodibenzofurans (Total) 0.0000561	
	Indeno(1,2,3-cd)pyrene 0.0252 (J)	
	Octachlorodibenzodioxin[1,2,3,4,6,7,8,9-] 0.0012	
	Octachlorodibenzofuran[1,2,3,4,6,7,8,9-] 0.00022	
	Pentachlorodibenzodioxin[1,2,3,7,8-] 0.000000204 (J)	
	Pentachlorodibenzodioxins (Total) 0.00000234	~ /
	Pentachlorodibenzofuran[1,2,3,7,8-] 0.00000101 (J) Pentachlorodibenzofuran[2,3,4,7,8-] 0.00000652	
	Pentachlorodibenzofurans (Totals) 0.0000449 (J)	
/	Phenanthrene 0.0317 (J)	
	Pyrene 0.0771	
	Tetrachlorodibenzofuran[2,3,7,8-] 0.00000377	
	Tetrachlorodibenzofurans (Totals) 0.000017	
	RE02-07-1573 2-2.6 ft QAL	
	Anthracene 0.00727 (J)	
	Benzo(a)anthracene 0.032 (J) Benzo(a)pyrene 0.0332 (J)	
	Benzo(b)fluoranthene 0.0424	
	Benzo(g,h,i)perylene 0.0222 (J)	
	Benzo(k)fluoranthene 0.0219 (J)	
	Chrysene 0.0386	
	Fluoranthene 0.0529	
	Heptachlorodibenzodioxin[1,2,3,4,6,7,8-] 0.000166	
	Heptachlorodibenzodioxins (Total) 0.00108 Heptachlorodibenzofuran[1,2,3,4,6,7,8-] 0.0000472	
	Heptachlorodibenzofuran[1,2,3,4,7,8,9-] 0.0000472 Heptachlorodibenzofuran[1,2,3,4,7,8,9-] 0.00000182 (J)	
	Heptachlorodibenzofurans (Total) 0.000235 6880	
	Hexachlorodibenzodioxin[1,2,3,4,7,8-] 0.000000738 (J)	
	Hexachlorodibenzodioxin[1,2,3,6,7,8-] 0.0000037	
	Hexachlorodibenzodioxin[1,2,3,7,8,9-] 0.00000091 (J)	
	Hexachlorodibenzodioxins (Total) 0.0000743	
	Hexachlorodibenzofuran[1,2,3,4,7,8-] 0.00000182 (J)	
	Hexachlorodibenzofuran[1,2,3,6,7,8-] 0.000000621 (J)	
	Hexachlorodibenzofuran[2,3,4,6,7,8-] 0.00000981 (J) . Hexachlorodibenzofurans (Total) 0.0000446	
	Indeno(1,2,3-cd)pyrene 0.0184 (J)	
	Octachlorodibenzodioxin[1,2,3,4,6,7,8,9-] 0.00123	
	Octachlorodibenzofuran[1,2,3,4,6,7,8,9-] 0.000248	
	Pentachlorodibenzodioxin[1,2,3,7,8-] 0.000000156 (J)	
	Pentachlorodibenzodioxins (Total) 0.0000007	
	Pentachlorodibenzofuran[2,3,4,7,8-] 0 00000155 (J)	
	Pentachlorodibenzofurans (Totals) 0.0000102 (J)	
	Phenanthrene 0.0212 (J)	
	Pyrene 0.0423 Tetrachlorodibenzofuran[2,3,7,8-] 0.000000867 (J)	
	Tetrachlorodibenzofurans (Totals) 0.00000232	
	Toluene 0.000465 (J)	
J	02-90	
(02-30	

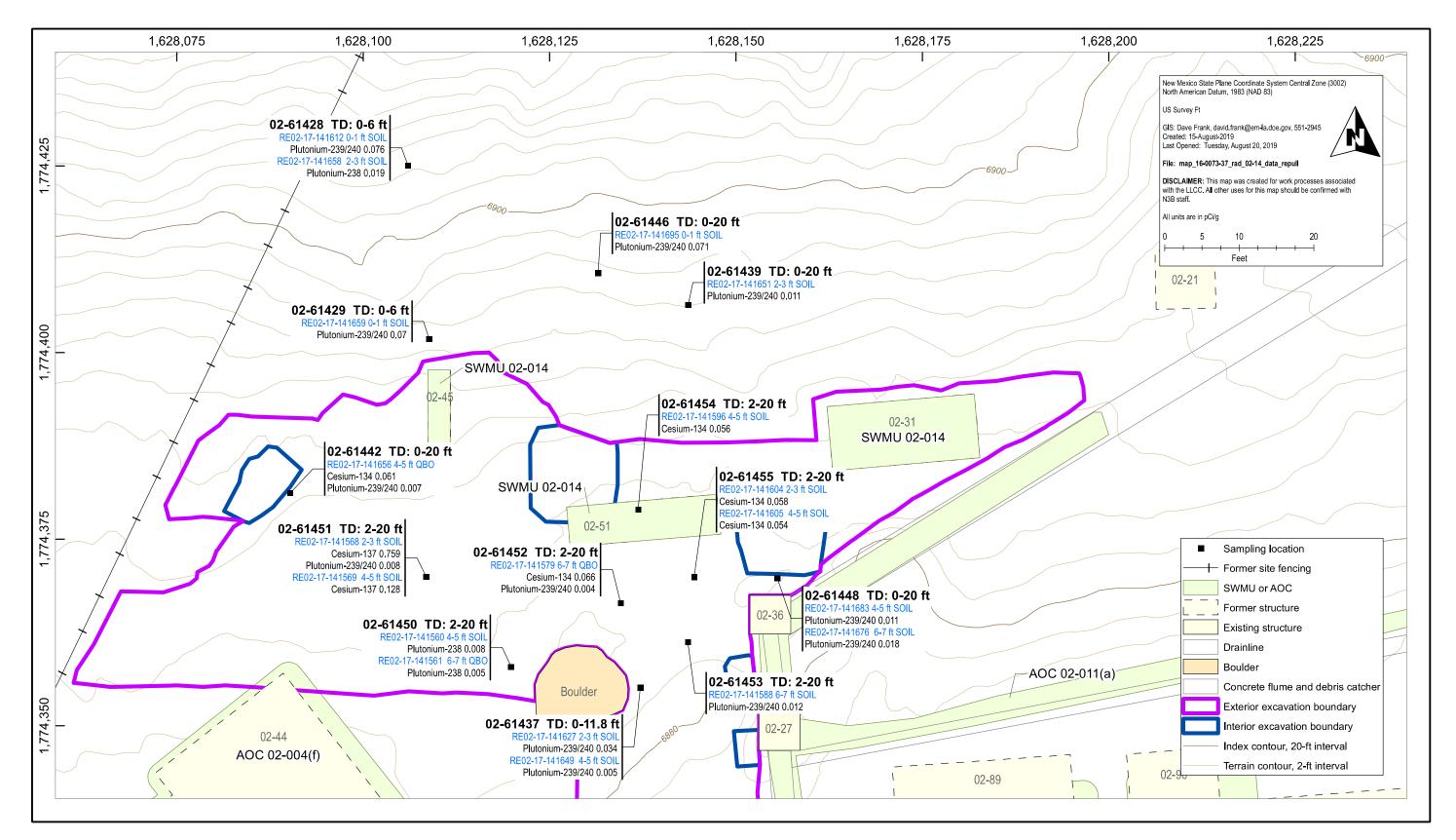


Figure 3.4-2 Radionuclides detected or detected above BVs/FVs at SWMU 02-014

Location ID	Easting (ft)	Northing (ft)
02-600387	1628200.95	1774392.37
02-600449	1628151.78	1774356.13
02-612451	1628151.78	1774360.13
02-612452	1628151.78	1774352.13
02-612453	1628147.78	1774356.13
02-613001	1628152.37	1774347.01
02-613002	1628151.78	1774364.13
02-613122	1628147.78	1774348.13
02-613124	1628147.78	1774352.13
02-613287	1628144.19	1774344.67
02-613288	1628144.48	1774365.63
02-613289	1628156.32	1774375.53
02-613292	1628179.77	1774384.08
02-613626	1628139.68	1774352.22
02-613627	1628139.24	1774371.95
02-613667	1628128.70	1774374.69
02-613668	1628132.69	1774348.76
02-613699	1628127.92	1774383.70
02-613700	1628135.67	1774352.38
02-613761	1628116.07	1774386.04
02-613762	1628116.07	1774393.22
02-61428	1628106.00	1774425.05
02-61429	1628108.86	1774401.79
02-61432	1628147.47	1774382.07
02-61435	1628098.51	1774371.75
02-61436	1628118.98	1774370.88
02-61437	1628137.16	1774355.10
02-61438	1628141.28	1774417.69
02-61439	1628143.58	1774406.38
02-61440	1628157.63	1774384.36
02-61441	1628165.80	1774378.78
02-61442	1628090.23	1774381.14
02-61443	1628087.15	1774367.30
02-61444	1628108.57	1774357.65
02-61445	1628122.45	1774351.33
02-61446	1628131.53	1774410.66
02-61447	1628141.00	1774394.41
02-61448	1628155.53	1774369.75
02-61450	1628119.84	1774357.92
02-61451	1628108.41	1774370.00

Table 2.2-1Surveyed Coordinates of Sample Locations at SWMU 02-014

Table 2.2-1 (continued)					
Location ID	Easting (ft)	Northing (ft)			
02-61452	1628134.53	1774366.40			
02-61453	1628143.55	1774361.22			
02-61454	1628136.89	1774378.94			
02-61455	1628144.37	1774369.91			
02-61474	1628147.87	1774337.53			
02-61475	1628134.91	1774331.46			
02-61476	1628115.98	1774350.53			
02-61477	1628091.65	1774346.90			
02-61478	1628095.26	1774363.51			
02-61479	1628080.82	1774358.13			
02-61480	1628070.92	1774380.57			
02-61481	1628082.09	1774394.89			
02-61482	1628099.42	1774396.68			
02-61483	1628122.09	1774402.79			
02-61486	1628175.20	1774361.63			
02-61487	1628177.53	1774383.54			
02-61488	1628164.78	1774393.19			
02-61489	1628177.37	1774392.79			
02-61490	1628161.70	1774409.30			
02-61491	1628176.65	1774405.29			
02-61492	1628157.65	1774417.01			
02-61493	1628168.42	1774418.46			
02-61494	1628185.18	1774411.65			
02-61526	1628195.96	1774398.92			
02-61528	1628115.36	1774326.09			
02-61529	1628099.68	1774323.29			
02-61530	1628071.55	1774346.83			
02-61537	1628099.45	1774356.78			
02-61538	1628077.29	1774354.58			
02-61539	1628074.25	1774359.06			
02-61540	1628073.85	1774365.97			
02-61541	1628086.43	1774383.55			
02-61542	1628185.49	1774387.19			
02-61543	1628185.81	1774394.23			
02-61544	1628079.96	1774386.23			
02-61545	1628069.32	1774365.16			
02-61547	1628077.23	1774350.25			
02-61548	1628086.82	1774388.51			
02-61549	1628083.30	1774380.59			

Table 2.2-1 (continued)

Location ID	Easting (ft)	Northing (ft)
02-61550	1628064.40	1774365.03
02-61551	1628064.79	1774359.02
02-61552	1628074.78	1774376.40
02-61553	1628070.73	1774361.03

Table 2.2-1 (continued)

Table 2.2–2
Field–Screening Results for Samples Collected at SWMU 02-014

				Beta/Gamma	
Location ID	Sample ID	Depth (ft)	Alpha (dpm)ª	(dpm)	VOCs (ppm)
02-613001	RELA-18-151143	6–7	35	3130	b
02-613001	RELA-18-151144	8–9	41	3110	—
02-613001	RE02-17-141631	10–11	13	2100	0.0
02-613001	RE02-17-141633	16–17	20	2030	0.0
02-613001	RE02-17-141634	19–20	6	2020	0.0
02-613287	RELA-18-151133	2–3	16	1271	—
02-613287	RELA-18-151134	4–5	60	2570	—
02-613287	RELA-18-151135	6–7	10	2510	—
02-613287	RE02-17-136051	8–9	27	1850	0.0
02-613289	RE02-17-141635	6–7	20	2030	0.0
02-613289	RELA-18-151137	6–7	21	2730	—
02-613289	RELA-18-151138	9–10	32	2410	—
02-613289	RELA-18-151139	11–11.25	16	2480	—
02-613667	RELA-18-151122	2–3	41	3230	—
02-613667	RELA-18-151123	4–5	35	3170	—
02-613667	RELA-18-151124	6–7	53	3230	—
02-613667	RELA-18-151125	8–9	41	3250	—
02-613668	RELA-18-151127	2–3	16	2510	—
02-613668	RELA-18-151128	4–5	37	2530	—
02-613668	RELA-18-151129	6–7	53	2750	—
02-613668	RELA-18-151130	8–9	32	2680	—
02-613762	RE02-17-141642	2–3	40	2070	0.0
02-613762	RELA-18-151117	2–3	37	2710	_
02-613762	RE02-17-141643	4–5	40	2290	0.0
02-613762	RELA-18-151118	4.25–5	53	2780	—
02-613762	RELA-18-151119	6–7	26	2780	—
02-613762	RE02-17-141644	6–7	40	2030	0.0

Location ID	Sample ID	Depth (ft)	Alpha (dpm)	Beta/Gamma (dpm)	VOCs (ppm)
02-613762	RELA-18-151120	8–9	21	2790	_
02-61428	RE02-17-135776	0–1	67	2480	0.0
02-61428	RE02-17-141612	0–1	67	2480	0.0
02-61428	RE02-17-135777	2–3	74	2570	0.0
02-61428	RE02-17-141658	2–3	74	2570	0.0
02-61428	RE02-17-141645	4–5	22	2330	0.0
02-61428	RE02-17-135778	4–5	22	2330	0.0
02-61428	RE02-17-135779	5–6	33	2560	0.0
02-61428	RE02-17-141621	5–6	33	2560	0.0
02-61429	RE02-17-135780	0–1	47	1812	0.0
02-61429	RE02-17-141659	0–1	47	1812	0.0
02-61429	RE02-17-141646	2–3	61	1840	0.0
02-61429	RE02-17-135781	2–3	61	1840	0.0
02-61429	RE02-17-135782	4–5	56	2380	0.0
02-61429	RE02-17-141622	4–5	56	2380	0.0
02-61429	RE02-17-141613	5–6	28	2360	0.0
02-61429	RE02-17-135783	5–6	28	2360	0.0
02-61432	RE02-17-141660	0–1	20	2350	0.0
02-61432	RE02-17-141623	2–3	33	2450	0.0
02-61432	RELA-18-151045	2–3	42	2480	
02-61432	RELA-18-151046	4–5	48	2530	_
02-61432	RE02-17-141647	4–5	33	2520	0.0
02-61432	RE02-17-141614	6–7	27	2660	0.0
02-61432	RELA-18-151047	6.75–7	24	2410	_
02-61432	RELA-18-151048	8-8.75	48	2770	_
02-61432	RE02-17-135853	11–12	27	2300	0.0
02-61432	RE02-17-135854	14–15	26.7	2630	0.0
02-61432	RE02-17-135855	16–17	33	2650	0.0
02-61432	RE02-17-135856	19–20	40	2810	0.0
02-61435	RE02-17-141615	0–1	33	2110	0.0
02-61435	RE02-17-141624	2–3	101	2260	0.0
02-61435	RELA-18-151057	2–3	37	2370	_
02-61435	RELA-18-151058	4–5	48	2400	 _
02-61435	RE02-17-141625	4–5	50	2200	0.0
02-61435	RELA-18-151059	6–7	48	2650	_
02-61435	RE02-17-141648	6–7	33	2340	0.0
02-61435	RE02-17-135893	11–12	16	2170	0.0
02-61435	RE02-17-135894	14–15	56	2320	0.0

Location ID	Sample ID	Depth (ft)	Alpha (dpm)	Beta/Gamma (dpm)	VOCs (ppm)
02-61435	RE02-17-135895	16–17	11	2320	0.0
02-61435	RE02-17-135896	19–20	11	2320	0.0
02-61436	RE02-17-141661	0–1	40	2150	0.0
02-61436	RE02-17-141626	2–3	92	1712	0.0
02-61436	RELA-18-151062	2–3	47	3220	_
02-61436	RELA-18-151063	4–5	53	3290	
02-61436	RE02-17-141616	4–5	28	1602	0.0
02-61436	RE02-17-141617	6–7	42	1726	0.0
02-61436	RELA-18-151064	6–7	35	3300	
02-61436	RE02-17-135901	8–9	49	1688	0.0
02-61436	RE02-17-135902	11–12	35	1753	0.0
02-61436	RE02-17-135903	14–15	42	1705	0.0
02-61436	RE02-17-135904	16–17	7	1527	0.0
02-61436	RE02-17-135905	19–20	42	1956	0.0
02-61437	RE02-17-141618	0–1	27	1923	0.0
02-61437	RE02-17-141627	2–3	34	1913	0.0
02-61437	RE02-17-144637	2–3	114	2300	—
02-61437	RE02-17-144638	4–5	87	2350	—
02-61437	RE02-17-141649	4–5	47	1920	0.0
02-61437	RE02-17-141662	6–7	34	1822	0.0
02-61437	RELA-18-151065	6–7	5	2570	—
02-61437	RELA-18-151066	8–9	16	2520	—
02-61437	RELA-18-151067	11–11.8	17	2820	_
02-61438	RE02-17-141663	0–1	56	2290	0.0
02-61438	RE02-17-135916	0–1	56	2290	0.0
02-61438	RE02-17-135917	2–3	73	2340	0.0
02-61438	RE02-17-141650	2–3	73	2340	0.0
02-61438	RE02-17-135918	4–5	16	2770	0.0
02-61438	RE02-17-141628	4–5	16	2270	0.0
02-61438	RE02-17-141619	6–7	44	2150	0.0
02-61438	RE02-17-135919	6–7	44	2150	0.0
02-61438	RE02-17-135920	8–9	28	2110	0.0
02-61438	RE02-17-135921	11–12	50	2190	0.0
02-61438	RE02-17-135922	14–15	27	2120	0.0
02-61438	RE02-17-135923	16–17	6	2190	0.0
02-61438	RE02-17-135924	19–20	20	1880	0.0
02-61439	RE02-17-135925	0–1	47	1902	0.0
02-61439	RE02-17-141664	0–1	47	1902	0.0

Table 2.2-2 (continu	ued)
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Location ID	Sample ID	Depth (ft)	Alpha (dpm)	Beta/Gamma (dpm)	VOCs (ppm)
02-61439	RE02-17-135926	2–3	68	2120	0.0
02-61439	RE02-17-141651	2–3	68	2120	0.0
02-61439	RE02-17-141629	4–5	34	2040	0.0
02-61439	RE02-17-135933	4–5	34	2040	0.0
02-61439	RE02-17-141620	6–7	6	1844	0.0
02-61439	RE02-17-135928	6–7	6	1849	0.0
02-61439	RE02-17-135929	8–9	34	2120	0.0
02-61439	RE02-17-135930	11–12	27	2060	0.0
02-61439	RE02-17-135931	14–15	54	1783	0.0
02-61439	RE02-17-135932	16–17	27	1884	0.0
02-61439	RE02-17-135927	19–20	13	2020	0.0
02-61440	RE02-17-141665	0–1	20	1952	0.0
02-61440	RE02-17-141652	2–3	20	2020	0.0
02-61440	RELA-18-151069	2–3	24	2680	_
02-61440	RELA-18-151070	4–5	48	2540	_
02-61440	RE02-17-141653	4–5	27	1851	0.0
02-61440	RE02-17-141666	6–7	6	1958	0.0
02-61440	RELA-18-151071	6–7	12	2520	_
02-61440	RELA-18-151072	8–9	18	2400	_
02-61440	RE02-17-135939	11–12	13	1952	0.0
02-61440	RE02-17-135940	14–15	20	1974	0.0
02-61440	RE02-17-135941	16–17	34	2120	0.0
02-61440	RE02-17-135942	19–20	6	2170	0.0
02-61441	RE02-17-141668	0–1	20	1847	0.0
02-61441	RE02-17-141667	2–3	13	1819	0.0
02-61441	RE02-17-144671	2–3	33	2570	_
02-61441	RE02-17-144672	4–5	87	2630	_
02-61441	RE02-17-141654	4–5	47	1973	0.0
02-61441	RE02-17-141655	6–7	13	1766	0.0
02-61441	RELA-18-151043	6–7	48	2500	_
02-61441	RELA-18-151073	8–8.5	18	2400	_
02-61441	RE02-17-135948	11–12	13	2070	0.0
02-61441	RE02-17-135949	14–15	34	1945	0.0
02-61441	RE02-17-135950	16–17	13	1847	0.0
02-61441	RE02-17-135951	19–20	20	2100	0.0
02-61442	RE02-17-141670	0–1	33	2220	0.0
02-61442	RE02-17-141669	2–3	50	2360	0.0
02-61442	RELA-18-151075	2–3	21	2270	_

Location ID	Sample ID	Depth (ft)	Alpha (dpm)	Beta/Gamma (dpm)	VOCs (ppm)
02-61442	RELA-18-151076	4–5	26	2330	_
02-61442	RE02-17-141656	4–5	78	2110	0.0
02-61442	RELA-18-151077	6–7	32	2420	_
02-61442	RE02-17-141657	6–7	44	2240	0.0
02-61442	RELA-18-151078	8–9	64	2470	—
02-61442	RE02-17-135957	11–12	39	2200	0.0
02-61442	RE02-17-135958	14–15	50	2510	0.0
02-61442	RE02-17-135959	16–17	73	2490	0.0
02-61442	RE02-17-135960	19–20	28	2450	0.0
02-61443	RELA-18-151084	11–12	10	2520	_
02-61443	RELA-18-151080	2–3	48	2470	—
02-61443	RE02-17-141678	2–3	20	2410	0.0
02-61443	RE02-17-141685	4–5	67	2300	0.0
02-61443	RELA-18-151081	4–5	64	2690	—
02-61443	RELA-18-151082	6–7	26	2720	_
02-61443	RE02-17-141785	6–7	60	2350	0.0
02-61443	RELA-18-151083	8–9	58	2670	_
02-61444	RE02-17-141693	0–1	57	1613	0.0
02-61444	RE02-17-141686	2–3	35	1640	0.0
02-61444	RELA-18-151086	2–3	47	3090	_
02-61444	RELA-18-151087	4–5	53	3340	_
02-61444	RELA-18-151088	6–7	59	3370	_
02-61444	RE02-17-141672	6–7	28	1733	0.0
02-61444	RELA-18-151089	8–8.85	59	3230	_
02-61444	RE02-17-136003	11–12	27	2000	0.0
02-61444	RE02-17-136004	14–15	6	1864	0.0
02-61444	RE02-17-136005	16–17	41	1994	0.0
02-61444	RE02-17-136006	19–20	27	1840	0.0
02-61445	RE02-17-141673	0–1	54	1682	0.0
02-61445	RE02-17-136007	0–1	54	1682	0.0
02-61445	RE02-17-136010	6–7	61	1949	0.0
02-61445	RE02-17-141694	6–7	61	1949	0.0
02-61445	RE02-17-136011	8–9	47	1970	0.0
02-61445	RE02-17-136012	11–12	13	1899	0.0
02-61445	RE02-17-136015	14–20	27	2000	_
02-61446	RE02-17-139388	0–1	41	1885	0.0
02-61446	RE02-17-141695	0–1	61	1949	0.0
02-61446	RE02-17-139389	2–3	54	1766	0.0

Location ID	Sample ID	Depth (ft)	Alpha (dpm)	Beta/Gamma (dpm)	VOCs (ppm)
02-61446	RE02-17-141688	2–3	54	1766	0.0
02-61446	RE02-17-141681	4-4.5	34	1812	0.0
02-61446	RE02-17-139390	4-4.5	34	1812	0.0
02-61446	RE02-17-139391	6–7	47	1996	0.0
02-61446	RE02-17-141674	6–7	47	1996	0.0
02-61446	RE02-17-139392	8–9	12	2530	0.0
02-61446	RE02-17-139393	11–12	27	2530	0.0
02-61446	RE02-17-139394	14–15	94	2270	0.0
02-61446	RE02-17-139395	16–17	33	2240	0.0
02-61446	RE02-17-139396	19–20	47	2390	0.0
02-61447	RE02-17-141696	0–1	27	1783	0.0
02-61447	RELA-18-151037	0–1	27	2650	
02-61447	RE02-17-141689	2–3	27	2150	0.0
02-61447	RELA-18-151038	2–3	21	2680	—
02-61447	RE02-17-141682	4–5	75	2011	0.0
02-61447	RELA-18-151039	4–5	74	2730	_
02-61447	RE02-17-141675	6–7	34	2050	0.0
02-61447	RELA-18-151040	6–7	8	2037	_
02-61447	RELA-18-151041	8–9	6	2025	_
02-61447	RE02-17-139402	11–12	40	1874	0.0
02-61447	RE02-17-139403	14–15	20	1866	0.0
02-61447	RE02-17-139404	16–17	34	2080	0.0
02-61447	RE02-17-139405	19–20	34	1923	0.0
02-61448	RE02-17-141697	0–1	6	1945	0.0
02-61448	RELA-18-151050	2–3	26	2490	
02-61448	RELA-18-151051	4–5	21	2660	_
02-61448	RE02-17-141683	4–5	54	1601	0.0
02-61448	RE02-17-141676	6–7	13	1647	0.0
02-61448	RELA-18-151052	6–7	26	2560	_
02-61448	RELA-18-151053	8–9	42	2670	_
02-61448	RELA-18-151054	11–12	32	2600	_
02-61448	RELA-18-151055	13–14	6	2024	 _
02-61448	RE02-17-139413	16–17	34	1903	0.0
02-61448	RE02-17-139414	19–20	27	2020	0.0
02-61450	RE02-17-141559	2–3	54	1633	0.0
02-61450	RELA-18-151091	2–3	29	3050	—
02-61450	RELA-18-151092	4–5	41	2970	1_
02-61450	RE02-17-141560	4–5	54	1791	0.0

Table 2.2-2 (continued)

Location ID	Sample ID	Depth (ft)	Alpha (dpm)	Beta/Gamma (dpm)	VOCs (ppm)
02-61450	RE02-17-141561	6–7	27	1899	0.0
02-61450	RE02-17-141562	8–9	34	1987	0.0
02-61450	RE02-17-141563	11–12	47	1840	0.0
02-61450	RE02-17-141564	14–15	82	1808	0.0
02-61450	RE02-17-141565	16–17	6	1899	0.0
02-61450	RE02-17-141566	19–20	61	1924	0.0
02-61451	RE02-17-141568	2–3	74	2570	0.0
02-61451	RELA-18-151094	2–3	41	3360	_
02-61451	RELA-18-151095	4-5	59	3370	
02-61451	RE02-17-141569	4–5	27	1877	0.0
02-61451	RE02-17-141570	6–7	13	1794	0.0
02-61451	RELA-18-151096	6.25–7	23	3280	<u> </u>
02-61451	RELA-18-151097	8–9	59	3290	
02-61451	RE02-17-141572	11–12	20	1988	0.0
02-61451	RE02-17-141573	14–15	13	1877	0.0
02-61451	RE02-17-141574	16–17	13	1784	0.0
02-61451	RE02-17-141575	19-20	34	2070	0.0
02-61452	RE02-17-141577	2–3	0	1747	0.0
02-61452	RELA-18-151099	2–3	24	2460	
02-61452	RELA-18-151100	4–5	48	2540	_
02-61452	RE02-17-141578	4–5	20	2010	0.0
02-61452	RE02-17-141579	6–7	13	1754	0.0
02-61452	RELA-18-151101	6–7	41	2380	_
02-61452	RELA-18-151102	8–9	36	2300	
02-61452	RE02-17-141581	11–12	13	1949	0.0
02-61452	RE02-17-141582	14–15	6	2020	0.0
02-61452	RE02-17-141583	16–17	20	1855	0.0
02-61452	RE02-17-141584	19–20	54	1909	0.0
02-61453	RE02-17-141586	2–3	54	1887	0.0
02-61453	RELA-18-151104	2–3	23	2359	_
02-61453	RELA-18-151105	4-5	31	2390	
02-61453	RE02-17-141587	4–5	20	1945	0.0
02-61453	RE02-17-141588	6–7	27	1967	0.0
02-61453	RE02-17-141692	6–7	27	1967	0.0
02-61453	RELA-18-151106	6–7	31	2960	<u> </u>
02-61453	RELA-18-151107	8–9	37	2365	1_
02-61453	RE02-17-141590	11–12	47	2000	0.0
02-61453	RE02-17-141592	16–17	20	1866	0.0

Location ID	Sample ID	Depth (ft)	Alpha (dpm)	Beta/Gamma (dpm)	VOCs (ppm)
02-61453	RE02-17-141593	19–20	13	1920	0.0
02-61454	RE02-17-141595	2–3	20	2020	0.0
02-61454	RELA-18-151109	2–3	36	2650	_
02-61454	RELA-18-151110	4–5	18	2640	_
02-61454	RE02-17-141596	4–5	27	1905	0.0
02-61454	RE02-17-141597	6–7	61	1920	0.0
02-61454	RE02-17-141598	8–9	0	2126	0.0
02-61454	RE02-17-141599	11–12	20	1916	0.0
02-61454	RE02-17-141600	14–15	1	2080	0.0
02-61454	RE02-17-136031	16–17	20	1895	0.0
02-61454	RE02-17-141602	19–20	13	2080	0.0
02-61455	RE02-17-141608	11–12	129	2000	0.0
02-61455	RE02-17-141609	14–15	13	1905	0.0
02-61455	RE02-17-141610	16–17	6	2000	0.0
02-61455	RE02-17-141611	19–20	27	2070	0.0
02-61455	RE02-17-141604	2–3	6	1981	0.0
02-61455	RELA-18-151112	2–3	36	2630	_
02-61455	RELA-18-151113	4–5	36	2510	_
02-61455	RE02-17-141605	4–5	40	1967	0.0
02-61455	RE02-17-141606	6–7	13	1949	0.0
02-61455	RELA-18-151114	6–7	30	2660	_
02-61455	RELA-18-151115	8–8.75	30	2530	_
02-61474	RE02-17-145060	2–3	20	1454	_
02-61474	RE02-17-145072	4–5	20	1686	_
02-61474	RE02-17-145084	6–7	34	1549	_
02-61475	RE02-17-145049	0–1	34	1388	_
02-61475	RE02-17-145061	2–3	41	1514	_
02-61475	RE02-17-145073	4–5	20	1486	_
02-61476	RE02-17-145050	0–1	13	1644	
02-61476	RE02-17-145062	2–3	13	1472	_
02-61476	RE02-17-145074	4–5	34	1552	_
02-61476	RE02-17-145086	6–7	20	1626	_
02-61476	RE02-17-145098	8–9	13	1430	 _
02-61476	RE02-17-145110	9–10	41	1742	 _
02-61477	RE02-17-145051	0–1	54	1556	_
02-61477	RE02-17-145063	2–3	20	1507	_
02-61478	RE02-17-145064	2–3	20	1423	_
02-61479	RE02-17-145065	2–3	6	1580	1_

Location ID	Sample ID	Depth (ft)	Alpha (dpm)	Beta/Gamma (dpm)	VOCs (ppm)
02-61479	RE02-17-145077	4–5	20	1563	_
02-61480	RE02-17-145054	0–1	41	1640	_
02-61480	RE02-17-145066	2–3	13	1735	_
02-61480	RE02-17-145078	4–5	27	1591	—
02-61480	RE02-17-145090	6–7	20	1491	_
02-61480	RE02-17-145102	8–8.5	34	1594	_
02-61481	RE02-17-145055	0–1	34	1871	—
02-61481	RE02-17-145067	2–3	13	1549	_
02-61481	RE02-17-145079	4–5	61	1640	—
02-61481	RE02-17-145091	6–7	20	1700	—
02-61481	RE02-17-145103	8–9	27	1703	_
02-61481	RE02-17-145115	9–10	47	1402	_
02-61482	RE02-17-145056	0–1	68	1573	—
02-61482	RE02-17-145068	2–3	136	1524	_
02-61482	RE02-17-145080	4–5	34	1672	_
02-61482	RE02-17-145092	6–7	34	1658	_
02-61482	RE02-17-145116	8–9	20	1419	_
02-61482	RE02-17-145104	9–10	34	1542	_
02-61483	RE02-17-145057	0–1	47	1780	_
02-61483	RE02-17-145069	2–3	34	1598	_
02-61483	RE02-17-145081	4–5	47	1577	_
02-61483	RE02-17-145093	6–7	41	1682	—
02-61483	RE02-17-145105	8–9	20	1700	_
02-61483	RE02-17-145117	9–10	54	1549	—
02-61486	RE02-17-145120	0–1	102	1482	_
02-61486	RE02-17-145130	2–3	41	1426	—
02-61487	RE02-17-145131	2–3	41	1770	—
02-61487	RE02-17-145141	4–5	47	1696	—
02-61487	RE02-17-145151	5–5.5	47	1698	—
02-61488	RE02-17-145132	2–3	39	2090	_
02-61488	RE02-17-145142	4-4.1	67	2130	_
02-61489	RE02-17-145133	2–3	56	2350	—
02-61489	RE02-17-145143	4–5	39	2370	_
02-61489	RE02-17-145153	6–7	41	1100	_
02-61489	RE02-17-145163	7–7.5	76	716	_
02-61490	RE02-17-145124	0–1	34	1381	_
02-61490	RE02-17-145134	2–3	20	1416	_
02-61491	RE02-17-145125	0–1	106	2480	_

Location ID	Sample ID	Depth (ft)	Alpha (dpm)	Beta/Gamma (dpm)	VOCs (ppm)
02-61491	RE02-17-145135	2–3	61	2610	_
02-61491	RE02-17-145145	4–5	33	2380	_
02-61492	RE02-17-145126	0–0.5	20	1566	_
02-61493	RE02-17-145127	0–1	61	1623	_
02-61493	RE02-17-145137	2–3	13	1777	—
02-61493	RE02-17-145147	4-4.5	27	1524	—
02-61494	RE02-17-145128	0–1	61	1559	_
02-61494	RE02-17-145138	2–3	82	1637	—
02-61526	RELA-18-151015	0–1	42	2540	—
02-61526	RELA-18-151020	2–3	58	2430	—
02-61526	RELA-18-151021	4–5	96	2390	—
02-61526	RELA-18-151022	6–7	16	2500	_
02-61528	RELA-18-151017	0–1	27	2390	—
02-61528	RELA-18-151023	2–3	22	2470	—
02-61528	RELA-18-151024	4–5	5	2590	
02-61529	RELA-18-151018	0–1	67	2470	_
02-61529	RELA-18-151027	2–3	50	2590	
02-61529	RELA-18-151028	4-4.65	39	2560	
02-61530	RELA-18-151019	0–1	56	2810	
02-61530	RELA-18-151030	2–3	37	2680	
02-61530	RELA-18-151031	4–5	50	2610	
02-61530	RELA-18-151032	6–7	45	2790	
02-61537	RELA-18-151036	11–12	42	1360	
02-61537	RELA-18-151033	2–3	28	2490	
02-61537	RELA-18-151034	4–5	33	2510	
02-61537	RELA-18-151035	6–7	42	2580	—
02-61537	RELA-18-151263	8–9	32	2580	
02-61538	RELA-18-161238	0–0.9	NDA ^c	1700	_
02-61538	RELA-18-161242	1–2	NDA	1600	
02-61539	RELA-18-161239	1–2	NDA	1500	
02-61540	RELA-18-161244	1–2	NDA	1500	
02-61541	RELA-18-164444	6–7	NDA	NDA	_
02-61542	RELA-18-161248	1–2	NDA	1200	 _
02-61543	RELA-18-161249	1–1.5	NDA	1400	 _
02-61544	RELA-18-164455	2–3	NDA	NDA	_
02-61544	RELA-18-164465	4.25–5	NDA	NDA	_
02-61544	RELA-18-164475	6–7	NDA	NDA	_
02-61545	RELA-18-164456	2–3	NDA	NDA	

Location ID	Sample ID	Depth (ft)	Alpha (dpm)	Beta/Gamma (dpm)	VOCs (ppm)
02-61545	RELA-18-164466	4–5	NDA	NDA	
02-61545	RELA-18-164476	6–7	NDA	NDA	
02-61547	RELA-18-164448	0–1	NDA	NDA	_
02-61547	RELA-18-164458	2–3	NDA	NDA	_
02-61547	RELA-18-164468	4–5	NDA	NDA	—
02-61547	RELA-18-164478	6–7	NDA	NDA	—
02-61548	RELA-18-164459	2–3	NDA	NDA	—
02-61548	RELA-18-164469	4-4.5	NDA	NDA	_
02-61548	RELA-18-164479	6–7	NDA	NDA	_
02-61549	RELA-18-164470	4–5	NDA	NDA	_
02-61549	RELA-18-164480	6–7	NDA	NDA	_
02-61550	RELA-18-164451	0–1	NDA	NDA	_
02-61550	RELA-18-164461	2–3	NDA	NDA	_
02-61550	RELA-18-164471	4–5	NDA	NDA	_
02-61550	RELA-18-164481	6–7	NDA	NDA	_
02-61551	RELA-18-164462	2–3	NDA	NDA	_
02-61551	RELA-18-164472	4–5	NDA	NDA	—
02-61551	RELA-18-164482	6–7	NDA	NDA	_
02-61552	RELA-18-164453	0–1	NDA	NDA	—
02-61552	RELA-18-164463	2–3	NDA	NDA	—
02-61552	RELA-18-164473	4–5	NDA	NDA	_
02-61552	RELA-18-164483	6–7	NDA	NDA	—
02-61553	RELA-18-164464	2–3	NDA	NDA	—
02-61553	RELA-18-164474	4–5	NDA	NDA	
02-61553	RELA-18-164484	6–7	NDA	NDA	_

Table 2.2-2 (continued)

^a dpm = Disintegrations per minute.

^b — = Measurement not performed.

^c NDA = No detectable activity.

Table 3.4-1Samples Collected and Analyses Requested at SWMU 02-014

Sample ID	Location ID	Depth (ft)	Media	Americium-241	Nitrate	Gamma-emitting Radionuclides	Tritium	Isotopic Plutonium	Isotopic Uranium	TAL Metals	PCBs	Perchlorate	Strontium-90	SVOCs	VOCs	Dioxins/Furans	Cyanide (Total)
RE02-07-1572	02-600387	0–0.5	ALLH	07-543 ^a	07-542	07-543	07-543	07-543	07-543	07-542	07-541	07-542	07-543	07-541	b	07-530	07-542
RE02-07-1573	02-600387	2–2.6	QAL	07-543	07-542	07-543	07-543	07-543	07-543	07-542	07-541	07-542	07-543	07-541	07-541	07-530	07-542
RE02-07-1878	02-600449	4.5–9.5	QAL	07-956	07-956	07-956	07-956	07-956	07-956	07-956	07-956	07-956	07-956	07-956	07-956	07-955	07-956
RE02-10-22130	02-600449	6–6.2	ALLH	—	—	—	—	—	—	—	10-4285	—	-	-	—	—	—
RE02-10-22133	02-612451	6–6.2	ALLH	—	—	_	—	—	—	—	10-4285	—	-	-	—	—	—
RE02-10-22136	02-612452	6–6.2	ALLH	_	—	—	_	_	_	_	10-4285	—	—	—	—	—	—
RE02-10-22137	02-612452	8–8.2	ALLH	—	—	—	_	_	_	—	10-4285	—	_	_	_	_	—
RE02-10-22139	02-612453	6–6.2	ALLH	_	—	_	_	_	_	_	10-4285	—	_	_	_	_	_
RE02-10-22140	02-612453	8–8.2	ALLH	—	—	—	—	_	_	—	10-4285	—	—	—	—	—	
RE02-10-26105	02-613001	6–6.5	ALLH	—	—	—	—	—	—	—	10-4454	—	 _	 _	—	—	—
RELA-18-151143	02-613001	6–7	ALLH	—	—	—	—	—	—	—	2018-1518	—	—	 _	—	—	—
RE02-10-26106	02-613001	7–7.5	ALLH	—	—	—	—	—	—	—	10-4454	—	—	—	—	—	—
RELA-18-151144	02-613001	8–9	ALLH	—	—	—	—	—	—	—	2018-1518	—	—	 _	—	—	_
RE02-17-141631	02-613001	10–11	QBO	_	—	_	_	_	—	—	2017-2020	—	-	-	_	_	_
RE02-17-141633	02-613001	16–17	QBO	_	—	_	_	_	—	—	2017-2020	—	-	-	_	_	
RE02-17-141634	02-613001	19–20	QBO	_	—	_	_	_	—	_	2017-2020	—	-	-	_	-	
RE02-10-26107	02-613002	6–6.5	ALLH	_	—	_	_	_	—	_	10-4454	—	-	-	_	_	
RE02-10-26108	02-613002	8–8.5	ALLH	_	—	_	_	_	—	_	10-4454	—	-	-	_	_	_
RE02-10-26638	02-613122	2–2.2	ALLH	_	—	_	_	_	_	_	10-4707	—	—	—	—	_	<u> </u>
RE02-10-26639	02-613122	4-4.2	ALLH	_	—	_	_	_	_	_	10-4707	—	—	—	—	_	<u> </u>
RE02-10-26640	02-613124	6–6.2	ALLH	_	_	_	_	_	_	_	10-4707	_	_	_	_	_	1
RE02-10-26641	02-613124	8–8.2	ALLH	_	_	_	_	_	_	_	10-4707	_	_	_	_	_	1
RELA-18-151133	02-613287	2–3	ALLH	_	_	_	_	_	_	_	2018-1518	_	_	_	_	_	1
RELA-18-151134	02-613287	4–5	ALLH	_	_	_	_	_	_	_	2018-1518	_	_	_	_	_	_
RE02-11-315	02-613287	6–6.2	ALLH	_	_	_	_	_	_	_	11-186	_	_	_	_	_	
RELA-18-151135	02-613287	6–7	ALLH	_		_			_		2018-1518	_	_	_		_	
RE02-11-316	02-613287	8–8.2	ALLH	—	—	<u> </u>	 		1_	_	11-186	—	_	—	<u> </u>	—	<u>+</u>
RE02-17-136051	02-613287	8–9	QBO	_	—	_	 	_	1_	_	2017-2087	—	_	—	<u> </u>	—	<u>+</u>
RE02-11-317	02-613288	6–6.2	ALLH	_	_	_	_	_	_	-	11-186	—	_	-	_	 _	_
RE02-11-318	02-613288	8–8.2	ALLH	_		_	_	<u> </u>	1_	_	11-186	—	_	_	1_	1_	1
RE02-11-320	02-613289	6–6.2	ALLH	_	_	_	_	_	_	_	11-209	—	_	_	_	1_	_

Table 3.4-1 (continued)

Sample ID	Location ID	Depth (ft)	Media	Americium-241	Nitrate	Gamma-emitting Radionuclides	Tritium	Isotopic Plutonium	Isotopic Uranium	TAL Metals	PCBs	Perchlorate	Strontium-90	SVOCs	VOCs	Dioxins/Furans	Cyanide (Total)
RE02-17-141635	02-613289	6–7	ALLH	—	—	2017-1985	—	2017-1985	2017-1985	2017-1985	—	_	2017-1985	—	—	—	_
RELA-18-151137	02-613289	6–7	ALLH	—	—	—	—	—	—	—	2018-1580	—	—	—	—	—	—
RELA-18-151138	02-613289	9–10	ALLH	—	—	—	—	—	—	—	2018-1580	—	—	—	—	—	—
RELA-18-151139	02-613289	11–11.25	ALLH	—	—	—	—	—	—	—	2018-1587	—	—	—	—	—	—
RE02-11-325	02-613292	4–4.2	ALLH	_	—	_	_	_	_	—	11-235	—	_	—	—	—	_
RE02-11-2218	02-613626	8–9	ALLH	_	—	—	_	_	_	—	11-541	—	—	_	—	—	_
RE02-11-2219	02-613626	10–11	ALLH	_	—	—	_	_	_	—	11-541	—	—	_	—	—	_
RE02-11-2220	02-613627	6–7	ALLH	_	_	_	_	_	—	_	11-541	_	_	_	_	_	_
RE02-11-2221	02-613627	8–9	ALLH	_	—	_	_	_	—	_	11-541	_	_	—	_	_	_
RE02-11-2222	02-613627	10–11	ALLH	_	—	_	_	_	_	_	11-686	_	_	_	_	_	_
RELA-18-151122	02-613667	2–3	ALLH	_	—	—	_	—	—	_	2018-1525	—	—	—	—	_	—
RELA-18-151123	02-613667	4–5	ALLH	_	—	—	—	_	—	—	2018-1525	—	—	—	_	_	—
RE02-11-2523	02-613667	6–6.2	ALLH	_	—	—	—	—	—	—	11-687	—	—	—	-	_	—
RELA-18-151124	02-613667	6–7	ALLH	_	—	—	—	—	—	—	2018-1525	—	—	—	-	_	—
RE02-11-2524	02-613667	8–8.2	ALLH	_	—	—	—	—	—	—	11-687	—	—	—	—	—	—
RELA-18-151125	02-613667	8–9	QBO	_	—	—	—	—	—	—	2018-1525	—	—	—	—	—	—
RE02-11-2525	02-613667	10–10.2	ALLH	—	—	—	—	—	—	—	11-687	—	—	_	—	—	—
RELA-18-151127	02-613668	2–3	ALLH	-	—	—	—	—	—	—	2018-1573	-	—	—	—	—	—
RELA-18-151128	02-613668	4–5	ALLH	_	—	—	—	—	—	—	2018-1573	—	—	—	—	—	—
RELA-18-151129	02-613668	6–7	ALLH	_	—	—	—	—	—	—	2018-1573	—	—	—	—	—	—
RE02-11-2526	02-613668	8–8.2	ALLH	_	—	—	—	—	—	—	11-729	—	—	—	—	—	—
RELA-18-151130	02-613668	8–9	ALLH	_	—	—	—	—	—	—	2018-1573	—	—	—	—	—	—
RE02-11-2527	02-613668	10–10.2	ALLH	—	—	—	—	—	—	—	11-729	—	—	_	—	—	—
RE02-11-2798	02-613699	12–12.2	QBT3	—	—	—	—	—	—	—	11-904	—	—	_	—	—	—
RE02-11-2799	02-613700	8–8.2	ALLH	—	—	—	—	—	—	—	11-904	—	—	—	—	—	—
RE02-11-2800	02-613700	10–10.2	ALLH	—	—	—	—	—	—	—	11-904	—	—	—	—	—	—
RE02-11-2801	02-613700	12–12.2	ALLH	—	—	—	—	—	—	—	11-904	—	—	_	—	—	—
RE02-11-3145	02-613700	14–14.2	ALLH	—	—	—	—	—	—	—	11-1006	—	—	—	—	—	—
RE02-11-3146	02-613761	6–6.2	ALLH	_	_		_		_	_	11-1009	_	_	_	_	_	_
RE02-11-3147	02-613761	8–8.2	ALLH	_	—		_		—	_	11-1009	_	—	_	_	_	_
RE02-11-3148	02-613761	10–10.2	ALLH	_	_	_	_	_	_	_	11-1009	_	_	_	_	_	_
RE02-11-3149	02-613761	12–12.2	QBT3	_	_	_	—	_	_	_	11-1009	—	—	_	_	_	

Table 3.4-1 (continued)

Sample ID	Location ID	Depth (ft)	Media	Americium-241	Nitrate	Gamma-emitting Radionuclides	Tritium	Isotopic Plutonium	Isotopic Uranium	TAL Metals	PCBs	Perchlorate	Strontium-90	SVOCs	VOCs	Dioxins/Furans	Cyanide (Total)
RE02-11-3150	02-613761	14–14.2	QBT3	_	—	_	—	_	_	—	11-1009	_	—	_	<u> </u>	—	—
RE02-17-141642	02-613762	2–3	ALLH	—	—	2017-2035	—	2017-2035	2017-2035	2017-2035	_	—	2017-2035	—	—	_	—
RELA-18-151117	02-613762	2–3	ALLH	—	—	_	—	—	—	—	2018-1580	_	_	—	—	_	—
RE02-17-141643	02-613762	4–5	QBO	—	—	2017-2035	—	2017-2035	2017-2035	2017-2035	—	_	2017-2035	—	_		—
RELA-18-151118	02-613762	4.25–5	QBO	_	—	_		_	—	_	2018-1580	—	—	—	—	—	—
RE02-17-141644	02-613762	6–7	QBO	—	_	2017-2035	_	2017-2035	2017-2035	2017-2035	—	_	2017-2035	_	_	_	_
RELA-18-151119	02-613762	6–7	QBO	_	_	_	_	—	_	_	2018-1580	_	_	_	_	_	_
RELA-18-151120	02-613762	8–9	QBO	—	—	—	—	—	-	_	2018-1580	—	-	—	—	-	-
RE02-17-135776	02-61428	0–1	ALLH	—	—	—	—	—	-	-	2017-1984	—	—	—	—	—	—
RE02-17-141612	02-61428	0–1	ALLH	_	—	2017-1981	—	2017-1981	2017-1981	2017-1981	—	_	2017-1981	_	—	-	-
RE02-17-135777	02-61428	2–3	ALLH	—	—	—	—	_	—	_	2017-1984	—	_	—	—	_	—
RE02-17-141658	02-61428	2–3	ALLH	_	_	2017-1981	_	2017-1981	2017-1981	2017-1981	—	—	2017-1981	—	_	_	_
RE02-17-141645	02-61428	4–5	ALLH	_	—	2017-1999	—	2017-1999	2017-1999	2017-1999	—	_	2017-1999	_	—	_	—
RE02-17-135778	02-61428	4–5	ALLH	_	—	—	_	_	—	_	2017-1997	_	—	_	_	—	_
RE02-17-135779	02-61428	5–6	ALLH	_	—	_	—	_	—	—	2017-1997	_	_	_	_	_	_
RE02-17-141621	02-61428	5–6	ALLH	_	—	2017-1999	—	2017-1999	2017-1999	2017-1999	—	_	2017-1999	_	_	_	_
RE02-17-135780	02-61429	0–1	ALLH	—	—	—	—	—	—	—	2017-1983	—	_	—	—	_	_
RE02-17-141659	02-61429	0–1	ALLH	—	—	2017-1985	—	2017-1985	2017-1985	2017-1985	—	—	2017-1985	—	—	_	_
RE02-17-141646	02-61429	2–3	ALLH	_	_	2017-1985	_	2017-1985	2017-1985	2017-1985	—	_	2017-1985	_	_	_	—
RE02-17-135781	02-61429	2–3	ALLH	_	_	_	_	_	_	_	2017-1983	_	_	_	_	_	—
RE02-17-135782	02-61429	4–5	ALLH	_	—	_	—	_	—	—	2017-1997	_	_	_	_	_	_
RE02-17-141622	02-61429	4–5	ALLH	_	—	2017-1999	—	2017-1999	2017-1999	2017-1999	—	_	2017-1999	_	_	_	_
RE02-17-141613	02-61429	5–6	ALLH	—	—	2017-1999	—	2017-1999	2017-1999	2017-1999	—	—	2017-1999	_	—	_	-
RE02-17-135783	02-61429	5–6	ALLH	—	—	_	—	—	—	—	2017-1997	—	_	_	—	_	-
RE02-17-141660	02-61432	0–1	ALLH	_	—	2017-1981	—	2017-1981	2017-1981	2017-1981	_	—	2017-1981	_	—	—	_
RE02-17-141623	02-61432	2–3	ALLH	_	—	2017-1981	—	2017-1981	2017-1981	2017-1981	_	—	2017-1981	_	—	—	_
RELA-18-151045	02-61432	2–3	ALLH	_	—	_	—	_	—	—	2018-1534	_	_	_	_	_	_
RELA-18-151046	02-61432	4–5	ALLH	—	—	—	—	—	—	—	2018-1534	—	-	—	—	-	—
RE02-17-141647	02-61432	4–5	ALLH	—	—	2017-1981	—	2017-1981	2017-1981	2017-1981	 _	—	2017-1981	—	—	-	<u> </u>
RE02-17-141614	02-61432	6–7	QBT3	_	—	2017-1981	—	2017-1981	2017-1981	2017-1981	—	—	2017-1981	_	—	-	<u> </u>
RELA-18-151047	02-61432	6.75–7	QBO	_	—	—	—	—	—	—	2018-1534	—	-	_	—	-	<u> </u>
RELA-18-151048	02-61432	8–8.75	QBO	—	—	-	—	-	-	—	2018-1534	-	_	—	—	-	-

Table 3.4-1 (continued)

Sample ID	Location ID	Depth (ft)	Media	Americium-241	Nitrate	Gamma-emitting Radionuclides	Tritium	Isotopic Plutonium	Isotopic Uranium	TAL Metals	PCBs	Perchlorate	Strontium-90	SVOCs	VOCs	Dioxins/Furans	Cyanide (Total)
RE02-17-135853	02-61432	11–12	ALLH	—	—	—	—	—	—	—	2017-1984	—	—	—	—	—	—
RE02-17-135854	02-61432	14–15	ALLH	_	—	_	—	_	—	_	2017-1984	—	—	—	—	_	—
RE02-17-135855	02-61432	16–17	ALLH	—	—	_	—	—	—	_	2017-1984	—	—	—	—	—	—
RE02-17-135856	02-61432	19–20	ALLH	—	—	_	—	—	—	_	2017-1984	—	—	—	—	—	—
RE02-17-141615	02-61435	0–1	ALLH	—	—	2017-1999	_	2017-1999	2017-1999	2017-1999	—	—	2017-1999	—	—	_	_
RE02-17-141624	02-61435	2–3	ALLH	—	_	2017-1999	_	2017-1999	2017-1999	2017-1999	—		2017-1999	_	_	_	_
RELA-18-151057	02-61435	2–3	ALLH	—	_	—	_	—	_	_	2018-1458		_	_	_	_	_
RELA-18-151058	02-61435	4–5	QBO	—	_	—	_	—	_	_	2018-1458		_	_	_	_	_
RE02-17-141625	02-61435	4–5	QBO	_	_	2017-1999	_	2017-1999	2017-1999	2017-1999	—	—	2017-1999	_	_	_	_
RELA-18-151059	02-61435	6–7	QBO	_	_	—	_	—	_	_	2018-1458	—	_	_	_	_	_
RE02-17-141648	02-61435	6–7	QBO	—	—	2017-1999	—	2017-1999	2017-1999	2017-1999	—	—	2017-1999	—	—	-	—
RE02-17-135893	02-61435	11–12	QBO	—	—	—	—	—	_	—	2017-1997	—	—	—	—	-	—
RE02-17-135894	02-61435	14–15	QBO	—	—	—	—	—	_	—	2017-1997	—	—	-	—	-	-
RE02-17-135895	02-61435	16–17	QBO	—	—	—	—	—	_	—	2017-1997	—	—	-	—	-	-
RE02-17-135896	02-61435	19–20	QBO	-	—	—	—	—	—	—	2017-1997	—	—	—	—	-	—
RE02-17-141661	02-61436	0–1	ALLH	—	—	2017-1932	—	2017-1932	2017-1932	2017-1932	—	—	2017-1932	—	—	—	—
RE02-17-141626	02-61436	2–3	ALLH	—	—	2017-1932	—	2017-1932	2017-1932	2017-1932	—	—	2017-1932	—	—	—	—
RELA-18-151062	02-61436	2–3	ALLH	—	—	—	—	—	—	_	2018-1501	_	—	—	—	—	—
RELA-18-151063	02-61436	4–5	ALLH	-	—	_	_	—	—	_	2018-1501	—	—	—	_	-	_
RE02-17-141616	02-61436	4–5	ALLH	—	—	2017-1932	—	2017-1932	2017-1932	2017-1932	—	-	2017-1932	—	—	—	—
RE02-17-141617	02-61436	6–7	QBO	—	—	2017-1932	—	2017-1932	2017-1932	2017-1932	—	—	2017-1932	—	—	-	—
RELA-18-151064	02-61436	6–7	QBO	-	—	—	—	—	—	—	2018-1501	—	—	—	—	-	—
RE02-17-135901	02-61436	8–9	QBO	—	—	—	—	—	—	_	2017-1931	—	—	—	—	—	—
RE02-17-135902	02-61436	11–12	QBO	—	—	—	—	—	—	_	2017-1931	—	—	—	—	—	—
RE02-17-135903	02-61436	14–15	QBO	-	—	_	_	—	—	_	2017-1931	—	—	—	_	-	_
RE02-17-135904	02-61436	16–17	QBO	-	—	-	—	-	—	—	2017-1931	—	—	-	—	—	—
RE02-17-135905	02-61436	19–20	QBO	—	—	—	—	—	—	_	2017-1931	_	—	—	—	—	—
RE02-17-141618	02-61437	0–1	ALLH	_	—	2017-2035	_	2017-2035	2017-2035	2017-2035	—	_	2017-2035	_	_	_	_
RE02-17-141627	02-61437	2–3	ALLH			2017-2035	_	2017-2035	2017-2035	2017-2035	_	_	2017-2035	_			_
RE02-17-144637	02-61437	2–3	ALLH	2017-2523	_	2017-2523	_	2017-2523	2017-2523	2017-2523	2017-2523	—	2017-2523	_	_	_	_
RE02-17-144638	02-61437	4–5	ALLH	2017-2523	_	2017-2523	_	2017-2523	2017-2523	2017-2523	2017-2523	—	2017-2523	_	_	_	_
RE02-17-141649	02-61437	4–5	ALLH	_	_	2017-2035	_	2017-2035	2017-2035	2017-2035	—	_	2017-2035	_		_	_

Table 3.4-1 (continued)

Sample ID	Location ID	Depth (ft)	Media	Americium-241	Nitrate	Gamma-emitting Radionuclides	Tritium	Isotopic Plutonium	Isotopic Uranium	TAL Metals	PCBs	Perchlorate	Strontium-90	SVOCs	VOCs	Dioxins/Furans	Cyanide (Total)
RE02-17-141662	02-61437	6–7	ALLH	—	—	2017-2080	_	2017-2080	2017-2080	2017-2080	_	_	2017-2080	_	—	_	—
RELA-18-151065	02-61437	6–7	ALLH	—	—	_	—	_	—	—	2018-1573	—	_	—	—	_	—
RELA-18-151066	02-61437	8–9	ALLH	—	—	_	—	—	—	—	2018-1573	—	—	—	—	_	—
RELA-18-151067	02-61437	11–11.8	QBO	—	—	_	—	—	—	—	2018-1573	—	—	—	—	_	—
RE02-17-141663	02-61438	0–1	ALLH	_	—	2017-1999	_	2017-1999	2017-1999	2017-1999	_	_	2017-1999	_	_	_	_
RE02-17-135916	02-61438	0–1	ALLH	_	—	—	_	—	—	—	2017-1997	_	—	_	_	_	_
RE02-17-135917	02-61438	2–3	ALLH	-	—	—	—	—	—	-	2017-1997	—	—	—	-	—	—
RE02-17-141650	02-61438	2–3	ALLH	—	—	2017-1999	—	2017-1999	2017-1999	2017-1999	—	—	2017-1999	—	—	_	—
RE02-17-135918	02-61438	4–5	QBO	—	—	—	_	—	—	—	2017-1997	_	_	—	—	_	_
RE02-17-141628	02-61438	4–5	QBO	—	—	2017-1999	_	2017-1999	2017-1999	2017-1999	—	_	2017-1999	—	—	_	_
RE02-17-141619	02-61438	6–7	QBO	—	—	2017-1999	_	2017-1999	2017-1999	2017-1999	—	_	2017-1999	—	—	_	_
RE02-17-135919	02-61438	6–7	QBO	_	—	_	_	—	—	—	2017-1997	—	—	—	_	_	_
RE02-17-135920	02-61438	8–9	QBO	—	—	_	_	—	—	—	2017-1998	—	_	—	—	_	_
RE02-17-135921	02-61438	11–12	QBO	—	—	—	_	—	—	—	2017-1998	—	—	—	—	_	_
RE02-17-135922	02-61438	14–15	QBO	_	—	_	_	—	—	—	2017-2020	_	_	—	_	_	_
RE02-17-135923	02-61438	16–17	QBO	_	—	_	_	—	—	—	2017-2020	_	_	—	_	_	_
RE02-17-135924	02-61438	19–20	QBO	—	—	—	—	—	—	—	2017-2020	_	—	—	—	—	_
RE02-17-135925	02-61439	0–1	ALLH	—	—	—	—	—	—	—	2017-2020	_	—	—	—	—	_
RE02-17-141664	02-61439	0–1	ALLH	—	—	2017-2021	—	2017-2021	2017-2021	2017-2021	_	_	2017-2021	—	—	_	_
RE02-17-135926	02-61439	2–3	ALLH	—	—	_	—	—	—	_	2017-2020	_	_	—	—	_	_
RE02-17-141651	02-61439	2–3	ALLH	—	—	2017-2021	_	2017-2021	2017-2021	2017-2021	_	_	2017-2021	—	_	_	_
RE02-17-141629	02-61439	4–5	QBO	—	—	2017-2021	—	2017-2021	2017-2021	2017-2021	_	_	2017-2021	—	—	_	_
RE02-17-135933	02-61439	4–5	QBO	-	—	—	—	—	—	—	2017-2020	_	—	—	—	—	-
RE02-17-141620	02-61439	6–7	QBO	—	—	2017-2021	—	2017-2021	2017-2021	2017-2021	_	_	2017-2021	—	—	—	_
RE02-17-135928	02-61439	6–7	QBO	—	—	_	—	—	—	_	2017-2020	_	_	—	—	_	_
RE02-17-135929	02-61439	8–9	QBO	—	—	_	—	—	—	_	2017-2020	_	_	—	—	_	_
RE02-17-135930	02-61439	11–12	QBO	-	—	—	—	—	—	—	2017-2020	_	—	—	—	—	-
RE02-17-135931	02-61439	14–15	QBO	—	—	—	—	—	—	_	2017-2020	_	_	—	—	—	
RE02-17-135932	02-61439	16–17	QBO	-	—	—	—	—	—	_	2017-2020	_	_	—	_	—	_
RE02-17-135927	02-61439	19–20	QBO	-	—	—	—	—	—	_	2017-2020	_	_	—	_	—	
RE02-17-141665	02-61440	0–1	ALLH	-	_	2017-2080	_	2017-2080	2017-2080	2017-2080	—	_	2017-2080	—	-	—	
RE02-17-141652	02-61440	2–3	ALLH	 _	-	2017-2080	—	2017-2080	2017-2080	2017-2080	—	—	2017-2080	 	 _	—	_

Table 3.4-1 (continued)

Sample ID	Location ID	Depth (ft)	Media	Americium-241	Nitrate	Gamma-emitting Radionuclides	Tritium	Isotopic Plutonium	Isotopic Uranium	TAL Metals	PCBs	Perchlorate	Strontium-90	SVOCs	VOCs	Dioxins/Furans	Cyanide (Total)
RELA-18-151069	02-61440	2–3	ALLH	—	—	_	—	—	—	—	2018-1572	—	—	—	-	_	—
RELA-18-151070	02-61440	4–5	ALLH	-	—	_	—	—	—	—	2018-1572	—	—	—	—	—	—
RE02-17-141653	02-61440	4–5	ALLH	_	_	2017-2080	—	2017-2080	2017-2080	2017-2080	—	—	2017-2080	—	—	—	—
RE02-17-141666	02-61440	6–7	ALLH	_	—	2017-2080	—	2017-2080	2017-2080	2017-2080	—	_	2017-2080	—	—	—	—
RELA-18-151071	02-61440	6–7	ALLH	_	_	_	—	—	—	_	2018-1572	—	—	_	_	_	_
RELA-18-151072	02-61440	8–9	ALLH	—	_	_	_		_	_	2018-1572	—	_	_	_	_	—
RE02-17-135939	02-61440	11–12	QBO	—	_	_			_	_	2017-2086	—	_	_	_	_	—
RE02-17-135940	02-61440	14–15	QBO	-	—	—	—	—	_	—	2017-2086	—	—	_	_	—	—
RE02-17-135941	02-61440	16–17	QBO	-	—	—	—	—	—	—	2017-2086	—	—	-	-	—	—
RE02-17-135942	02-61440	19–20	QBO	-	_	_	—	—	—	_	2017-2086	—	—	—	—	—	_
RE02-17-141668	02-61441	0–1	ALLH	-	_	2017-1985	_	2017-1985	2017-1985	2017-1985	—	—	2017-1985	_	_	—	_
RE02-17-141667	02-61441	2–3	ALLH	_	_	2017-1985	_	2017-1985	2017-1985	2017-1985	—	—	2017-1985	_	_	_	_
RE02-17-144671	02-61441	2–3	ALLH	2017-2523		2017-2523	_	2017-2523	2017-2523	2017-2523	2017-2523	_	2017-2523	_	_	_	—
RE02-17-144672	02-61441	4–5	ALLH	2017-2523	_	2017-2523	—	2017-2523	2017-2523	2017-2523	2017-2523	—	2017-2523	_	_	_	_
RE02-17-141654	02-61441	4–5	ALLH	_	_	2017-1985	_	2017-1985	2017-1985	2017-1985	—	_	2017-1985	_	_	_	_
RE02-17-141655	02-61441	6–7	QBO	_	_	2017-1985	_	2017-1985	2017-1985	2017-1985	—	_	2017-1985	_	_	_	_
RELA-18-151043	02-61441	6–7	QBO	—	—	—	—	—	—	—	2018-1551	—	—	—	—	—	—
RELA-18-151073	02-61441	8–8.5	QBO	—	—	—	—	—	—	—	2018-1551	—	—	—	—	—	—
RE02-17-135948	02-61441	11–12	QBO	_	_	_	_	_	—	_	2017-1983	_	—	_	_	_	—
RE02-17-135949	02-61441	14–15	QBO	_	_	_	_	_	—	_	2017-1983	_	—	_	_	_	—
RE02-17-135950	02-61441	16–17	QBO	_	_	_	_	—	_	_	2017-1983	_	—	_	_	_	_
RE02-17-135951	02-61441	19–20	QBO	_	_	_	_	—	_	_	2017-1983	_	—	_	_	_	_
RE02-17-141670	02-61442	0–1	ALLH	—	—	2017-1999	—	2017-1999	2017-1999	2017-1999	—	—	2017-1999	—	—	—	—
RE02-17-141669	02-61442	2–3	ALLH	—	—	2017-1999	—	2017-1999	2017-1999	2017-1999	—	—	2017-1999	—	—	—	—
RELA-18-151075	02-61442	2–3	ALLH	_	_	_	—	_	_	_	2018-1458	—	—	_	_	_	_
RELA-18-151076	02-61442	4–5	QBO	_	—	_	—	—	—	—	2018-1458	_	—	_	_	_	_
RE02-17-141656	02-61442	4–5	QBO	_	—	2017-1999	—	2017-1999	2017-1999	2017-1999	—	—	2017-1999	—	—	_	_
RELA-18-151077	02-61442	6–7	QBO	-	—	—	—	<u> </u>	_	—	2018-1458	_	—	_	—	_	—
RE02-17-141657	02-61442	6–7	QBO	-	—	2017-1999	—	2017-1999	2017-1999	2017-1999	_	<u> </u>	2017-1999	_	_	_	_
RELA-18-151078	02-61442	8–9	QBO	_	_	—	—	—	—	—	2018-1458	_	—	_	_	_	—
RE02-17-135957	02-61442	11–12	QBO	-	—	—	—	—	_	—	2017-1998	_	—	_	_	_	_
RE02-17-135958	02-61442	14–15	QBO	-	—	-	—	-	_	_	2017-1998	_	—	_	_	—	_

Table 3.4-1 (continued)

Sample ID	Location ID	Depth (ft)	Media	Americium-241 Nitrate	Gamma-emitting Radionuclides	Tritium	lsotopic Plutonium	lsotopic Uranium	TAL Metals	PCBs	Perchlorate	Strontium-90	SVOCs	vocs	Dioxins/Furans	Cyanide (Total)
RE02-17-135959	02-61442	16–17	QBO				_			2017-1998	_		_			_
RE02-17-135960	02-61442	19–20	QBO		_	_	 	 	_	2017-1998	_	_	_	_	 	
RELA-18-151080	02-61443	2–3	ALLH		_	_	_	_		2018-1441	_	_	_	_	_	_
RE02-17-141678	02-61443	2–3	ALLH		2017-1981	—	2017-1981	2017-1981	2017-1981	_	_	2017-1981	—	_	—	_
RE02-17-141685	02-61443	4–5	QBO	_ _	2017-1981	—	2017-1981	2017-1981	2017-1981	—	_	2017-1981	—	_	—	_
RELA-18-151081	02-61443	4–5	QBO		—	_	—	—	_	2018-1441	_	_	—	_	_	_
RE02-17-141785	02-61443	6–7	QBO	-	2017-1981	_	2017-1981	2017-1981	2017-1981	—	-	2017-1981	—	_	_	_
RELA-18-151082	02-61443	6–7	QBO	-	_	_	—	_	_	2018-1441	-	_	—	_	_	_
RELA-18-151083	02-61443	8–9	QBO	-	—	_	—	—	_	2018-1441	_	—	—	—	—	_
RELA-18-151084	02-61443	11–12	QBO		—	—	—	—	—	2018-1441	_	_	—	_	—	_
RE02-17-141693	02-61444	0–1	ALLH		2017-1932	—	2017-1932	2017-1932	2017-1932	—	_	2017-1932	—	_	—	_
RE02-17-141686	02-61444	2–3	ALLH		2017-1932	_	2017-1932	2017-1932	2017-1932	—	_	2017-1932	—	_	—	_
RELA-18-151086	02-61444	2–3	ALLH		—	_	—	—	_	2018-1501	_	_	—	_	—	_
RELA-18-151087	02-61444	4–5	ALLH		—	—	—	—	—	2018-1501	—	—	—	—	—	—
RELA-18-151088	02-61444	6–7	ALLH		—	—	—	—	—	2018-1501	-	—	—	—	—	—
RE02-17-141672	02-61444	6–7	ALLH		2017-1932	—	2017-1932	2017-1932	2017-1932	—	—	2017-1932	—	—	—	—
RELA-18-151089	02-61444	8–8.85	ALLH		—	—	—	—	—	2018-1501	—	—	—	—	—	—
RE02-17-136003	02-61444	11–12	QBO		—	—	—	—	—	2017-1944	-	-	—	—	—	_
RE02-17-136004	02-61444	14–15	QBO		—	—	—	—	—	2017-1944	-	—	—	—	—	—
RE02-17-136005	02-61444	16–17	QBO	— —	—	_	—	_	_	2017-1944	—	—	—	—	_	_
RE02-17-136006	02-61444	19–20	QBO	— —	—	_	—	_	_	2017-1944	—	—	—	—	_	_
RE02-17-141673	02-61445	0–1	ALLH		2017-1945	_	2017-1945	2017-1945	2017-1945	—	—	2017-1945	_	—	—	_
RE02-17-136007	02-61445	0–1	ALLH		—	_	-	-	—	2017-1944	-	-	—	—	—	—
RE02-17-136010	02-61445	6–7	QBO	— —	_	_	_	_	_	2017-1944	—	—	_	—	_	_
RE02-17-141694	02-61445	6–7	QBO	— —	2017-1945	_	2017-1945	2017-1945	2017-1945	—	—	2017-1945	—	—	_	_
RE02-17-136011	02-61445	8–9	QBO		—	_	-	-	—	2017-1944	-	-	—	—	-	—
RE02-17-136012	02-61445	11–12	QBO		—	_	-	-	—	2017-1944	-	-	—	—	—	—
RE02-17-136015	02-61445	14–20	QBO							2017-1970	_					_
RE02-17-139388	02-61446	0–1	ALLH		_					2017-1982	—				_	_
RE02-17-141695	02-61446	0–1	ALLH	— —	2017-1985	—	2017-1985	2017-1985	2017-1985	_	—	2017-1985			_	_
RE02-17-139389	02-61446	2–3	ALLH		_	_	—	—	_	2017-1982	—	—	—	_	—	—
RE02-17-141688	02-61446	2–3	ALLH	-	2017-1985	-	2017-1985	2017-1985	2017-1985	—	_	2017-1985	—		<u> </u>	

Table 3.4-1 (continued)

Sample ID	Location ID	Depth (ft)	Media	Americium-241	Nitrate	Gamma-emitting Radionuclides	Tritium	Isotopic Plutonium	Isotopic Uranium	TAL Metals	PCBs	Perchlorate	Strontium-90	SVOCs	VOCs	Dioxins/Furans	Cyanide (Total)
RE02-17-141681	02-61446	4-4.5	QBO	—	—	2017-1985	—	2017-1985	2017-1985	2017-1985	—	—	2017-1985	—	—	—	—
RE02-17-139390	02-61446	4-4.5	QBO	_	—	_	—		—	_	2017-1982	_	—	—	—	—	—
RE02-17-139391	02-61446	6–7	ALLH	_	—	_	—		—	_	2017-1984	_	—	—	—	—	—
RE02-17-141674	02-61446	6–7	QBO	—	—	2017-1981	—	2017-1981	2017-1981	2017-1981	—	_	2017-1981	—	—	—	—
RE02-17-139392	02-61446	8–9	ALLH	—	—	—	_	—	—	—	2017-1984	—	—	—	_	_	—
RE02-17-139393	02-61446	11–12	ALLH	—	—	—	—	—	—	—	2017-1984	—	—	—	_	_	—
RE02-17-139394	02-61446	14–15	QBO	_		_		_	_	_	2017-1984	—	_	_	_	_	—
RE02-17-139395	02-61446	16–17	QBO	—	—	—	—	—	_	—	2017-1984	—	—	_	_	—	—
RE02-17-139396	02-61446	19–20	ALLH	—	—	—	—	—	—	—	2017-1984	-	—	—	—	—	—
RE02-17-141696	02-61447	0–1	ALLH	_	—	2017-2021	—	2017-2021	2017-2021	2017-2021	—	—	2017-2021	—	—	_	_
RELA-18-151037	02-61447	0–1	ALLH	_	—	—	—	—	—	_	2018-1596	—	—	—	—	—	_
RE02-17-141689	02-61447	2–3	ALLH	_	—	2017-2021	—	2017-2021	2017-2021	2017-2021	—	—	2017-2021	—	—	_	_
RELA-18-151038	02-61447	2–3	ALLH	—	—	—	—	—	—	_	2018-1596	_	—	_	—	—	_
RE02-17-141682	02-61447	4–5	ALLH	—	—	2017-2021	—	2017-2021	2017-2021	2017-2021	—	—	2017-2021	—	—	—	—
RELA-18-151039	02-61447	4–5	QBO	_	—	—	—	—	_	_	2018-1596	_	—	_	_	—	_
RE02-17-141675	02-61447	6–7	ALLH	_	—	2017-2021	—	2017-2021	2017-2021	2017-2021	—	—	2017-2021	—	—	_	_
RELA-18-151040	02-61447	6–7	QBO	—	—	—	—	—	—	_	2018-1596	_	—	_	—	—	_
RELA-18-151041	02-61447	8–9	QBO	—	—	—	—	—	—	_	2018-1596	_	—	_	—	—	_
RE02-17-139402	02-61447	11–12	QBO	—	_	—	—	—	_	_	2017-2037	—	_	_	_	—	_
RE02-17-139403	02-61447	14–15	QBO	_	—	—	—	—	_	_	2017-2037	_	—	_	_	—	_
RE02-17-139404	02-61447	16–17	QBO	_	—	—	—	—	_	_	2017-2037	_	—	_	_	—	_
RE02-17-139405	02-61447	19–20	QBO	_	—	—	—	—	_	_	2017-2037	_	—	_	_	—	_
RE02-17-141697	02-61448	0–1	ALLH	—	—	2017-1985	—	2017-1985	2017-1985	2017-1985	—	—	2017-1985	_	—	—	_
RELA-18-151050	02-61448	2–3	ALLH	—	—	—	—	—	—	_	2018-1587	_	—	_	—	—	_
RELA-18-151051	02-61448	4–5	ALLH	_	—	—	—	—	_	_	2018-1587	_	—	_	_	—	_
RE02-17-141683	02-61448	4–5	ALLH	—	_	2017-1985	—	2017-1985	2017-1985	2017-1985	—	—	2017-1985	_	_	_	_
RE02-17-141676	02-61448	6–7	ALLH	—	—	2017-1985	_	2017-1985	2017-1985	2017-1985	—	—	2017-1985	_	_	_	_
RELA-18-151052	02-61448	6–7	ALLH	_	—	—	_	—	_	_	2018-1587	_	—	_	_	_	_
RELA-18-151053	02-61448	8–9	ALLH	—	—	—	—	—	—	—	2018-1587	_	—	—	—	—	_
RELA-18-151054	02-61448	11–12	ALLH	_	—	_	_	_	_		2018-1587	_	—	_	_	_	_
RELA-18-151055	02-61448	13–14	ALLH	_	_	_	_	_	_		2018-1596	_	_	_	_	_	_
RE02-17-139413	02-61448	16–17	QBO	_	—	_	_	_	—	_	2017-1982	_	—	—	_		—

Table 3.4-1 (continued)

Sample ID	Location ID	Depth (ft)	Media	Americium-241	Nitrate	Gamma-emitting Radionuclides	Tritium	Isotopic Plutonium	Isotopic Uranium	TAL Metals	PCBs	Perchlorate	Strontium-90	SVOCs	vocs	Dioxins/Furans	Cyanide (Total)
RE02-17-139414	02-61448	19–20	QBO	—	—	_	_	_	_	—	2017-1982	—	_	_	_	—	_
RE02-17-141559	02-61450	2–3	ALLH	_	—	2017-2078	_	2017-2078	2017-2078	2017-2078	—	_	2017-2078	_	_	_	_
RELA-18-151091	02-61450	2–3	ALLH	_	—	—	_	—	_	—	2018-1518	_	_	_	_	_	_
RELA-18-151092	02-61450	4–5	ALLH	_	—	_	—	_	—	—	2018-1518	_	_	_	—	_	—
RE02-17-141560	02-61450	4–5	ALLH	—	—	2017-2078	_	2017-2078	2017-2078	2017-2078	_	—	2017-2078	_	_	_	_
RE02-17-141561	02-61450	6–7	QBO	_	—	2017-2078	_	2017-2078	2017-2078	2017-2078	2017-2087	_	2017-2078	_	_	_	_
RE02-17-141562	02-61450	8–9	QBO	_	—	—	_	—	_	—	2017-2087	_	_	_	_	_	_
RE02-17-141563	02-61450	11–12	QBO	_	—	—	_	—	_	—	2017-2087	_	_	_	_	_	_
RE02-17-141564	02-61450	14–15	QBO	_	—	_		—	_	_	2017-2087	_	_	_	_	_	_
RE02-17-141565	02-61450	16–17	QBO	—	—	—		—	_	_	2017-2087	—	—	_	_	—	_
RE02-17-141566	02-61450	19–20	QBO	_	—	_		—	_	_	2017-2087	_	_	_	_	_	_
RE02-17-141568	02-61451	2–3	ALLH		—	2017-2080		2017-2080	2017-2080	2017-2080	—	_	2017-2080	_	_	_	_
RELA-18-151094	02-61451	2–3	ALLH	_	—	_		—	_	_	2018-1501	—	_	_	_	_	_
RELA-18-151095	02-61451	4–5	ALLH	_	—	—	_	—	_	—	2018-1501	_	_	_	_	_	_
RE02-17-141569	02-61451	4–5	ALLH	_	—	2017-2080		2017-2080	2017-2080	2017-2080	_	_	2017-2080	_	_	_	_
RE02-17-141570	02-61451	6–7	QBO		—	2017-2080		2017-2080	2017-2080	2017-2080	—	_	2017-2080	_	_	_	_
RELA-18-151096	02-61451	6.25–7	QBO	—	—	—	—	—	_	—	2018-1501	-	-	_	_	-	-
RELA-18-151097	02-61451	8–9	QBO	—	—	—	—	—	_	—	2018-1501	-	-	_	_	-	—
RE02-17-141572	02-61451	11–12	QBO	—	—	—	—	—	_	—	2017-2081	—	-	_	_	—	—
RE02-17-141573	02-61451	14–15	QBO	_	—	—	—	—	_	—	2017-2081	—	—	_	—	_	_
RE02-17-141574	02-61451	16–17	QBO		—	_	_	—	_	_	2017-2087	_	—	_	_	_	_
RE02-17-141575	02-61451	19–20	QBO	_	—	_		—	_	—	2017-2087	_	—	_	_	_	_
RE02-17-141577	02-61452	2–3	ALLH	—	—	2017-2080	—	2017-2080	2017-2080	2017-2080	_	-	2017-2080	_	_	-	-
RELA-18-151099	02-61452	2–3	ALLH	—	—	—	—	—	_	—	2018-1558	-	-	_	_	-	—
RELA-18-151100	02-61452	4–5	ALLH	—	—	—	—	—	—	—	2018-1558	—	-	_	_	—	—
RE02-17-141578	02-61452	4–5	ALLH	—	—	2017-2080	—	2017-2080	2017-2080	2017-2080	—	—	2017-2080	_	_	—	—
RE02-17-141579	02-61452	6–7	QBO	—	—	2017-2080	—	2017-2080	2017-2080	2017-2080	—	—	2017-2080	—	—	—	—
RELA-18-151101	02-61452	6–7	QBO	_	_	—	_	_	_	_	2018-1558	_	_	_	_	_	_
RELA-18-151102	02-61452	8–9	QBO	_	—	_	_	_	_	—	2018-1558	_	—	_	_	_	_
RE02-17-141581	02-61452	11–12	QBO	_	—	_	_	_	_	—	2017-2081	_	—	_	_	_	_
RE02-17-141582	02-61452	14–15	QBO	_	_	_	_	_	_	_	2017-2081	_	_	_	_	_	_
RE02-17-141583	02-61452	16–17	QBO	_	—	—	—	—	—	—	2017-2081	—	—	—	—	—	—

Table 3.4-1 (continued)

Sample ID	Location ID	Depth (ft)	Media	Americium-241	Nitrate	Gamma-emitting Radionuclides	Tritium	Isotopic Plutonium	Isotopic Uranium	TAL Metals	PCBs	Perchlorate	Strontium-90	SVOCs	VOCs	Dioxins/Furans	Cyanide (Total)
RE02-17-141584	02-61452	19–20	QBO	—	—	_	—	—	—	—	2017-2081	—	—	—	—	—	—
RE02-17-141586	02-61453	2–3	ALLH	<u> </u>	—	2017-2021	—	2017-2021	2017-2021	2017-2021	—	—	2017-2021	—	—	—	—
RELA-18-151104	02-61453	2–3	ALLH	—	—	—	—	—	—	_	2018-1572	—	—	—	—	—	—
RELA-18-151105	02-61453	4–5	ALLH	—	—	—	—	—	—	—	2018-1572	—	—	—	—	—	—
RE02-17-141587	02-61453	4–5	ALLH		—	2017-2021	—	2017-2021	2017-2021	2017-2021	—	_	2017-2021	—	_		—
RE02-17-141588	02-61453	6–7	ALLH	—	—	2017-2021	_	2017-2021	2017-2021	2017-2021	—	—	2017-2021	—	—	—	—
RELA-18-151106	02-61453	6–7	ALLH	_	—	—	_	—	_	_	2018-1572	—	_	_	_	_	—
RELA-18-151107	02-61453	8–9	ALLH	—	—	—	—	—	—	—	2018-1572	—	—	—	_	_	—
RE02-17-141590	02-61453	11–12	QBO	—	—	—	—	—	_	—	2017-2020	—	—	—	—	—	—
RE02-17-141592	02-61453	16–17	QBO	—	_	—	_	—	_	_	2017-2020	—	—	—	—	_	_
RE02-17-141593	02-61453	19–20	QBO	-	—	—	—	—	—	—	2017-2020	—	—	—	—	—	—
RE02-17-141595	02-61454	2–3	ALLH	-	—	2017-2035	—	2017-2035	2017-2035	2017-2035	—	-	2017-2035	—	—	-	—
RELA-18-151109	02-61454	2–3	ALLH	-	—	—	—	—	—	_	2018-1534	—	—	—	—	—	_
RELA-18-151110	02-61454	4–5	ALLH	—	—	—	—	—	—	_	2018-1534	—	—	—	—	—	—
RE02-17-141596	02-61454	4–5	ALLH	_	_	2017-2035	_	2017-2035	2017-2035	2017-2035	—	—	2017-2035	—	_	_	_
RE02-17-141597	02-61454	6–7	QBO	-	—	2017-2035	—	2017-2035	2017-2035	2017-2035	2017-2037	—	2017-2035	—	_	_	_
RE02-17-141598	02-61454	8–9	QBO	_	—	—	—	_	—	_	2017-2037	—	—	—	_	—	_
RE02-17-141599	02-61454	11–12	QBO	_	—	—	—	_	—	_	2017-2037	—	—	—	_	—	_
RE02-17-141600	02-61454	14–15	QBO	_	_	—	_	—	_	_	2017-2037	_	—	—	_	—	_
RE02-17-136031	02-61454	16–17	QBO	_	_	—	_	—	_	_	2017-2038	_	—	—	_	—	_
RE02-17-141602	02-61454	19–20	QBO	_	_	—	_	—	_	_	2017-2037	_	—	—	_	—	_
RE02-17-141604	02-61455	2–3	ALLH	_	_	2017-2035	_	2017-2035	2017-2035	2017-2035	—	—	2017-2035	—	_	_	_
RELA-18-151112	02-61455	2–3	ALLH	_	_	—	—	_	—	_	2018-1551	—	—	—	_	_	_
RELA-18-151113	02-61455	4–5	ALLH	_	_	—	—	_	—	_	2018-1551	—	—	—	_	—	_
RE02-17-141605	02-61455	4–5	ALLH	_	_	2017-2035	_	2017-2035	2017-2035	2017-2035	—	—	2017-2035	—	_	_	_
RE02-17-141606	02-61455	6–7	ALLH	_	_	2017-2035	_	2017-2035	2017-2035	2017-2035	—	—	2017-2035	—	_	_	_
RELA-18-151114	02-61455	6–7	ALLH	_	—	—	_	—	_	_	2018-1551	_	—	—	_	_	_
RELA-18-151115	02-61455	8–8.75	ALLH	_	—	—	_	—	_	_	2018-1551	_	—	—	_	_	_
RE02-17-141608	02-61455	11–12	QBO	_	—	—	—	—	—	—	2017-2036	_	—	—	—	—	_
RE02-17-141609	02-61455	14–15	QBO	—	—	_	_	-	_		2017-2036	_	_	_	_	_	_
RE02-17-141610	02-61455	16–17	QBO	_	_	_	_	-	_		2017-2036	_	_	_	_	_	_
RE02-17-141611	02-61455	19–20	QBO	_	_	_	_	_	—	_	2017-2036	_	—	—	_		—

Table 3.4-1 (continued)

Sample ID	Location ID	Depth (ft)	Media	Americium-241	Nitrate	Gamma-emitting Radionuclides	Tritium	Isotopic Plutonium	Isotopic Uranium	TAL Metals	PCBs	Perchlorate	Strontium-90	SVOCs	VOCs	Dioxins/Furans	Cyanide (Total)
RE02-17-145060	02-61474	2–3	ALLH	—	—	-	—	—	— ·	_	2017-2647	—	—	—	_	_	—
RE02-17-145072	02-61474	4–5	ALLH	—	—	-	—	—	— ·	_	2017-2647	—	—	—	_	_	—
RE02-17-145084	02-61474	6–7	ALLH	—	—	—	_	—			2017-2647	_	-	_	—	_	—
RE02-17-145049	02-61475	0–1	ALLH	_	—	—	_	_			2017-2647		—	—	_		—
RE02-17-145061	02-61475	2–3	ALLH	_	_	_	_	_	_		2017-2647		—	_	_		_
RE02-17-145073	02-61475	4–5	ALLH	_	_	—	_	_	_		2017-2647		—	_	_		_
RE02-17-145050	02-61476	0–1	ALLH	_	_	—	_	_	_		2017-2678		—	_	_		_
RE02-17-145062	02-61476	2–3	ALLH	_	—	—	_	_			2017-2678		—	—	_		_
RE02-17-145074	02-61476	4–5	ALLH	_	—	—	_	_	_ ·		2017-2678		—	—	_		_
RE02-17-145086	02-61476	6–7	QBO	_	—	—	_	_	_	_	2017-2678		—	_	_		—
RE02-17-145098	02-61476	8–9	QBO	_	—	—	_	_	_	_	2017-2678		—	_	_		_
RE02-17-145110	02-61476	9–10	QBO		_	—	_	_	_		2017-2678		—	_	_		_
RE02-17-145051	02-61477	0–1	ALLH	_	—	—	_	_			2017-2647		—	—	_		_
RE02-17-145063	02-61477	2–3	ALLH	_	_	—	_	_	_		2017-2647		—	_	_		_
RE02-17-145064	02-61478	2–3	ALLH	_	_	—	_	_			2017-2647	_	—	_	_	_	_
RE02-17-145065	02-61479	2–3	ALLH	_	—	—	_	_			2017-2647	_	—	_	_	_	_
RE02-17-145077	02-61479	4–5	ALLH	_	_	—	_	_	_		2017-2647		—	_	_		_
RE02-17-145054	02-61480	0–1	ALLH	_	_	—	_	_	_		2017-2667		—	_	_		—
RE02-17-145066	02-61480	2–3	ALLH	_	—	—	_	_	_		2017-2667		—	_	_		_
RE02-17-145078	02-61480	4–5	ALLH	_	_	_	_	_	_		2017-2667		—	_	_		_
RE02-17-145090	02-61480	6–7	ALLH	_	_	_	_	_	_		2017-2667		—	_	_		_
RE02-17-145102	02-61480	8–8.5	QBT3	_	_	—	_	_	_		2017-2667		—	_	_		_
RE02-17-145055	02-61481	0–1	ALLH	—	—	-	—	_		_	2017-2667	_	—	—	_	_	—
RE02-17-145067	02-61481	2–3	ALLH	_	_	—	_	_	_		2017-2667		—	_	_		_
RE02-17-145079	02-61481	4–5	ALLH	—	—	-	—	—	— ·	_	2017-2667	—	—	—	_	_	—
RE02-17-145091	02-61481	6–7	ALLH	—	—	-	—	—	— ·	_	2017-2667	—	—	—	_	_	—
RE02-17-145103	02-61481	8–9	QBO	—	—	—	—	—			2017-2667	_	—	—	—	_	—
RE02-17-145115	02-61481	9–10	QBO	_	—	_	_	—			2017-2667		—	—		_	_
RE02-17-145056	02-61482	0–1	ALLH	_	—	—	_	—		_	2017-2667		—	_	_	_	_
RE02-17-145068	02-61482	2–3	ALLH	_	—	_	_	_		_	2017-2667		—	_	_	_	_
RE02-17-145080	02-61482	4–5	QBO	_	—	_	_	_	·	_	2017-2667		_	_	_	_	_
RE02-17-145092	02-61482	6–7	QBT3	_	—	_	_	_			2017-2667	_	—	_			

Table 3.4-1 (continued)

Sample ID	Location ID	Depth (ft)	Media	Americium-241	Nitrate	Gamma-emitting Radionuclides	Tritium	Isotopic Plutonium	Isotopic Uranium	TAL Metals	PCBs	Perchlorate	Strontium-90	SVOCs	VOCs	Dioxins/Furans	Cyanide (Total)
RE02-17-145116	02-61482	8–9	QBO	_	_	_	_	_	_	_	2017-2667		_	_	_	_	—
RE02-17-145104	02-61482	9–10	QBO	_	_	_	—	_	—	_	2017-2667	—	—	_	_	_	—
RE02-17-145057	02-61483	0–1	ALLH	_	_	_	_	_	_	_	2017-2678	_	_	_	_	_	—
RE02-17-145069	02-61483	2–3	ALLH	—	—	_	_	—		—	2017-2678		—	—	—	_	_
RE02-17-145081	02-61483	4–5	ALLH	—	—	_	_	—	—	—	2017-2678	—	—	—	—	_	—
RE02-17-145093	02-61483	6–7	QBO	_	_	_	_	_	_	_	2017-2678	_	_	_	_	_	—
RE02-17-145105	02-61483	8–9	QBO	_	_	_	_	—	—	_	2017-2678	_	—	_	_	_	—
RE02-17-145117	02-61483	9–10	QBO	_	_	_	_	_	_	_	2017-2678		_	_	_	_	—
RE02-17-145120	02-61486	0–1	ALLH	_	_	_	_	_	_	_	2017-2647	_	_	_	_	_	_
RE02-17-145130	02-61486	2–3	ALLH	_	_	_	_	_	_	_	2017-2647		_	_	_	_	—
RE02-17-145131	02-61487	2–3	ALLH	_	_	_	_	_	_	_	2017-2667		_	_	_	_	—
RE02-17-145141	02-61487	4–5	ALLH	—	—	_	_	—	—	—	2017-2667	_	—	_	—	—	—
RE02-17-145151	02-61487	5–5.5	QBO	_	_	_	_	_	_	_	2017-2667	_	_	_	_	_	—
RE02-17-145132	02-61488	2–3	QBO	_	_	_	_	_	_	_	2017-2695	_	_	_	_	_	—
RE02-17-145142	02-61488	4-4.1	QBO	_	_	_	_	_	_	_	2017-2695	_	_	_	_	_	_
RE02-17-145133	02-61489	2–3	ALLH	_	_	_	_	_	_	_	2017-2695	_	_	_	_	_	_
RE02-17-145143	02-61489	4–5	ALLH	—	—	_	—	—	—	—	2017-2695	—	—	-	—	—	—
RE02-17-145153	02-61489	6–7	QBO	—	—	_	—	—	—	—	2017-2710	—	—	-	—	—	—
RE02-17-145163	02-61489	7–7.5	QBO	—	—	_	—	—	—	—	2017-2710	—	—	-	—	—	—
RE02-17-145124	02-61490	0–1	ALLH	—	—	—	—	—	—	—	2017-2647	—	—	-	—	—	—
RE02-17-145134	02-61490	2–3	ALLH	_	_	_	_	_	_	_	2017-2647	_	_	_	_	_	_
RE02-17-145125	02-61491	0–1	ALLH	—	—	_	_	—	—	—	2017-2695	_	—	_	—	—	_
RE02-17-145135	02-61491	2–3	ALLH	—	—	—	_	—	—	—	2017-2695	—	—	—	—	_	—
RE02-17-145145	02-61491	4–5	QBO	_	—	_	—	_	_	_	2017-2695	_	_	_	_	—	—
RE02-17-145126	02-61492	0–0.5	ALLH	—	—	_	—	—	—	—	2017-2647	—	—	-	—	—	—
RE02-17-145127	02-61493	0–1	ALLH	—	—	—	—	—	—	—	2017-2647	—	—	-	—	—	—
RE02-17-145137	02-61493	2–3	ALLH	—	—	—	_	—	—	—	2017-2647	—	—	—	—	_	—
RE02-17-145147	02-61493	4-4.5	QBO	_	_	_	_		_	_	2017-2647	—	_	_	_	_	_
RE02-17-145128	02-61494	0–1	ALLH	_	_	_	_			_	2017-2667	—	_	_	_	_	_
RE02-17-145138	02-61494	2–3	ALLH	_	—	_	_	_	_	_	2017-2667	—	_	_	_	_	_
RELA-18-151015	02-61526	0–1	ALLH	_	_	_	_	_	_	_	2018-1441	—	_	_	_	_	_
RELA-18-151020	02-61526	2–3	ALLH	—	_	_	_	_		_	2018-1441		—	_	—	_	

Table 3.4-1 (continued)

Sample ID	Location ID	Depth (ft)	Media	Americium-241 Nitrate	Gamma-emitting Radionuclides	Tritium	Isotopic Plutonium	Isotopic Uranium	TAL Metals	PCBs	Perchlorate	Strontium-90	SVOCs	vocs	Dioxins/Furans	Cyanide (Total)
RELA-18-151021	02-61526	4–5	ALLH	— —	—	—	_	—	—	2018-1441	_	_	—	—	_	—
RELA-18-151022	02-61526	6–7	QCT		—	—	—	—	—	2018-1458	—	—	—	—	—	—
RELA-18-151017	02-61528	0–1	ALLH	— —	_	—	_	_	_	2018-1422	_	_	_	—	—	_
RELA-18-151023	02-61528	2–3	ALLH	— —	_	_	—	_	_	2018-1422	_	—	—	_	—	—
RELA-18-151024	02-61528	4–5	ALLH	— —	_	_	_	_	_	2018-1422	_	_	_	_	_	_
RELA-18-151018	02-61529	0–1	ALLH		—	_	_	_	_	2018-1422	_	_	_	—	—	_
RELA-18-151027	02-61529	2–3	ALLH	— —	_	_	_	_	_	2018-1422	_	_	_	_	_	_
RELA-18-151028	02-61529	4-4.65	ALLH	— —	_	—	_	_	_	2018-1422	_	_	_	—	—	_
RELA-18-151019	02-61530	0–1	ALLH		—	—	_	—	—	2018-1422	—	_	_	—	—	_
RELA-18-151030	02-61530	2–3	ALLH	— —	—	_	_	_	—	2018-1422	_	_	_	—	_	_
RELA-18-151031	02-61530	4–5	QBO	— —	—	_	_	_	—	2018-1422	_	_	_	—	_	_
RELA-18-151032	02-61530	6–7	QBO		—	—	—	—	—	2018-1422	_	—	—	—	_	_
RELA-18-151033	02-61537	2–3	ALLH	— —	_	—	_	_	_	2018-1422	_	_	_	—	—	_
RELA-18-151034	02-61537	4–5	ALLH	— —	_	_	_	_	_	2018-1422	_	_	_	_	—	—
RELA-18-151035	02-61537	6–7	ALLH	— —	_	_	_		_	2018-1441	_	_	_	_	—	_
RELA-18-151263	02-61537	8–9	QBO		—	_	_	_	_	2018-1441	_	_	—	—	—	_
RELA-18-151036	02-61537	11–12	QBO	— —	_	_	_	_	_	2018-1441	_	_	_	_	—	—
RELA-18-161238	02-61538	0–0.9	ALLH	— —	_	_	_	_	_	N3B-2018-3962-1	_	_	_	_	—	—
RELA-18-161242	02-61538	1–2	ALLH	— —	—	_	_	_	—	N3B-2018-3962-1	_	_	_	—	_	_
RELA-18-161239	02-61539	1–2	ALLH	— —	_	_	_	_	_	N3B-2018-3962-1	_	_	_	_	_	_
RELA-18-161244	02-61540	1–2	ALLH	— —	_	_	_	_	_	N3B-2018-3962-1	_	_	_	_	_	_
RELA-18-164444	02-61541	6–7	QBT1G	— —	_	_	_	_	_	N3B-2019-380	_	_	_	—	_	_
RELA-18-161248	02-61542	1–2	ALLH		—	—	—	—	—	N3B-2018-3962-1	—	—	—	—	—	—
RELA-18-161249	02-61543	1–1.5	ALLH	— —	_	_	_	_	_	N3B-2018-3962-1	_	_	_	_	—	—
RELA-18-164455	02-61544	2–3	ALLH	— —	—	_	_	_	—	N3B-2019-391	_	_	_	—	_	_
RELA-18-164465	02-61544	4.25–5	QBT1G		—	—	—	—	—	N3B-2019-391	—	—	—	—	_	_
RELA-18-164475	02-61544	6–7	QBT1G		—	—	—	—	—	N3B-2019-391	—	—	—	—	—	_
RELA-18-164456	02-61545	2–3	ALLH		_	_	_	_	_	N3B-2019-412	_	_	_	_	_	_
RELA-18-164466	02-61545	4–5	ALLH	— —	_	—	_	—	—	N3B-2019-412	_	_	_	_	_	_
RELA-18-164476	02-61545	6–7	ALLH	— —	_	—	_	—	—	N3B-2019-412	_	_	_	_	_	_
RELA-18-164448	02-61547	0–1	ALLH	— —	_	_	_	_	_	N3B-2019-391	_	_	_	_	_	_
RELA-18-164458	02-61547	2–3	ALLH	— —	—	_			—	N3B-2019-391			—	—	_	—

Table 3.4-1 (continued)

Sample ID	Location ID	Depth (ft)	Media	Americium-241	Nitrate	Gamma-emitting Radionuclides	Tritium	Isotopic Plutonium	Isotopic Uranium	TAL Metals	PCBs	Perchlorate	Strontium-90	SVOCs	vocs	Dioxins/Furans	Cyanide (Total)
RELA-18-164468	02-61547	4–5	ALLH	_	_	_	_	_	—	_	N3B-2019-391	_	_	_	—	—	_
RELA-18-164478	02-61547	6–7	ALLH	_	_	_	_	_	—	_	N3B-2019-391	_	_	_	—	—	_
RELA-18-164459	02-61548	2–3	ALLH	_	_	_	_	_	—	_	N3B-2019-380	_	_	_	_	_	—
RELA-18-164469	02-61548	4-4.5	ALLH	_	_	—	_	—	—	_	N3B-2019-380	_	_	—	—	—	—
RELA-18-164479	02-61548	6–7	QBT1G	—	_	—	—	—	—	—	N3B-2019-380	_	—	—	-	—	-
RELA-18-164470	02-61549	4–5	QBT1G	—	—	—	—	—	—	—	N3B-2019-391	_	—	—	—	—	—
RELA-18-164480	02-61549	6–7	QBT1G	_	_	—	—	—	_	—	N3B-2019-391	_	—	—	_	_	_
RELA-18-164451	02-61550	0–1	ALLH	_	_	—	—	_	_	—	N3B-2019-391	_	—	_	_	_	_
RELA-18-164461	02-61550	2–3	ALLH	—	_	—	_		_	—	N3B-2019-391	_	—	_	-	_	_
RELA-18-164471	02-61550	4–5	ALLH	_	_	_	—	—	_	_	N3B-2019-412	_	—	—	_	—	_
RELA-18-164481	02-61550	6–7	ALLH	_	_	_	—	—	_	_	N3B-2019-412	_	—	—	_	—	_
RELA-18-164462	02-61551	2–3	ALLH	_	_	_	—	—	_	_	N3B-2019-412	_	—	—	_	—	_
RELA-18-164472	02-61551	4–5	ALLH	_	_	—	—	_	_	—	N3B-2019-412	_	—	_	_	_	_
RELA-18-164482	02-61551	6–7	ALLH	_	_	—	—	_	_	—	N3B-2019-412	_	—	_	_	_	_
RELA-18-164453	02-61552	0–1	ALLH	—	_	—	_		_	—	N3B-2019-391	_	—	_	-	_	_
RELA-18-164463	02-61552	2–3	ALLH	_	_	—	—	_	—	—	N3B-2019-391	_	—	_	_	_	_
RELA-18-164473	02-61552	4–5	ALLH	_	_	—	—	—	—	—	N3B-2019-391	_	—	_	_	_	_
RELA-18-164483	02-61552	6–7	QBT1G	—	-	-	—	_	—	—	N3B-2019-391	_	—	—	-	-	_
RELA-18-164464	02-61553	2–3	ALLH	—	_	-	—	—	_	—	N3B-2019-412	_	—	—	-	-	-
RELA-18-164474	02-61553	4–5	ALLH	—	—	—	—	—	—	—	N3B-2019-412	_	—	—	—	_	_
RELA-18-164484	02-61553	6–7	ALLH	—	_	-	—	—	—	—	N3B-2019-412	_	—	—	-	 _	<u> </u>

^a Analytical request number.

^b — = Analysis not requested.

 Table 3.4-2

 Inorganic Chemicals Detected or Detected above BVs at SWMU 02-014

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Field Sample ID	Location ID	Depth (ft)	Media	Aluminu	Arsenic	Barium	Cadmium	Calcium	Chromium	Copper	Iron	Magnesi
Qbt1 g, Qct, Qbo I	Background Value ^a	1	1	3560	0.56	25.7	0.4	1900	2.6	3.96	3700	739
Qbt 2, 3, 4 Backgr	-			7340	2.79	46	1.63	2200	7.14	4.66	14,500	1690
Soil Background				29,200	8.17	295	0.4	6120	19.3	14.7	21,500	4610
Industrial SSL ^b				1,290,000	35.9	255,000	1110	40,700,000	505 ^c	51,900	908,000	5,680,000
Recreational SSL ^c	l			619,000	42.9	124,000	457	na ^e	281 ^c	24,800	434,000	na
Residential SSL ^b				78,000	7.07	15,600	70.5	13,000,000	96.6 ^c	3130	54,800	20,900,000
RE02-07-1572	02-600387	0–0.5	SOIL	f	_	—	—	—	—	_	—	—
RE02-07-1573	02-600387	2–2.6	QAL	_	_	_	0.497 (U)	—	_	_	_	_
RE02-07-1878	02-600449	4.5–9.5	QAL	—	_	—	0.517 (U)	—	—	—	—	—
RE02-17-141643	02-613762	4–5	QBO	10,800	0.735 (J)	57.1	—	7260	3.16	—	4860	1270
RE02-17-141644	02-613762	6–7	QBO	5020	—	—	—	—	2.74	—	—	—
RE02-17-141646	02-61429	2–3	SOIL	—	—	—	—	—	—	—	—	—
RE02-17-141660	02-61432	0–1	SOIL	—	—	—	—	—	—	—	—	—
RE02-17-141614	02-61432	6–7	QBT3	—	—	—	—	—	—	—	—	—
RE02-17-141615	02-61435	0–0.1	SOIL	—	—	—	—	—	—	47.5	—	—
RE02-17-141625	02-61435	4–5	QBO	6440	0.662 (J)	29.9	—	—	—	—	5250	808
RE02-17-141648	02-61435	6–7	QBO	9180	0.855 (J)	47.5	—	2130	3.46	7.2	7560	1560
RE02-17-141661	02-61436	0–1	SOIL	—	_	—	—	—	—	_	—	—
RE02-17-141626	02-61436	2–3	SOIL	_		_	0.5 (U)	—	_	_	—	—
RE02-17-141617	02-61436	6–7	QBO	—	0.647 (J)	—	0.5 (U)	—	14.5	—	3810	—
RE02-17-141618	02-61437	0–1	SOIL	—	_	—	—	—	—	—	—	—
RE02-17-144637	02-61437	2–3	SOIL	—	_	—	—	—	—	—	—	—
RE02-17-144638	02-61437	4–5	SOIL	—	_	—	—	—	—	—	—	—
RE02-17-141662	02-61437	6–7	SOIL	—	_	—	—	—	—	—	—	—
RE02-17-141663	02-61438	0–1	SOIL	—	—	—	—	—	—	—	—	—
RE02-17-141628	02-61438	4–5	QBO	15,000	0.94 (J)	96.1	—	—	4.08	4.91	5700	1340
RE02-17-141619	02-61438	6–7	QBO	10,900	0.821 (J)	43.8	—	—	3	—	5580	1710
RE02-17-141651	02-61439	2–3	SOIL	—	—	—	—	—	—	—	—	—
RE02-17-141629	02-61439	4–5	QBO	9010	—	60.2	—	—	2.79	—	—	—
RE02-17-141620	02-61439	6–7	QBO	9420	0.628 (J)	49.3	—	—	2.81	—	3760	—
RE02-17-141668	02-61441	0–1	SOIL	—		—	—	—	_	—	—	—
RE02-17-144671	02-61441	2–3	SOIL	—		_	—	—	_	—	—	—
RE02-17-141654	02-61441	4–5	SOIL	—		_	—	—	_	—	—	—
RE02-17-144672	02-61441	4–5	SOIL	—	_	—	—	—	—	—	—	—

Table 3.4-2 (continued)

							,		1		1	1
Field Sample ID	Location ID	Depth (ft)	Media	Aluminum	Arsenic	Barium	Cadmium	Calcium	Chromium	Copper	Lon	Magnesium
Qbt1 g, Qct, Qbo E	Background Value ^a			3560	0.56	25.7	0.4	1900	2.6	3.96	3700	739
Qbt 2, 3, 4 Backgro	ound Value ^a			7340	2.79	46	1.63	2200	7.14	4.66	14,500	1690
Soil Background V	/alue ^a			29,200	8.17	295	0.4	6120	19.3	14.7	21,500	4610
Industrial SSL ^b				1,290,000	35.9	255,000	1110	40,700,000	505 ^c	51,900	908,000	5,680,000
Recreational SSL ^d				619,000	42.9	124,000	457	na ^e	281 ^c	24,800	434,000	na
Residential SSL ^b				78,000	7.07	15,600	70.5	13,000,000	96.6 ^c	3130	54,800	20,900,000
RE02-17-141655	02-61441	6–7	QBO	—	0.583 (J)	—	—	—	—	—	4040	—
RE02-17-141670	02-61442	0–1	SOIL	—	—	—	—	—	_	—	—	—
RE02-17-141656	02-61442	4–5	QBO	4450	0.604 (J)	25.9	—	—	—	—	5360	—
RE02-17-141657	02-61442	6–7	QBO	6880	0.762 (J)	43.2	—	—	2.85	—	5460	882
RE02-17-141685	02-61443	4–5	QBO	—	0.652 (J)	25.8	—	1950	_	—	4120	—
RE02-17-141785	02-61443	6–7	QBO	4560	0.653 (J)	29.8	—	—	—	—	5010	—
RE02-17-141693	02-61444	0–1	SOIL	—	—	—	—	—	_	—	—	_
RE02-17-141672	02-61444	6–7	SOIL	—	—	—	0.5 (U)	—	_	—	—	—
RE02-17-141673	02-61445	0–1	SOIL	—	—	—	—	—	_	—	—	_
RE02-17-141694	02-61445	6–7	QBO	3670	0.621 (J)	34.5	—	—	—	—	4320	—
RE02-17-141688	02-61446	2–3	SOIL	—	—	—	—	—	_	—	—	—
RE02-17-141681	02-61446	4-4.5	QBO	6520	0.591 (J)	40	—	—	2.62	—	—	871
RE02-17-141674	02-61446	6–7	QBO	9250	0.722 (J)	72.9	—	—	—	—	—	—
RE02-17-141696	02-61447	0–1	SOIL	—	—	—	—	—	_	—	—	—
RE02-17-141689	02-61447	2–3	SOIL	—	—	—	—	—	_	—	—	—
RE02-17-141682	02-61447	4–5	SOIL	—	—	—	—	—	—	—	—	—
RE02-17-141675	02-61447	6–7	SOIL	—	—	—	—	—	—	—	—	—
RE02-17-141697	02-61448	0–1	SOIL	—	—	—	—	—	_	_	—	—
RE02-17-141683	02-61448	4–5	SOIL	—	—	—	—	—	—	—	—	—
RE02-17-141561	02-61450	6–7	QBO	4070	—	25.8	—	—	—	—	4800	—
RE02-17-141570	02-61451	6–7	QBO	6250	0.594 (J)	31	—	—	3.27	_	6030	851
RE02-17-141579	02-61452	6–7	QBO	5380	0.564 (J)	—	—	—	—	—	5650	819
RE02-17-141586	02-61453	2–3	SOIL	—	—	—	—	—	—	—	—	—
RE02-17-141588	02-61453	6–7	SOIL	—	—	—	—	—	_	_	—	—
RE02-17-141595	02-61454	2–3	SOIL	—	—	—	—	—	_	_	—	—
RE02-17-141596	02-61454	4–5	SOIL	—	—	—	0.5 (U)	—	_	_	—	—
RE02-17-141597	02-61454	6–7	QBO	5760	0.733 (J)	34.5	_	_	2.96	_	6050	785

Table 3.4-2 (continued)

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Field Sample ID	Location ID	Depth (ft)	Media	Mangan	Mercury	Nickel	Nitrate	Perchlorate	Selenium	Thallium	Vanadium	Zinc
Qbt1 g, Qct, Qbo I	Background Value ^a			189	0.1	2	na	na	0.3	1.22	4.59	40
Qbt 2, 3, 4 Backgr	ound Value ^a			482	0.1	6.58	na	na	0.3	1.1	17	63.5
Soil Background	Value ^a			671	0.1	15.4	na	na	1.52	0.73	39.6	48.8
Industrial SSL ^b				160,000	389	25,700	2,080,000	908	6490	13	6530	389,000
Recreational SSL ^o	d			14,800	186	12,400	991,000	434	3100	6.19	3100	186,000
Residential SSL ^b				10,500	23.5	1560	125,000	54.8	391	0.782	394	23,500
RE02-07-1572	02-600387	0–0.5	SOIL	—	—	—	1.71 (J-)	—	—	—	—	78.2
RE02-07-1573	02-600387	2–2.6	QAL	—	—	—	1.77 (J-)	0.000813 (J)	—	—	—	—
RE02-07-1878	02-600449	4.5–9.5	QAL	—	—	—	—	—	—	-	—	—
RE02-17-141643	02-613762	4–5	QBO	—	—	2.76	NA ^g	NA	0.462 (U)	—	4.95	—
RE02-17-141644	02-613762	6–7	QBO	—	—	2.26	NA	NA	—	—	—	—
RE02-17-141646	02-61429	2–3	SOIL	—	—	—	NA	NA	2.5 (U)	—	—	—
RE02-17-141660	02-61432	0–1	SOIL	—	—	_	NA	NA	2.5 (U)	—	—	—
RE02-17-141614	02-61432	6–7	QBT3	—	—	_	NA	NA	1.11 (J)	—	—	—
RE02-17-141615	02-61435	0–0.1	SOIL	—	0.538 (J)	—	NA	NA	_	—	—	58.7
RE02-17-141625	02-61435	4–5	QBO	—	_	2.32	NA	NA	_	—	4.62	—
RE02-17-141648	02-61435	6–7	QBO	232	_	3.64	NA	NA	0.576 (J)	_	6.43	_
RE02-17-141661	02-61436	0–1	SOIL	—	0.127 (J)	_	NA	NA	_	—	—	54.6
RE02-17-141626	02-61436	2–3	SOIL	_		_	NA	NA	_	_	_	_
RE02-17-141617	02-61436	6–7	QBO	251	_	2.21	NA	NA	0.697 (U)	2.5 (U)	_	_
RE02-17-141618	02-61437	0–1	SOIL	—	0.209 (J)	—	NA	NA	_	—	—	—
RE02-17-144637	02-61437	2–3	SOIL	_		—	NA	NA	_	—	—	_
RE02-17-144638	02-61437	4–5	SOIL	_		_	NA	NA	_	_	_	_
RE02-17-141662	02-61437	6–7	SOIL	_	0.116 (J)	_	NA	NA	_	_	_	_
RE02-17-141663	02-61438	0–1	SOIL	_		—	NA	NA	2.5 (U)	—	—	_
RE02-17-141628	02-61438	4–5	QBO	213		4.37	NA	NA	_	—	6.34	_
RE02-17-141619	02-61438	6–7	QBO	—	_	3.32	NA	NA	 _	—	5.83	—
RE02-17-141651	02-61439	2–3	SOIL	—	_	_	NA	NA	2.5 (U)	—	—	—
RE02-17-141629	02-61439	4–5	QBO	_	_	2.36	NA	NA	2.5 (U)	_	_	_
RE02-17-141620	02-61439	6–7	QBO	_	_	2.56	NA	NA	2.5 (U)	_	_	_
RE02-17-141668	02-61441	0–1	SOIL	_	_	—	NA	NA	2.5 (U)	-	_	—
RE02-17-144671	02-61441	2–3	SOIL	_	_	_	NA	NA	_	_	_	_
RE02-17-141654	02-61441	4–5	SOIL	_	_	_	NA	NA	2.5 (U)	-	_	—

Table 3.4-2 (continued)

Field Sample ID	Location ID	Depth (ft)	Media	Manganese	Mercury	Nickel	Nitrate	Perchlorate	Selenium	Thallium	Vanadium	Zinc
Qbt1 g, Qct, Qbo I	Background Value ^a	1	1	189	0.1	2	na	na	0.3	1.22	4.59	40
Qbt 2, 3, 4 Backgr	ound Value ^a			482	0.1	6.58	na	na	0.3	1.1	17	63.5
Soil Background	Value ^a			671	0.1	15.4	na	na	1.52	0.73	39.6	48.8
Industrial SSL ^b				160,000	389	25,700	2,080,000	908	6490	13	6530	389,000
Recreational SSL ^c	1			14,800	186	12,400	991,000	434	3100	6.19	3100	186,000
Residential SSL ^b				10,500	23.5	1560	125,000	54.8	391	0.782	394	23,500
RE02-17-144672	02-61441	4–5	SOIL	—	—	—	NA	NA	—	—	—	—
RE02-17-141655	02-61441	6–7	QBO	—	—	—	NA	NA	2.5 (U)	2.5 (U)	—	_
RE02-17-141670	02-61442	0–1	SOIL	—	—	—	NA	NA	2.5 (U)	—	—	_
RE02-17-141656	02-61442	4–5	QBO	—	—	—	NA	NA	0.476 (J)	—	—	—
RE02-17-141657	02-61442	6–7	QBO	242	—	2.65	NA	NA	2.5 (U)	—	5.74	—
RE02-17-141685	02-61443	4–5	QBO	—	—	—	NA	NA	0.363 (J)	—	—	—
RE02-17-141785	02-61443	6–7	QBO	—	—	2.1	NA	NA	2.5 (U)	—	—	—
RE02-17-141693	02-61444	0–1	SOIL	—	0.289 (J)	—	NA	NA	—	—	—	—
RE02-17-141672	02-61444	6–7	SOIL	—	—	—	NA	NA	—	2.5 (U)	—	—
RE02-17-141673	02-61445	0–1	SOIL	—	0.434 (J)	—	NA	NA	—	—	—	—
RE02-17-141694	02-61445	6–7	QBO	240	—	—	NA	NA	0.42 (J)	—	—	—
RE02-17-141688	02-61446	2–3	SOIL	—	—	—	NA	NA	2.5 (U)	—	—	—
RE02-17-141681	02-61446	4-4.5	QBO	—	—	2.31	NA	NA	2.5 (U)	—	—	
RE02-17-141674	02-61446	6–7	QBO	—	—	2.06	NA	NA	0.464 (J)	-	—	—
RE02-17-141696	02-61447	0–1	SOIL	—	—	—	NA	NA	2.5 (U)	—	—	—
RE02-17-141689	02-61447	2–3	SOIL	—	—	—	NA	NA	2.5 (U)	—	—	—
RE02-17-141682	02-61447	4–5	SOIL	—	—	—	NA	NA	2.5 (U)	—	—	
RE02-17-141675	02-61447	6–7	SOIL	—	—	—	NA	NA	2.5 (U)	—	—	
RE02-17-141697	02-61448	0–1	SOIL	—	—	—	NA	NA	—	—	—	105
RE02-17-141683	02-61448	4–5	SOIL	—	—	—	NA	NA	2.5 (U)	—	—	—
RE02-17-141561	02-61450	6–7	QBO	210	—	—	NA	NA	0.403 (U)	—	—	—
RE02-17-141570	02-61451	6–7	QBO	229	—	2.48	NA	NA	2.5 (U)	—	4.86	—
RE02-17-141579	02-61452	6–7	QBO	—	—	—	NA	NA	0.445 (U)	—	—	—
RE02-17-141586	02-61453	2–3	SOIL	_	—	_	NA	NA	2.5 (U)	—	—	_

Table 3.4-2 (continued)

Field Sample ID	Location ID	Depth (ft)	Media	Manganese	Mercury	Nickel	Nitrate	Perchlorate	Selenium	Thallium	Vanadium	Zinc
Qbt1 g, Qct, Qbo	Background Value ^a	3		189	0.1	2	na	na	0.3	1.22	4.59	40
Qbt 2, 3, 4 Backgr	ound Value ^a			482	0.1	6.58	na	na	0.3	1.1	17	63.5
Soil Background	Value ^a			671	0.1	15.4	na	na	1.52	0.73	39.6	48.8
Industrial SSL ^b				160,000	389	25,700	2,080,000	908	6490	13	6530	389,000
Recreational SSL	d			14,800	186	12,400	991,000	434	3100	6.19	3100	186,000
Residential SSL ^b				10,500	23.5	1560	125,000	54.8	391	0.782	394	23,500
RE02-17-141588	02-61453	6–7	SOIL	—	—	—	NA	NA	2.5 (U)	—	—	—
RE02-17-141595	02-61454	2–3	SOIL	—	—	—	NA	NA	—	2.5 (U)	—	—
RE02-17-141596	02-61454	4–5	SOIL	—	—	—	NA	NA	—	2.5 (U)	—	—
RE02-17-141597	02-61454	6–7	QBO	274	—	2.45	NA	NA	0.657 (U)	—	5.84	—

Notes: Results are in mg/kg. Data qualifiers are defined in Appendix A.

^a BVs are from LANL (1998, 059730).

 $^{\rm b}\,$ SSLs are from NMED (2017, 602273), unless otherwise noted.

^c SSLs are for total chromium.

^d SSLs are from LANL (2017, 602581).

^e na = Not available.

 f — = Not detected or not detected above BV.

^g NA = Not analyzed.

Table 3.4-3Organic Chemicals other than PCBs Detected at SWMU 02-014

Sample ID	Location ID	Depth (ft)	Media	Anthracene	Benzo(a)anthracene	Benzo(a)pyrene	Benzo(b)fluoranthene	Benzo(g,h,i)perylene	Benzo(k)fluoranthene	Chrysene	Fluoranthene	Heptachlorodibenzodioxin[1,2,3,4,6,7,8-]	Heptachlorodibenzodioxins (Total)	Heptachlorodibenzofuran[1,2,3,4,6,7,8-]	Heptachlorodibenzofuran[1,2,3,4,7,8,9-]
Industrial SSL ^a				253,000	32.3	23.6	32.3	25,300 ^b	323	3230	33,700	na ^c	na	na	na
Recreational SSL ^d				86,300	88.8	8.88	88.8	8630 ^b	888	8880	11,500	na	na	na	na
Residential SSL ^a				17400	1.53	1.12	1.53	1740 ^b	15.3	1530	2320	na	na	na	na
RE02-07-1572	02-600387	0–0.5	SOIL	0.00903 (J)	0.0516	0.0551	0.0702	0.0271 (J)	0.041	0.0598	0.0984	0.00016	0.001	0.0000475	0.00000235 (J)
RE02-07-1573	02-600387	2–2.6	QAL	0.00727 (J)	0.032 (J)	0.0332 (J)	0.0424	0.0222 (J)	0.0219 (J)	0.0386	0.0529	0.000166	0.00108	0.0000472	0.00000182 (J)
RE02-07-1878	02-600449	4.5–9.5	QAL	e	—	—	_	—	—	_	—	0.00000134 (J)	0.00000429	0.00000545	0.00000799 (J)

Table 3.4-3 (continued)

Sample ID	Location ID	Depth (ft)	Media	Heptachlorodibenzofurans (Total)	Hexachlorodibenzodioxin[1,2,3,4,7,8-]	Hexachlorodibenzodioxin[1,2,3,6,7,8-]	Hexachlorodibenzodioxin[1,2,3,7,8,9-]	Hexachlorodibenzodioxins (Total)	Hexachlorodibenzofuran[1,2,3,4,7,8-]	Hexachlorodibenzofuran[1,2,3,6,7,8-]	Hexachlorodibenzofuran[1,2,3,7,8,9-]	Hexachlorodibenzofuran[2,3,4,6,7,8-]	Hexachlorodibenzofurans (Total)	Indeno(1,2,3-cd)pyrene
Industrial SSL ^a				na	na	na	na	na	na	na	na	na	na	32.3
Recreational SSL ^c				na	na	na	na	na	na	na	na	na	na	88.8
Residential SSL ^a				na	na	na	na	na	na	na	na	na	na	1.53
RE02-07-1572	02-600387	0–0.5	SOIL	0.000205	0.00000071 (J)	0.00000332	0.0000013 (J)	0.0000622	0.00000486	0.00000166 (J)	0.0000005 (J)	0.00000273	0.0000561	0.0252 (J)
RE02-07-1573	02-600387	2–2.6	QAL	0.000235	0.000000738 (J)	0.0000037	0.0000091 (J)	0.0000743	0.00000182 (J)	0.00000621 (J)	—	0.00000981 (J)	0.0000446	0.0184 (J)
RE02-07-1878	02-600449	4.5–9.5	QAL	0.0000109	—	—	_	-	0.00000641	0.00000263	0.00000051 (J)	0.00000430	0.0000454	_

Table 3.4-3 (continued)

Sample ID	Location ID	Depth (ft)	Media	Octachlorodibenzodioxin[1,2,3,4,6,7,8,9-]	Octachlorodibenzofuran[1,2,3,4,6,7,8,9-]	Pentachlorodibenzodioxin[1,2,3,7,8-]	Pentachlorodibenzodioxins (Total)	Pentachlorodibenzofuran[1,2,3,7,8-]	Pentachlorodibenzofuran[2,3,4,7,8-]	Pentachlorodibenzofurans (Totals)	Phenanthrene	Pyrene	Tetrachlorodibenzofuran[2,3,7,8-]	Tetrachlorodibenzofurans (Totals)	Toluene
Industrial SSL ^a				na	na	na	na	na	na	na	25,300	25,300	0.00243	na	61,100
Recreational SSL ^c				na	na	na	na	na	na	na	8630	8630	0.00297	na	47,600
Residential SSL ^a				na	na	na	na	na	na	na	1740	1740	0.00049	na	5220
RE02-07-1572	02-600387	0–0.5	SOIL	0.0012	0.00022	0.00000204 (J)	0.00000234	0.00000101 (J)	0.00000652	0.0000449 (J)	0.0317 (J)	0.0771	0.00000377	0.000017	NA ^f
RE02-07-1573	02-600387	2–2.6	QAL	0.00123	0.000248	0.000000156 (J)	0.0000007	—	0.00000155 (J)	0.0000102 (J)	0.0212 (J)	0.0423	0.00000867 (J)	0.00000232	0.000465 (J)
RE02-07-1878	02-600449	4.5–9.5	QAL	0.0000134	0.0000057	—	—	0.0000019 (J)	0.0000128	0.000073	1-	—	0.00000639	0.0000324	—

Notes: Results are in mg/kg. Data qualifiers are defined in Appendix A.

^a SSLs are from NMED (2017, 602273), unless otherwise noted.

^b Pyrene used as a surrogate based on structural similarity.

^c na = Not available.

^d SSLs are from LANL (2017, 602581).

^e — = Not detected.

^f NA = Not analyzed.

Table 3.4-4 PCBs Detected at SWMU 02-014

Sample ID	Location ID	Depth (ft)	Media	Aroclor-1254	Aroclor-1260
Industrial SSL ^a				11	11.1
Recreational SSL ^b	5.53	10.3			
Residential SSL ^a	1.14	2.43			
RE02-07-1572	02-600387	0–0.5	SOIL	0.234	0.553
RE02-07-1573	02-600387	2–2.6	QAL	0.0532	0.105
RE02-07-1878	02-600449	4.5–9.5	QAL	c	0.171
RE02-10-22130	02-600449	6–6.2	SOIL	_	0.69
RE02-10-22133	02-612451	6–6.2	SOIL	_	4
RE02-10-22136	02-612452	6–6.2	SOIL	_	1.1
RE02-10-22137	02-612452	8–8.2	SOIL	_	2.14
RE02-10-22139	02-612453	6–6.2	SOIL	_	0.377
RE02-10-22140	02-612453	8–8.2	SOIL	_	0.191
RE02-10-26105	02-613001	6–6.5	SOIL	_	0.536
RELA-18-151143	02-613001	6–7	SOIL	_	5.15
RE02-10-26106	02-613001	7–7.5	SOIL	—	5.48
RELA-18-151144	02-613001	8–9	SOIL	_	1.47
RE02-10-26107	02-613002	6–6.5	SOIL	—	7.98
RE02-10-26108	02-613002	8–8.5	SOIL	—	1.67
RE02-10-26638	02-613122	2–2.2	SOIL	—	0.328
RE02-10-26639	02-613122	4-4.2	SOIL	0.0407 (J)	0.178
RE02-10-26640	02-613124	6–6.2	SOIL	—	2.03
RE02-10-26641	02-613124	8–8.2	SOIL	0.17	1.24
RELA-18-151133	02-613287	2–3	SOIL	—	0.485
RELA-18-151134	02-613287	4–5	SOIL	—	0.769
RE02-11-315	02-613287	6–6.2	SOIL	—	13.9
RELA-18-151135	02-613287	6–7	SOIL	—	1.92
RE02-11-316	02-613287	8–8.2	SOIL	—	12.6
RE02-17-136051	02-613287	8–9	QBO	—	0.586
RE02-11-317	02-613288	6–6.2	SOIL	—	2.48
RE02-11-318	02-613288	8–8.2	SOIL	—	1.07
RE02-11-320	02-613289	6–6.2	SOIL		3.39
RELA-18-151137	02-613289	6–7	SOIL	_	0.767
RELA-18-151138	02-613289	9–10	SOIL		10.8
RELA-18-151139	02-613289	11–11.25	SOIL	—	20.7

Table 3.4-4 (continued)

Sample ID	Location ID	Depth (ft)	Media	Aroclor-1254	Aroclor-1260
Industrial SSL ^a				11	11.1
Recreational SSL ^b				5.53	10.3
Residential SSL ^a				1.14	2.43
RE02-11-325	02-613292	4-4.2	SOIL	—	0.825
RE02-11-2218	02-613626	8–9	SOIL	0.197	1.03
RE02-11-2219	02-613626	10–11	SOIL	—	3.3
RE02-11-2220	02-613627	6–7	SOIL	1.39	7.29
RE02-11-2221	02-613627	8–9	SOIL	—	4.16
RE02-11-2222	02-613627	10–11	SOIL	—	1.89
RELA-18-151122	02-613667	2–3	SOIL	—	0.199
RELA-18-151123	02-613667	4–5	SOIL	—	0.107
RE02-11-2523	02-613667	6–6.2	SOIL	7.11	13.3
RELA-18-151124	02-613667	6–7	SOIL	—	0.0347
RE02-11-2524	02-613667	8–8.2	SOIL	3.74	6.3
RELA-18-151125	02-613667	8–9	QBO	—	0.013
RE02-11-2525	02-613667	10–10.2	SOIL	4.93	7.73
RELA-18-151127	02-613668	2–3	SOIL	—	0.552
RELA-18-151128	02-613668	4–5	SOIL	—	0.0465
RELA-18-151129	02-613668	6–7	SOIL	—	0.144
RE02-11-2526	02-613668	8–8.2	SOIL	—	1.52
RELA-18-151130	02-613668	8–9	SOIL	—	0.0694
RE02-11-2527	02-613668	10–10.2	SOIL	—	3.21
RE02-11-2798	02-613699	12–12.2	QBT3	0.636 (J)	3.34
RE02-11-2799	02-613700	8–8.2	SOIL	0.0505	0.258
RE02-11-2800	02-613700	10–10.2	SOIL	0.0617	0.327
RE02-11-2801	02-613700	12–12.2	SOIL	0.256 (J)	1.19
RE02-11-3145	02-613700	14–14.2	SOIL	0.779	2.27
RE02-11-3146	02-613761	6–6.2	SOIL	—	10.7
RE02-11-3147	02-613761	8–8.2	SOIL	—	3.85
RE02-11-3148	02-613761	10–10.2	SOIL	—	1.53
RE02-11-3149	02-613761	12–12.2	QBT3	_	1.75
RE02-11-3150	02-613761	14–14.2	QBT3	_	0.0445
RELA-18-151117	02-613762	2–3	SOIL	—	1.82
RELA-18-151118	02-613762	4.25–5	QBO	_	0.0673
RELA-18-151119	02-613762	6–7	QBO	_	0.0464
RELA-18-151120	02-613762	8–9	QBO	_	0.0432

Table 3.4-4 (continued)

Sample ID	Location ID	Depth (ft)	Media	Aroclor-1254	Aroclor-1260
Industrial SSL ^a				11	11.1
Recreational SSL ^b				5.53	10.3
Residential SSL ^a				1.14	2.43
RE02-17-135776	02-61428	0–1	SOIL	—	0.0227
RE02-17-135778	02-61428	4–5	SOIL	—	0.00645 (J)
RE02-17-135780	02-61429	0–1	SOIL	—	0.333
RE02-17-135781	02-61429	2–3	SOIL	—	0.0457
RE02-17-135782	02-61429	4–5	SOIL	—	0.0106
RE02-17-135783	02-61429	5–6	SOIL	—	0.0133
RELA-18-151045	02-61432	2–3	SOIL	0.00811	0.0169
RELA-18-151046	02-61432	4–5	SOIL	—	0.128
RELA-18-151047	02-61432	6.75–7	QBO	—	0.974
RELA-18-151048	02-61432	8–8.75	QBO	0.0618	0.116
RE02-17-135854	02-61432	14–15	SOIL	_	0.00381 (J)
RE02-17-135855	02-61432	16–17	SOIL	0.0193	—
RE02-17-135856	02-61432	19–20	SOIL	0.00388 (J)	—
RELA-18-151057	02-61435	2–3	SOIL	—	0.00467
RELA-18-151058	02-61435	4–5	QBO	_	0.00253 (J)
RELA-18-151059	02-61435	6–7	QBO	—	0.00755
RE02-17-135893	02-61435	11–12	QBO	—	0.0184
RE02-17-135894	02-61435	14–15	QBO	—	0.00879
RELA-18-151062	02-61436	2–3	SOIL	—	0.264
RELA-18-151063	02-61436	4–5	SOIL	—	0.0586
RELA-18-151064	02-61436	6–7	QBO	—	0.288
RE02-17-144637	02-61437	2–3	SOIL	—	1.41
RE02-17-144638	02-61437	4–5	SOIL	—	1.9
RELA-18-151065	02-61437	6–7	SOIL	—	0.324
RELA-18-151066	02-61437	8–9	SOIL	—	0.036
RELA-18-151067	02-61437	11–11.8	QBO	—	0.0106
RE02-17-135916	02-61438	0–1	SOIL	—	0.0228
RE02-17-135917	02-61438	2–3	SOIL	—	0.00532 (J)
RE02-17-135919	02-61438	6–7	QBO	_	0.00369 (J)
RE02-17-135920	02-61438	8–9	QBO	—	0.00417 (J)
RE02-17-135921	02-61438	11–12	QBO	_	0.00314 (J)
RE02-17-135922	02-61438	14–15	QBO	_	0.00422 (J)
RE02-17-135923	02-61438	16–17	QBO	_	0.00368 (J)

Sample ID	Location ID	Depth (ft)	Media	Aroclor-1254	Aroclor-1260
Industrial SSL ^a		_ • • • • • • • • • • • • • • • • • • •		∢	∢
Recreational SSL ^b				5.53	10.3
Residential SSL ^a				1.14	2.43
RE02-17-135925	02-61439	0–1	SOIL	_	0.0928
RE02-17-135926	02-61439	2–3	SOIL	_	0.0183
RE02-17-135928	02-61439	6–7	QBO	_	0.00459 (J)
RE02-17-135929	02-61439	8–9	QBO	_	0.00411 (J)
RE02-17-135930	02-61439	11–12	QBO	_	0.00441 (J)
RE02-17-135931	02-61439	14–15	QBO	—	0.0111
RE02-17-135932	02-61439	16–17	QBO	—	0.00586 (J)
RE02-17-135927	02-61439	19–20	QBO	—	0.0207
RELA-18-151069	02-61440	2–3	SOIL	—	0.423
RELA-18-151070	02-61440	4–5	SOIL	—	0.257
RELA-18-151071	02-61440	6–7	SOIL	—	3.75 (J)
RELA-18-151072	02-61440	8–9	SOIL	—	0.0545
RE02-17-135939	02-61440	11–12	QBO	_	0.0511
RE02-17-135940	02-61440	14–15	QBO	_	0.00316 (J)
RE02-17-144671	02-61441	2–3	SOIL	_	16.1
RE02-17-144672	02-61441	4–5	SOIL	_	0.919
RELA-18-151043	02-61441	6–7	QBO	_	0.0636
RELA-18-151073	02-61441	8–8.5	QBO	_	0.0277
RE02-17-135948	02-61441	11–12	QBO	_	0.0123
RE02-17-135949	02-61441	14–15	QBO	_	0.0077
RELA-18-151075	02-61442	2–3	SOIL	_	0.888
RELA-18-151076	02-61442	4–5	QBO	_	0.911
RELA-18-151077	02-61442	6–7	QBO	_	3.4
RELA-18-151078	02-61442	8–9	QBO	_	0.15
RE02-17-135957	02-61442	11–12	QBO	_	0.0193
RE02-17-135958	02-61442	14–15	QBO	—	0.0221
RELA-18-151080	02-61443	2–3	SOIL	—	0.824
RELA-18-151081	02-61443	4–5	QBO	—	0.21
RELA-18-151082	02-61443	6–7	QBO	—	0.028
RELA-18-151083	02-61443	8–9	QBO	—	0.0434
RELA-18-151084	02-61443	11–12	QBO	—	0.016
RELA-18-151086	02-61444	2–3	SOIL	—	0.0227
RELA-18-151087	02-61444	4–5	SOIL	—	0.0356

Sample ID	Location ID	Depth (ft)	Media	Aroclor-1254	Aroclor-1260
Industrial SSL ^a				11	11.1
Recreational SSL ^b				5.53	10.3
Residential SSL ^a				1.14	2.43
RELA-18-151088	02-61444	6–7	SOIL	_	0.0283
RELA-18-151089	02-61444	8–8.85	SOIL	—	1.55
RE02-17-136003	02-61444	11–12	QBO	—	0.00645 (J)
RE02-17-136004	02-61444	14–15	QBO	—	0.00938
RE02-17-136005	02-61444	16–17	QBO	—	0.0117
RE02-17-136006	02-61444	19–20	QBO	—	0.00667 (J)
RE02-17-136007	02-61445	0—1	SOIL	—	0.0869
RE02-17-136010	02-61445	6–7	QBO	—	0.0225
RE02-17-136011	02-61445	8–9	QBO	—	0.00381 (J)
RE02-17-136015	02-61445	14-20	QBO	—	0.0746
RE02-17-139388	02-61446	0–1	SOIL	—	0.0593
RE02-17-139389	02-61446	2–3	SOIL	—	0.00725
RE02-17-139390	02-61446	4-4.5	QBO	—	0.0132
RE02-17-139391	02-61446	6–7	SOIL	—	0.0227
RE02-17-139392	02-61446	8–9	SOIL	—	0.00911
RE02-17-139393	02-61446	11–12	SOIL	—	0.0164
RE02-17-139394	02-61446	14–15	QBO	—	0.004 (J)
RE02-17-139395	02-61446	16–17	QBO	—	0.00974
RE02-17-139396	02-61446	19–20	SOIL	—	0.0064 (J)
RELA-18-151037	02-61447	0—1	SOIL	3.93	8.91
RELA-18-151038	02-61447	2–3	SOIL	0.174	0.348
RELA-18-151039	02-61447	4–5	QBO	0.0461	0.112
RELA-18-151040	02-61447	6–7	QBO	0.15	0.319
RE02-17-139403	02-61447	14—15	QBO	—	0.141
RE02-17-139404	02-61447	16–17	QBO	—	0.0663
RE02-17-139405	02-61447	19–20	QBO	—	0.174
RELA-18-151050	02-61448	2–3	SOIL	—	0.0562
RELA-18-151051	02-61448	4–5	SOIL	—	0.0103
RELA-18-151052	02-61448	6–7	SOIL	—	0.0233
RELA-18-151053	02-61448	8–9	SOIL	—	0.00685
RELA-18-151054	02-61448	11–12	SOIL	—	0.591
RELA-18-151055	02-61448	13–14	SOIL	—	0.0707
RE02-17-139413	02-61448	16–17	QBO		0.00424 (J)

Sample ID	Location ID	Depth (ft)	Media	Aroclor-1254	Aroclor-1260
Industrial SSL ^a				11	11.1
Recreational SSL ^b				5.53	10.3
Residential SSL ^a				1.14	2.43
RE02-17-139414	02-61448	19–20	QBO	_	0.0186
RELA-18-151091	02-61450	2–3	SOIL	_	0.0224
RELA-18-151092	02-61450	4–5	SOIL	_	0.00351 (J)
RE02-17-141562	02-61450	8–9	QBO	_	0.0131
RE02-17-141564	02-61450	14–15	QBO	_	0.0701
RE02-17-141565	02-61450	16–17	QBO	_	0.0201
RELA-18-151094	02-61451	2–3	SOIL	_	0.067
RELA-18-151095	02-61451	4–5	SOIL	—	0.0382
RELA-18-151096	02-61451	6.25–7	QBO	_	0.0205
RELA-18-151097	02-61451	8–9	QBO	_	0.0244
RE02-17-141572	02-61451	11–12	QBO	_	0.0113
RE02-17-141573	02-61451	14–15	QBO	_	0.0396
RE02-17-141574	02-61451	16–17	QBO	_	0.0132
RELA-18-151099	02-61452	2–3	SOIL	—	21
RELA-18-151100	02-61452	4–5	SOIL	—	0.827
RELA-18-151101	02-61452	6–7	QBO	_	0.307
RELA-18-151102	02-61452	8–9	QBO	—	0.0167
RE02-17-141582	02-61452	14–15	QBO	—	0.218
RE02-17-141583	02-61452	16–17	QBO	_	0.0329
RE02-17-141584	02-61452	19–20	QBO	—	0.174
RELA-18-151104	02-61453	2–3	SOIL	—	2.34 (J)
RELA-18-151105	02-61453	4–5	SOIL	—	0.0121
RELA-18-151106	02-61453	6–7	SOIL	—	0.372
RELA-18-151107	02-61453	8–9	SOIL	—	0.0095
RE02-17-141590	02-61453	11–12	QBO	—	0.0405
RE02-17-141592	02-61453	16–17	QBO	—	0.0832
RE02-17-141593	02-61453	19–20	QBO	—	0.0522
RELA-18-151109	02-61454	2–3	SOIL		0.022
RELA-18-151110	02-61454	4–5	SOIL		0.00136 (J)
RE02-17-141597	02-61454	6–7	QBO	—	0.00933
RE02-17-141598	02-61454	8–9	QBO		0.00346 (J)
RE02-17-141599	02-61454	11–12	QBO	_	0.00921
RE02-17-136031	02-61454	16–17	QBO	_	0.0101

Sample ID	Location ID	Depth (ft)	Media	Aroclor-1254	Aroclor-1260
Industrial SSL ^a				11	11.1
Recreational SSL ^b				5.53	10.3
Residential SSL ^a				1.14	2.43
RELA-18-151112	02-61455	2–3	SOIL	—	0.00263 (J)
RELA-18-151113	02-61455	4–5	SOIL		0.00898
RELA-18-151114	02-61455	6–7	SOIL	—	0.035
RELA-18-151115	02-61455	8–8.75	SOIL	—	0.0112
RE02-17-141608	02-61455	11–12	QBO	—	0.172 (J+)
RE02-17-141609	02-61455	14–15	QBO		0.0531 (J+)
RE02-17-145060	02-61474	2–3	SOIL	—	1.67
RE02-17-145072	02-61474	4–5	SOIL	—	5.53
RE02-17-145084	02-61474	6–7	SOIL	—	3.54
RE02-17-145049	02-61475	0–1	SOIL	—	0.452
RE02-17-145061	02-61475	2–3	SOIL	—	0.073
RE02-17-145073	02-61475	4–5	SOIL	—	0.0395
RE02-17-145050	02-61476	0–1	SOIL	—	1.29
RE02-17-145062	02-61476	2–3	SOIL	—	0.0178
RE02-17-145074	02-61476	4–5	SOIL	—	0.0478
RE02-17-145086	02-61476	6–7	QBO	—	0.053
RE02-17-145098	02-61476	8–9	QBO	—	0.1
RE02-17-145110	02-61476	9–10	QBO	—	0.023
RE02-17-145051	02-61477	0–1	SOIL	—	5
RE02-17-145063	02-61477	2–3	SOIL	—	1.21
RE02-17-145064	02-61478	2–3	SOIL	—	2.41
RE02-17-145065	02-61479	2–3	SOIL	—	1.25
RE02-17-145077	02-61479	4–5	SOIL	—	1.57
RE02-17-145054	02-61480	0–1	SOIL	—	0.976
RE02-17-145066	02-61480	2–3	SOIL	—	0.0653
RE02-17-145078	02-61480	4–5	SOIL		0.179
RE02-17-145090	02-61480	6–7	SOIL		0.063
RE02-17-145102	02-61480	8–8.5	QBT3		0.101
RE02-17-145055	02-61481	0–1	SOIL		0.551
RE02-17-145067	02-61481	2–3	SOIL	—	0.192
RE02-17-145079	02-61481	4–5	SOIL	—	0.12
RE02-17-145091	02-61481	6–7	SOIL	—	0.0876
RE02-17-145103	02-61481	8–9	QBO	—	0.0505

Sample ID	Location ID	Depth (ft)	Media	Aroclor-1254	Aroclor-1260
Industrial SSL ^a				11	11.1
Recreational SSL ^b				5.53	10.3
Residential SSL ^a				1.14	2.43
RE02-17-145115	02-61481	9–10	QBO	_	0.0243
RE02-17-145056	02-61482	0–1	SOIL	_	11.3
RE02-17-145068	02-61482	2–3	SOIL	—	0.642
RE02-17-145080	02-61482	4–5	QBO	—	0.77
RE02-17-145092	02-61482	6–7	QBT3	—	0.471
RE02-17-145116	02-61482	8–9	QBO	—	0.456
RE02-17-145104	02-61482	9–10	QBO	—	0.641
RE02-17-145057	02-61483	0–1	SOIL	—	0.132
RE02-17-145069	02-61483	2–3	SOIL	—	0.0309
RE02-17-145081	02-61483	4–5	SOIL	—	0.0079
RE02-17-145093	02-61483	6–7	QBO	—	0.0107
RE02-17-145105	02-61483	8–9	QBO	—	0.00545
RE02-17-145117	02-61483	9–10	QBO	—	0.00353 (J)
RE02-17-145120	02-61486	0–1	SOIL	—	0.3
RE02-17-145130	02-61486	2–3	SOIL	—	0.333
RE02-17-145131	02-61487	2–3	SOIL	—	0.768
RE02-17-145141	02-61487	4–5	SOIL	—	1.59
RE02-17-145151	02-61487	5–5.5	QBO	—	0.476
RE02-17-145132	02-61488	2–3	QBO	—	1.6
RE02-17-145142	02-61488	4-4.1	QBO	—	2.25
RE02-17-145133	02-61489	2–3	SOIL	—	0.456
RE02-17-145143	02-61489	4–5	SOIL	—	0.422
RE02-17-145153	02-61489	6–7	QBO	—	0.0512
RE02-17-145163	02-61489	7–7.5	QBO	—	0.733
RE02-17-145124	02-61490	0–1	SOIL	0.0366	0.0572
RE02-17-145134	02-61490	2–3	SOIL	0.0284	0.0449
RE02-17-145125	02-61491	0–1	SOIL	—	0.016
RE02-17-145135	02-61491	2–3	SOIL	—	0.00456
RE02-17-145145	02-61491	4–5	QBO	—	0.00623
RE02-17-145126	02-61492	0–0.5	SOIL	0.0418	0.0792
RE02-17-145127	02-61493	0–1	SOIL	0.0248	0.043
RE02-17-145137	02-61493	2–3	SOIL	0.00632	0.011
RE02-17-145147	02-61493	4-4.5	QBO	0.00327 (J)	0.006

Sample ID	Location ID	Depth (ft)	Media	Aroclor-1254	Aroclor-1260
Industrial SSL ^a				11	11.1
Recreational SSL ^b				5.53	10.3
Residential SSL ^a				1.14	2.43
RE02-17-145128	02-61494	0–1	SOIL	0.0344	0.066
RE02-17-145138	02-61494	2–3	SOIL	_	0.00525
RELA-18-151015	02-61526	0–1	SOIL	_	0.531
RELA-18-151020	02-61526	2–3	SOIL	_	0.0232
RELA-18-151021	02-61526	4–5	SOIL	_	0.0439
RELA-18-151022	02-61526	6–7	QCT	—	0.00279 (J)
RELA-18-151017	02-61528	0–1	SOIL	—	0.0791
RELA-18-151023	02-61528	2–3	SOIL	—	0.0491
RELA-18-151024	02-61528	4–5	SOIL	—	0.00672
RELA-18-151018	02-61529	0–1	SOIL	—	3.77
RELA-18-151027	02-61529	2–3	SOIL	0.149	0.0754
RELA-18-151028	02-61529	4-4.65	SOIL	0.0401	0.0451
RELA-18-151019	02-61530	0–1	SOIL	—	1.68
RELA-18-151030	02-61530	2–3	SOIL	—	0.1
RELA-18-151031	02-61530	4–5	QBO	—	0.0251
RELA-18-151032	02-61530	6–7	QBO	—	0.0375
RELA-18-151033	02-61537	2–3	SOIL	—	23.9
RELA-18-151034	02-61537	4–5	SOIL	—	7.09
RELA-18-151035	02-61537	6–7	SOIL	—	0.614
RELA-18-151263	02-61537	8–9	QBO	—	1.44
RELA-18-151036	02-61537	11–12	QBO	—	2.48
RELA-18-161238	02-61538	0-0.9	SOIL	—	17.3
RELA-18-161242	02-61538	1–2	SOIL	—	1.1
RELA-18-161239	02-61539	1–2	SOIL	—	5.28
RELA-18-161244	02-61540	1–2	SOIL	—	12.3
RELA-18-164444	02-61541	6–7	QBT1G	—	0.963
RELA-18-161248	02-61542	1–2	SOIL	—	1.43
RELA-18-161249	02-61543	1–1.5	SOIL	—	6.77
RELA-18-164455	02-61544	2–3	SOIL	—	1.1
RELA-18-164465	02-61544	4.25–5	QBT1G	—	0.00375
RELA-18-164475	02-61544	6–7	QBT1G	—	0.413
RELA-18-164456	02-61545	2–3	SOIL	—	2.03
RELA-18-164466	02-61545	4–5	SOIL	—	0.446

Table 3.4-4 (continued)

Sample ID	Location ID	Depth (ft)	Media	Aroclor-1254	Aroclor-1260
Industrial SSL ^a				11	11.1
Recreational SSL ^b				5.53	10.3
Residential SSL ^a		1		1.14	2.43
RELA-18-164476	02-61545	6–7	SOIL	—	0.041
RELA-18-164448	02-61547	0–1	SOIL		0.921
RELA-18-164458	02-61547	2–3	SOIL	—	0.0299
RELA-18-164468	02-61547	4–5	SOIL	_	0.104
RELA-18-164478	02-61547	6–7	SOIL	—	0.00849
RELA-18-164459	02-61548	2–3	SOIL	—	0.684
RELA-18-164469	02-61548	4-4.5	SOIL	—	2.98
RELA-18-164479	02-61548	6–7	QBT1G	—	0.578
RELA-18-164470	02-61549	4–5	QBT1G	—	4.17
RELA-18-164480	02-61549	6–7	QBT1G	_	3.65
RELA-18-164451	02-61550	0–1	SOIL	—	5.35
RELA-18-164461	02-61550	2–3	SOIL	—	0.115
RELA-18-164471	02-61550	4–5	SOIL	_	0.157
RELA-18-164481	02-61550	6–7	SOIL	_	0.0342
RELA-18-164462	02-61551	2–3	SOIL	—	0.155
RELA-18-164472	02-61551	4–5	SOIL	_	0.169
RELA-18-164482	02-61551	6–7	SOIL	_	0.0631
RELA-18-164453	02-61552	0–1	SOIL	_	1.18
RELA-18-164463	02-61552	2–3	SOIL	_	0.031
RELA-18-164473	02-61552	4–5	SOIL	_	0.0205
RELA-18-164483	02-61552	6–7	QBT1G	_	0.0248
RELA-18-164464	02-61553	2–3	SOIL	_	0.834
RELA-18-164474	02-61553	4–5	SOIL	_	0.828
RELA-18-164484	02-61553	6–7	SOIL	_	1.25

Notes: Results are in mg/kg. Data qualifiers are defined in Appendix A.

^a SSLs are from NMED (2017, 602273), unless otherwise noted.

^b SSLs are from LANL (2017, 602581).

^c — = Not detected.

Table 3.4-5Radionuclides Detected or Detected above BV/FV at SWMU 02-014

Field Sample ID	Location ID	Depth (ft)	Media	Cesium-134	Cesium-137	Plutonium-238	Plutonium-239/240
Soil Background	Value/Fallout Value	a		na ^b	1.65	0.023	0.054
Qbt 1g, Qct, Qbo	Background Value/	Fallout Value ^a		na	na	na	na
Industrial SAL ^c				17	41	1300	1200
Recreational SAL	C			150	370	1400	1300
Residential SAL ^c	_			5	12	84	79
RE02-17-141612	02-61428	0–1	SOIL	d	—	—	0.076
RE02-17-141658	02-61428	2–3	SOIL	—	—	0.019	—
RE02-17-141659	02-61429	0–1	SOIL	—	—	—	0.07
RE02-17-141627	02-61437	2–3	SOIL	—	—	—	0.034
RE02-17-141649	02-61437	4–5	SOIL	—	—	_	0.005
RE02-17-141651	02-61439	2–3	SOIL	—	—	—	0.011
RE02-17-141656	02-61442	4–5	QBO	0.061	—	—	0.007
RE02-17-141695	02-61446	0–1	SOIL	—	—	—	0.071
RE02-17-141683	02-61448	4–5	SOIL	—	—	—	0.011
RE02-17-141676	02-61448	6–7	SOIL	—	—	—	0.018
RE02-17-141560	02-61450	4–5	SOIL	—	—	0.008	—
RE02-17-141561	02-61450	6–7	QBO	—	—	0.005	—
RE02-17-141568	02-61451	2–3	SOIL	—	0.759	—	0.008
RE02-17-141569	02-61451	4–5	SOIL	—	0.128	—	—
RE02-17-141579	02-61452	6–7	QBO	0.066	—	—	0.004
RE02-17-141588	02-61453	6–7	SOIL	—	—	_	0.012

Field Sample ID	Location ID	Depth (ft)	Media	Cesium-134	Cesium-137	Plutonium-238	Plutonium-239/240
Soil Background	Soil Background Value/Fallout Value ^a			na ^b	1.65	0.023	0.054
Qbt1g, Qct, Qbo E	Background Value/F	allout Value ^a		na	na	na	na
Industrial SAL ^c				17	41	1300	1200
Recreational SAL	c			150	370	1400	1300
Residential SAL ^c	Residential SAL ^c		5	12	84	79	
RE02-17-141596	02-61454	4–5	SOIL	0.056	—	_	—
RE02-17-141604	02-61455	2–3	SOIL	0.058	_	_	—
RE02-17-141605	02-61455	4–5	SOIL	0.054	—		—

Notes: Results are in pCi/g. Data qualifiers are defined in Appendix A.

^a BVs/FVs are from LANL (1998, 059730).

^b na = Not available.

^c SALs are from LANL (2015, 600929).

^d — = Not detected or not detected above BV/FV.

Appendix A

Acronyms and Abbreviations, Metric Conversion Table, and Data Qualifier Definitions

A-1.0 ACRONYMS AND ABBREVIATIONS

AK	acceptable knowledge
ALARA	as low as reasonably achievable
AOC	area of concern
ATSDR	Agency for Toxic Substances and Disease Registry
bgs	below ground surface
BV	background value
CCV	continuing calibration verification
COC	chain of custody
Consent Order	Compliance Order on Consent
COPC	chemical of potential concern
DAF	dilution attenuation factor
D&D	decontamination and decommissioning
DL	detection limit
DOE	Department of Energy (U.S.)
dpm	disintegrations per minute
Eh	oxidation-reduction potential
EPA	Environmental Protection Agency (U.S.)
EPC	exposure point concentration
EQL	estimated quantitation limit
ESL	ecological screening level
FV	fallout value
GPS	global-positioning system
HI	hazard index
HQ	hazard quotient
ICS	interference check sample
ICV	initial calibration verification
IR	investigation report
IS	internal standard
K _d	soil-water partition coefficient
K _{oc}	organic carbon-water partition coefficient
Kow	octanol-water partition coefficient
LANL	Los Alamos National Laboratory

LAL	lower acceptance limit
LCS	laboratory control sample
LiDAR	light detection and imaging
LLW	low-level waste
LOAEL	lowest observed adverse effect level
MDC	minimum detectable concentration
MDL	method detection limit
mmHg	millimeters of mercury
MS	matrix spike
N3B	Newport News Nuclear BWXT-Los Alamos, LLC
NDA	no detectable activity
NMED	New Mexico Environment Department
NOAEL	no observed adverse effect level
OWR	Omega West Reactor
PAH	polycyclic aromatic hydrocarbon
PCB	polychlorinated biphenyl
QA	quality assurance
QC	quality control
%R	percent recovery
%RSD	percent relative standard deviation
RCT	radiological control technician
RESRAD	Residual Radioactivity (computer code)
RfD	reference dose
RPD	relative percent difference
RTK	real-time kinematic
SAL	screening action level
SCL	sample collection log
SF	slope factor
SMO	Sample Management Office
SOP	standard operating procedure
SSL	soil screening level
SVOC	semivolatile organic compound
SWMU	solid waste management unit
T&E	threatened and endangered

ТА	technical area
TAL	target analyte list
TCDD[2,3,7,8]	2,3,7,8-tetrachlorodibenzo-p-dioxin
TEC	toxicity equivalency concentration
TEF	toxic equivalency factor
TSCA	Toxic Substances Control Act
UAL	upper acceptance limit
UCL	upper confidence limit
VOC	volatile organic compound
WBR	water boiler reactor
WCSF	waste characterization strategy form

A-2.0 METRIC CONVERSION TABLE

Multiply SI (Metric) Unit	by	To Obtain US Customary Unit		
kilometers (km)	0.622	miles (mi)		
kilometers (km)	3281	feet (ft)		
meters (m)	3.281	feet (ft)		
meters (m)	39.37	inches (in.)		
centimeters (cm)	0.03281	feet (ft)		
centimeters (cm)	0.394	inches (in.)		
millimeters (mm)	0.0394	inches (in.)		
micrometers or microns (µm)	0.0000394	inches (in.)		
square kilometers (km ²)	0.3861	square miles (mi ²)		
hectares (ha)	2.5	acres		
square meters (m ²)	10.764	square feet (ft ²)		
cubic meters (m ³)	35.31	cubic feet (ft ³)		
kilograms (kg)	2.2046	pounds (lb)		
grams (g)	0.0353	ounces (oz)		
grams per cubic centimeter (g/cm3)	62.422	pounds per cubic foot (lb/ft ³)		
milligrams per kilogram (mg/kg)	1	parts per million (ppm)		
micrograms per gram (µg/g)	1	parts per million (ppm)		
liters (L)	0.26	gallons (gal.)		
milligrams per liter (mg/L)	1	parts per million (ppm)		
degrees Celsius (°C)	9/5 + 32	degrees Fahrenheit (°F)		

A-3.0 DATA QUALIFIER DEFINITIONS

Data Qualifier	Definition
U	The analyte was analyzed for but not detected.
J	The analyte was positively identified, and the associated numerical value is estimated to be more uncertain than would normally be expected for that analysis.
J+	The analyte was positively identified, and the result is likely to be biased high.
J-	The analyte was positively identified, and the result is likely to be biased low.
UJ	The analyte was not positively identified in the sample, and the associated value is an estimate of the sample-specific detection or quantitation limit.
R	The data are rejected as a result of major problems with quality assurance/quality control parameters.

Appendix B

Field Methods

B-1.0 INTRODUCTION

This appendix summarizes field methods implemented during the 2018–2019 sampling and remediation activities at Solid Waste Management Unit (SWMU) 02-014 within the Middle Los Alamos Canyon Aggregate Area at Los Alamos National Laboratory (LANL or the Laboratory). Descriptions of field methods for previous investigations at SWMU 02-014 are included in the Phase II Investigation Report for Middle Los Alamos Canyon Aggregate, Revision 2 (N3B 2018, 700091). Table B-1.0-1 summarizes the field investigation methods, and the following sections provide more detailed descriptions of these methods.

B-2.0 FIELD-SCREENING METHODS

This section summarizes the field-screening methods used during the investigation activities. Field-screening results are presented in Table 2.2-2 of the addendum.

All soil/tuff samples were screened for radioactivity before submittal to the Sample Management Office (SMO). Samples submitted to the SMO were screened in the field by radiological control technicians (RCTs) using a ThermoFisher Model SHP-380, with Eberline Model E600 Geiger Counter, for detection of low-energy radiation. Gross alpha and beta/gamma screening measurements were recorded on the sample collection log/chain-of-custody (SCL/COC) forms and are reported in Table 2.2-2.

The outside of sealed sample containers, as well as sampling tools and equipment, were screened using smears and counted by the Ludlum Model 3030 Alpha Beta Sample Counter for detection of removable alpha and beta contamination.

B-3.0 FIELD-SCREENING INSTRUMENT CALIBRATION

The RCT calibrated field screening instruments daily before local background levels for radioactivity were measured. Instruments were calibrated using plutonium-239 and chloride-36 sources for alpha and beta emissions, respectively. All calibrations met the manufacturer's specifications and specifications in the applicable radiation detection instrument manual.

B-4.0 SURFACE AND SUBSURFACE SAMPLING

This section summarizes the methods used for collecting surface and subsurface samples.

B-4.1 Sample Collection

Surface and shallow subsurface samples were collected using a stainless-steel hand auger to collect material within the prescribed sampling intervals. For samples collected at depths greater than 3 ft, 4-in. polyvinyl chloride pipe was decontaminated and inserted into the hole to prevent hole collapse and cross-contamination of samples. A stainless-steel bowl and scoop were used to capture the sample from the hand-auger bucket. The sample was then transferred to sterile sample collection jars. Samples were appropriately labeled, sealed with custody seals, and documented before transporting to the SMO. Samples were transported to the SMO for processing and shipment to off-site contract analytical laboratories. The SMO personnel reviewed and approved the SCLs and accepted custody of the samples.

B-4.2 Subsurface Sampling Methods

Subsurface samples were collected in accordance with approved subcontractor procedures technically equivalent to standard operating procedure (SOP) ER-SOP-20069, "Soil, Tuff, and Sediment Sampling." Borehole samples were collected in a stainless-steel split-spoon core-barrel sampler that retrieved core in 2.5-ft intervals. The samples collected, listed by location and depth, are provided in tables for each site in the addendum to the Phase II investigation report.

Core retrieved from the subsurface was field screened for radioactivity and was visually inspected and logged. Following inspection, the 2.5-ft core section to be sampled was removed from the core barrel and placed in a stainless-steel bowl and homogenized. The material was crushed, if necessary, with a decontaminated rock hammer and stainless-steel spoon to allow core material to fit into sample containers.

No samples were collected for volatile organic compound (VOC) analysis. The sample collection tools were decontaminated immediately before each sample was collected in accordance with an approved subcontractor procedure technically equivalent to ER-SOP-5061, "Field Decontamination of Equipment" (section B-5.4).

B-4.3 Quality Control Samples

Quality assurance/quality control samples were collected in accordance with an approved subcontractor procedure technically equivalent to ER-SOP-20235, R0, "Sample Containers, Preservation, and Field Quality Control." The quality control samples included field duplicates and field rinsate blanks. Field duplicate samples were collected from the same material as a regular investigation sample and submitted for the same analyses. Field duplicate samples were collected at a frequency of at least 1 duplicate sample for every 10 samples submitted to the SMO. A total of 6 field duplicate samples were collected.

Field rinsate blanks were collected to evaluate field decontamination procedures. Rinsate blanks were collected by rinsing sampling equipment (i.e., auger buckets, sampling bowls, and scoops) with deionized water after decontamination. The rinsate water was collected in a sample container and submitted to the SMO. A total of four field rinsate samples were collected.

B-4.4 Decontamination of Sampling Equipment

The split-spoon core barrels and all other sampling equipment that came (or could have come) in contact with sample material were decontaminated after each core was retrieved and logged. Decontamination included wiping the equipment with Fantastik and paper towels. The drilling equipment was decontaminated before mobilization of the drill rig to another borehole to avoid cross-contamination between samples and borehole locations. Residual material adhering to equipment was removed using dry decontamination methods such as the use of wire brushes and scrapers. Decontamination activities were performed in accordance with an approved subcontractor procedure technically equivalent to ER-SOP-5061, "Field Decontamination of Equipment." Field rinsate blank samples were collected in accordance with an approved procedure technically equivalent to ER-SOP-5059, "Field Quality Control Samples."

B-5.0 SOIL REMEDIATION

B-5.1 Radiological Controls

Heavy equipment entering and leaving the site was monitored for radioactivity using smear samples. Heavy equipment screening smears were taken by the RCT and counted by a Ludlum Model 3030 Alpha Beta Sample Counter for detection of removable alpha and beta contamination. A ThermoFisher Model SHP-380, with Eberline Model E600 Geiger Counter, was used for direct screening of alpha and beta contamination of personnel after completing daily work, as requested. Any tools and personnel entering an excavation were screened out by an RCT.

B-5.2 Soil Excavation

Between November 7, 2018, and December 17, 2018, approximately 282 yd³ of polychlorinated biphenyl– (PCB-) contaminated soil was removed from the excavation sites at SWMU 02-014. Soil was removed from the entire excavation area to a depth of 1 ft and deeper excavation was performed at five interior areas to depths of 4.5 ft to 10.5 ft. Excavation areas are shown on Plate 1.

The planned extent and depths of excavations were initially identified based on the results of previous sampling for PCBs. Following collection and analysis of confirmation samples, several areas were identified where excavation had to be extended vertically within the interior of the original area (interior areas) or laterally beyond the original excavation area. Excavation was determined to be complete upon confirmation of excavation depth and extent using several methods. Both the Topcon HiPer V real-time kinematic (RTK) global positioning system (GPS) unit and visual measurements were used to confirm the 1-ft excavation depth and extent. RTK GPS and visual tape measurements were used to confirm depth extent of most of the deep, interior excavations. Elevation was recorded at the original surface and at the bottom of the excavation depth was reached. Points along the edges were often slightly higher than the central depth because of sloping during the excavation process and some sloughing of surface material. For the deepest interior excavation, a laser tape measure was affixed to a 10 ft × 2 in. × 4 in. wood beam to safely check the depth of excavation. In addition, 10-ft, 10.5-ft, and 11-ft lengths were marked with tape on the excavator arm for a second visual check.

Both a Yanmar mini-excavator and Volvo excavator were used for excavation as well as the placement and tamping of backfill and base course materials. Excavated material was placed in 5.18-yd³ soft-sided IP-1 bags with 6-mil polypropylene built-in inner liners. After an IP-1 bag was filled, the inner liners of the bags were sealed per manufacturer's instructions and discussions with the waste management coordinator and Field Execution Team lead. Once sealed, IP-1 bags were secured to a telehandler and moved to Technical Area 41 (TA-41) where they were staged.

During excavation of deep interior area I-1, two asbestos-wrapped pipes (assumed to be decommissioned gas and water lines based on conversations with LANL Utilities Management) were exposed within the first 1–2 feet below ground surface (bgs). A "pause work" commenced the afternoon of November 19, 2018, and went to December 3, 2018. During this time, an asbestos abatement plan and integrated work document addendum were developed. The approved asbestos abatement plan was then implemented and the exposed pipes and wrapping were removed from the excavation and properly packaged and labeled. Abatement took place from December 4, 2018, through December 6, 2018. About 1 yd³ of waste (pipe, asbestos wrapping, contact waste materials) is estimated to have been generated in the process.

Restoration at SWMU 02-014 began on December 18, 2018, and was completed on January 28, 2019. Fill was first placed into the deep excavation areas. Material was spread and compressed by the Volvo excavator in 6–8-in. lifts to 1 ft below grade (same level as the 1-ft deep excavation) and tamped after each lift using the approximately 1-yd³ bucket. Visual and tape measure methods were used to confirm that fill was deposited to 4–6 in. below grade. Base course was then placed on top of the fill. Visual observation and the Topcon RTK GPS system were used to confirm that base course was deposited as close to original grade as possible.

B-6.0 GEODETIC SURVEYING

RTK GPS surveying was conducted using Topcon HiPer V Navigation Satellite System antennas coupled with a Topcon FC-5000 Data Collector Controller. This system was used to stake sampling locations, locations to be left unexcavated (i.e., uncontaminated locations), locations excavated, planned excavation boundaries, and pre- and post-excavation topographic elevations. Light detection and ranging (LiDAR) was also used for pre- and post-excavation topographic elevation data collection, specifically for excavations too deep to safely check via RTK GPS.

If a planned sampling location needed to be offset more than 1 ft because of surface or subsurface obstructions, the relocated point was re-surveyed. Surveyed coordinates for sampling locations of samples submitted to the SMO are presented in Table 2.2-1 of the addendum.

B-7.0 WASTE STORAGE AND DISPOSAL

All excavated media at SWMU 02-014 was placed in 5.18-yd³ soft-sided IP-1 bags. IP-1 bags were positioned on pallets for loading and staging. After the bag was sealed, an RCT screened all sides of the bag before releasing the bag for staging at TA-41. Temporary storage was within a posted radiological waste storage area at TA-41. All waste containers that were staged at TA-41 were covered with tarps for additional protection from the elements.

All waste was managed in accordance with the project waste characterization strategy form (WCSF) and an approved subcontractor procedure technically equivalent to ER-DIR-SOP-10021, R1, "Characterization and Management of Environmental Programs Waste." Contact waste was stored in labeled 1-gal. plastic bags in the radioactive waste accumulation area on-site until it could be transferred to a 5.18-yd³ soft-sided IP-1 bag.

B-8.0 DEVIATIONS FROM THE WORK PLAN

The 2018–2019 sampling and remediation activities at SWMU 02-014 were not in the approved Phase II investigation work plan for Middle Los Alamos Canyon Aggregate Area (LANL 2009, 105073; NMED 2009, 105595). The need for these activities was identified as a result of the Phase II sampling performed at Area of Concern 02-011(a) and the subsequent identification of a new SWMU.

B-9.0 REFERENCES

The following reference list includes all documents cited in this report. Parenthetical information following each reference provides the author(s), publication date, and ERID, ESHID, or EMID. This information is also included in text citations. ERIDs were assigned by the Laboratory's Associate Directorate for Environmental Management (IDs through 599999); ESHIDs were assigned by the Laboratory's Associate Directorate for Environment, Safety, and Health (IDs 600000 through 699999); and EMIDs are assigned by Newport News Nuclear BWXT-Los Alamos, LLC (N3B) (IDs 700000 and above). IDs are used to locate documents in N3B's Records Management System and in the Master Reference Set. The NMED Hazardous Waste Bureau and N3B maintain copies of the Master Reference Set. The set ensures that NMED has the references to review documents. The set is updated when new references are cited in documents.

- LANL (Los Alamos National Laboratory), February 2009. "Phase II Investigation Work Plan for Middle Los Alamos Canyon Aggregate Area, Revision 1," Los Alamos National Laboratory document LA-UR-09-1206, Los Alamos, New Mexico. (LANL 2009, 105073)
- N3B (Newport News Nuclear BWXT-Los Alamos, LLC), September 2018. "Phase II Investigation Report for Middle Los Alamos Canyon Aggregate Area, Revision 2," Newport News Nuclear BWXT-Los Alamos, LLC, document EM2018-0039, Los Alamos, New Mexico. (N3B 2018, 700091)
- NMED (New Mexico Environment Department), March 25, 2009. "Approval, Middle Los Alamos Canyon Aggregate Area Phase II Work Plan, Revision 1," New Mexico Environment Department letter to D. Gregory (DOE LASO) and D. McInroy (LANL) from J.P. Bearzi (NMED-HWB), Santa Fe, New Mexico. (NMED 2009, 105595)

Method	Summary
Hand-Auger Sampling	This method is typically used for sampling soil or sediment at depths of less than 10–15 ft, but in some cases may be used to collect samples of weathered or nonwelded tuff. The method involves hand-turning a stainless-steel bucket auger (typically 3–4 in. inside diameter), creating a vertical hole that can be advanced to the desired sampling depth. When the desired depth was reached, the auger was decontaminated before advancing the hole through the sampling depth. The sample material was transferred from the auger bucket to a stainless-steel sampling bowl before the various required sample containers were filled.
Split-Spoon Core- Barrel Sampling	A stainless-steel core barrel was advanced using a hollow-stem auger drilling rig. The core barrel extracted a continuous length of soil and/or rock. The split-spoon core barrel is a cylindrical barrel split lengthwise so the two halves can be separated to expose the core sample. If necessary, pieces small enough to fit into the sample container were removed from the core using a decontaminated rock hammer or stainless-steel spoon. The section of core in the core barrel was then screened for radioactivity. A portion of the core was then collected as a discrete sample from the desired depth for remaining analyses.
Handling, Packaging, and Shipping of Samples	Field team members sealed and labeled samples before packing them to ensure the sample containers and the containers used for transport were free of external contamination. Field team members packaged all samples to minimize the possibility of breakage during transport. After all environmental samples were collected, packaged, and preserved, a field team member transported them to the SMO. The SMO arranged to ship the samples to the analytical laboratories.
Sample Control and Field Documentation	The collection, screening, and transport of samples were documented on standard forms generated by the SMO. These included SCLs and sample container labels. SCLs were completed at the time of sample collection, and the logs were signed by the sampler and a reviewer who verified the logs for completeness and accuracy. Corresponding labels were initialed and applied to each sample container, and custody seals were placed around each sample container. SCLs were completed and signed to verify that the samples were not left unattended.
Field Quality Control Samples	 Field quality control samples were collected as follows: <i>Field Duplicates</i>: At a frequency 10%; collected at the same time as a regular sample and submitted for the same analyses. <i>Equipment Rinsate Blank</i>: At a frequency of 10%; collected by rinsing decontaminated sampling equipment with deionized water, which was collected in a sample container and submitted for laboratory analysis.
Field Decontamination of Drilling and Sampling Equipment	Dry decontamination was used to minimize the generation of liquid waste. Dry decontamination included the use of a wire brush or other tool to remove soil or other material adhering to the sampling equipment, followed by use of a commercial cleaning agent (nonacid, waxless cleaners) and paper wipes.
Containers and Preservation of Samples	Specific requirements/processes for sample containers, preservation techniques, and holding times are based on U.S. Environmental Protection Agency guidance for environmental sampling, preservation, and quality assurance. Specific requirements for each sample were printed on the SCL provided by the SMO (size and type of container [e.g., glass, amber glass, and polyethylene]). All samples were preserved by placing them with ice in insulated containers to maintain a temperature of 4°C.

Table B-1.0-1Summary of Field Investigation Methods

Method	Summary
Coordinating and Evaluating Geodetic Surveys	Geodetic surveys focused on obtaining survey data of acceptable quality to use during project investigations. Geodetic surveys were conducted with Topcon HiPer V Navigation Satellite System Antennas coupled with a Topcon FC-5000 Data Collection Controller. All coordinates were expressed as State Plane Coordinate System 83, NM Central, U.S. feet. All elevation data were reported relative to the National Geodetic Vertical Datum of 1983.
Management of Environmental Restoration Project Waste, Waste Characterization	Waste was managed, characterized, and stored in accordance with an approved WCSF that documented the site history, field activities, and characterization approach for each waste stream managed. Waste characterization complied with on- or off-site waste acceptance criteria. All stored waste was marked with appropriate signage and labels. Drummed waste was stored on pallets to prevent deterioration of containers. A waste storage area was established before waste was generated. Waste storage areas were located in controlled areas of the Laboratory to prevent unauthorized personnel from inadvertently adding or managing wastes. Each container of waste generated was individually labeled with waste classification, item identification number, and radioactivity category (if applicable), immediately following containerization. All waste was segregated by classification and compatibility to prevent cross-contamination. Management of waste is described in Appendix C.

Table B-1.0-1 (continued)

Appendix C

Waste Management

C-1.0 INTRODUCTION

This appendix describes management of the waste generated during the 2018–2019 investigation and remediation activities at solid waste management unit (SWMU) 02-014 within the Middle Los Alamos Canyon Aggregate Area of Los Alamos National Laboratory (the Laboratory) and during sampling activities conducted in 2017 to characterize the area requiring remediation. In general, waste generated during the investigation and remediation activities was managed in accordance with approved subcontractor procedures technically equivalent to standard operating procedure (SOP) EP-DIR-SOP-10021, R1, "Characterization and Management of Environmental Program Waste." This procedure incorporates the requirements of applicable U.S. Environmental Protection Agency and New Mexico Environment Department regulations, and U.S. Department of Energy orders.

Waste characterization strategy forms (WCSFs) were prepared to address characterization approaches, on-site management, and final disposition options for wastes. Analytical data and information on wastes generated during previous investigations and/or acceptable knowledge (AK) were used to complete the WCSF. WCSF EP2016-0128 was prepared in October 2016 to address Middle Los Alamos Canyon Aggregate Area site characterization activities. WCSF EM2018-0021 was prepared in July 2018 to address SWMU 02-014 remediation activities. The WCSFs are provided in Attachment C-1 (on CD).

The selection of waste containers was based on appropriate U.S. Department of Transportation requirements, waste types, and estimated volumes of wastes to be generated. Immediately following containerization, each waste container was individually labeled with a unique identification number and with information regarding waste classification, contents, and radioactivity, if applicable.

Wastes were staged in clearly marked, appropriately constructed waste accumulation areas. Waste accumulation area postings, regulated storage duration, and inspection requirements were based on the type of waste and its classification. Container and storage requirements were detailed in the WCSFs and approved before waste was generated.

To the extent possible, investigation and remediation activities were conducted in a manner that minimized the generation of waste.

C-2.0 WASTE STREAMS

The waste streams generated and managed during the investigation are described below and are summarized in Table C-2.0-1. The waste stream numbers correspond with those identified in the WCSFs. Waste streams 1, 2, 5, and 6 are the same for the two WCSFs.

- WCSF Waste Stream #1: Contact Waste—This waste stream is composed of solid waste generated during investigation and removal activities that has come into contact with contaminated environmental media and equipment. This includes, but is not limited to, personal protective equipment (e.g., gloves), plastic sheeting (e.g., tarps, liners), plastic and glass sample bottles, disposable sampling supplies (e.g., filters, tubing, plastic bags), and dry decontamination wastes (e.g., paper items). Less than 1 yd³ of contact waste was generated and was combined with excavated media.
- WCSF Waste Streams #2, #5, and #6—No municipal solid waste (waste stream #2), petroleumcontaminated soils (waste stream #5), or decontamination fluids (waste stream #6) were generated.

- WCSF EP2016-0128 Waste Stream #3: Drill cuttings—This waste stream consisted of sediment, soil, and rock removed during mechanical auger drilling. Approximately 4 yd³ of drill cuttings was generated during this investigation and stored in 55-gal. drums.
- WCSF EP2016-0128 Waste Stream #4/WCSF EM2018-0021 Waste Stream # 3: Environmental Media—This waste stream consists of contaminated soil, sediment, and tuff excavated to remove media that exceeds polychlorinated biphenyl cleanup levels. Approximately 282 yd³ of this waste was generated. This waste stream also includes media from surface and subsurface sampling.
- WCSF EM2018-0021 Waste Stream #4: Excavated Debris—This waste stream consists of residual manmade debris encountered during soil excavation. Debris encountered during the soil excavation includes asbestos-wrapped pipe. Approximately 1 yd³ of this waste was generated.

 Table C-2.0-1

 Summary of Waste Generation and Management

WCSF EP2016-0128 Waste Stream #	WCSF EM2018-0021 Waste Stream #	Waste Stream	Waste Type	Volume	Characterization Method	On-Site Management	Disposition
1	1	Contact waste	LLW ^a TSCA ^b	<1 yd3	AK and analytical results of site characterization	Plastic bags, 5.18 yd ³ IP-1 hard-side bags	Energy Solutions, Clive UT
2	2	Municipal solid waste	n/a ^c	0	n/a	n/a	n/a
3	n/a	Drill cuttings	Industrial	4 yd ³	Direct sampling	55-gal. drum	Waste Control Specialists, Andrews TX
4	3	Environmental media	LLW TSCA	282 yd ³	AK and analytical results of site characterization	5.18 yd ³ IP-1 hard-side bags	Energy Solutions, Clive UT
n/a	4	Excavated debris	LLW TSCA	1 yd ³	AK and analytical results of site characterization	5.18 yd ³ IP-1 hard-side bags	Energy Solutions, Clive UT
5	5	Petroleum- contaminated soils	n/a	0	n/a	n/a	n/a
6	6	Decontamination fluids	n/a	0	n/a	n/a	n/a

^a LLW = Low-level waste.

^b TSCA = Toxic Substances Control Act.

^c n/a = Not applicable.

Attachment C-1

Waste Characterization Strategy Forms (on CD included with this document)

Appendix D

Analytical Program

D-1.0 INTRODUCTION

This appendix discusses the analytical methods and data-quality review for samples collected during investigations at Solid Waste Management Unit (SWMU) 02-014 within the Middle Los Alamos Canyon Aggregate Area at Los Alamos National Laboratory (LANL or the Laboratory). All decision-level data presented in the investigation report addendum are evaluated, including data from samples collected in 2007, 2010, 2017, and 2018. Additionally, this appendix gives a summary of the effects of data-quality issues on the acceptability of the analytical data.

The results of the quality assurance/quality control (QA/QC) procedures were used to estimate the accuracy, bias, and precision of the analytical measurements. Samples for QC include method blanks, matrix spikes (MSs), laboratory control samples (LCSs), internal standards, initial calibration verifications (ICVs) and continuing calibration verifications (CCVs), surrogates, and tracers.

The type and frequency of laboratory QC analyses are described in the statements of work for analytical laboratories. Other QC factors, such as sample preservation and holding times, were also assessed in accordance with the requirements outlined in Standard Operating Procedure (SOP) 5056, "Sample Containers and Preservation."

The following SOPs were used for data validation:

- SOP-5161, "Routine Validation of Volatile Organic Compound (VOC) Analytical Data"
- SOP-5162, "Routine Validation of Semivolatile Organic Compound (SVOC) Analytical Data"
- SOP-5163, "Routine Validation of Organochlorine Pesticide (PEST) and Polychlorinated Biphenyl (PCB) Analytical Data"
- SOP-5165, "Routine Validation of Metals Analytical Data"
- SOP-5166, "Routine Validation of Gamma Spectroscopy, Chemical Separation Alpha Spectrometry, Gas Proportional Counting, and Liquid Scintillation Analytical Data"
- SOP-5169, "Routine Validation of Dioxin Furan Analytical Data (EPA Method 1618 and SW-846 EPA Method 8290)"
- SOP-5191, "Routine Validation of LC/MS/MS Perchlorate Analytical Data" (SW-846 EPA Method 6850)

Routine data validation was performed for each data package (referred to by a request number), and analytical data were reviewed and evaluated based on U.S. Environmental Protection Agency (EPA) National Functional Guidelines, where applicable (EPA 1994, 048639; EPA 1999, 066649). As a result of the data validation and assessment efforts, qualifiers are assigned to the analytical records as appropriate. The data qualifier definitions are provided in Appendix A. Sample collection logs (SCLs) and chain of custody forms (COCs) are provided in Appendix E (on CD included with this document). The analytical data, instrument printouts, and data validation reports are provided in Appendix E.

D-2.0 ANALYTICAL DATA ORGANIZATION

Historical data evaluated in this report were collected during Resource Conservation and Recovery Act facility investigations, other corrective actions, and other investigations. All historical investigation samples were submitted to and analyzed by approved off-site laboratories. These data are determined to be of sufficient quality for decision-making purposes and have been reviewed and revalidated to current QA standards.

D-3.0 INORGANIC CHEMICAL ANALYSES

A total of 92 samples (plus 8 field duplicates) collected at SWMU 02-014 were analyzed for inorganic chemicals. Ninety-two samples (plus 8 field duplicates) were analyzed for target analyte list (TAL) metals; three of these samples were also analyzed for nitrate, perchlorate, and total cyanide. The analytical methods used for inorganic chemicals are listed in Table D-1.0-1.

Tables in the addendum to the investigation report summarize all samples collected and the analyses requested for the investigation of SWMU 02-014. All analyses conducted during the investigation are presented in Appendix E (on CD included with this document).

D-3.1 Inorganic Chemical QA/QC Samples

QA/QC samples are used to produce measures of the reliability of the data. The results of the QA/QC analyses performed on a sample provide confidence about whether the analyte is present and whether the concentration reported is accurate. To assess the accuracy and precision of inorganic chemical analyses, this investigation included analyses of LCSs, preparation blanks, MSs, laboratory duplicate samples, interference check samples (ICSs), and serial dilution samples. Each of these QA/QC sample types is described briefly in the paragraphs below.

The LCS serves as a monitor of the overall performance of each step during the analysis, including sample digestion. For inorganic chemicals in soil or tuff, LCS percent recoveries (%R) should fall within the control limits of 75% to 125% (LANL 1995, 049738; LANL 2000, 071233; LANL 2008, 109962).

The preparation blank is an analyte-free matrix to which all reagents are added in the same volumes or proportions as those used in the environmental sample processing; it is extracted and analyzed in the same manner as the corresponding environmental samples. Preparation blanks are used to measure bias and potential cross-contamination. All inorganic chemical results should be below the method detection limit (MDL).

MS samples assess the accuracy of inorganic chemical analyses. These samples are designed to provide information about the effect of the sample matrix on the sample preparation procedures and analytical technique. The MS acceptance criterion is 75% to 125% recovery, inclusive, for all spiked analytes (LANL 1995, 049738; LANL 2000, 071233; LANL 2008, 109962).

Laboratory duplicate samples assess the precision of inorganic chemical analyses. All relative percent differences (RPDs) between the sample and laboratory duplicate should be ±35% for soil (LANL 1995, 049738; LANL 2000, 071233; LANL 2008, 109962).

The ICSs assess the accuracy of the analytical laboratory's interelement and background correction factors used for inductively coupled plasma emission spectroscopy. The ICS %R should be within the acceptance range of 80% to 120%. The QC acceptance limits are ±20%.

Serial dilution samples measure potential physical or chemical interferences and correspond to a sample dilution ratio of 1:5. The chemical concentration in the undiluted sample must be at least 50 times the MDL (100 times for inductively coupled plasma mass spectroscopy) for valid comparison. For sufficiently high concentrations, the RPD should be within 10%.

D-3.2 Data-Quality Results for Inorganic Chemicals

The majority of the analytical results for inorganic chemicals were either not assigned a qualifier or qualified as not detected (U) because the analytes were not detected by the respective analytical methods. These data do not have any quality issues associated with the values presented.

D-3.2.1 Maintenance of COC

SCL/COC forms were maintained properly for all samples analyzed for inorganic chemicals (see Appendix E, on CD included with this document).

D-3.2.2 Sample Documentation

All samples analyzed for inorganic chemicals were properly documented on SCL/COC forms in the field (see Appendix E, on CD included with this document).

D-3.2.3 Sample Dilutions

Some samples were diluted for inorganic chemical analyses. No qualifiers were applied to any inorganic chemical sample results because of dilutions.

D-3.2.4 Sample Preservation

Preservation criteria were met for all samples analyzed for inorganic chemicals.

D-3.2.5 Holding Times

Holding-time criteria were met for all samples analyzed for inorganic chemicals.

D-3.2.6 ICVs and CCVs

ICV and CCV criteria were met for all samples analyzed for inorganic chemicals.

D-3.2.7 Interference Check Sample and/or Serial Dilutions

One vanadium result was qualified as estimated (J) because the serial dilution sample RPD exceeded criteria.

D-3.2.8 Laboratory Duplicate Samples

One barium result was qualified as estimated (J) because the duplicate result exceeded the RPD requirements.

A total of 20 TAL metals results were qualified as estimated (J) because a duplicate sample was not prepared and/or analyzed with the samples for unspecified reasons.

D-3.2.9 Blanks

A total of 193 TAL metals results were qualified as not detected (U) because the sample result was less than 5 times the concentration of the related analyte in the method blank.

D-3.2.10 MS Samples

Six TAL metals results were qualified as estimated and biased high (J+) because the spike percent recovery value is less than 30%.

One calcium result and two nitrate results were qualified as estimated and biased low (J-) because the MS %R value was less than the lower acceptance limit (LAL) but greater than 30%.

Three antimony results were qualified as estimated not detected (UJ) because the MS %R value was less than the LAL but greater than 30%.

One aluminum result was qualified as estimated and biased high (J+) because of MS recovery problems.

D-3.2.11 LCS Recoveries

No inorganic chemical results were qualified because of LCS recovery.

D-3.2.12 Detection Limits

A total of 388 TAL metals results, 1 nitrate result, and 1 perchlorate result were qualified as estimated (J) because the sample result was reported as detected between the instrument detection limit and the estimated detection limit.

D-3.2.13 Rejected Results

No inorganic chemical results were qualified as rejected (R).

D-4.0 ORGANIC CHEMICAL ANALYSES

A total of 353 samples (plus 21 field duplicates) collected at SWMU 02-014 were analyzed for organic chemicals. A total of 353 samples (plus 21 field duplicates) were analyzed for polychlorinated biphenyls (PCBs); 2 of these samples were also analyzed for volatile organic compounds (VOCs); and 3 samples were also analyzed for semivolatile organic compounds (SVOCs) and for dioxins and furans. The analytical methods used for organic chemicals are listed in Table D-1.0-1.

Tables within the addendum to the investigation report summarize all samples collected at SWMU 02-014 and the analyses requested. All organic chemical results are provided on CD in Appendix E.

D-4.1 Organic Chemical QA/QC Samples

QA/QC samples are used to produce measures of the reliability of the data. The results of the QA/QC analyses performed on a sample provide confidence about whether the analyte is present and whether the concentration reported is accurate. To assess the accuracy and precision of organic chemical analyses, this investigation included calibration verifications and the analysis of LCSs, method blanks, MSs, surrogates, and internal standards (ISs). Each of these QA/QC sample types is described briefly in the paragraphs below.

Calibration verification is the establishment of a quantitative relationship between the response of the analytical procedure and the concentration of the target analyte. There are two aspects of calibration verification: initial and continuing. The initial calibration verifies the accuracy of the calibration curve as well as the individual calibration standards used to perform the calibration. The continuing calibration ensures that the initial calibration is still holding and correct as the instrument is used to process samples. The continuing calibration also serves to determine that analyte identification criteria such as retention times and spectral matching are being met.

The LCS is a sample of a known matrix that has been spiked with compounds that are representative of the target analytes, and it serves as a monitor of overall performance on a "controlled" sample. The LCS is the primary demonstration, on a daily basis, of the ability to analyze samples with good qualitative and quantitative accuracy. The LCS recoveries should within the method-specific acceptance criteria.

A method blank is an analyte-free matrix to which all reagents are added in the same volumes or proportions as those used in the environmental sample processing; it is extracted and analyzed in the same manner as the corresponding environmental samples. Method blanks are used to assess the potential for sample contamination during extraction and analysis. All target analytes should be below the contract-required detection limit in the method blank.

MS samples are used to measure the ability to recover prescribed analytes from a native sample matrix and consist of aliquots of the submitted samples spiked with a known concentration of the target analyte(s). Spiking typically occurs before sample preparation and analysis. The spike sample recoveries should be between the LAL and the upper acceptance limit (UAL).

A surrogate compound (surrogate) is an organic compound used in the analyses of target analytes that is similar in composition and behavior to the target analytes but not normally found in environmental samples. Surrogates are added to every blank, sample, and spike to evaluate the efficiency with which analytes are recovered during extraction and analysis. The recovery percentage of the surrogates must be within specified ranges or the sample may be rejected or assigned a qualifier.

ISs are chemical compounds added to every blank, sample, and standard extract at a known concentration. They are used to compensate for (1) analyte concentration changes that might occur during storage of the extract, and (2) quantitation variations that can occur during analysis. ISs are used as the basis for quantitation of target analytes. The %R for ISs should be within the range of 50% to 200%.

D-4.2 Data-Quality Results for Organic Chemicals

The majority of the analytical results for organic chemicals were either not assigned a qualifier or qualified as not detected (U) because the analytes were not detected by the respective analytical methods. These data do not have any quality issues associated with the values presented.

One dioxin/furan result was qualified as estimated (J) because the analytical laboratory qualified the detected result as estimated.

D-4.2.1 Maintenance of COC

SCL/COC forms were maintained properly for all samples analyzed for organic chemicals (see Appendix E, on CD included with this document).

D-4.2.2 Sample Documentation

All samples analyzed for organic chemicals were properly documented on the SCL in the field (see Appendix E, on CD included with this document).

D-4.2.3 Sample Dilutions

Some samples were diluted for organic chemical analyses. No qualifiers were applied to any organic chemical sample results because of dilutions.

D-4.2.4 Sample Preservation

Preservation criteria were met for all samples analyzed for organic chemicals.

D-4.2.5 Holding Times

Holding-time criteria were met for all samples analyzed for organic chemicals.

D-4.2.6 ICVs and CCVs

Ten SVOC and eight VOC results were qualified as estimated not detected (UJ) because the initial calibration curve exceeded the percent relative standard deviation (%RSD) criteria and/or the associated multipoint calibration correlation coefficient was less than 0.995.

D-4.2.7 Surrogate Recoveries

No organic chemical results were qualified because of surrogate recovery.

D-4.2.8 IS Responses

No organic chemical results were qualified because of IS response.

D-4.2.9 Blanks

Three PCB results and two dioxin/furan results were qualified as estimated (J) because the sample concentration was greater than 5 times the amount in the method blank.

D-4.2.10 MS Samples

No organic chemical results were qualified because of MS analyses.

D-4.2.11 Laboratory Duplicate Samples

Laboratory duplicates collected for organic chemical analyses indicated acceptable precision for all samples.

D-4.2.12 LCS Recoveries

Two PCB results were qualified as estimated and biased high (J+) because the LCS recovery was greater than the UAL.

Three SVOC results were qualified as estimated not detected (UJ) because the LCS documentation is missing.

D-4.2.13 Detection Limits

A total of 11 SVOC results, 1 VOC result, 30 PCB results, and 19 dioxin/furan results were qualified as estimated (J) because the sample result was reported as detected between the instrument detection limit and the estimated detection limit.

D-4.2.14 Rejected Data

No organic chemical results were qualified as rejected (R).

D-5.0 RADIONUCLIDE ANALYSES

A total of 92 samples (plus 7 field duplicates) collected at SWMU 02-014 were analyzed for radionuclides. A total of 92 samples (plus 7 field duplicates) were analyzed for gamma-emitting radionuclides, isotopic plutonium, isotopic uranium, and strontium-90. Seven of these samples were also analyzed for americium-241 and three samples were also analyzed for tritium. The analytical methods used for radionuclides are listed in Table D-1.0-1.

Tables in the addendum to the investigation report summarize all samples collected at SWMU 02-014 and the analyses requested. All radionuclide results are provided on CD (Appendix E).

D-5.1 Radionuclide QA/QC Samples

To assess the accuracy and precision of radionuclide analyses, this investigation included analyses of LCSs, method blanks, MS samples, laboratory duplicate samples, and tracers. Each of these QA/QC sample types is described briefly in the paragraphs below.

The LCS serves as a monitor of the overall performance of each step during the analysis, including sample digestion. For radionuclides in soil or tuff, LCS %R should fall between the control limits of 80% and 120%.

A method blank is an analyte-free matrix to which all reagents are added in the same volumes or proportions as those used in the environmental sample processing; it is analyzed in the same manner as the corresponding environmental samples. Method blanks are used to assess the potential for sample contamination during analysis. All radionuclide results should be below the minimum detectable concentration (MDC).

MS samples assess the accuracy of radionuclide analyses. These samples are designed to provide information about the effect of the sample matrix on the sample preparation procedures and analytical technique. The MS acceptance criterion is 75% to 125% recovery.

Tracers are radioisotopes added to a sample for the purposes of monitoring losses of the target analytes. The tracer is assumed to behave in the same manner as the target analyte. The tracer recoveries should fall between the LAL and UAL.

Laboratory duplicate samples assess the precision of radionuclide analyses. All RPDs between the sample and laboratory duplicate should be $\pm 35\%$ for soil (LANL 1995, 049738; LANL 2000, 071233; LANL 2008, 109962).

D-5.2 Data-Quality Results for Radionuclides

Approximately one-fifth (202) of the analytical results for radionuclides were not assigned a qualifier. These data do not have any quality issues associated with the values presented.

The majority of results (623) were qualified as not detected (U) because the associated sample activity was less than or equal to the minimum detectable activity.

D-5.2.1 Maintenance of COC

SCL/COC forms were maintained properly for all samples (see Appendix E, on CD included with this document).

D-5.2.2 Sample Documentation

All samples were properly documented on the SCL/COC forms in the field (see Appendix E, on CD included with this document).

D-5.2.3 Sample Dilutions

Some samples were diluted for radionuclide analyses. No qualifiers were applied to any radionuclide sample results because of dilutions.

D-5.2.4 Sample Preservation

Preservation criteria were met for all samples analyzed for radionuclides.

D-5.2.5 Holding Times

Holding-time criteria were met for all samples analyzed for radionuclides.

D-5.2.6 Method Blanks

No radionuclide results were qualified because of method blank analyses.

A total of 14 isotopic uranium results were qualified as not detected (U) because the sample results were equal to or less than 5 times the activity of the related analyte in the method blank.

A total of 14 isotopic uranium results were qualified as estimated (J) and 66 isotopic uranium results were qualified as estimated and biased high (J+) because the sample activities were greater than 5 times the amount in the method blank.

D-5.2.7 MS Samples

No radionuclide results were qualified because of MS recovery.

D-5.2.8 Tracer Recoveries

Two isotopic uranium results were qualified as estimated and biased low (J-) because the tracer receiver was less than the LAL but equal to or greater than 10%.

Three isotopic uranium results were qualified as estimated and biased high (J+) because the tracer recovery was greater than the UAL.

D-5.2.9 LCS Recoveries

No radionuclide results were qualified because of LCS recovery.

One strontium-90 result was qualified as estimated and biased high (J+) because the LCS recovery was greater than the UAL.

D-5.2.10 Laboratory Duplicate Sample Recoveries

No radionuclide results were qualified because of laboratory duplicate sample recovery.

D-5.2.11 Rejected Data

Two cesium-134 results and one cesium-137 result were qualified as rejected (R) because the minimum detectable activity and/or total propagated uncertainty documentation was missing.

D-6.0 REFERENCES

The following reference list includes documents cited in this appendix. Parenthetical information following each reference provides the author(s), publication date, and ERID, ESHID, or EMID. This information is also included in text citations. ERIDs were assigned by the Laboratory's Associate Directorate for Environmental Management (IDs through 599999); ESHIDs were assigned by the Laboratory's Associate Directorate for Environment, Safety, and Health (IDs 600000 through 699999); and EMIDs are assigned by Newport News Nuclear BWXT-Los Alamos, LLC (N3B) (IDs 700000 and above). IDs are used to locate documents in N3B's Records Management System and in the Master Reference Set. The NMED Hazardous Waste Bureau and N3B maintain copies of the Master Reference Set. The set ensures that NMED has the references to review documents. The set is updated when new references are cited in documents.

- EPA (U.S. Environmental Protection Agency), February 1994. "USEPA Contract Laboratory Program National Functional Guidelines for Inorganic Data Review," EPA-540/R-94/013, Office of Emergency and Remedial Response, Washington, D.C. (EPA 1994, 048639)
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Table D-1.0-1
Inorganic Chemical, Organic Chemical, and
Radionuclide Analytical Methods for Samples Collected at SWMU 02-014

Analytical Method	Analytical Description	Analytical Suite							
Inorganic Chemicals									
EPA 300.0	Ion chromatography	Anions (nitrate)							
EPA SW-846: 6010B/6010C	Inductively coupled plasma emission spectroscopy—atomic emission spectroscopy	Aluminum, antimony, arsenic, barium, beryllium, calcium, cadmium, cobalt, chromium, copper, iron, lead, magnesium, manganese, nickel, potassium, selenium, silver, sodium, thallium, uranium, vanadium, and zinc (TAL metals)							
EPA SW-846:6020/6020B	Inductively coupled plasma mass spectrometry	Aluminum, antimony, arsenic, barium, beryllium, calcium, cadmium, cobalt, chromium, copper, iron, lead, magnesium, manganese, nickel, potassium, selenium, silver, sodium, thallium, vanadium, and zinc (TAL metals)							
EPA SW-846:9012A	Automated colorimetric/off-line distillation	Total cyanide							
EPA SW-846:6850	Liquid chromatography–mass spectrometry/mass spectrometry/	Perchlorate							
EPA SW-846:7471A	Cold vapor atomic absorption	Mercury							
Organic Chemicals									
EPA SW-846: 8082	Gas chromatography	PCBs							
EPA SW-846:8260B	Gas chromatography–mass spectrometry	VOCs							
EPA SW-846:8270C	Gas chromatography–mass spectrometry	SVOCs							
EPA SW-846:8290	High-resolution gas chromatography/high-resolution mass spectrometry	Dioxins/furans							
Radionuclides									
EPA 901.1	Gamma spectroscopy	Cesium-134, cesium-137, cobalt-60, sodium-22							
HASL Method 300:AM-241 HASL Method 300:ISOPU HASL Method 300:ISOU	Chemical separation alpha spectrometry	Americium-241 Isotopic plutonium Isotopic uranium							
EPA 905.0	Gas proportional counting	Strontium-90							
EPA 906.0	Liquid scintillation	Tritium							

Appendix E

Analytical Suites and Results and Analytical Reports (on CD included with this document)

Appendix F

Risk Assessments

CONTENTS

F-1.0	INTRO	DUCTION	F-1
F-2.0	BACK	GROUND	F-1
	F-2.1	Site Description and Operational History	F-1
		F-2.1.1 SWMU 02-014	
	F-2.2	Investigation Sampling	F-2
	F-2.3	Determination of COPCs	F-2
F-3.0	CONC	EPTUAL SITE MODEL	F-2
	F-3.1	Receptors and Exposure Pathways	F-2
	F-3.2	Environmental Fate and Transport	F-3
		F-3.2.1 Inorganic Chemicals	F-4
		F-3.2.2 Organic Chemicals	F-5
		F-3.2.3 Radionuclides	F-6
	F-3.3	Exposure Point Concentration Calculations	F-7
F-4.0	HUMA	N HEALTH RISK-SCREENING EVALUATIONS	F-7
	F-4.1	Human Health SSLs	F-8
	F-4.2	Results of Human Health Screening Evaluation	F-8
		F-4.2.1 SWMU 02-014	F-8
	F-4.3	Vapor Intrusion Pathway	F-9
		F-4.3.1 SWMU 02-014	F-9
	F-4.4	Essential Nutrients	F-10
	F-4.5	Uncertainty Analysis	F-10
		F-4.5.1 Data Evaluation and COPC Identification Process	F-10
		F-4.5.2 Exposure Evaluation	F-10
		F-4.5.3 Toxicity Evaluation	F-11
		F-4.5.4 Additive Approach	F-12
	F-4.6	Interpretation of Human Health Risk Screening Results	F-12
		F-4.6.1 SWMU 02-014	F-12
F-5.0	ECOL	OGICAL RISK EVALUATION TA-02 CORE AREA	F-13
F-6.0	CONC	LUSIONS	F-13
	F-6.1	Human Health Risk	F-13
F-7.0	REFE	RENCES	F-14

Figures

Figure F-3.1-1	Conceptual site model for Solid Waste Management Unit 02-014	F_17
Figure F-3. I-1	Conceptual site model for Solid Waste Management Onit 02-014	

Tables

Table F-2.3-1	EPCs at SWMU 02-014 for the Industrial and Recreational Scenarios	F-19
Table F-2.3-2	EPCs at SWMU 02-014 for the Residential Scenario	F-20
Table F-3.2-1	Physical and Chemical Properties of Inorganic COPCs at SWMU 02-014	F-22
Table F-3.2-2	Physical and Chemical Properties of Organic COPCs at SWMU 02-014	F-22
Table F-3.2-3	Physical and Chemical Properties of Radionuclide COPCs at SWMU 02-014	F-23
Table F-3.3-1	TEFs Used for Calculating TCDD-Equivalent Concentrations	F-23
Table F 4.1-1	Exposure Parameters Used to Calculate Chemical SSLs for the Industrial, Recreational, and Residential Scenarios	F-24
Table F-4.1-2	Parameter Values Used to Calculate Radionuclide SALs for the Residential Scenario	F-25
Table F-4.1-3	Parameter Values Used to Calculate Radionuclide SALs for the Industrial and Construction Worker Scenarios	F-26
Table F-4.1-4	Parameter Values Used to Calculate Radionuclide SALs for the Recreational Scenario	F-26
Table F-4.2-1	Industrial Carcinogenic Screening Evaluation for SWMU 02-014	F-27
Table F-4.2-2	Industrial Noncarcinogenic Screening Evaluation for SWMU 02-014	F-27
Table F-4.2-3	Industrial Radionuclide Screening Evaluation for SWMU 02-014	F-28
Table F-4.2-4	Recreational Carcinogenic Screening Evaluation for SWMU 02-014	F-28
Table F-4.2-5	Recreational Noncarcinogenic Screening Evaluation for SWMU 02-014	F-28
Table F-4.2-6	Recreational Radionuclide Screening Evaluation for SWMU 02-014	F-29
Table F-4.2-7	Residential Carcinogenic Screening Evaluation for SWMU 02-014	F-29
Table F-4.2-8	Residential Noncarcinogenic Screening Evaluation for SWMU 02-014	F-30
Table F-4.2-9	Residential Radionuclide Screening Evaluation for SWMU 02-014	F-30
Table F-4.2-10	Construction Worker Noncarcinogenic Screening Evaluation for SWMU 02-014	F-31
Table F-4.4-1	Essential Nutrient Screening Assessment	F-32

Attachments

Attachment F-1	Dioxin and Furan Toxicity Equivalency Factor Calculations
	(on CD included with this document)

Attachment F-2 ProUCL Files (on CD included with this document)

F-1.0 INTRODUCTION

This appendix presents the results of the human health risk-screening evaluations conducted in support of the environmental characterization of Solid Waste Management Unit (SWMU) 02-014, located in the northern portion of Los Alamos National Laboratory (LANL or the Laboratory). The evaluations of potential risk at this SWMU are based on decision-level data from the 2007, 2010, 2017, and 2018 investigations.

F-2.0 BACKGROUND

A brief description of SWMU 02-014, which was assessed for potential risks and dose in this addendum, is presented below.

F-2.1 Site Description and Operational History

SWMU 02-014 is located within Technical Area 02 (TA-02) at the Laboratory. TA-02 was used to house a series of research reactors from 1943 to 2003 when decontamination and decommissioning (D&D) of the site occurred. The main reactor building (02-1) was constructed in 1943. It housed five separate nuclear reactors: three iterations of water boiler reactors located on the east side of the building, one plutoniumfueled reactor (the Clementine Reactor), and the Omega West Reactor (OWR). A number of facilities were constructed over the years to support the TA-02 research activities. TA-02 was active from 1943 to 1993 (WD-3 2003, 082646, pp. 1–2). Various remedial actions, such as soil removal and D&D, were conducted in the bottom of Los Alamos Canyon, including at TA-02, after the Cerro Grande fire. These actions were taken to reduce the risk of contaminants dispersing from post-fire floods. Approximately 54 yd³ of soil contaminated with cesium-137 was removed in 2000, following an extensive field survey for gross-gamma radiation (LANL 2001, 070352). The OWR and associated structures underwent D&D in 2002 and 2003 (WD-3 2003, 082646). After all structures at TA-02 were removed, field radiological surveys were conducted to confirm that surface contamination release limits were not exceeded (WD-3 2003, 082646, pp. 18–19). The land was returned to its original contour and reseeded (WD-3 2003, 082646, pp. 1–2). The road accessing the reactor site is controlled by the Laboratory via a locked gate.

F-2.1.1 SWMU 02-014

SWMU 02-014 consists of three former electrical transformer stations (structures 02-31, 02-45, and 02-51) that served buildings in TA-02. This site was not identified as a SWMU or AOC in the 1990 SWMU report. (LANL 1990, 007511). This site was identified during efforts to discover the source of polychlorinated biphenyl (PCB) contamination identified during investigation sampling at storm drain area of concern (AOC) 02-011(a)(ii). Historical records, including engineering drawings and photographs, were reviewed and three potential sources of PCBs were identified. Former structure 02-31 was an electrical transformer station located 40 ft behind building 02-1. The transformer station was built in 1944 and was removed in 1950. Former structure 02-45 was built in 1954 to serve building 02-44. The transformer structure consisted of three transformers mounted across two telephone poles approximately 14 ft above the ground. The transformer station was replaced with another transformer station (structure 02-51). Former structure 02-51 was an electrical transformer station located approximately 20 ft southwest of former structure 02-31 and 20 ft southeast of former structure 02-45. Historical records indicated PCB-containing transformer oil had been used at this former transformer station. Structure 02-51 was constructed in 1961 and demolished in 2003. Soil at SWMU 02-014 was remediated in 2018.

F-2.2 Investigation Sampling

The final data set used to identify chemicals of potential concern (COPCs) for SWMU 02-014 and used in this appendix to evaluate the potential risks to human health and the environment are the qualified analytical results from the 2007–2018 investigations. Only those data determined to be of decision-level quality following the data-quality assessment (Appendix D) are included in the final data set evaluated in this appendix.

F-2.3 Determination of COPCs

Section 5.0 of the Phase II investigation report (Phase II IR) (N3B 2018, 700091) summarizes the COPC selection process. Only COPCs detected above background (inorganic chemicals and naturally occurring radionuclides); with detection limits greater than background values (BVs) (inorganic chemicals); and detected organic chemicals, inorganic chemicals with no BVs, and fallout radionuclides were retained. The industrial scenario and the recreational scenario used data for samples collected from 0.0 to 1.0 ft below ground surface (bgs). The residential scenario used data for samples collected from 0.0 to 10.0 ft bgs. However, sampling depths often overlapped because of multiple investigations; therefore, samples with a starting depth less than the lower bound of the interval were included in the risk-screening assessments for a given scenario as appropriate.

Tables F-2.3-1 and F-2.3-2 summarize the COPCs evaluated for potential risk at SWMU 02-014. Some of the COPCs identified in this addendum may not be evaluated for potential risk under one or more scenarios because they were not detected within the specified depth intervals associated with a given scenario.

F-3.0 CONCEPTUAL SITE MODEL

The primary mechanisms of release related to historical contaminant sources are described in detail in the approved investigation work plan for the Middle Los Alamos Canyon Aggregate Area (LANL 2006, 092571.12; NMED 2006, 095416). Releases from the sites at TA-02 may have occurred as a result of air emissions, surface releases, subsurface leaks, or effluent discharges. Previous sampling results indicated contamination from inorganic chemicals, organic chemicals, and radionuclides (LANL 2008, 101669.12).

F-3.1 Receptors and Exposure Pathways

The primary exposure pathway for human receptors is surface soil and subsurface soil/tuff that may be brought to the surface through intrusive activities. Migration of contamination to groundwater through the vadose zone is unlikely given the depth to regional groundwater (more than 500 ft bgs). Human receptors may be exposed through direct contact with soil or suspended particulates by ingestion, inhalation, dermal contact, and external irradiation pathways. Direct contact exposure pathways from subsurface contamination to human receptors are complete for the resident. The exposure pathways are the same as those for surface soil. Sources, exposure pathways, and receptors are shown in the conceptual site model (Figure F-3.1-1). Complete exposure pathways having soil/tuff as a secondary source are addressed in section F-4.2 and volatilization to air is addressed in section F-4.3.

New Mexico Environment Department (NMED) guidance (NMED 2017, 602273) requires that sites larger than 2 acres be evaluated to determine if beef ingestion is a plausible and complete exposure pathway. SWMU 02-014 is smaller than 2 acres. In addition, grazing is not allowed on Laboratory property. Therefore, further evaluation of the beef ingestion pathway is not necessary.

The Middle Los Alamos Canyon Aggregate Area is primarily a former industrial area, and all sites are on Laboratory property. None of the sites are active and therefore they currently provide habitat for ecological receptors. Weathering of tuff is the only viable natural process that may result in the exposure of receptors to COPCs in tuff. However, because of the slow rate of weathering expected for tuff, exposure to COPCs in tuff is negligible, although it is included in the assessments. Exposure pathways to subsurface contamination below 10.0 ft (human health) are not complete and would be complete only if contaminated soil or tuff were excavated and brought to the surface.

F-3.2 Environmental Fate and Transport

The evaluation of environmental fate addresses the chemical processes affecting the persistence of chemicals in the environment, and the evaluation of transport addresses the physical processes affecting mobility along a migration pathway. Migration into soil and tuff depends on precipitation or snowmelt, soil moisture content, depth of soil, soil hydraulic properties, and properties of the COPCs. Migration into and through tuff also depends on the unsaturated flow properties of the tuff and the presence of joints and fractures.

The most important factor with respect to the potential for COPCs to migrate to groundwater is the presence of saturated conditions. Downward migration in the vadose zone is also limited by a lack of hydrostatic pressure as well as the lack of a source for the continued release of contamination. Without sufficient moisture and a source, there is little or no potential for migration of materials through the vadose zone to groundwater.

Contamination at depth is addressed in the discussion of nature and extent in the Phase II IR (N3B 2018, 700091). Results from the deepest samples collected at most sites showed either no detected concentrations of COPCs or low- to trace-level concentrations of only a few inorganic, radionuclide, and/or organic COPCs in tuff. The limited extent of contamination is related to the absence of the key factors that facilitate migration, as discussed above. Given how long the contamination has been present in the subsurface, the physical and chemical properties of the COPCs, and the lack of saturated conditions, the potential for contaminant migration to groundwater is very low.

NMED guidance (NMED 2017, 602273) contains screening levels that consider the potential for contaminants in soil to result in groundwater contamination. These screening levels consider equilibrium partitioning of contaminants among solid, aqueous, and vapor phases and account for dilution and attenuation in groundwater through the use of dilution attenuation factors (DAFs). These DAF soil screening levels (SSLs) may be used to identify chemical concentrations in soil that have the potential to contaminate groundwater (EPA 1996, 059902). Screening contaminant concentrations in soil against these DAF SSLs does not, however, provide an indication of the potential for contaminants to migrate to groundwater. The assumptions used in the development of these DAF SSLs include an assumption of uniform contaminant concentrations from the contaminant source to the water table (i.e., migration to groundwater is assumed to have already occurred). Furthermore, this assumption is inappropriate for cases such as SWMU 02-014, where sampling has shown that contamination is vertically bounded near the surface and the distance from the surface to the water table is large. For these reasons, screening of contaminant concentrations in soil against the DAF SSLs was not performed.

The relevant release and transport processes of the COPCs are a function of chemical-specific properties that include the relationship between the physical form of the constituents and the nature of the constituent transport processes in the environment. Specific properties include the degree of saturation and the potential for ion exchange (barium and other inorganic chemicals) or sorption and the potential for natural bioremediation. The transport of volatile organic compounds (VOCs) occurs primarily in the vapor phase by diffusion or advection in subsurface air.

Current potential transport mechanisms that may lead to exposure include

- dissolution and/or particulate transport of surface contaminants during precipitation and runoff events,
- airborne transport of contaminated surface soil,
- continued dissolution and advective/dispersive transport of chemical contaminants contained in subsurface soil and tuff as a result of past operations,
- disturbance of contaminants in shallow soil and subsurface tuff by Laboratory operations, and
- disturbance and uptake of contaminants in shallow soil by plants and animals.

Contaminant distributions indicate that after the initial deposition of contaminants from operational activities and historical remediation efforts, elevated levels of COPCs tend to remain concentrated in the vicinity of the original release points. The primary potential release and transport mechanisms identified for the site in the Middle Los Alamos Canyon Aggregate Area addendum include direct discharge; precipitation, sorption, and mechanical transport; dissolution and advective transport in water; and volatilization, diffusion, and dispersion. Less significant transport mechanisms include wind entrainment and, given the asphalt pavement covering most sites, dispersal of surface soil and uptake of contaminants from soil and water by biota.

Gas or vapor-phase contaminants such as VOCs are likely to volatilize to the atmosphere from nearsurface soil and sediment and/or migrate by diffusion through air-filled pores in the vadose zone. Migration of vapor-phase contaminants from tuff into ambient air may occur by diffusion or advection driven by barometric pressure changes.

F-3.2.1 Inorganic Chemicals

In general, and particularly in a semiarid climate, inorganic chemicals are not highly soluble or mobile in the environment, although there are exceptions. The physical and chemical factors that determine the distribution of inorganic COPCs within the soil and tuff at SWMU 02-014 include the soil-water partition coefficient (K_d) of the inorganic chemicals, the pH of the soil, soil characteristics (such as sand or clay content), and the redox potential (Eh). The interaction of these factors is complex, but the K_d values provide a general assessment of the potential for migration through the subsurface; chemicals with higher K_d values are less likely to be mobile than those with lower ones. Chemicals with K_d values greater than 40 are very unlikely to migrate through soil towards the water table (Kincaid et al. 1998, 093270). Table F-3.2-1 presents the K_d values and water solubility for the inorganic COPCs at SWMU 02-014. Based on this criterion, the following COPCs have a low potential to mobilize and migrate through soil and the vadose zone: aluminum, barium, chromium, manganese, mercury, nickel, thallium, and vanadium. The K_d values for arsenic, copper, iron, and selenium are less than 40 and may indicate a greater potential to mobilize and migrate through soil and the vadose zone beneath the sites. A K_d is not available for perchlorate.

It is important to note that other factors besides the K_d values (e.g., speciation in soil, oxidation-reduction potential, pH, and soil mineralogy) also play significant roles in the likelihood that inorganic chemicals will migrate. The COPCs with K_d values less than 40 are discussed further below. Information about the fate and transport properties of inorganic chemicals was obtained from individual chemical profiles published by the Agency for Toxic Substances and Disease Registry (ATSDR) (ATSDR 1997, 056531, and http://www.atsdr.cdc.gov/toxpro2).

Arsenic may undergo a variety of reactions, including oxidation-reduction reactions, ligand exchange, precipitation, and biotransformation. Arsenic forms insoluble complexes with iron, aluminum, and magnesium oxides found in soil and in this form, arsenic is relatively immobile. However, under low pH and reducing conditions, arsenic can become soluble and may potentially leach into groundwater or result in runoff of arsenic into surface waters. Arsenic is expected to have low mobility under the environmental conditions (neutral to alkaline soil pH and oxidizing near-surface conditions) present in the Middle Los Alamos Canyon Aggregate Area.

Copper movement in soil is determined by physical and chemical interactions with the soil components. Most copper deposited in soil will be strongly adsorbed and remains in the upper few centimeters of soil. Copper will adsorb to organic matter, carbonate minerals, clay minerals, or hydrous iron, and manganese oxides. In most temperate soil, pH, organic matter, and ionic strength of the soil solutions are the key factors affecting adsorption. Soil in the area is neutral to slightly alkaline, so the leaching of copper is not a concern at this site. Copper binds to soil much more strongly than other divalent cations, and the distribution of copper in the soil solution is less affected by pH than other metals. Copper is expected to be bound to the soil and move in the system by water transport of soil particles as opposed to dissolved species.

Iron is naturally occurring in soil and tuff and may be relatively mobile under reducing conditions. Iron is sensitive to soil pH conditions, occurring in two oxidation states, iron(III), the insoluble oxidized form, and iron(II), the reduced soluble form. Most iron in well-drained neutral-to-alkaline soil is present as precipitates of iron(III) hydroxides and oxides. With time, these precipitates are mineralized and form various iron minerals, such as lepidcrocite, hematite, and goethite. Iron is not expected to be mobile in the neutral to slightly alkaline, well-drained soil at the Middle Los Alamos Canyon Aggregate Area.

Perchlorate is somewhat soluble in water and may migrate with water molecules in saturated soil. As noted above, the subsurface material beneath the sites has low moisture content, which inhibits the mobility of perchlorate as well as most other inorganic chemicals.

Selenium is not often found in the environment in its elemental form but is usually combined with sulfide minerals or with silver, copper, lead, and nickel minerals. In soil, pH and Eh are determining factors in the transport and partitioning of selenium. In soil with a pH of greater than 7.5, selenates, which have high solubility and a low tendency to adsorb onto soil particles, are the major selenium species and are very mobile. The soil pH at most sites in the Middle Los Alamos Canyon Aggregate Area is neutral to slightly alkaline, indicating that selenium is not likely to migrate.

F-3.2.2 Organic Chemicals

Table F-3.2-2 presents the physical and chemical properties (organic carbon-water partition coefficient $[K_{oc}]$, logarithm to the base 10 octanol/water partition coefficient $[\log K_{ow}]$, solubility, and vapor pressure) of the organic COPCs identified at SWMU 02-014. The physical and chemical properties of organic chemicals are important when evaluating their fate and transport. The following physiochemical property information illustrates some aspects of the fate and transport of COPCs at SWMU 02-014. The information is summarized from Ney (1995, 058210).

Water solubility may be the most important chemical characteristic used to assess mobility of organic chemicals. The higher the water solubility of a chemical, the more likely it is to be mobile and the less likely it is to accumulate, bioaccumulate, volatilize, or persist in the environment. A highly soluble chemical (water solubility greater than 1000 mg/L) is prone to biodegradation and metabolism that may detoxify the parent chemical. No chemicals detected at SWMU 02-014 in the Middle Los Alamos Canyon Aggregate Area have water solubilities greater than 1000 mg/L.

The lower the water solubility of a chemical, especially below 10 mg/L, the more likely it will be immobilized by adsorption. Chemicals with lower water solubilities are more likely to accumulate or bioaccumulate and persist in the environment, are slightly prone to biodegradation, and are metabolized in plants and animals. The chemicals identified as having water solubilities less than 10 mg/L are the polycyclic aromatic hydrocarbons (PAHs), PCBs, and 2,3,7,8-tetrachlorodibenzodioxin (2,3,7,8-TCDD).

Vapor pressure is a characteristic used to evaluate the tendency of organic chemicals to volatize. Chemicals with vapor pressure greater than 0.01 millimeters of mercury (mmHg) are likely to volatilize and, therefore, concentrations at the site are reduced over time; vapors of these chemicals are more likely to travel toward the atmosphere and not migrate towards groundwater. Toluene has vapor pressure greater than 0.01 mmHg.

Chemicals with vapor pressures less than 0.000001 mmHg are less likely to volatilize and, therefore, tend to remain immobile. Benzo(a)pyrene, benzo(a)anthracene, benzo(b)fluoranthene, benzo(g,h,i)perylene, benzo(k)fluoranthene, chrysene, indeno(1,2,3-cd)pyrene, and 2,3,7,8-TCDD have vapor pressures less than 0.000001 mmHg.

The K_{ow} is an indicator of a chemical's potential to bioaccumulate or bioconcentrate in the fatty tissues of living organisms. The unitless K_{ow} value is an indicator of water solubility, mobility, sorption, and bioaccumulation. The higher the K_{ow} above 1000, the greater the affinity the chemical has for bioaccumulation/bioconcentration in the food chain, the greater the potential for sorption in the soil, and the lower the mobility (Ney 1995, 058210).

No COPCs have a K_{ow} greater than 1000. A K_{ow} of less than 500 indicates high water solubility; mobility; little to no affinity for bioaccumulation; and degradability by microbes, plants, and animals. PAHs; PCBs; 2,3,7,8-TCDD; and toluene have a K_{ow} much less than 500.

The K_{oc} measures the tendency of a chemical to adsorb to organic carbon in soil. K_{oc} values above 500 L/kg indicate a strong tendency to adsorb to soil, leading to low mobility (NMED 2017, 602273). Most organic chemicals detected have K_{oc} values above 500 L/kg, indicating a very low potential to migrate toward groundwater. The only organic chemical with a K_{oc} value less than 500 L/kg is toluene.

The PAHs; PCBs; and 2,3,7,8-TCDD are the least mobile and the most likely to bioaccumulate. Toluene is more soluble and volatile and is more likely to travel toward the atmosphere and not migrate toward groundwater. Because the organic chemicals detected were at low concentrations and extent is defined, they are not likely to migrate to groundwater.

F-3.2.3 Radionuclides

Radionuclides are generally not highly soluble or mobile in the environment, particularly in the semiarid climate of the Laboratory. The physical and chemical factors that determine the distribution of radionuclides within soil and tuff are the K_d, the pH of the soil and other soil characteristics (e.g., sand or clay content), and the Eh. The interaction of these factors is complex, but K_d values provide a general assessment of the potential for migration through the subsurface: chemicals with higher K_d values are less likely to be mobile than those with lower values. Radionuclides with K_d values greater than 40 are very unlikely to migrate through soil towards the water table (Kincaid et al. 1998, 093270).

Table F-3.2-3 gives physical and chemical properties of the radionuclide COPCs identified at sites in SWMU 02-014. Based on K_d values, cesium-134, cesium-137, plutonium-238, and plutonium-239/240 have a very low potential to migrate towards groundwater at SWMU 02-014.

F-3.3 Exposure Point Concentration Calculations

The exposure point concentrations (EPCs) represent upper-bound concentrations of COPCs. For comparison with risk-screening levels, the upper confidence limit (UCL) of the arithmetic mean was calculated when possible and used as the EPC. The UCLs were calculated using all available decision-level data within the depth range of interest. If an appropriate UCL of the mean could not be calculated, the maximum detected concentration of the COPC was used as the EPC. The summary statistics, including the EPC for each COPC for the human health risk-screening assessments and the distribution used for the calculation, are presented in Tables F-2.3-1 and F-2.3-2.

The EPCs for the dioxin and furan congeners are the sums of the detected congeners weighted by the toxic equivalency factors (TEFs) (NMED 2017, 602273); the sum is expressed as the 2,3,7,8-TCDD- equivalent concentration. The toxic equivalency factors used are presented in Table F-3.3-1. The results of the TEF calculations for each site where dioxins/furans are COPCs are presented in Attachment F-1 and the 2,3,7,8-TCDD-equivalent concentrations (95% UCLs or maximum concentrations) are presented in the section F-4.2 tables.

The UCLs of the mean concentrations were calculated using the U.S. Environmental Protection Agency (EPA) ProUCL 5.1.002 software (EPA 2015, 601725), which is based on EPA guidance (EPA 2002, 085640). Consistent with the ProUCL v5.1 Technical Guide, a minimum of eight samples and five detections are needed to calculate UCLs (EPA 2015, 601724). The ProUCL program calculates 95%, 97.5%, and 99% UCLs and recommends a distribution and UCL. The 95% UCL for the recommended calculation method was used as the EPC. The ProUCL software performs distributional tests on the data set for each COPC and calculates the most appropriate UCL based on the distribution of the data set. Environmental data may have a normal, lognormal, or gamma distribution but are often nonparametric (no definable shape to the distribution). The ProUCL documentation strongly recommends against using the maximum detected concentration for the EPC. The maximum detected concentration was used to represent the EPC for COPCs only when there were too few detections to calculate a UCL. Input and output data files for ProUCL calculations are provided on CD as Attachment F-2.

F-4.0 HUMAN HEALTH RISK-SCREENING EVALUATIONS

The human health risk-screening assessment for SWMU 02-014 evaluated residential, industrial, and recreational exposure scenarios. SWMU 02-014 was screened for the residential scenario using data from 0.0 to 10.0 ft bgs and screened for industrial and recreational scenarios using data from 0.0 to 1.0 ft bgs, where available. The human health risk-screening assessments compared either the 95% UCL of the mean concentration or the maximum detected concentration of each COPC with SSLs for chemicals.

For most constituents, the residential exposure scenario is the most protective, and the residential scenario is used for evaluating whether a site is appropriate for corrective action complete without controls. Although based on current and foreseeable land use, potential exposure to construction workers is not expected at SWMU 02-014, sites being recommended for corrective action complete without controls must not pose an unacceptable risk to construction workers. For some constituents, the construction worker SSL is less than the residential SSL and the residential exposure scenario may not also be protective of construction workers. Therefore, sites posing no potential unacceptable risk under the residential scenario are evaluated to determine whether the residential exposure scenario is also protective of construction workers. If not, the construction worker scenario is evaluated to determine whether the site could be recommended for corrective action complete without controls.

F-4.1 Human Health SSLs and SALs

Human health risk-screening assessments were conducted using SSLs for the industrial and residential scenarios obtained from NMED guidance (NMED 2017, 602273). The NMED SSLs are based on a target hazard quotient (HQ) of 1 and a target cancer risk of 1×10^{-5} (NMED 2017, 602273). Recreational SSLs were obtained from Laboratory guidance (LANL 2017, 602581) and are based on the same target risk levels as the NMED SSLs. Surrogate chemicals were also used for some COPCs without an SSL based on structural similarity or because the COPC is a breakdown product (NMED 2003, 081172). Exposure parameters used to calculate the industrial, recreational, and residential SSLs are presented in Table F-4.1-1.

Radionuclide screening action levels (SALs) were used for comparison with radionuclide COPC EPCs and were derived using the Residual Radioactivity (RESRAD) model, Version 7.0 (LANL 2015, 600929). The SALs are based on a 25-mrem/yr dose as authorized by U.S. Department of Energy (DOE) Order 458.1. Exposure parameters used to calculate the SALs are presented in Tables F-4.1-2, F-4.1-3, and F-4.1-4.

F-4.2 Results of Human Health Screening Evaluation

The EPC of each COPC was compared with the SSLs for the industrial, recreational, and residential scenarios, as appropriate. For carcinogenic chemicals, the EPCs were divided by the SSL and multiplied by 1×10^{-5} . The sum of the carcinogenic risks was compared with the NMED target cancer risk level of 1×10^{-5} . For noncarcinogenic chemicals, an HQ was generated for each COPC by dividing the EPC by the SSL. The HQs were summed to generate a hazard index (HI). The HI was compared with the NMED target HI of 1. The radionuclide EPCs were divided by the SAL and multiplied by 25 mrem/yr. The total doses were compared with the DOE target level of 25 mrem/yr, as authorized by DOE Order 458.1.The results are presented in Tables F-4.2-1 to F-4.2-10 and are described below for SWMU 02-014.

Sites posing no unacceptable risk under the residential scenario may be recommended for corrective action complete without controls if the residential scenario is also protective of construction workers. For SWMU 02-014, the following COPCs have construction worker SSLs less than residential SSLs: aluminum, barium, chromium, manganese, and nickel. The residential EPC for each of these COPCs was compared with the construction worker SSL. The ratio of the residential EPC for SWMU 02-014 to the construction worker SSL (i.e., the maximum HQ) was 0.14 for aluminum, 0.01 for barium, 0.02 for chromium, 0.44 for manganese, and 0.003 for nickel. Thus, manganese is the only COPC that could potentially pose an unacceptable construction worker risk. The construction worker scenario was evaluated to determine whether this site can be recommended for corrective action complete without controls. Because manganese is associated with noncarcinogenic risk, only the construction worker HI was evaluated; the residential scenario is protective of construction worker cancer risk at all sites. Concerning radionuclides, all construction worker SALs are equal to or greater than residential SALs and the residential scenario is protective of construction workers for all sites.

F-4.2.1 SWMU 02-014

The results of the risk-screening assessment for the industrial scenario are presented in Tables F-4.2-1, F-4.2-2, and F-4.2-3. The total excess cancer risk for the industrial scenario is 5×10^{-6} , which is less than the NMED target risk level of 1×10^{-5} (NMED 2017, 602273). The industrial HI is 0.06, which is less than the NMED target HI of 1 (NMED 2017, 602273). The total dose is 0.0009 mrem/yr, which is less than the target dose of 25 mrem/yr as authorized by DOE Order 458.1.

The results of the risk-screening assessment for the recreational scenario are presented in Tables F-4.2-4, F-4.2-5, and F-4.2-6. The total excess cancer risk for the recreational scenario is 5×10^{-6} , which is less than the NMED target risk level of 1×10^{-5} (NMED 2017, 602273). The recreational HI is 0.2, which is less than the NMED target HI of 1 (NMED 2017, 602273). The total dose is 0.0008 mrem/yr, which is less than the target dose of 25 mrem/yr as authorized by DOE Order 458.1.

The results of the risk-screening assessment for the residential scenario are presented in Tables F-4.2-7, F-4.2-8, and F-4.2-9. The total excess cancer risk for the residential scenario is 1×10^{-5} , which is equal to the NMED target risk level of 1×10^{-5} (NMED 2017, 602273). The residential HI is 0.6, which is less than the NMED target HI of 1 (NMED 2017, 602273). The total dose is 2 mrem/yr, which is less than the target dose of 25 mrem/yr as authorized by DOE Order 458.1.

Manganese is a COPC at SWMU 02-014 and may potentially pose an unacceptable noncarcinogenic risk to the construction worker. Construction worker risk for SWMU 02-014 was evaluated using the EPCs for the residential scenario (Table F-2.3-2) since both residential and construction worker scenarios consider the 0.0-ft to 10.0-ft-bgs interval. The noncarcinogenic risk screening results for the construction worker at SWMU 02-014 are presented in Table F-4.2-10. The construction worker HI is 0.8, which is less than the NMED target of 1 (NMED 2017, 602273). The primary contributor to construction worker noncarcinogenic risk is manganese. The residential scenario is protective of the construction worker for cancer risk and total dose.

F-4.3 Vapor Intrusion Pathway

NMED guidance (NMED 2017, 602273) requires an evaluation of the vapor-intrusion pathway. The vapor intrusion pathway of VOCs into a building was evaluated where appropriate. The evaluation can be qualitative for a potentially complete pathway if the following criteria are met:

- Volatile and toxic compounds are minimally detected,
- Concentrations are below NMED's vapor-intrusion screening levels for soil-gas and/or groundwater. There is no suspected source(s) for volatile and toxic compounds, and
- Concentrations are decreasing with depth (for soil).

Because only bulk soil data are available for this site, the vapor-intrusion screening levels are not applicable for the evaluation. The vapor-intrusion pathway was qualitatively evaluated as part of the residential scenario. Among the factors considered for the vapor-intrusion pathway to be relevant to human health risk is the current extent of structures and their proximity to the VOC source. One may also consider if construction of buildings is possible or proposed in the reasonably foreseeable future.

F-4.3.1 SWMU 02-014

SWMU 02-014 consists of three former electrical transformer stations (structures 02-31, 02-45, and 02-51) that served buildings in TA-02. This site was not identified as a SWMU or AOC in the 1990 SWMU report. (LANL 1990, 007511). This site was identified during efforts to discover the source of PCB contamination identified during investigation sampling at storm drain AOC 02-011(a)(ii). Historical records, including engineering drawings and photographs, were reviewed and three potential sources of PCBs were identified. Former structure 02-31 was an electrical transformer station located 40 ft behind building 02-1. The transformer station was built in 1944 and was removed in 1950. Former structure 02-45 was built in 1954 to serve building 02-44. The transformer structure consisted of three transformers mounted across two telephone poles approximately 14 ft above the ground. The transformer station was replaced with another transformer station (structure 02-51). Former structure 02-51 was an electrical transformer structure 02-51 was an electrical transformer structure 02-51 was an electrical transformer structure 02-51 was mounted across two telephone poles approximately 14 ft above the ground. The transformer station was replaced with

located approximately 20 ft southwest of former structure 02-31 and 20 ft southeast of former structure 02-45. Historical records indicated PCB-containing transformer oil had been used at this former transformer station. Structure 02-51 was constructed in 1961 and demolished in 2003. Soil at SWMU 02-014 was remediated in 2018.

One VOC, toluene was minimally detected at this site (one detection in two samples) with a maximum concentration of 0.000465 mg/kg that was less than the nondetected concentration (0.00107 mg/kg) in the second sample. Furthermore, the site description does not indicate that solvents were used, so no sources of VOCs are present and VOCs were only detected minimally. In addition, the structures have been removed, soil has been remediated, and the site is inactive. The vapor-intrusion pathway is therefore potentially complete based on NMED guidance (NMED 2017, 602273) and no additional evaluation is necessary.

F-4.4 Essential Nutrients

NMED has SSLs for evaluation of essential nutrients (NMED 2017, 602273). Magnesium was identified as a COPC at SWMU 02-014 and the maximum detected concentration of magnesium was compared with the appropriate NMED SSL. The results of the comparison found the maximum magnesium concentration to be substantially less than the SSL as presented in Table F-4.4-1. Further evaluation of magnesium at this site is not necessary.

F-4.5 Uncertainty Analysis

F-4.5.1 Data Evaluation and COPC Identification Process

A primary uncertainty associated with the COPC identification process is the possibility that a chemical may be inappropriately identified as a COPC when it is actually not a COPC or that a chemical may not be identified as a COPC when it actually should be identified as a COPC. Inorganic chemicals are appropriately identified as COPCs because only the chemicals detected or that have detection limits above background are retained for further analysis. There are no established BVs for organic chemicals, and all detected organic chemicals are identified as COPCs and are retained for further analysis unless shown to be from a source not related to the site. Other uncertainties may include errors in sampling, laboratory analysis, and data analysis. However, because concentrations used in the risk-screening evaluations include those detected below the estimated quantitation limits and nondetections above BVs, data evaluation uncertainties are expected to have little effect on the risk-screening results.

F-4.5.2 Exposure Evaluation

The current and reasonably foreseeable future land use is industrial. To the degree actual activity patterns are not represented by those activities assumed by the industrial scenario, uncertainties are introduced in the assessment, and the evaluation presented in this assessment overestimates potential risk. An individual may be subject to exposures in a different manner than the exposure assumptions used to derive the industrial SSLs. For the site evaluated, individuals might not be on-site at present or in the future for that frequency and duration. The industrial assumptions for the SSLs are that the potentially exposed individual is outside on-site for 8 hr/day, 225 days/yr, and 25 yr (NMED 2017, 602273). The residential SSLs are based on exposure of 24 hr/day, 350 days/yr, and 30 yr (NMED 2017, 602273). As a result, the industrial and residential scenarios evaluated at SWMU 02-014 likely overestimate the exposure and risk.

A number of assumptions are made relative to exposure pathways, including input parameters, completeness of a given pathway, the contaminated media to which an individual may be exposed, and intake rates for different routes of exposure. In the absence of site-specific data, the exposure assumptions used were consistent with default values (NMED 2017, 602273). When several upper-bound values (as are found in NMED 2017, 602273) are combined to estimate exposure for any one pathway, the resulting risk estimate can exceed the 99th percentile, and therefore, can exceed the range of risk that may be reasonably expected. Also, the assumption that residual concentrations of chemicals in the tuff are available and result in exposure in the same manner as if they were in soil overestimates the potential exposure and risk to receptors.

Uncertainty is introduced in the concentration aggregation of data for estimating the EPCs at a site. Risk from a single location or area with relatively high COPC concentrations may be underestimated by using a representative sitewide value. The use of a UCL is intended to provide a protective upper-bound (i.e., conservative) COPC concentration and is assumed to be representative of the average exposure to a COPC across the entire site. Potential risk and exposure from a single location or area with relatively high COPC concentrations may be overestimated if a representative sitewide value is used. The use of the maximum detected concentration for the EPC overestimates the exposure to contamination because receptors are not consistently exposed to the maximum detected concentration across the site. The use of EPCs similar to BVs will overestimate risk due to site-related contamination since much of the risk will be associated with background concentrations.

SWMU 02-014

The residential total excess cancer risk at SWMU 02-014 was approximately 1×10^{-5} . Aroclor-1260 (9.47 x 10⁻⁶), arsenic (1.13 x 10⁻⁶), and 2,3,7,8-TCDD (1.35 x 10⁻⁶), make up a majority of this risk. The residential EPCs for arsenic (0.798 mg/kg) and 2,3,7,8-TCDD (6.6E-06 mg/kg) are less than their respective SSLs of 7.07 mg/kg and 4.9E-05 mg/kg. The Aroclor-1260 EPC (2.3 mg/kg) is slightly less than the SSL (2.43 mg/kg).

F-4.5.3 Toxicity Evaluation

The primary uncertainty associated with the SSLs is related to the derivation of toxicity values used in their calculation. Toxicity values (reference doses [RfDs] and slope factors [SFs]) were used to derive the SSLs used in this risk-screening evaluation (NMED 2017, 602273). Uncertainties were identified in five areas with respect to the toxicity values: (1) extrapolation from other animals to humans, (2) interindividual variability in the human population, (3) the derivation of RfDs and SFs, (4) the chemical form of the COPC, and (5) the use of surrogate chemicals.

Extrapolation from Animals to Humans. The SFs and RfDs are often determined by extrapolation from animal data to humans, which may result in uncertainties in toxicity values because differences exist in chemical absorption, metabolism, excretion, and toxic responses between animals and humans. Differences in body weight, surface area, and pharmacokinetic relationships between animals and humans are taken into account to address these uncertainties in the dose-response relationship. However, conservatism is usually incorporated in each of these steps, resulting in the overestimation of potential risk.

Individual Variability in the Human Population. For noncarcinogenic effects, the degree of variability in human physical characteristics is important both in determining the risks that can be expected at low exposures and in defining the no observed adverse effect level (NOAEL). The NOAEL uncertainty factor approach incorporates a 10-fold factor to reflect individual variability within the human population that can

contribute to uncertainty in the risk evaluation; this factor of 10 is generally considered to result in a conservative estimate of risk to noncarcinogenic COPCs.

Derivation of RfDs and SFs. The RfDs and SFs for different chemicals are derived from experiments conducted by different laboratories that may have different accuracy and precision that could lead to an overestimation or underestimation of the risk. The uncertainty associated with the toxicity factors for noncarcinogens is measured by the uncertainty factor, the modifying factor, and the confidence level. For carcinogens, the weight of evidence classification indicates the likelihood that a contaminant is a human carcinogen. Toxicity values with high uncertainties may change as new information is evaluated.

Chemical Form of the COPC. COPCs may be bound to the environment matrix and not available for absorption into the human body. However, the COPCs are assumed to be bioavailable. This assumption can lead to an overestimation of the total risk.

Use of Surrogate Chemicals. The use of surrogates for chemicals that do not have EPA-approved or provisional toxicity values also contributes to uncertainty in the risk assessment. A surrogate was used to provide an SSL for benzo(g,h,i)perylene based on structural similarity. The overall impact of surrogates on the risk assessment is minimal because these COPCs were detected infrequently and at low concentrations.

F-4.5.4 Additive Approach

For noncarcinogens, the effects of exposure to multiple chemicals are generally unknown, and possible interactions could be synergistic or antagonistic, resulting in either an overestimation or underestimation of the potential risk. Additionally, RfDs used in the risk calculations typically are not based on the same endpoints with respect to severity, effects, or target organs. Therefore, the potential for noncarcinogenic effects may be overestimated for individual COPCs that act by different mechanisms or by different modes of action but are addressed additively.

F-4.6 Interpretation of Human Health Risk Screening Results

F-4.6.1 SWMU 02-014

Industrial Scenario

The total excess cancer risk for the industrial scenario is 5×10^{-6} , which is less than the NMED target risk level of 1×10^{-5} (NMED 2017, 602273). The industrial HI is 0.06, which is less than the NMED target HI of 1 (NMED 2017, 602273). The total dose is 0.0009 mrem/yr, which is less than the target dose of 25 mrem/yr as authorized by DOE Order 458.1. The total dose for the industrial scenario is equivalent to a total risk of 4×10^{-9} , based on conversion from dose using RESRAD Version 7.0 (LANL 2015, 600929).

Recreational Scenario

The total excess cancer risk for the recreational scenario is 5×10^{-6} , which is less than the NMED target risk level of 1×10^{-5} (NMED 2017, 602273). The recreational HI is 0.2, which is less than the NMED target HI of 1 (NMED 2017, 602273). The total dose is 0.0008 mrem/yr, which is less than the target dose of 25 mrem/yr as authorized by DOE Order 458.1. The total dose for the recreational scenario is equivalent to a total risk of 2×10^{-9} , based on conversion from dose using RESRAD Version 7.0 (LANL 2015, 600929).

Residential Scenario

The total excess cancer risk for the residential scenario is 1×10^{-5} , which is equivalent to the NMED target risk level of 1×10^{-5} (NMED 2017, 602273). The residential HI is 0.6, which is less than the NMED target HI of 1 (NMED 2017, 602273). The total dose is 2 mrem/yr, which is less than the target dose of 25 mrem/yr as authorized by DOE Order 458.1. The total dose for the residential scenario is equivalent to a total risk of 3×10^{-5} , based on conversion from dose using RESRAD Version 7.0 (LANL 2015, 600929).

The residential exposure scenario is also protective of construction workers.

F-5.0 ECOLOGICAL RISK EVALUATION TA-02 CORE AREA

Ecological risk was evaluated collectively for sites within the TA-02 core area, including SWMU 02-014, in the "Phase II Investigation Report for Middle Los Alamos Canyon Aggregate Area, Revision 2" (N3B 2018, 700091). Based on evaluations of the minimum ESLs, HI analyses, potential effects to populations (individuals for threatened and endangered species), lowest observed adverse effect level analyses, the relationship of detected concentrations and screening levels to background concentrations, and results of site-specific ecological risk studies conducted within the TA-02 core area, the Phase II IR concluded that no potential ecological risk sexist for the TA-02 core area, which includes SWMU 02-014. Therefore, further evaluation of ecological risk was not performed for this Phase II IR addendum.

F-6.0 CONCLUSIONS

F-6.1 Human Health Risk

The total excess cancer risks were less than the target risk level of 1×10^{-5} for the industrial and recreational scenarios at SWMU 02-014. The residential risks at SWMU 02-014 were equivalent to the target risk level of 1×10^{-5} .

The HIs were less than the target HI of 1 for the industrial, recreational, and residential scenarios at SWMU 02-014.

The total doses were below the target dose limit of 25 mrem/yr as authorized by DOE Order 458.1 for the industrial, recreational, and residential scenarios at SWMU 02-014. The total doses were equivalent to total risks of 4×10^{-9} for the industrial scenario, 2×10^{-9} for the recreational scenario, and 3×10^{-5} for the residential scenario, based on conversion from dose using RESRAD Version 7.0 (LANL 2015, 600929).

There was no potential noncarcinogenic risk to construction workers, and the residential scenario demonstrated protection of construction workers for carcinogenic risk and dose.

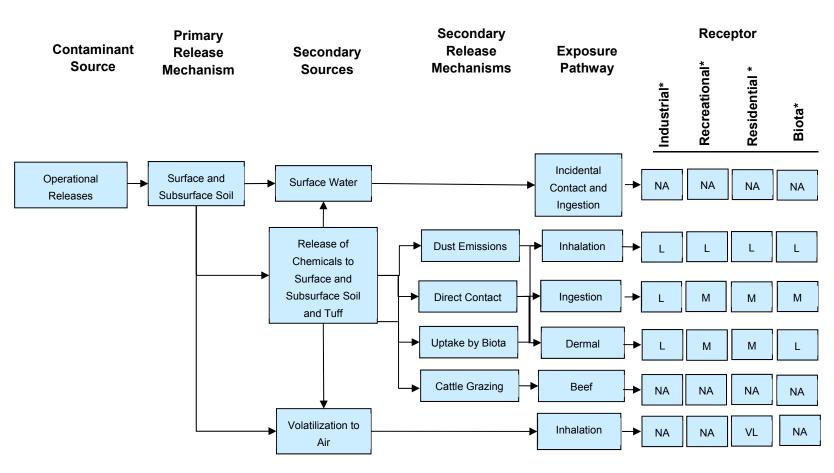
SWMU 02-014 in the Middle Los Alamos Canyon Aggregate Area is not accessible by the public and is not planned for release by DOE in the foreseeable future. Therefore, an as low as reasonably achievable (ALARA) evaluation for radiological exposure to the public is not currently required. Should DOE's plans for releasing this area change, an ALARA evaluation will be conducted at that time.

F-7.0 REFERENCES

The following reference list includes documents cited in this report. Parenthetical information following each reference provides the author(s), publication date, and ERID, ESHID, or EMID. This information is also included in text citations. ERIDs were assigned by the Laboratory's Associate Directorate for Environmental Management (IDs through 599999); ESHIDs were assigned by the Laboratory's Associate Directorate for Environment, Safety, and Health (IDs 600000 through 699999); and EMIDs are assigned by Newport News Nuclear BWXT-Los Alamos, LLC (N3B) (IDs 700000 and above). IDs are used to locate documents in N3B's Records Management System and in the Master Reference Set. The NMED Hazardous Waste Bureau and N3B maintain copies of the Master Reference Set. The set ensures that NMED has the references to review documents. The set is updated when new references are cited in documents.

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*Notes: Very low (VL), low (L), and moderate (M) designations indicate the pathway is a potentially complete pathway and is evaluated in the risk assessments. Not applicable (NA) indicates the pathway is incomplete and is not evaluated in the risk assessments.

Figure F-3.1-1 Conceptual site model for Solid Waste Management Unit 02-014

	Number	Number					
СОРС	of Analyses	of Detects	Minimum Concentration	Maximum Concentration	Distribution	EPC	EPC Method
	7 mary 505	Detects	ooneentration	ooncentration	Distribution		El o metriod
Inorganic Chemicals (mg/kg)		1		Γ	1	T	
Copper	17	16	2.38	47.5	Lognormal	11.7	95% Percentile Bootstrap
Mercury	17	17	0.013	0.538	Lognormal	0.284	95% Chebyshev (Mean, Sd)
Selenium	17	9	0.119	2.5 (U)	Normal	0.66	95% KM (t)
Organic Chemicals (mg/kg)							
Anthracene	1	1	0.00903	0.00903	n/a*	0.00903	Maximum detected concentration
Aroclor-1254	29	6	0.00114 (U)	3.93	Nonparametric	0.787	95% KM (Chebyshev)
Aroclor-1260	29	29	0.016	17.3	Gamma	3.91	95% Adjusted Gamma
Benzo(a)anthracene	1	1	0.0516	0.0516	n/a	0.0516	Maximum detected concentration
Benzo(a)pyrene	1	1	0.0551	0.0551	n/a	0.0551	Maximum detected concentration
Benzo(b)fluoranthene	1	1	0.0702	0.0702	n/a	0.0702	Maximum detected concentration
Benzo(g,h,i)perylene	1	1	0.0271	0.0271	n/a	0.0271	Maximum detected concentration
Benzo(k)fluoranthene	1	1	0.041	0.041	n/a	0.041	Maximum detected concentration
Chrysene	1	1	0.0598	0.0598	n/a	0.0598	Maximum detected concentration
Fluoranthene	1	1	0.0984	0.0984	n/a	0.0984	Maximum detected concentration
Indeno(1,2,3-cd)pyrene	1	1	0.0252	0.0252	n/a	0.0252	Maximum detected concentration
Phenanthrene	1	1	0.0317	0.0317	n/a	0.0317	Maximum detected concentration
Pyrene	1	1	0.0771	0.0771	n/a	0.0771	Maximum detected concentration
TCDD[2,3,7,8-] equivalent	1	1	0.0000066	0.0000066	n/a	0.0000066	Maximum detected concentration
Radionuclides (pCi/g)							
Plutonium-239/240	17	9	0 (U)	0.076	Gamma	0.0414	95% KM Adjusted Gamma

Table F-2.3-1 EPCs at SWMU 02-014 for the Industrial and Recreational Scenarios

Note: Data qualifiers are defined in Appendix A.

* n/a = Not applicable.

F-19

 Table F-2.3-2

 EPCs at SWMU 02-014 for the Residential Scenario

COPC	Number of Analyses	Number of Detects	Minimum Concentration	Maximum Concentration	Distribution	EPC	EPC Method		
norganic Chemicals (mg/kg)									
Aluminum	92	92	268	15,000	Nonparametric	5990	95% Chebyshev (Mean, Sd)		
Arsenic	92	92	0.321	2.38	Nonparametric	0.799	95% Student's-t		
Barium	92	92	14.1	96.1	Nonparametric	36.8	95% Student's-t		
Chromium	92	89	1.05 (U)	14.5	Nonparametric	3.32	95% KM (t)		
Copper	92	89	1.02	47.5	Nonparametric	5.81	95% KM (Chebyshev)		
Iron	92	92	285	12,800	Nonparametric	5300	95% Student's-t		
Manganese	92	92	103	305	Normal	204	95% Student's-t		
Mercury	92	90	0.003	0.538	Nonparametric	0.0697	95% KM (Chebyshev)		
Nickel	92	90	0.774	8.48	Nonparametric	2.39	95% KM (t)		
Perchlorate	3	1	0.000813	0.00214 (U)	n/a ^a	0.000813	Maximum detected concentration		
Selenium	92	39	0.119	2.5 (U)	Normal	0.459	95% KM (t)		
Thallium	92	28	0.027	2.5 (U)	Lognormal	0.0807	95% Percentile Bootstrap		
Vanadium	92	92	1.69	19.6	Nonparametric	4.87	95% Student's-t		
Organic Chemicals (mg/kg)			·		-	·			
Anthracene	3	2	0.00727	0.0356 (U)	n/a	0.00903	Maximum detected concentration		
Aroclor-1254	262	24	0.00112 (U)	7.11	Lognormal	0.132	95% Percentile Bootstrap		
Aroclor-1260	262	255	0.00123 (U)	23.9	Nonparametric	2.3	95% KM (Chebyshev)		
Benzo(a)anthracene	3	2	0.032	0.0516	n/a	0.0516	Maximum detected concentration		
Benzo(a)pyrene	3	2	0.0332	0.0551	n/a	0.0551	Maximum detected concentration		
Benzo(b)fluoranthene	3	2	0.0356 (U)	0.0702	n/a	0.0702	Maximum detected concentration		
Benzo(g,h,i)perylene	3	2	0.0222	0.0356 (U)	n/a	0.0271	Maximum detected concentration		
Benzo(k)fluoranthene	3	2	0.0219	0.041	n/a	0.041	Maximum detected concentration		
Chrysene	3	2	0.0356 (U)	0.0598	n/a	0.0598	Maximum detected concentration		

СОРС	Number of Analyses	Number of Detects	Minimum Concentration	Maximum Concentration	Distribution	EPC	EPC Method
Fluoranthene	3	2	0.0356 (U)	0.0984	n/a	0.0984	Maximum detected concentration
Indeno(1,2,3-cd)pyrene	3	2	0.0184	0.0356 (U)	n/a	0.0252	Maximum detected concentration
Phenanthrene	3	2	0.0212	0.0356 (U)	n/a	0.0317	Maximum detected concentration
Pyrene	3	2	0.0356 (U)	0.0771	n/a	0.0771	Maximum detected concentration
TCDD[2,3,7,8-] equivalent	3	3	0.00000418	0.0000066	n/a	0.0000066	Maximum detected concentration
Toluene	2	1	0.000465	0.00107 (U)	n/a	0.000465	Maximum detected concentration
Radionuclides (pCi/g)							
Cesium-134	90	5	-0.07 (U)	0.171 (U)	n/a	0.066 ^b	Maximum detected concentration
Cesium-137	91	8	-0.028 (U)	0.759	n/a	0.759 ^b	Maximum detected concentration
Plutonium-238	92	3	-0.00976 (U)	0.019	n/a	0.019	Maximum detected concentration
Plutonium-239/240	92	18	-0.0471 (U)	0.076	n/a	0.076 ^b	Maximum detected concentration

Note: Data qualifiers are defined in Appendix A.

^a n/a = Not applicable.

^b Value used is the maximum detected concentration because ProUCL recommended value is negative.

СОРС	K _d a (cm³/g)	Water Solubility ^a (g/L)
Aluminum	1500	Insoluble
Arsenic	29	Insoluble
Barium	41	Insoluble
Chromium	850	Insoluble
Copper	35	Insoluble
Iron	25	Insoluble
Manganese	65	Insoluble
Mercury	52	Insoluble
Nickel	65	Insoluble
Perchlorate	na ^b	245
Selenium	5	Insoluble
Thallium	71	Insoluble
Vanadium	1000	Insoluble

Table F-3.2-1Physical and Chemical Properties of Inorganic COPCs at SWMU 02-014

^a Information from <u>http://rais.ornl.gov/cgi-bin/tools/TOX_search</u>.

^b na = Not available.

Table F-3.2-2Physical and Chemical Properties of OrganicCOPCs at SWMU 02-014

COPC	Water Solubility* (mg/L)	Organic Carbon Coefficient K _{oc} * (L/kg)	Log Octanol-Water Partition Coefficient Kow*	Vapor Pressure* (mm Hg at 25°C)
Anthracene	4.34E-02	1.64E+04	4.45E+00	6.53E-06
Aroclor-1254	4.30E-02	1.30E+05	6.50E+00	7.71E-05
Aroclor-1260	1.44E-02	3.50E+05	7.55E+00	4.05E-05
Benzo(a)anthracene	9.40E-03	1.77E+05	5.76+00	2.1E-07
Benzo(a)pyrene	1.62E-03	5.87E+05	6.13E+00	5.49E-09
Benzo(b)fluoranthene	1.50E-03	5.99E+05	5.78E+00	5.00E-07
Benzo(g,h,i)perylene	2.60E-04	1.95E+06	6.63E+00	1.00E-10
Benzo(k)fluoranthene	8.00E-04	5.87E+05	6.11E+00	9.65E-10
Chrysene	2.00E-03	1.80E+05	5.81E+00	6.23E-09
Fluoranthene	2.60E-01	5.54E+04	5.16E+00	9.22E-06
Indeno(1,2,3-cd)pyrene	1.90E-04	1.95E+06	6.70E+00	1.25E-10
Phenanthrene	1.15E+00	1.67E+04	4.46E+00	1.21E-04
Pyrene	1.35E-01	5.43E+04	4.88E+00	4.50E-06
Tetrachlorodibenzodioxin[2,3,7,8-]	2.00E-04	2.49E+05	6.80E+00	1.50E-09
Toluene	5.26E+02	2.34E+02	2.73E+00	2.84E+01

* Information from http://rais.ornl.gov/cgi-bin/tools/TOX_search, unless noted otherwise.

 Table F-3.2-3

 Physical and Chemical Properties of Radionuclide COPCs at SWMU 02-014

COPC	Soil-Water Partition Coefficient, Kd ^a (cm ³ /g)	Water Solubility ^b (g/L)
Cesium-134	1000	Insoluble
Cesium-137	1000	Insoluble
Plutinium-238	4500	Insoluble
Plutonium-239/240	4500	Insoluble

^a Information from <u>http://rais.ornl.gov/cgi-bin/tools/TOX_search</u>.

^b Denotes reference information from <u>http://www.epa.gov/superfund/sites/npl/hrsres/tools/scdm.htm</u>.

Table F-3.3-1 TEFs Used for Calculating TCDD-Equivalent Concentrations

Dioxin and Furan Congeners	TEFs*
TCDD[2,3,7,8-]	1
Pentachlorodibenzodioxin[1,2,3,7,8-]	1
Hexachlorodibenzodioxin[1,2,3,4,7,8-]	0.1
Hexachlorodibenzodioxin[1,2,3,6,7,8-]	0.1
Hexachlorodibenzodioxin[1,2,3,7,8,9-]	0.1
Heptachlorodibenzodioxin[1,2,3,4,6,7,8-]	0.01
Octachlorodibenzodioxin[1,2,3,4,6,7,8,9-]	0.0003
Tetrachlorodibenzofuran[2,3,7,8-]	0.1
Pentachlorodibenzofuran[1,2,3,7,8-]	0.03
Pentachlorodibenzofuran[2,3,4,7,8-]	0.3
Hexachlorodibenzofuran[1,2,3,4,7,8-]	0.1
Hexachlorodibenzofuran[1,2,3,6,7,8-]	0.1
Hexachlorodibenzofuran[1,2,3,7,8,9-]	0.1
Hexachlorodibenzofuran[2,3,4,6,7,8-]	0.1
Heptachlorodibenzofuran[1,2,3,4,6,7,8-]	0.01
Heptachlorodibenzofuran[1,2,3,4,7,8,9-]	0.01
Octachlorodibenzofuran[1,2,3,4,6,7,8,9-]	0.0003

*TEFs from NMED (2017, 602273).

Parameters	Industrial Values	Recreational Values	Residential Values
Target HQ	1	1	1
Target cancer risk	10 ⁻⁵	10 ⁻⁵	10 ⁻⁵
Averaging time (carcinogen/mutagen)	70 yr × 365 days	70 yr × 365 days	70 yr × 365 days
Averaging time (noncarcinogen)	ED × 365 days	Exposure duration × 365 days	ED × 365 days
Skin absorption factor	$SVOC^a = 0.1$	SVOC = 0.1	SVOC = 0.1
	Chemical-specific	Chemical-specific	Chemical-specific
Adherence factor-child	n/a ^b	0.2 mg/cm ²	0.2 mg/cm ²
Body weight-child	n/a	31 kg	15 kg (0–6 yr of age)
Cancer slope factor–oral (chemical-specific)	(mg/kg-day) ⁻¹	(mg/kg-d) ⁻¹	(mg/kg-d) ⁻¹
Inhalation unit risk (chemical-specific)	(µg/m³)	(µg/m³)	(µg/m³)
Exposure frequency	225 days/yr	200 days/yr	350 days/yr
Exposure time	8 hr/day	1 hr/day	24 hr/day
Exposure duration-child	n/a	6 yr (6 to <12 yr of age)	6 yr ^c
Age-adjusted ingestion factor for carcinogens	n/a	n/a	36,750 mg/kg
Age-adjusted ingestion factor for mutagens	n/a	n/a	25,550 mg/kg
Soil ingestion rate-child	n/a	91 mg/d	200 mg/d
Particulate emission factor	6.61 × 10 ⁹ m ³ /kg	6.61 × 10 ⁹ m ³ /kg	6.61 × 10 ⁹ m³/kg
Reference dose-oral (chemical-specific)	(mg/kg-day)	(mg/kg-day)	(mg/kg-day)
Reference dose– inhalation (chemical- specific)	(mg/kg-day)	(mg/kg-day)	(mg/kg-day)
Exposed surface area– child	n/a	4030 cm ²	2690 cm²/day
Age-adjusted skin contact factor for carcinogens	n/a	n/a	112,266 mg/kg
Age-adjusted skin contact factor for mutagens	n/a	n/a	166,833 mg/kg
Volatilization factor for soil (chemical-specific)	(m ³ /kg)	(m ³ /kg)	(m ³ /kg)
Body weight-adult	80 kg	80 kg	80 kg
Exposure duration ^d	25 yr	26 yr (20 yr carcinogens)	30 yr ^e

 Table F-4.1-1

 Exposure Parameters Used to Calculate Chemical SSLs

 for the Industrial, Recreational, and Residential Scenarios

Parameters	Industrial Values	Recreational Values	Residential Values
Adherence factor-adult	0.12 mg/cm ²	0.07 mg/cm ²	0.07 mg/cm ²
Soil ingestion rate-adult	100 mg/day	30 mg/day	100 mg/day
Exposed surface area- adult	3470 cm²/day	6032 cm ²	6032 cm²/day

Table F-4.1-1 (continued)

Note: Parameter values from NMED (2017, 602273) and LANL (2017, 602581).

^a SVOC = Semivolatile organic compound.

^b n/a = Not applicable.

 $^{\rm c}$ The child exposure duration for mutagens is subdivided into 0–2 yr and 2–6 yr.

^d Exposure duration for lifetime resident is 26 yr. For carcinogens, the exposures are combined for child (6 yr) and adult (20 yr).

^e The adult exposure duration for mutagens is subdivided into 6–16 yr and 16–30 yr.

Table F-4.1-2

Parameter Values Used to Calculate Radionuclide SALs for the Residential Scenario

Parameter	Residential, Child	Residential, Adult
Inhalation rate (m ³ /yr)	4712 ^a	7780 ^b
Mass loading (g/m ³)	1.5 × 10 ^{-7¢}	1.5 × 10 ^{-7c}
Outdoor time fraction	0.0926 ^d	0.0934 ^e
Indoor time fraction	0.8656 ^f	0.8648 ^g
Soil ingestion (g/yr)	73 ^h	36.5 ⁱ

^a Calculated as 12.9 m³/day × 365.25 day/yr, where 12.9 m³/day is the mean upper percentile daily inhalation rate of a child (EPA 2011, 208374, Table 6-1).

^b Calculated as 21.3 m³/day × 365.25 day/yr, where 21.3 m³/day is the mean upper percentile daily inhalation rate of an adult from 21 to less than 61 yr old (EPA 2011, 208374, Table 6-1).

^c Calculated as $(1/6.6 \times 10^9 \text{ m}^3/\text{kg}) \times 1000 \text{ g/kg}$, where $6.6 \times 10^9 \text{ m}^3/\text{kg}$ is the particulate emission factor (NMED 2015, 600915).

^d Calculated as (2.32 hr/day × 350 day/yr)/8766 hr/yr, where 2.32 hr/day (139 min) is the largest amount of time spent outdoors for child age groups between 1 to less than 3 mo and 3 to less than 6 yr (EPA 2011, 208374, Table 16-1) and is comparable with the adult time spent outdoors at a residence.

^e Calculated as (2.34 hr/day × 350 day/yr)/8766 hr/yr, where 4.68 hr/day is the average total time spent outdoors for adults age 18 to less than 65 yr in all environments (EPA 2011, 208374, Table 16-1); 50% of this value (2.34 hr/day) was applied to time spent outdoors at a residence and is similar to mean time outdoors at a residence for this age group (EPA 2011, 208374, Table 16-22).

^f Calculated as [(24 hr/day–2.32 hr/day) × 350 day/yr]/8766 hr/yr.

^g Calculated as [(24 hr/day–2.34 hr/day) × 350 day/yr]/8766 hr/yr.

^h The soil ingestion rate compensates for the time-based occupancy factor applied by RESRAD (LANL 2015, 600929 in calculating exposure from the soil ingestion pathway. Calculated as [0.2 g/day × 350 day/yr]/[indoor + outdoor time fractions], where 0.2 g/day is the upper percentile site-related daily child soil ingestion rate (NMED 2015, 600915; EPA 2011, 208374, Table 5-1).

¹ The soil ingestion rate compensates for the time-based occupancy factor applied by RESRAD (LANL 2015, 600929 in calculating exposure from the soil ingestion pathway. Calculated as [0.1 g/day × 350 day/yr]/[indoor + outdoor time fractions], where 0.1 g/day is the site-related daily adult soil ingestion rate (NMED 2015, 600915).

Table F-4.1-3 Parameter Values Used to Calculate Radionuclide SALs for the Industrial and Construction Worker Scenarios

Parameter	Industrial, Adult	Construction Worker, Adult
Inhalation rate (m ³ /yr)	7780 ^a	7780 ^a
Mass loading (g/m ³)	1.51 × 10 ^{-7b}	4.76 × 10 ^{-7c}
Outdoor time fraction	0.2053 ^d	0.2282 ^e
Indoor time fraction	O ^f	0
Soil ingestion (g/yr)	109.6 ^g	362 ^h

^a Calculated as [21.3 m³/day × 365.25 day/yr], where 21.3 m³/day is the upper percentile daily inhalation rate of an adult from 21 to less than 61 yr old (EPA 2011, 208374, Table 6-1).

^b Calculated as $(1/6.6 \times 10^9 \text{ m}^3/\text{kg}) \times 1000 \text{ g/kg}$, where $6.6 \times 10^9 \text{ m}^3/\text{kg}$ is the particulate emission factor (NMED 2015, 600915).

^c Calculated as $(1/2.1 \times 10^6 \text{ m}^3/\text{kg}) \times 1000 \text{ g/kg}$, where $2.1 \times 10^6 \text{ m}^3/\text{kg}$ is the particulate emission factor (NMED 2015, 600915).

^d Calculated as (8 hr/day × 225 day/yr)/8766 hr/yr, where 8 hr/day is an estimate of the average length of the workday and 225 day/yr is the exposure frequency (NMED 2015, 600915).

^e Calculated as (8 hr/day × 250 day/yr)/8766 hr/yr, where 8 hr/day is an estimate of the average length of the workday and 250 day/yr is the exposure frequency (NMED 2015, 600915).

^f The commercial/industrial worker is defined as someone who "spends most of the work day conducting maintenance or manual labor activities outdoors" (NMED 2015, 600915).

^g The soil-ingestion rate compensates for the time-based occupancy factor applied by RESRAD (LANL 2015, 600929) in calculating exposure from the soil-ingestion pathway. Calculated as [0.1 g/day × 225 day/yr]/[indoor + outdoor time fractions], where 0.1 g/day is the site-related daily adult soil-ingestion rate (NMED 2015, 600915).

^h The soil-ingestion rate compensates for the time-based occupancy factor applied by RESRAD (LANL 2015, 600929) in calculating exposure from the soil-ingestion pathway. Calculated as [0.33 g/day × 250 day/yr]/[indoor + outdoor time fractions], where 0.33 g/day is the site-related daily adult soil-ingestion rate (NMED 2015, 600915).

Parameter	Recreational, Child	Recreational, Adult
Inhalation rate (m ³ /yr)	15,250 ^a	19,460 ^b
Mass loading (g/m ³)	1.5 × 10 ^{-7¢}	1.5 × 10 ^{-7¢}
Outdoor time fraction	0.0228 ^d	0.0228 ^d
Indoor-time fraction	0	0
Soil ingestion (g/yr)	797 ^e	244 ^f

Table F-4.1-4

Parameter Values Used to Calculate Radionuclide SALs for the Recreational Scenario

^a Calculated as (0.029 m³/min × 60 min/hr × 24 hr/day × 365.25 day/yr), where 0.029 m³/min is the upper percentile child inhalation rate for moderate activity for 6 to <11 yr old (EPA 2011, 208374, Table 6-2).

^b Calculated as (0.037 m³/min × 60 min/hr × 24 hr/day × 365.25 day/yr), where 0.037 m³/min is the age-weighted upper percentile adult inhalation rate for moderate activity (12 to 35 yr) (EPA 2011, 208374, Table 6-2).

^c Calculated as (1/6.6 × 10⁹ m³/kg) × 1000 g/kg, where 6.6 × 10⁹ m³/kg is the particulate emission factor used for residential and industrial scenarios (NMED 2015, 600915).

^d Calculated as (1 hr/day × 200 day/yr)/8766 hr/yr, where 1 hr/day is the exposure time for a recreational adult or child and 200 day/yr is the exposure frequency (LANL 2015, 600929).

^e The soil ingestion rate is defined to compensate for the time-based occupancy factor applied by RESRAD (LANL 2015, 600929) in calculating exposure from the soil ingestion pathway. One hundred percent of daily soil ingestion is protectively assumed to occur during outdoor activity. Calculated as [(0.2 g/day/2.2 hr/day) × 1 hr/day × 200 day/yr)]/[indoor + outdoor time fractions], where 2.2 hr/day is the mean time spent outdoors per d for a 6 to <11 yr old child (EPA 2011, 208374, Table 16-1), and where 0.2 g/day is the upper-bound child soil ingestion rate (EPA 2011, 208374, Table 5-1; NMED 2015, 600915).</p>

^f Calculated as [(0.1 g/day/3.6 hr/day) × 1 hr/day × 200 day/yr)]/[indoor + outdoor time fractions], where 3.6 hr/day is the mean time spent outdoors per d for an adult (12 to 35 yr) (EPA 2011, 208374, Table 16-1) and where 0.1 g/day is the adult soil ingestion rate (NMED 2015, 600915).

СОРС	EPC (mg/kg)	Industrial SSL* (mg/kg)	Cancer Risk
Aroclor-1254	0.787	11	7.15E-07
Aroclor-1260	3.91	11.1	3.53E-06
Benzo(a)anthracene	0.0516	32.3	1.6E-08
Benzo(a)pyrene	0.0551	23.6	2.33E-08
Benzo(b)fluoranthene	0.0702	32.3	2.17E-08
Benzo(k)fluoranthene	0.041	323	1.27E-09
Chrysene	0.0598	3230	1.85E-10
Indeno(1,2,3-cd)pyrene	0.0252	32.3	7.8E-09
TCDD[2,3,7,8-] equivalent	0.0000066	0.000238	2.77E-07
Total Excess Cancer Risk			5E-06

 Table F-4.2-1

 Industrial Carcinogenic Screening Evaluation for SWMU 02-014

* SSLs from NMED (2017, 602273).

Table F-4.2-2

Industrial Noncarcinogenic Screening Evaluation for SWMU 02-014

COPC	EPC (mg/kg)	Industrial SSL ^a (mg/kg)	HQ
Copper	11.7	51,900	2.25E-04
Mercury	0.284	389	7.3E-04
Selenium	0.66	6490	1.02E-04
Aroclor-1254	0.787	16.4	4.8E-02
Anthracene	0.00903	253,000	3.57E-08
Benzo(g,h,i)perylene	0.0271	25,300 ^b	1.07E-06
Fluoranthene	0.0984	33,700	2.92E-06
Phenanthrene	0.0317	25,300	1.25E-06
Pyrene ^b	0.0771	25,300	3.05E-06
TCDD[2,3,7,8-] equivalent	0.000066	0.000808	8.17E-03
		HI	0.06

^a SSLs from NMED (2017, 602273).

^b Pyrene used as a surrogate based on structural similarity.

СОРС	EPC (pCi/g)	Industrial SAL* (pCi/g)	Dose (mrem/yr)
Plutonium-239/240	0.0414	1200	8.63E-04
		Total Dose	0.0009

 Table F-4.2-3

 Industrial Radionuclide Screening Evaluation for SWMU 02-014

* SALs from LANL (2015, 600929).

Table F-4.2-4 Recreational Carcinogenic Screening Evaluation for SWMU 02-014

СОРС	EPC (mg/kg)	Recreational SSL* (mg/kg)	Cancer Risk	
Aroclor-1254	0.787	10	7.87E-07	
Aroclor-1260	3.91	10.3	3.8E-06	
Benzo(a)anthracene	0.0516	88.8	5.81E-09	
Benzo(a)pyrene	0.0551	8.88	6.2E-08	
Benzo(b)fluoranthene	0.0702	88.8	7.91E-09	
Benzo(k)fluoranthene	0.041	888	4.62E-10	
Chrysene	0.0598	8880	6.73E-11	
Indeno(1,2,3-cd)pyrene	0.0252	88.8	2.84E-09	
TCDD[2,3,7,8-] equivalent	0.0000066	0.000297	2.22E-07	
Total Excess Cancer Risk 5E-06				

* SSLs from LANL (2017, 602581).

	Table F-4	.2-5	
Recreational Noncarc	inogenic Screen	ing Evaluation for SV	VMU 02-014

СОРС	EPC (mg/kg)	Recreational SSL ^a (mg/kg)	HQ
Copper	11.7	24,800	4.72E-04
Mercury	0.284	186	1.49E-03
Selenium	0.66	3100	2.13E-04
Aroclor-1254	0.787	5.53	1.42E-01
Anthracene	0.00903	86,300	1.05E-07
Benzo(a)pyrene	0.0551	86	6.41E-04
Benzo(g,h,i)perylene	0.0271	8630 ^b	3.14E-06
Fluoranthene	0.0984	11,500	8.56E-06
Phenanthrene	0.0317	8630	3.67E-06
Pyrene	0.0771	8630	8.93E-06
TCDD[2,3,7,8-] equivalent	0.0000066	0.00034	1.94E-02
	·	H	0.2

^a SSLs from LANL (2017, 602581).

^b Pyrene used as a surrogate based on structural similarity

Recreational Radio	Table F- nuclide Screeni		SWMU 02-014
	EPC	Recreational SAL*	

COPC	EPC (pCi/g)	Recreational SAL* (pCi/g)	Dose (mrem/yr)
Plutonium-239/240	0.0414	1300	7.96E-04
		Total Dose	0.0008

* SALs from LANL (2015, 600929).

Table F-4.2-7
Residential Carcinogenic Screening Evaluation for SWMU 02-014

COPC	EPC (mg/kg)	Residential SSL* (mg/kg)	Cancer Risk
Arsenic	0.799	7.07	1.13E-06
Chromium	3.32	96.6	3.44E-07
Nickel	2.39	595,000	4.02E-11
Aroclor-1254	0.132	2.43	5.43E-07
Aroclor-1260	2.3	2.43	9.47E-06
Benzo(a)anthracene	0.0516	1.53	3.37E-07
Benzo(a)pyrene	0.0551	1.12	4.92E-07
Benzo(b)fluoranthene	0.0702	1.53	4.59E-07
Benzo(k)fluoranthene	0.041	15.3	2.68E-08
Chrysene	0.0598	153	3.91E-09
Indeno(1,2,3-cd)pyrene	0.0252	1.53	1.65E-07
TCDD[2,3,7,8-] equivalent	0.0000066	0.000049	1.35E-06
	т	otal Excess Cancer Risl	< 1E-05

* SSLs from NMED (2017, 602273).

COPC	EPC (mg/kg)	Residential SSL ^a (mg/kg)	HQ
Aluminum	5990	78,000	7.68E-02
Arsenic	0.799	13	6.15E-02
Barium	36.8	15,600	2.36E-03
Chromium	3.32	45,200	7.35E-05
Copper	5.81	3130	1.86E-03
Iron	5300	54,800	9.67E-02
Manganese	204	10,500	1.94E-02
Mercury	0.0697	23.5	2.97E-03
Nickel	2.39	1560	1.53E-03
Perchlorate	0.000813	54.8	1.48E-05
Selenium	0.459	391	1.17E-03
Thallium	0.0807	0.78	1.03E-01
Vanadium	4.87	394	1.24E-02
Aroclor-1254	0.132	1.14	1.16E-01
Anthracene	0.00903	17,400	5.19E-07
Benzo(g,h,i)perylene	0.0271	1740 ^b	1.56E-05
Fluoranthene	0.0984	2320	4.24E-05
Phenanthrene	0.0317	1740	1.82E-05
Pyrene	0.0771	1740	4.43E-05
TCDD[2,3,7,8-] equivalent	0.0000066	0.000051	1.3E-01
Toluene	0.000465	5220	8.91E-08
		Н	0.6

 Table F-4.2-8

 Residential Noncarcinogenic Screening Evaluation for SWMU 02-014

^a SSLs from NMED (2017, 602273).

^b Pyrene used as a surrogate based on structural similarity.

Table F-4.2-9

Residential Radionuclide Screening Evaluation for SWMU 02-014

COPC	EPC (pCi/g)	Residential SAL* (pCi/g)	Dose (mrem/yr)
Cesium-134	0.066	5	3.30E-01
Cesium-137	0.759	12	1.58E+00
Plutonium-238	0.019	84	5.65E-03
Plutonium-239/240	0.076	79	2.41E-02
		Total Dose	2

* SALs from LANL (2015, 600929).

COPC	EPC (mg/kg)	Construction Worker SSL (mg/kg)ª	HQ
Aluminum	5990	41,400	1.45E-01
Arsenic	0.799	41.2	1.94E-02
Barium	36.8	4390	8.38E-03
Chromium	3.32	134	2.48E-02
Copper	5.81	14,200	4.09E-04
Iron	5300	248,000	2.14E-02
Manganese	204	464	4.40E-01
Mercury	0.0696	77.1	9.03E-04
Nickel	2.39	753	3.17E-03
Perchlorate	0.000813	248	3.28E-06
Selenium	0.459	1750	2.62E-04
Thallium	0.0807	3.54	2.28E-02
Vanadium	4.87	614	7.93E-03
Aroclor-1254	0.132	4.91	2.69E-02
Anthracene	0.00903	75,300	1.20E-07
Benzo(g,h,i)perylene	0.0271	7530 ^b	3.60E-06
Fluoranthene	0.0984	10,000	9.84E-06
Phenanthrene	0.0317	7530	4.21E-06
Pyrene	0.0771	7530	1.02E-05
TCDD[2,3,7,8-] equivalent	0.0000066	0.00023	2.87E-02
Toluene	0.000465	14,000	3.32E-08
		HI	0.8

 Table F-4.2-10

 Construction Worker Noncarcinogenic Screening Evaluation for SWMU 02-014

^a SSLs from NMED (2017, 602273).

^b Pyrene used as a surrogate based on structural similarity.

 Table F-4.4-1

 Essential Nutrient Screening Assessment

			Maximum Concentration	SSL	
SWMU / AOC	Scenario	COPC	(mg/kg)	(mg/kg)*	Ratio
02-014	Residential	Magnesium	1710	20,900,000	8.2E-05

* SSLs from NMED (2017, 602273).

Attachment F-1

Dioxin and Furan Toxicity Equivalency Factor Calculations (on CD included with this document)

Attachment F-2

ProUCL Files (on CD included with this document)

Appendix G

Box Plots and Statistical Results

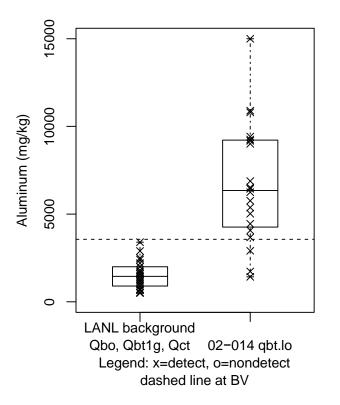


Figure G-1 Box plot for aluminum in tuff at SWMU 02-014

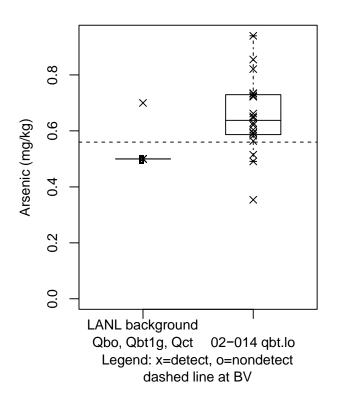


Figure G-2 Box plot for arsenic in tuff at SWMU 02-014

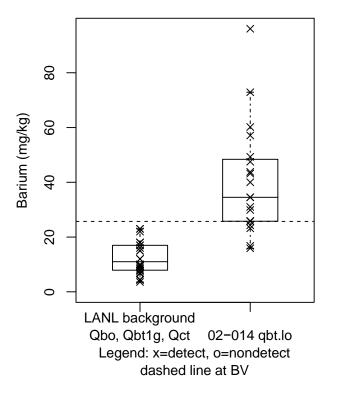


Figure G-3 Box plot for barium in tuff at SWMU 02-014

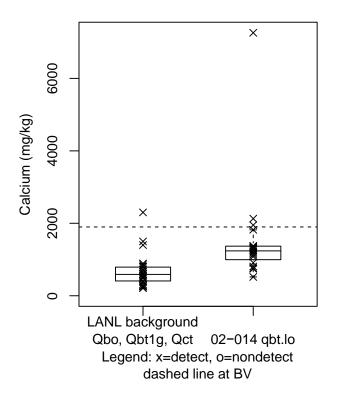


Figure G-4 Box plot for calcium in tuff at SWMU 02-014

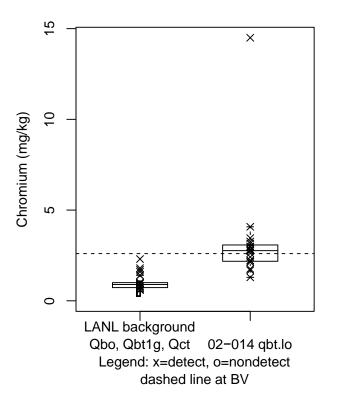


Figure G-5 Box plot for chromium in tuff at SWMU 02-014

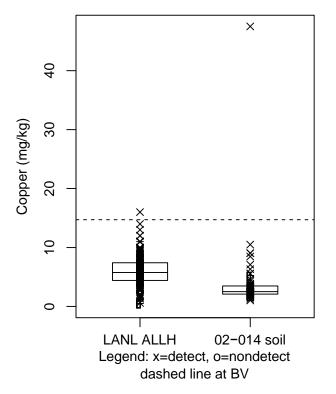


Figure G-6 Box plot for copper in soil at SWMU 02-014

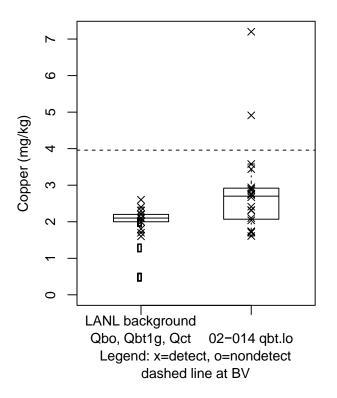


Figure G-7 Box plot for copper in tuff at SWMU 02-014

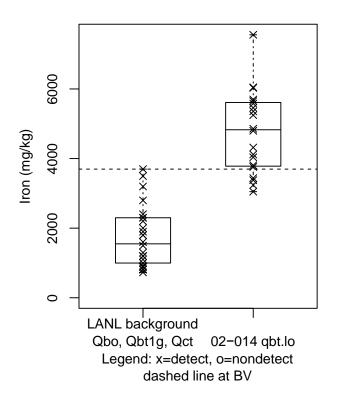


Figure G-8 Box plot for iron in tuff at SWMU 02-014

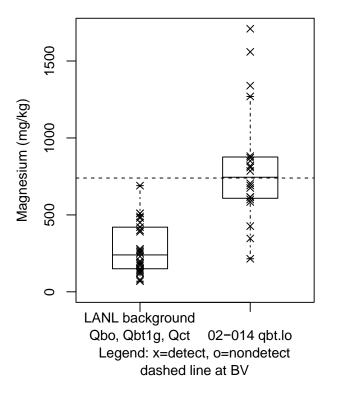


Figure G-9 Box plot for magnesium in tuff at SWMU 02-014

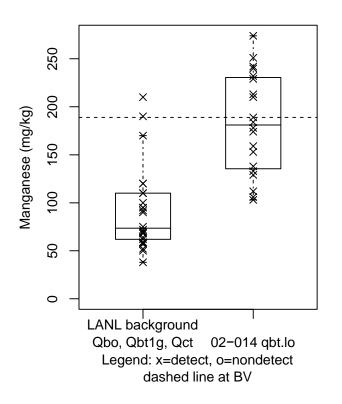


Figure G-10 Box plot for manganese in tuff at SWMU 02-014

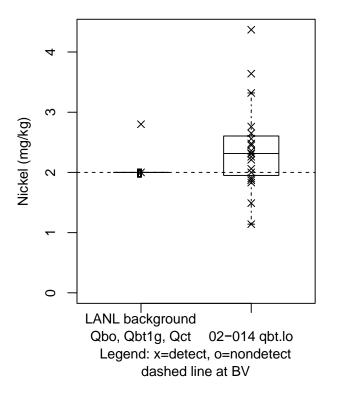


Figure G-11 Box plot for nickel in tuff at SWMU 02-014

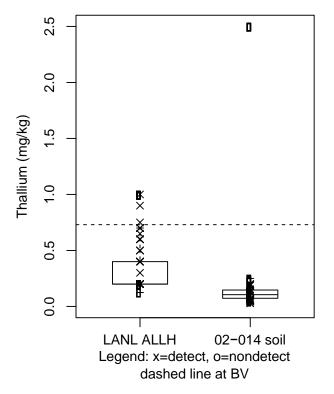


Figure G-12 Box plot for thallium in soil at SWMU 02-014

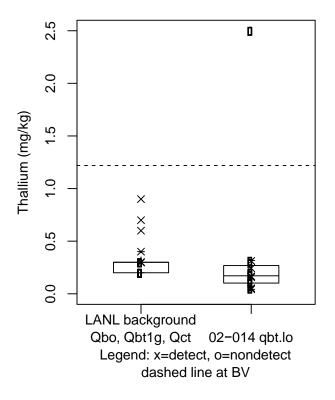


Figure G-13 Box plot for thallium in tuff at SWMU 02-014

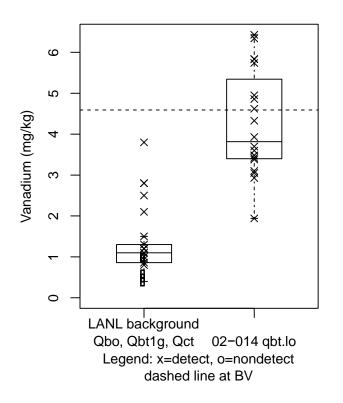


Figure G-14 Box plot for vanadium in tuff at SWMU 02-014

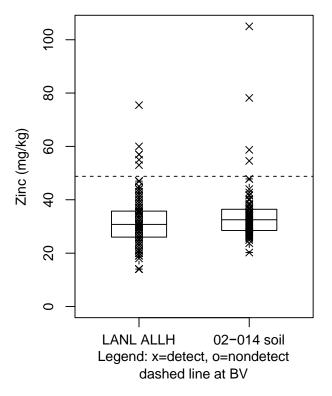


Figure G-15 Box plot for zinc in soil at SWMU 02-014

Analyte	Gehan Test p-Value	Quantile Test p-Value	Slippage p-Value	COPC?
Aluminum	<0.001	<0.001	n/a*	Yes
Arsenic	n/a	0.00374	0.00171	Yes
Barium	<0.001	<0.001	n/a	Yes
Calcium	<0.001	0.131	0.444	No
Chromium	<0.001	<0.001	n/a	Yes
Copper	<0.001	<0.001	n/a	Yes
Iron	<0.001	<0.001	n/a	Yes
Magnesium	<0.001	<0.001	n/a	Yes
Manganese	<0.001	<0.001	n/a	Yes
Nickel	n/a	0.00312	0.0751	Yes
Thallium	n/a	1.00	0.184	No
Vanadium	<0.001	<0.001	n/a	Yes

 Table G-1

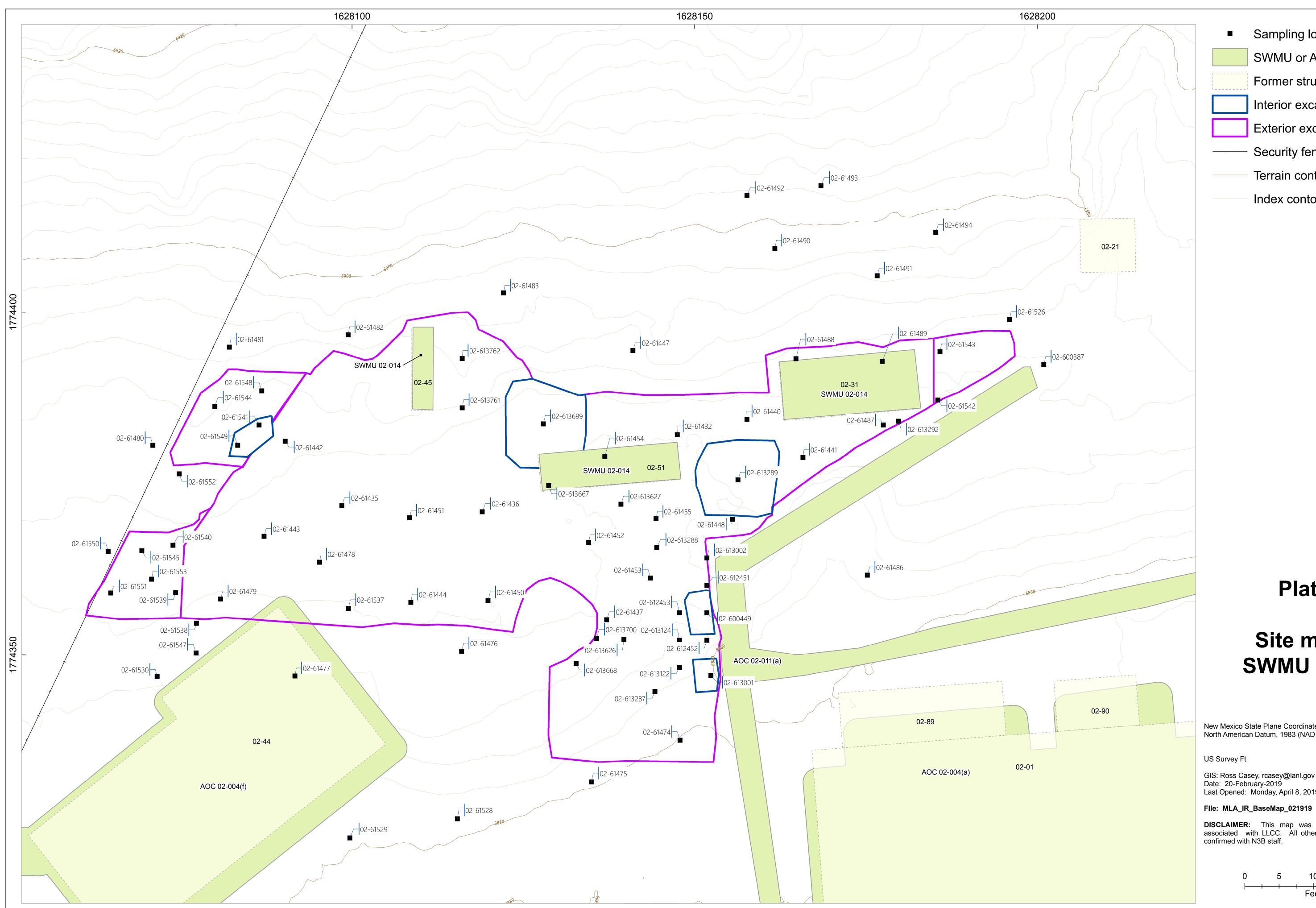
 Results for Statistical Tests for Inorganic Chemicals in Tuff at SWMU 02-014

*n/a = Not applicable.

Table G-2Results for Statistical Tests for Inorganic Chemicals in Soil at SWMU 02-014

Analyte	Gehan Test p-Value	Quantile Test p-Value	Slippage p-Value	COPC?
Copper	1.00	1.00	n/a	No
Thallium	n/a	n/a	0.0239	Yes
Zinc	0.0143	0.212	0.0828	No

*n/a = Not applicable.



Sampling location SWMU or AOC Former structures Interior excavation boundary Exterior excavation boundary Security fence Terrain contour, 20 ft interval

Index contour, 2 ft interval

Plate 1

Site map of **SWMU 02-014**

New Mexico State Plane Coordinate System Central Zone (3002) North American Datum, 1983 (NAD 83)

US Survey Ft

GIS: Ross Casey, rcasey@lanl.gov Date: 20-February-2019 Last Opened: Monday, April 8, 2019



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