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## Justification for Class III Permit Modification March 2005 SWMU 136 Operable Unit 1303 Building 907 Septic Systems at Techincal Area II

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#### This work supported by the United States Department of Energy under contract DE-AC04-94AL85000.

# SWMU 136 Building 907 Septic System and HE Drain System (TA-II)





Environmental Restoration Project

#### Site History

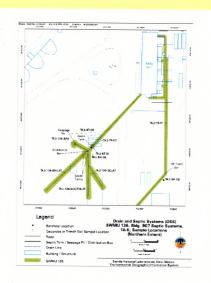
- SWMU 136, Building 907 Septic System and HE Drain System, is located in the northern portion of TA-II, and covers approximately 0.46 acres.
- Building 907 was constructed in 1948 and used for a variety of purposes including nuclear weapon assembly, HE development and testing, film developing, and laser research.
- The building had one septic system and one HE waste water drain system.
- The septic system discharged to three septic tanks, a seepage pit, and a drainfield southwest of the building. The septic system was connected to the COA sanitary sewer system in 1992.
- The HE drain system, including an HE particulate catch box, extended south then southwest of the building, passing under Building 906, and discharged into a 200-ft open ditch. When weapon assembly operations were discontinued in the late 1950s, HE residues were no longer washed into the Building 907 floor drains.
- Building 907 was decontaminated and demolished in late 2002.
- The HE catch box reportedly partially collapsed in 1991 or 1992. It was sampled in December 2001 and in June 2003 was abandoned by backfilling.

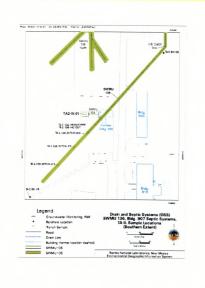
#### Depth to Groundwater

 The regional aquifer is approximately 545 ft bgs, and the perched aquifer (not a source of drinking water) is approximately 300 ft bgs.

#### Constituents of Concern

- VOCs
- SVOCs
- PCBsHE compounds
- Metals
- Radionuclides





#### Investigations

- In 1987, all septic tanks and drainfields throughout TA-II, III, and V were grouped together as part of the RFA. SWMU 136 was listed because industrial wastes could have been discharged to the sanitary sewer system.
- In 1991, a study was conducted for SWMU 136 that included background information reviews and personnel interviews.
- In October and November 1993, soil samples were collected from a trench excavated across the sanitary system drain line and a trench excavated across the HE drain line system.
- In December 1993 the area surrounding Building 907 was included in a geophysical (electromagnetic) survey. No anomalies related to buried material, other than underground utilities, were identified.
- In November-December 1993, and January-February 1994, passive soil-vapor surveys were conducted in TA-II. The majority of compounds detected were chlorinated solvents and petroleum hydrocarbons. The highest detections were in the southern part of TA-II.
- In March and November 1994, soil samples were collected from a borehole drilled near the first septic tank, from a borehole drilled near the HE catch box, and from a borehole drilled near the end of the HE drain line outfall.
- In August 1995, soil samples were collected from two boreholes drilled adjacent to the first septic tank.
- In November and December 1996, three boreholes were drilled at TA-II and sampled for soil vapor to approximately 100 ft bgs. Two of the borings were completed as active soil-vapor monitoring wells. The long-term monitoring results indicate an apparent widespread presence of VOCs in soil vapor; the concentrations were low and remained steady with no apparent periodicity.
- In early 2000, portions of the septic system were excavated with a backhoe to confirm the positions and dimensions of the septic tanks, seepage pit, drain line laterals, and HE outfall drain pipe.
- In March and August 2000, a backhoe was used to collect soil samples from three locations along the HE outfall ditch.
- In August 2000, soil samples were collected from boreholes drilled beneath the seepage pit and drainfield laterals.
- In December 2001, samples from the partially collapsed HE catch box contents were collected. Because
  the material in the catch box appeared to be predominantly native soil, removal was not required by
  NMED.

#### Summary of Data Used for NFA Justification

- All confirmatory soil sample analytical results, with the exception of the HE catch box samples, were used
  for characterizing the site, for performing the risk assessment, and for justification for the NFA proposal.
- VOCs, SVOCs, and one HE compound were detected in these samples. Numerous metals were detected above background values. Two radionuclides and tritium were detected above background activities, and one radionuclide had an MDA above background activities. No PCBs or cyanide were detected in these soil samples.

#### Recommended Future Land Use

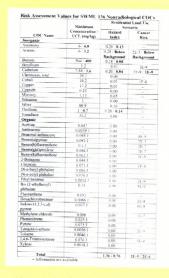
· Industrial land use was established for this site.



This partially excavated drainfield lateral piping at SWMU 136. The pipe is 4 in. diameter vitrified clay with unsealed joints between pipe sections to permit drainage. Drain lines were typically buried in gravel-filled trenches. Drain line trench depths varied between 8 and 10 ft below grade at this site.

#### Results of Risk Analysis

- Risk assessment results for the residential scenario are calculated per NMED risk assessment guidance as presented in "Supplemental Risk Document Supporting Class 3 Permit Modification Process" (SNL October 2003)
- Because COCs were present in concentrations greater than background-screening levels or because
  constituents were present that did not have background-screening numbers, it was necessary to perform
  a risk assessment for the site. The risk assessment analysis evaluated the potential for adverse health
  effects for the residential land-use scenario.
- The maximum concentration for lead was 58.4 mg/kg. NMED guidance for lead screening concentrations for construction and industrial land-use scenarios are 750 and 1500 mg/kg respectively. The EPA screening guidance value for a residential land-use scenario is 400 mg/kg. The maximum concentration for lead at this site is less than all the screening values; therefore, lead was eliminated from further consideration in the human health risk assessment.
- The residential HI (1.36) is above the NMED guidance. The total estimated excess cancer risk for the residential land-use scenario, 2E-5, is above the NMED guideline. However using the UCL of the average concentrations for the main contributors to excess cancer risk [antimony, arsenic (below background), barium, cadmium, and thallium] reduces the total HI to 0.76 and the total estimated excess cancer risk to 2E-6. Thus, by using realistic concentrations in the risk calculations that more accurately depict actual site conditions, the total risk calculations are below NMED guidelines.
- The residential land-use scenario incremental TEDE was 3.1E-1 mrem/yr, which is below the EPA numerical guideline of 75 mrem/yr. Therefore, SWMU 136 is eligible for unrestricted radiological release.
- Using the SNL predictive ecological risk assessment methodology, the ecological risk for SWMU 136 is predicted to be low.
- In conclusion, human health and ecological risks are acceptable per NMED guidance. Thus, SWMU 136 is proposed for CAC without institutional controls.



#### For More Information Contact

U.S. Department of Energy Sandia Site Office Environmental Restoration Mr. John Gould Telephone (505) 845-6089 Sandia National Laboratories Environmental Restoration Project Task Leader: Brenda Langkopf Telephone (505) 284-3272



## Sandia National Laboratories

## Justification for Class III Permit Modification March 2005

SWMU 136 Operable Unit 1303 Building 907 Septic Systems at Technical Area II

NFA Originally Submitted August 1995
Comment Response October 1996
NOD Response January 2000
RSI Response June 2004
Soil Vapor Sampling June 2004

Environmental Restoration Project



United States Department of Energy Sandia Site Office



#### Department of Energy

Albuquerque Operations Office Kirtland Area Office P. O. Box 5400 Albuquerque, New Mexico 87185-5400

AUG 2 8 1995

#### CERTIFIED MAIL - RETURN RECEIPT REQUESTED

Mr. David Neleigh, Chief
New Mexico and Federal Facilities Section
RCRA Permits Branch
U. S. Environmental Protection Agency, Region VI
1445 Ross Avenue, Suite 1200
Dallas, TX 75202-2733

Dear Mr. Neleigh:

Enclosed are copies of the second set of No Further Action (NFA) proposals for 23 solid waste management units (SWMUs) from the Resource Conservation and Recovery Act (RCRA) Hazardous and Solid Waste Amendments (HSWA) Final Permit for Sandia National Laboratories/New Mexico (SNL/NM), ID No. NM5890110518.

Copies of these proposals are also being submitted for comment to the New Mexico Environment Department (NMED), Hazardous and Radioactive Materials Bureau. The Class 3 permit modification process will be initiated after regulatory comments are addressed.

If you have any questions, please contact John Gould at (505) 845-6089 or Mark Jackson at (505) 845-6288.

Sincerely,

Michael J. Zamorski Acting Area Manager

**Enclosures** 

cc w/enclosures: T. Trujillo, AL, ERD L. Aker, AIP (2 copies) W. Cox, SNL, MS 1147 cc w/o enclosures:

- M. Jackson, KAO
- J. Johnsen, KAO-AiP
- C. Soden, AL, EPD
- N. Morlock, EPA, Region VI
- T. Roybal, SNL, MS 1147
- M. Davis, SNL, MS 1147
- T. Vandenberg, SNL, MS 0141
- E. Krauss, SNL, MS 0141



## Sandia National Laboratories / New Mexico

## PROPOSAL FOR NO FURTHER ACTION ENVIRONMENTAL RESTORATION PROJECT SITE 136, BUILDING 907 SEPTIC SYSTEM OPERABLE UNIT 1303

**June 1995** 

Environmental Restoration Project



United States Department of Energy Albuquerque Operations Office

## PROPOSAL FOR NO FURTHER ACTION

Site 136, Building 907 Septic System Operable Unit 1303

SANDIA NATIONAL LABORATORIES/NEW MEXICO

#### 1. Introduction

#### 1.1 ER Site Identification Number and Name

Sandia National Laboratories/New Mexico (SNL/NM) is proposing an administrative no further action (NFA) decision based on confirmatory sampling for Environmental Restoration (ER) Site 136, Building 907 Septic System, Operable Unit (OU) 1303. The Resource Conservation and Recovery Act (RCRA) Facility Assessment (RFA) grouped all septic tanks and leachfields found throughout Technical Area (TA) II, III, and V together. The Building 907 Septic System and all other associated systems were given RFA number 79 [Environmental Protection Agency (EPA 1987)]. ER Site 136 was identified as the Building 907 Septic System in the Hazardous and Solid Waste Amendment (HSWA) Module IV (EPA 1993) of the SNL/NM RCRA Hazardous Waste Management Facility Permit (NM5890110518) (EPA 1992).

#### 1.2 SNL/NM Administrative NFA Based on Confirmatory Sampling Process

This proposal for a determination of an administrative NFA decision based on confirmatory sampling has been prepared using the criteria presented in Section 4.5.3. of the SNL/NM Program Implementation Plan (PIP) (SNL/NM 1995). Specifically, this proposal "contains information demonstrating that there are no releases of hazardous waste (including hazardous constituents) from solid waste management units (SWMU) at the facility that may pose a threat to human health or the environment" [as proposed in the Code of Federal Regulations (CFR) Section 40 Part 264.51(a)(2)] (EPA 1990). The HSWA Module IV contains the same requirements for an NFA demonstration:

Based on the results of the RFI (RCRA Facility Investigation) and other relevant information, the Permittee may submit an application to the Administrative Authority for a Class III permit modification under 40 CFR 270.42(c) to terminate the RFI/CMS (corrective measures study) process for a specific unit. This permit modification application must contain information demonstrating that there are no releases of hazardous waste including hazardous constituents from a particular SWMU at the facility that pose threats to human health and/or the environment, as well as additional information required in 40 CFR 270.42(c) (EPA 1993).

In requesting an administrative NFA decision based on confirmatory sampling for ER Site 136, Building 907 Septic System, this proposal is using existing administrative/archival information to satisfy permit requirements. This unit is eligible for an administrative with confirmatory sampling NFA proposal based on one or more of the following criteria taken from the RCRA Facility Assessment Guidance (EPA 1986):

Criterion A: The unit has never contained constituents of concern

Criterion B: The unit has design and/or operating characteristics that effectively prevent releases to the environment

Criterion C: The unit clearly has not released hazardous waste or constituents into the environment

Specifically, ER Site 136 is being proposed for an administrative NFA decision based on confirmatory sampling because the SWMU clearly has not released hazardous waste or constituents into the environment (Criterion C).

#### 1.3 Local Setting

SNL/NM occupies 2,829 acres of land owned by the Department of Energy (DOE), with an additional 14,920 acres of land provided by land-use permits with Kirtland Air Force Base (KAFB), the United States Forest Service (USFS), the State of New Mexico, and the Isleta Indian Reservation. SNL/NM has been involved in nuclear weapons research, component development, assembly, testing, and other nuclear activities since 1945.

ER Site 136 (Figure 1) is owned by DOE. The site is located in the central to north-central portion of TA-II. TA-II, one of five technical areas within SNL/NM, is diamond-shaped, approximately 1,450 feet on a side, and encompasses 45 acres. The center of TA-II is approximately 3,000 feet south of TA-I, the location for most administrative and research activities. TA-II abuts TA-IV to the south. TA-II is surrounded by a 10-foot high chain link fence, with a security guarded gate at the west corner. In earlier years, guard towers were located at each corner; now only the west entrance tower remains. TA-II currently contains 22 buildings, 27 high explosives (HE) bunkers of various sizes, and four mobile offices (MOs).

TA-II lies west of the basin-bounding fault complex and northwest of the Tijeras Arroyo fault, which are the two main structural features of the Albuquerque Basin. The geologic materials consist of thick alluvial sediments which overlie deep bedrock. An alluvial fan and piedmont colluvium overlie Santa Fe Group strata. The Santa Fe deposits are estimated to be approximately 3,000 feet thick beneath TA-II (Hawley and Haase 1992). Detailed descriptions of the regional geology are in the PIP and in the annual Site-Wide Hydrogeologic Characterization Project (SWHCP) 1993 Annual Report (SNL/NM 1993).

Previous SWHCP soil surveys and 1993 surficial mapping activities provide general soil characteristics for TA-II. Soil associated with the escarpments of the Tijeras Arroyo is poorly developed, such as the Bluepoint-Kokan Association (Hacker 1977). Areas underlain by this soil series, however, locally contain well-developed calcic horizons, which are the remnants of the Tijeras, Wink, and Madurez soil originally developed on older surficial deposits. The Bluepoint-Kokan soil reflects erosion of older soil and, therefore, is characterized by discontinuous soil horizons. The heterogeneity would be expected to strongly influence the location and rates of infiltration and geochemical interactions between soil and percolating water (SNL/NM 1993). TA-II is characterized as having an average surface soil permeability of approximately 0.1 inch per hour (SNL/NM 1993).

No perennial surface-water bodies are present within TA-II or in the immediate vicinity of the area. However, a large ephemeral surface drainage, the Tijeras Arroyo, is located directly southeast of TA-II. TA-II is located outside the 100- and 500-year floodplains of the Tijeras Arroyo.

Depth of regional ground water in the vicinity of TA-II is approximately 540 feet, with shallower water-bearing units present at approximately 305 to 315 feet. In the shallower saturated zones, the ground water gradient is to the south-southeast at 0.016 ft/ft. No water supply wells are present within TA-II.

#### 2. History of the SWMU

#### 2.1 Sources of Supporting Information

In preparation to request an administrative NFA decision based on confirmatory sampling for ER site 136, a background study was conducted to collect available and relevant site information. Background information sources included existing records and reports of site activity. In addition, interviews were conducted with SNL/NM staff and contractors familiar with site operational history. The study was completely documented and has provided traceable references which sustain the integrity of this proposal.

The following information sources were available for use in the evaluation of ER site 136:

- Interviews were combined and summarized in three reports (Anonymous no date; Haines, Kelly, and Cochran 1991; and Byrd 1991).
- Radiation and organic vapor surveys were performed over the sanitary and the HE drain trench areas. Soil samples were collected from three horizons from two trench sample locations.
- The Site-Wide Hydrogeologic Characterization Project 1993 Annual Report (SNL/NM 1993).
- Sequential historical aerial photographs from 1951 to 1992 for the specifically prescribed area of ER Site 136 (Ebert 1994).
- A passive soil vapor survey (SVS) investigation was conducted in the vicinity of the Building 907 Septic System [Northeast Research Institute (NERI 1994)].
- One borehole was drilled and soil samples were collected near the outfall of the HE drainline for Building 907. Two additional boreholes were drilled and soil samples were collected; one along the sanitary line near the seepage pit and the second along the HE drain trench near the HE catch box.

Utilizing this information, a brief history of ER Site 136 and a discussion of all relevant evidence regarding past waste practices and releases at the site have been prepared and are presented in this proposal for an administrative NFA decision based on confirmatory sampling.

#### 2.2 Previous Audits, Inspections, and Findings

The RCRA RFA grouped all septic tanks and leachfields found throughout TA-II, TA-III, and TA-V together. The Building 907 Septic System and all associated systems were given RFA number 79 (EPA 1987). The Building 907 Septic System was listed as an SWMU because sanitary wastes were not separated from industrial wastes; therefore, hazardous wastes may have been discharged to septic tanks and leachfields. Site 136 includes both the sanitary and the HE septic lines.

The 1987 RCRA RFA is summarized below.

The wastes managed at this location include sanitary and industrial wastes including trichloroethylene (TCE), toluene, and methanol. Septic tank contents were discharged to leachfields. Release controls do not appear to have been present. There is no history of releases at this location. The potential for air contamination resulting from ER Site 136 is low because the wastes were discharged to underground septic tanks then to leachfields. The potential for soil contamination is high because the wastes were released to leachfields. The potential for surface water and ground water contamination was not determined in the RFA. Because sanitary wastes were disposed in tanks and leached through surface soils, there is a potential for subsurface gas generation.

The 1987 Comprehensive Environmental Assessment and Response Program (CEARP) Phase 1 (DOE 1987) grouped the septic systems for Building 904 and 907 together in a brief description. The 1987 CEARP Phase 1 is summarized below.

Industrial and sanitary wastes from TA-II buildings are discharged through a system of septic tanks and drainfields. Buildings 904 and 907 (TA-II) were formerly used for weapon assembly and high-explosives development and processing work. The buildings' drain systems may be contaminated with high explosives and solvents from process washdown. The floor drains in these buildings emptied to the septic system through a trap containing burlap bags. High explosives retained in the burlap were burned in a pit. There was a Comprehensive Environmental Response, Compensation and Liability Act (CERCLA) finding of Uncertain for Federal Facility Site Discovery and Identification Findings (FFSDIF), Preliminary Assessment (PA), and Pollutant Standards Index (PSI); insufficient information is available to calculate a hazard ranking system (HRS) migration mode score.

#### 2.3 Historical Operations

Building 907, currently the Explosives Application Facility, was constructed in 1948 and is located in the northern portion of TA-II (Figure 1). The building was initially designed for

the final assembly of nuclear weapons. An earthen berm on the south side of the building was originally built to isolate the building from surrounding areas from potential accidental explosions.

During the early 1950s, 8 to 10 employees worked three eight-hour shifts per day assembling and packaging nuclear weapons. Compacted HE may have been occasionally dropped and/or broken during assembly. Floor sweeping was not allowed in Building 907 because it could potentially generate static electricity. Therefore, explosives materials and debris were flushed down the floor drain with large quantities of water and discharged into the Building 907 HE system. A mixture of kerosene and water may have been used to clean the floor. The floor drains appear to have been designed to discharge large volumes of water, and may have received significant amounts of water containing kerosene and small quantities of explosive materials.

In the mid-1950s, bomb assembly operations were discontinued, and the building was converted into an explosives testing and development facility. These operations involved mixing small quantities of explosives in separate handling areas. According to interviews, explosives residues were not flushed down the floor drains, but small amounts of cleaning solvents may have been discharged to the sanitary system. Work areas and components were typically cleaned with solvents, including isopropyl alcohol, toluene, petroleum distillates, nitromethane, acetone, and methanol. Carbon tetrachloride was used to clean the work tables and components. Long-time employees indicated that they sometimes experienced the sensation of drunkenness on the carbon tetrachloride vapors. It was used liberally (more than 2 to 3 gallons a month) until about 1951, when it was replaced by TCE because the carbon tetrachloride caused headaches.

Between 1980 and 1985, Building 907 contained a darkroom for processing X-ray and high-speed black-and-white film. During this time, developing solutions and rinse water were discharged through a sink and into the Building 907 sanitary system. After 1985, only rinse water was discharged to the sanitary system, which was shut down in 1992.

From 1982 to 1992, cooling water from several lasers also was discharged to the Building 907 sanitary system. No contaminants are known to have been discharged with the cooling water.

Aerial photographs show vegetation, 20 to 30 feet in width, in the areas above the sanitary and HE drainlines. Between 1948 and 1992, an estimated 13 million gallons of effluent may have been discharged to the septic system.

To summarize, based on historical records and interviews, contaminants that could have been discharged into the Building 907 Septic System include:

- Explosive materials such as HE compounds (e.g., cyclo-1,3,5-trimethylene-2,4,6-trinitramine (RDX), octahydro-1,3,5,7-tetranitro-1,3,5,7-tetrazocine (HMX), nitroguanidine, and nitrocellulose), plastic-bonded explosives (PBX), nitromethane, and Baratol
- Cleaning solvents such as methanol, acetone, isopropyl alcohol, toluene,

petroleum distillates, TCE, and carbon tetrachloride

- Photochemicals, including silver, cadmium, chromium, and cyanide
- Metals, including arsenic, barium, copper, lead, manganese, mercury, selenium, silver, zinc

#### 3. Evaluation of Relevant Evidence

#### 3.1 Unit Characteristics

Building 907 was constructed with two drain systems: a sanitary system and an HE system (Figure 2). The systems encompass 800 feet of sanitary and HE drainlines.

The HE system was designed and constructed to drain the assembly bays, and passes through a catch box about 200 feet south of Building 907. The catch box was designed to collect HE particulates by mechanical filtration of the discharge water. Sometime in 1991 or 1992, the soil above the catch box collapsed. In 1992, a corrugated metal cover was constructed over the catch box to divert rain water. The HE system extends an additional 600 feet to the southwest from the catch box, passing under Building 906. The discharge drains into a 200-foot-long, open ditch west-southwest of Building 906.

The sanitary septic system extends about 250 feet southwest from the southwest corner of Building 907, and consists of a 900-gallon septic tank, an 8.5-foot deep by 5-foot wide seepage pit, and a leachfield consisting of about 150 feet of perforated pipe in trenches. The floor drains in the building reportedly may have been cross-connected between the sanitary system and the HE system.

#### 3.2 Operating Practices

To reduce the potential for HE introduction into the environment, HE collected in the catch box were removed periodically and were burned at the Explosive Burn Pit area, ER Site 114.

#### 3.3 Presence or Absence of Visual Evidence

Because ER Site 136 is located underground, no visual evidence was obtained to determine that contamination has not occurred from this site to the environment.

#### 3.4 Results of Previous Sampling/Surveys

On December 5 and 6, 1991, surface radiation and organic vapor surveys were performed over the site areas. The organic vapor survey was performed using an HNu photoionization detector (PID) held at waist height and field calibrated to read equivalent benzene concentration. No readings were detected above background levels. The radiation survey was performed using a Bicron 2000 gamma detector at waist height, and an ASP-1 survey meter with an HP-260 G-M pancake probe at ground level for beta-gamma detection. The

objectives of the survey were to determine both the general area gamma radiation levels and the presence of surface contamination. The radiation levels varied from approximately 0.02 to 0.03 milliroentgens per hour (mR/hr). Surface readings varied from 80 to 140 counts per minute (cpm), which are within the range of area background. Several swipe surveys were also performed on the ground and personnel were monitored for contamination. No elevated activities were identified.

Changes in vegetation that appeared to be related to septic line discharge were identified through the interpretation and digital mapping of vegetation from sequential historical aerial photographs for the specifically prescribed area of ER Site 136 (Ebert 1994). Vegetation was identified along both the sanitary and HE septic lines between 1964 and 1992.

Between October 28 and November 1, 1993, soil samples were collected from two trenches excavated across the HE drain trench and sanitary sewerlines (Trenches 1 and 2, Figure 2). The soil samples were collected from three depths at each trench: at the surface, immediately above, and below the pipe. Trench soil samples were analyzed for VOCs, SVOCs, HE compounds, tritium, metals, and radioisotopes. No constituents of concern were reported over the Sandia site-wide upper tolerance limit (UTL) background concentration and/or the proposed RCRA Subpart S action level (Table 1). With the exception of tritium, identified in Trench 2 at 360 and 260 pCi/l, at 0.5 and 5.0 feet depth, respectively, no anomalous activities were identified when compared to natural background activities.

The raw data, along with QA/QC documentation, are readily available and can be viewed in the Environmental Operations (EO) Records Center. A summary of the trench soil data is presented in Table 1 and includes the maximum concentration of the constituents of concern, the site-wide UTL background concentrations (IT 1994), and the proposed RCRA Subpart S action levels, as appropriate and available.

#### 3.5 Assessment of Gaps in Information

Identified data gaps required that a more comprehensive analysis of volatile organic compounds (VOCs) and semivolatile organic compounds (SVOCs) along the drainlines be accomplished by a soil gas survey. This more comprehensive investigation was needed to locate and qualify the nature and extent of potential organic contamination. Location-specific soil sampling and analysis for organics, inorganics, and radionuclides were deemed necessary at the outfalls, seepage pits, and HE catch box to provide supplementary confirmation of the soil gas survey results and to quantify contamination at potential source areas.

#### 3.6 Confirmatory Sampling

Two investigations were determined to be necessary to fill the data gaps (see Section 3.5). The results for the investigations are presented below. The Sampling and Analysis Plans for the two borehole investigations are located in Appendix A. The raw data, along with QA/QC documentation, are readily available and can be viewed in the EO Records Center. A

summary of the borehole data is presented in Table 2 and includes the maximum concentrations of the constituents of concern, the site-wide background UTLs, and the proposed RCRA Subpart S action levels, as appropriate and available.

Between November 11 and December 4, 1993, a passive SVS investigation was conducted in the vicinity of the Building 907 Septic System. Except for one isolated detection of benzene, toluene, ethylbenzene, total xylenes (BTEX) at the Building 907 seepage pit and one isolated detection of TCE south of the seepage pit, no other VOCs or SVOCs were identified in soil vapor collected during the SVS investigation. The BTEX and TCE levels are considered extremely low and probably do not reflect contamination in the vicinity of the seepage pit and leachfield areas (NERI 1994). Further explanation is contained in the 1994 NERI report which has been included as a separate report with the submittal of this NFA proposal.

On March 10, 1994, Borehole TA2-BH-05 was drilled near the outfall of the HE drainline for Building 907 (Figure 2). This borehole was located based on results of the passive SVS. The borehole was drilled to a total depth of 30.75 feet BGL. Soil samples were collected at depths of 4, 9, 13.5, 18.75, and 27.2 feet BGL. The samples were analyzed for tritium, metals, HE compounds, radioisotopes, total uranium, VOCs, and SVOCs. No VOCs were detected in any of the soil samples except for acetone and methylene chloride. Acetone and methylene chloride are common laboratory contaminants and were both reported significantly below the proposed RCRA Subpart S action level. The common laboratory contaminant bis (2-ethylhexyl) phthalate was the only SVOC detected [up to 350 micrograms per kilogram (µg/kg)]. No HE compounds or radionuclides were detected in any of the borehole soil samples. No metals exceeded their respective SNL/NM site-wide calculated UTL background concentrations (IT 1994). Manganese and vanadium UTL background concentrations were not calculated, but reported concentrations were below the proposed RCRA Subpart S action levels (Table 2).

On November 2 and 6, 1994, Boreholes TA2-BH-08 and TA2-BH-09 were drilled near Building 907 (Figure 2). Borehole TA2-BH-08 was drilled along the sanitary sewerline near the seepage pit. Borehole TA2-BH-09 was drilled along the HE drain trench near the HE catch box. Soil samples were collected from each borehole at depths of 5, 10, 15, 20, 30, 40, and 50 feet BGL. The soil samples were analyzed at off-site laboratories for VOCs, SVOCs, HE, total metals, cyanide, tritium, and gamma spectroscopy.

No VOCs, HE, cyanide, or mercury were identified above instrument detection limits. The only SVOC detected was the common laboratory contaminant bis (2-ethylhexyl) phthalate in Borehole TA2-BH-05 with a maximum concentration of 350  $\mu$ g/kg at a depth of 14.6 feet. To support the conclusion that this constituent was a result of laboratory contamination, the passive SVS identified no SVOCs in the vicinity of Building 907, and no SVOCs were identified as potential constituents of concern for this site (Section 2.3).

Metals exceeding the SNL/NM site-wide UTL background concentrations (IT 1994) were: barium [UTL = 407.9 milligrams per kilogram (mg/kg)], detected at a maximum of 944 mg/kg (10-foot depth) in Borehole TA2-BH-09, less than the proposed RCRA Subpart S action level value of 6000 mg/kg; beryllium (UTL = 0.8 mg/kg), detected at a maximum of 1.2 mg/kg (31-foot depth) in Borehole TA2-BH-08 compared to the proposed Subpart S

action level of 0.2 mg/kg; copper (UTL = 16.7 mg/kg), detected at 17.2 mg/kg (31-foot depth) in Borehole TA2-BH-08, no action level calculated as copper is an important essential element; nickel (UTL = 15.4 mg/kg), detected at 18.4 mg/kg (31-foot depth) in Borehole TA2-BH-08, less than the proposed RCRA Subpart S action level of 2000 mg/kg; and zinc (UTL = 46.7 mg/kg), detected at 58.8 mg/kg (31-foot depth) in Borehole TA2-BH-08, less than the proposed Subpart S action level of 20,000 mg/kg. Radiological data identified no elevated activities compared to background.

Site-wide UTL background concentrations were not calculated for arsenic, manganese or vanadium. However, proposed RCRA Subpart S action levels were available. Arsenic was identified at a maximum concentration of 5.7 mg/kg (10-foot depth) in Borehole TA2-BH-09, less than the proposed RCRA Subpart S action level of 20 mg/kg. Manganese was identified at a maximum concentration of 548 mg/kg (31-foot depth) in Borehole TA2-BH-08, less than the proposed RCRA Subpart S action level of 10,000 mg/kg (in food). Vanadium was identified at a maximum concentration of 33.2 mg/kg (31-foot depth) in Borehole TA2-BH-08, less than the proposed RCRA Subpart S action level of 600 mg/kg.

Although beryllium was detected slightly above the SNL/NM site-wide background UTL, the value of 1.2 mg/kg could be within background concentrations since the UTL is based on the 95 percent tolerance coefficient; i.e. 5 percent of the background data will exist above the UTL.

#### 3.7 Rationale for Pursuing a Confirmatory Sampling NFA Decision

A comparison of analytical results to SNL/NM background levels and RCRA proposed Subpart S action levels shows that all chemical constituents of concern are either within background concentration levels and/or below the prescribed action level. The results of the surface surveys and soil sampling and analytical results indicate that there has been no release of hazardous constituents from this site that may pose a threat to human health and/or the environment.

#### 4. Conclusion

ER Site 136 is being proposed for an administrative NFA decision based on confirmatory sampling because the evidence cited above demonstrates that the SWMU clearly has not released hazardous wastes or constituents into the environment (Criterion C) (see Section 1.2). Therefore, no threat to human health or the environment exists.

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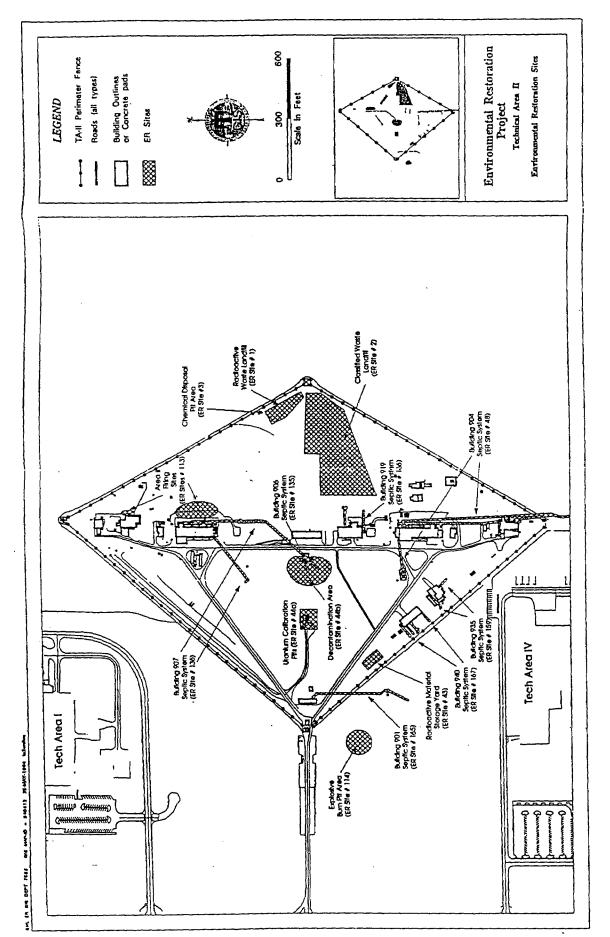
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Map Showing Technical Area II and the Locations of Environmental Restoration Sites Figure 1.

No Further Action Proposal (Site 136)

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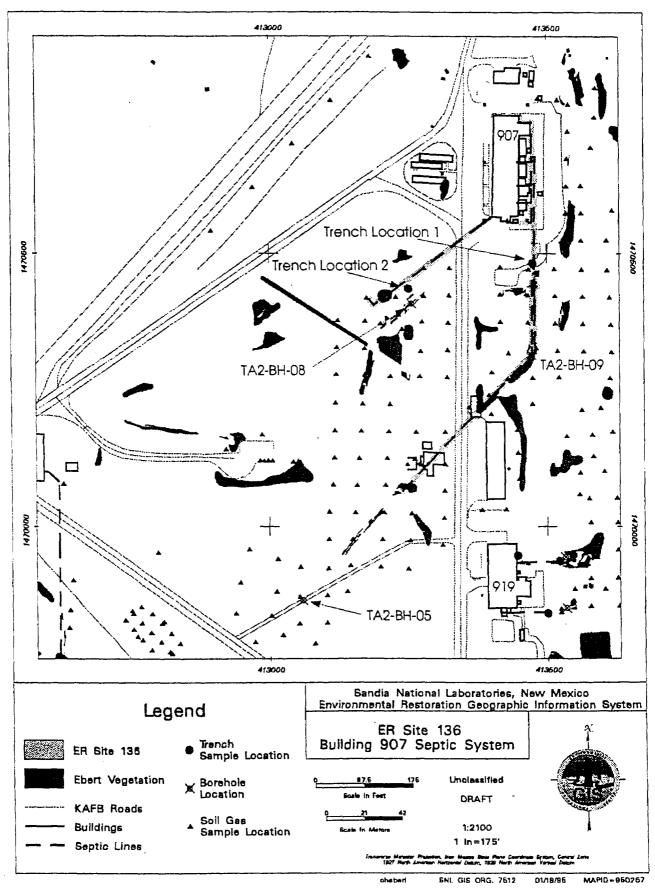


Figure 2. Map Showing ER Site 136 Building 907 Septic System

Table 1. Site 136, Building 907 Septic System, Data Summary of Soil Samples Collected from Trenches 1 and 2

Parameter	Trench 1 maximum concen. mg/kg-mtls ug/kg-org.	Trench 2 maximum concen. mg/kg-mtls ug/kg-org.	Site-wide UTL background concen. mg/kg-mtls ug/kg-org.	RCRA Subpart S Action Level mg/kg-mtls ug/kg-org.
Antimony	ND	ND	NC	30
Arsenic	2.5	3.4	NC	20
Barium	104	101	407.9	6,000
Beryllium	.36	.45	.8	.2
Cadmium	.68	ND	3.5	80
Chromium	6.3	6.5	22.9	80,000
Cobalt	3.8	5.0	21ª	NC
Copper	12.2	11.6	16.7	NC
Lead	14.5	10.2	15	NC
Manganese	171	225	NC	10,000
Mercury	ND	ND	NC	20
Nickel	5.1	6.7	15.4	2,000
Selenium	ND	ND	NC	400
Silver	3.4	ND	4.0	400
Thallium	ND	ND	NC	6.92
Vanadium	16.6	24	NC	600
Zinc	34.1	37	46.7	20,000
Acetone	12	ND	NC	8,000
Carbon Tetra- chloride	ND	ND	NC	5

Table 1. Site 136, Building 907 Septic System, Data Summary of Soil Samples Collected from Trenches 1 and 2 (Concluded)

Parameter	Trench 1 maximum concen. mg/kg-mtls ug/kg-org.	Trench 2 maximum concen. mg/kg-mtls ug/kg-org.	Site-wide UTL background concen. mg/kg-mtls ug/kg-org.	RCRA Subpart S Action Level mg/kg-mtls ug/kg-org.
Methyl ethyl ketone	ND	ND	NC	50,000
Methylene chloride	6B	8B	NC	90
Trichloro- ethene	ND	7	NC	60
Toluene	6.6	13	NC	20,000
Xylene	7	12	NC	200,000
HMX	ND	ND	NC	NC
RDX	ND	ND	NC	NC

#### **Notes**

ND = Not detected.

N/A = Not applicable.

NC = Not calculated.

B = Parameter detected in blank.

a = A site-wide UTL was not calculated for cobalt. However, a UTL was calculated for the Tijeras Arroyo sites which are adjacent to TA-II. The UTL for Tijeras Arroyo was used in this NFA proposal.

Aluminum, calcium, iron, magnesium, potassium and sodium were excluded from the table due to natural abundance.

Site 136, Building 907 Septic System, Data Summary of Soil Samples Collected from Borehole 05, Borehole 08 and Borehole 09 Table 2.

Parameter	BH-05	Depth	BH-08	Depth	BH-09	Depth	Site-Wide	RCRA
	max.	(ft)	max.	(ft)	max.	(ft)	OTL	Subpart S
	concen.		concen.		concen.		backgnd	Action
							concen.	Tevel
	mg/kg- metale		mg/kg- metals		metals		mø/ko-	mo/ko-
	no/ko-		nσ/kg-		ng/kg-		metals	metals
	organic		organic		organic		ug/kg-	ug/kg-
							organic	organic
Antimony	QN	N/A	ND	N/A	ND	N/A	NC	30
Arsenic	3.8	30	4.8	11	5.7	10	NC	20
Barium	150	30	174	16.5	944	10	407.9	6,000
Beryllium	.26	21.2	1.2	31	.46	28	8.	.2
Cadmium	.71	30	ON	N/A	ND	N/A	3.5	80
Chromium	9.3	6, 21.2	15.5	31	9.5	28	22.9	80,000
Cobalt	6.9	14.6	10.2	31	5.5	28	21ª	NC
Copper	10.8	21.2	17.2	31	11.3	28	16.7	NC
Cyanide	SN	N/A	ON	N/A	ND	N/A	NC	2,000
Lead	5.3	30	8.2	31	5.8	28	15	NC
Manganesc	281	11	548	31	284	28	NC	10,000
Mercury	ND	N/A	ND	N/A	ND	N/A	NC	20
Nickel	7.9	30	18.4	31	10.5	28	15.4	2,000

Site 136, Building 907 Septic System, Data Summary of Soil Samples Collected from Borehole 05, Borehole 08 and Borehole 09 (Continued) Table 2.

Parameter	BH-05	Depth	BH-08	Depth (ft)	BH-09 max.	Depth (ft)	Site-Wide UTL	RCRA Subpart S
-	concen.		concen.	· · · · · · · · · · · · · · · · · · ·	concen.		backgnd concen.	Action Level
	mg/kg- metals		mg/kg- metals		mg/kg- metals		mg/kg-	mg/kg-
	ug/kg-		ug/kg-		ug/kg-		metals	metals
	organic		organic		organic		ug/kg- organic	ug/kg- organic
Selenium	QN	N/A	ND	N/A	QN	N/A	NC	400
Silver	ND	N/A	ND	N/A	ND	N/A	4.0	400
Thallium	QN	N/A	ND	N/A	ND	N/A	NC	6.92
Vanadium	31.7	9	33.2	31	24	28	NC	009
Zinc	30.8	21.2	58.8	31	37.9	28	46.7	20,000
Acetone	22B	15.5	NS	N/A	NS	N/A	NC	8,000
bis(2ethylhe- xyl)phthalate	350	14.6	ND	N/A	ND	N/A	NC	50
Methylene Chloride	5.9B	30.2	ND	N/A	ND	N/A	NC	90
HMX	ND	N/A	ND	N/A	ND	N/A	NC	NC
RDX	ND	N/A	ND	N/A	ND	N/A	NC	NC

Table 2. Site 136, Building 907 Septic System, Data Summary of Soil Samples Collected from Borehole 05, Borehole 08 and Borehole 09 (Concluded)

#### Notes

ND = not detected.

N/A = not applicable.

NS = Not sampled.

NC = Not calculated.

B = Parameter detected in blank.

a = A site-wide UTL was not calculated for cobalt. However, a UTL was calculated for the Tijeras Arroyo sites which are adjacent to TA-II. The UTL for Tijeras Arroyo was used in this NFA proposal.

Aluminum, calcium, iron, magnesium, potassium and sodium were excluded from the table due to natural abundance.

#### APPENDIX A

Confirmatory Sampling and Analysis Plan

## Workplan for Drilling at Technical Area 2

#### Introduction

Beginning on March 1, 1994, drilling will be conducted at four locations at Technical Area II (TA-2) (Figure 1) to collect preliminary data for the RFI Workplan. The scope of work will be divided into two phases. Phase I involves drilling three deep boreholes and eleven shallow boreholes in the vicinity of TA-2. Phase II involves advancing one and possibly two of the deep boreholes to first water and completing each as monitor wells. All Phase I boreholes will be drilled with a Barber drill rig using sonic resonance. All Phase II boreholes will be drilled with a Dresser XT-70E drill rig using mud-rotary punch coring. During both Phase I and Phase II, continuous core will be collected from each borehole with a core barrel. The scope of work for Phase I involves:

- Drilling and sampling nine boreholes to 6 ft BGL and two boreholes to 12 ft BGL in the vicinity of the former High Explosives Burn Pit (HEBP) area;
- Drilling and sampling one borehole to 150 ft below ground level (BGL) near the Building 906 dry well;
- Drilling and sampling one borehole to 100 ft BGL near the Building 935 retention tank and dry well; and
- Drilling and sampling one borehole to 150 ft BGL in the Classified Waste Landfill (CWL).

The scope of work for Phase II involves:

- Advancing the 150-ft deep borehole drilled near the Building 906 dry well to first water and completing the borehole as a monitor well; and
- If time and budget permit, the 150-ft deep borehole drilled in the CWL will be advanced to first water and completed as a monitor well.

The Phase II boreholes and monitor wells will be advanced and installed, respectively, with the XT-70E Dresser drill rig after it has completed work at the Chemical Waste Landfill (expected to be April 1994).

#### The purposes of Phase I are to:

- Drill and sample shallow boreholes with the Barber drill rig at the former HEBP area;
- Drill and sample pilot boreholes with the Barber drill rig prior to using the larger Dresser drill rig; and
- Determine the nature and vertical extent of contamination, if any, beneath the designated ER sites prior to well installation.

#### The purposes of Phase II are to:

- Advance the boreholes at Building 906 and the CWL (if time and budget permit) to first water using the Dresser XT-70E drill rig;
- Collect soil samples from the two deep boreholes; and
- Complete the two boreholes as monitor wells in the first water-bearing zone encountered (assumed to be at 300 ft BGL.

The Dresser XT-70E drill rig will not be available until about mid- to late April. Therefore, the smaller Barber drill rig will be used to drill pilot boreholes at Building 906 and in the vicinity of the CWL, and drill and sample boreholes in the former HEBP area and one at Building 935. The Dresser drill rig is better suited for drilling deep boreholes than the Barber drill rig. In addition, mud-rotary wireline punch coring can be used with the Dresser drill rig, which is more effective and reliable than sonic coring at depths greater than about 150 ft.

The soil analytical data collected during Phase I will be used to determine the nature and extent of potential contamination at the four drilling locations. These data also will be used to evaluate waste management issues regarding the use of mud-rotary drilling and well installation methods if the subsurface soil is contaminated. Geophysical logging will be conducted after well completion and will consist of EM Induction (EMI) resistivity or resistivity (which ever is available), natural gamma, and neutron. Neutron logging will not be conducted in the open boreholes to avoid the potential for losing the radiation source in the borehole.

Details of the drilling and sampling at the four locations at TA-2 are described separately below. Management of investigation-derived waste (IDW) is presented in the Waste Management Plan for Investigation-Derived Waste at Technical Area II.

#### **Project Personnel**

A site-specific Health and Safety Plan (HASP) has been developed for the planned drilling activities at TA-2. The Technical Task Leader for the drilling activities is Rarilee Conway of Department 7582. The Site Safety Officer (SSO) for the drilling activities is Michael Wade and/or a designated Health Physics (HP) Technician or person from IT Corporation (IT). The boreholes will be drilled by Water Development Corporation (WDC) of Woodland, California, and logged by Michael Wade, Rarilee Conway, John Copland, and/or Tom Tharp. WDC will provide equipment for drilling (Drill rod, PVC pipe, etc.). Jerry Mercer (Department 9333) will serve as a consultant who will be "on call" during drilling operations to assist with any technical and/or other drilling situations that may occur during drilling. He will also assist with ordering necessary drilling supplies and assisting with coordinating drilling activities.

Two technicians/geologists from IT will assist with soil and ground-water sampling, filling out chain-of-custody (COC) forms, health and safety monitoring, obtaining field sampling equipment and sample jars, delivering samples to the Sample Management Office (SMO), and any other related field work. IT will also provide field-screening for VOCs with a PID or FID (i.e., sample equipment, core, etc.). Sandia will provide a full-time, Certified Industrial Hygienist (CIH) and HP Technician to monitor drilling and sampling equipment and soil and samples for health and safety reasons. At Buildings 906 and 935 and at the CWL, the HP will be available for the first 50 ft of drilling (i.e., the primary zone of potential contamination). The HP will let the field team know when it is no longer necessary for continued radiation field-screening. Potential project personnel and their phone numbers are listed in the site-specific HASP.

If any laboratory questions arise regarding sample containers, sample quantity, holding times, etc., please contact the following people:

Bob Friberg, Sample Coordinator at TMA/Eberline (or Jim Lozito): 505-345-3461. Ellen LaRiviere, ENSECO: 303-421-6611; ext. 308.

Samples collected for volatile organic compounds (VOCs), semi-volatile organic compounds (SVOCs), high explosives (HE) compounds, and Target Analyte Listed

(TAL) metals will be submitted to ENSECO unless otherwise notified. Radiological samples (total uranium, tritium, and photon emitters (gamma scan), will be submitted to TMA/Eberline. In addition, soil will be collected in a plastic marinelli beaker and screened by Amir Monagheghi at the Health Physics Laboratory, Department 7715 (Radioactive Protection & Measurement Department) for radiological screening. These samples will be collected at every interval where soil samples are collected for pre-laboratory screening purposes. Equipment rinse blanks will also be collected and submitted to the analytical laboratories at various times in the field, such as after completing a borehole. The frequency of collecting equipment blanks will be determined in the field by the sampling team.

The backgrounds for the Environmental Restoration (ER) sites associated with this project are described separately in the Draft RFI Workplan and are available on request. The HASP provides additional information concerning potential contaminants of concern (COCs) at each site. Details of the Phase I and Phase II sampling and analysis plans (SAPs) for each of the four locations are presented below.

#### Phase I Drilling and Soil Sampling

Phase I activities involve drilling 14 boreholes from 6 ft to 150 ft deep at four locations at TA-2. These activities are described separately below. Because of ongoing activities at TA-2, drilling during the weekend is necessary in the CWL. To avoid the potential for shrapnel and debris striking the drill rig, drilling must take place at the CWL during Friday, March 4 through Monday, March 7. Therefore, Phase 1 drilling will begin at the former HEBP area on Tuesday March 1 to avoid the potential to stop drilling in a deep borehole before it is finished (such as at Building 906 or 935).

The estimated schedule for Phase 1 drilling at TA-2 is as follows, and assumes that no major drill rig or health and safety issues occur during fieldwork:

Proposed schedule for drilling at TA-2 in March 1994.

Estimated Drilling Dates	Location/Activity
Week Prior to March 1	Drillers and any field personnel take RAD 101 on Tuesday, February 22. Mobilize drill rig to TA-2; set-up at former HEBP area.
Tuesday, March 1	Start drilling shallow 11 boreholes (6 ft to 12 ft deep) at the former HEBP area. Backfill boreholes with grout.
Thursday, March 3 - afternoon	Decontaminate drill rod; mob and set-up at CWL.
Friday, March 4	Start to drill 150-ft pilot borehole at CWL.
Monday, March 7	Backfill borehole with grout, if necessary.  Decontaminate drill rod; mob and set-up at Building 906.
Tuesday, March 8	Start to drill 150-ft pilot borehole at Building 906.
Thursday, March 11	Possibly complete pilot borehole at Building 906. Backfill with grout, if necessary. End of first field cycle.
Tuesday, March 15	Decontaminate drill rig and equipment; mob and set-up at Building 935. Start to drill 100-ft borehole at Building 935.
Friday, March 18	Complete borehole at Building 935. Backfill with grout. Decontaminate drill rig and equipment. TA-2 Phase 1 drilling complete. Drill rig mobilizes to LWDS site.

The total estimated drilling days (including decontamination and mobilization) are 14. If Phase 1 drilling takes longer, Phase 2 drilling time will be reduced, and potentially only one monitor well will be installed (Building 906) instead of two. In addition, drilling schedules could change depending on activities or security issues at TA-2 or the availability of drill rigs. For additional information, Attachment A presents draft site background information about Buildings 906 and 935 and the Classified Waste Landfill.

#### Former High Explosives Burn Pit Area

Extensive field sampling has been conducted in the vicinity of the former HEBP area to locate two former burn pits and to determine the vertical and lateral extent of any potential contamination (Figure 2). Between September 3rd and 22nd, 1992, a total of 185 boreholes were drilled with a CME-75 drill rig to 6 ff BGL. The boreholes were

drilled on a 4-ft grid pattern to achieve a 95 percent confidence level for locating the two former burn pits. As part of the investigation, the two former burn pits were informally named Burn Pit Y (BPY) and Burn Pit Z (BPZ) (IT, 1993) (Figures 2 and 3). A total of 370 soil samples were collected (one each at depths of 3 ft and 6 ft at each borehole location). The soil samples were analyzed for total RCRA metals and HE compounds. The soil samples were not analyzed for VOCs or SVOCs. However, all soil samples were field screened for VOCs with a PID and for alpha, beta, and gamma radiation using alpha scintillation and G-M probes. No VOCs or radioisotopes were detected above background levels during field sampling (IT, 1993). Details of this investigation are discussed in IT (1993) and are summarized below.

Figures 2 and 3 show the locations of soil samples collected from boreholes BPY-11,4 through BPY-11,6 which contained elevated metals concentrations (IT, 1993). The highest elevated concentrations of the metals in soil from these boreholes are:

- Barium at 23,800 mg/kg (6-ft sample from borehole BPY-11,5);
- Cadmium at 16.8 mg/kg (6-ft sample from borehole BPY-11,5);
- Chromium at 9.7 mg/kg (6-ft sample from borehole BPY-11,5); and
- Lead at 110 mg/kg (6-ft sample from borehole BPY-11,4).

Only the soil sample collected from 6 ft in borehole BPY-11,5 contained detectable concentrations of HE. At this location, RDX was detected at a concentration of 5.9 mg/kg, and HMX was detected at 3.1 mg/kg (IT, 1993). No other HE compounds were detected in this or any other sample collected in the vicinity of the former HEBP area; the detection limit was 1 mg/kg (IT, 1993).

The purpose the additional sampling described in this document is to collect data to:

- 1) Identify the lateral extent of barium and any associated COCs in the immediate vicinity east of boreholes BPY-11,4 through BPY-11,6 (i.e., between Burn Pit Y and Burn Pit Z);
- 2) Determine the vertical extent of barium and any associated COCs at depths greater than 6 ft in boreholes BPY-11,4 and BPY-11,5; and
- 3) Support a proposal for No Further Action (NFA) or for evaluating remedial alternatives at the HEBP area, if necessary.

The sampling for the HEBP area involves:

- Collecting soil samples at 3-ft and 6-ft depths in the area between boreholes BPY-11,4 through BPY-11,6 (Burn Pit Y) and Burn Pit Z. If any COCs are detected above background concentrations at the 6 ft depth, additional soil samples will be collected vertically and laterally as part of the RFI Workplan activities for TA-2 until no COCs are identified significantly above background concentrations;
- Collecting soil samples at the 9 ft and 12 ft depths at borehole locations BPY-11,4 and BPY-11,5; and
- Analyzing the soil samples for HE compounds and metals.

To fully determine the lateral extent of barium and any associated COCs between Burn Pit Y and Burn Pit Z, soil samples will be collected at 3-ft and 6-ft depths from boreholes using a 4-ft sampling grid. The boreholes will be drilled with the Barber drill rig using continuous sonic coring. The 4-ft sampling grid established during the September 1992 investigation will be extended east toward Burn Pit Z.

As shown on Figures 2 and 3, three boreholes each will be drilled east of boreholes BPY-11,4, BPY-11,5, and BPY-11,6 (i.e., 9 boreholes total). Because of the extensive soil sampling conducted in this area in September 1992, the boreholes and soil samples in this investigation will be identified using the same name convention as in September 1992, except that all samples will be preceded by TA-2 (e.g., TA2-BPY-12,5-3.0). As shown in Tables 1 and 2, the soil samples will be analyzed for TAL metals by EPA Methods 6010 and 7000 and HE compounds by USATHAMA Method. Only one sample needs to be collected from each depth, but can be used for all analyses to reduce sample collection time and management. If any potential COCs are detected in soil samples collected from the easternmost locations, additional boreholes will be drilled to the western edge of Burn Pit Z as part of the TA-2 RFI Workplan activities (since analytical results will not be available for up to 2 months).

At boreholes BPY-11,4 and BPY-11,5, soil samples will be collected at 9-ft and 12-ft depths to evaluate the vertical extent of elevated metals concentrations (i.e., primarily barium). The samples collected for metals will be analyzed by EPA Methods 6010 and 7000 (Table 1). In addition, soil samples collected from boreholes

BPY-11,4 and BPY-11,5 also will be analyzed for HE compounds by the USATHAMA Method for HE compounds. This HE analysis has a 7-day holding time, so samples must be delivered to the SMO as soon as possible (Table 1). If any potential COCs are detected in soil samples collected from 12 ft deep in either BPY-11,4 or BPY-11,5, the borehole will be advanced and sampled below 12 ft deep as part of the TA-2 RFI Workplan activities (since analytical results will not be available for up to 2 months).

All drilling equipment and drill core material will be field-screened for VOCs with a PID or FID and for radioisotopes using alpha scintillation and G-M probes. If any potential COCs are identified above background levels, the samples will be submitted for analysis. All soil samples collected will submitted for off-site analysis; samples also will be submitted for QA/QC (duplicates and Matrix Spike/Matrix Spike Duplicate, etc.) (Tables 1 and 2). In addition, all samples should be preserved on ice. The drilling geologist may collect additional soil samples in more permeable zones or submit more samples for QA/QC analysis if determined necessary in the field. In addition, ground water samples should be collected from any perched water-bearing zone(s) encountered during drilling, although no perched ground water is expected at these shallow locations. The samples should be collected, submitted, and analyzed for the potential COCs as listed in Tables 1 and 2. To minimize any potential cross-contamination, all sampling equipment should be decontaminated according to PRO 91-27 - General Equipment Decontamination (i.e., a mixture of water and Alconox soap followed by deionized water).

After each borehole has been drilled to the total depth (i.e., 6 ft or 12 ft BGL), each will be backfilled with grout. No additional drilling is planned in the vicinity of the former HEBP area for Phase II. This site is planned for an NFA proposal.

Table 1. Summary of analyses for soil samples to be collected from boreholes drilled in the immediate vicinity of Burn Pit Y (BPY), former HEBP area, Technical Area 2 (see Figure 5 for the locations of boreholes).

Borehole Locations	Sample Type or QA/QC Type	Sample Depth (in ft)	TAL Metals <sup>a</sup>	HEp	Sample Container Type <sup>c</sup>	Number of containers <sup>d</sup>
Lateral Sample Grid - BPY Row 12						
TA2-BPY-12,4-3.0	Subsurface soil	3	1	1	500 ml	1
TA2-BPY-12,4-6.0	Subsurface soil	6	1	1	500 ml	1
TA2-BPY-12,4-6.0	MS/MSD - Include on COC for BPY-12,4-6.0	6				
TA2-BPY-12,5-3.0	Subsurface soil	3	1	1	500 ml	1
TA2-BPY-12,5-6.0	Subsurface soil	6	1	1	500 ml	1
TA2-BPY-12,6-3.0	Subsurface soil	3	1	1	500 ml	1
TA2-BPY-12,6-6.0	Subsurface soil	6	1	1	500 ml	1
TA2-BPY-12,6-6.0	Duplicate of TA2- BPY-12,9-6.0	6	1	1	500 ml	1
Total			7	7	7	7 total
			analyses	analyses	containers	containers
Lateral Sampling Grid - BPY Row 13						
TA2-BPY-13,4-3.0	Subsurface soil	3	1	1	500 ml	1
TA2-BPY-13,4-6.0	Subsurface soil	6	1	1	500 ml	1
TA2-BPY-13,5-3.0	Subsurface soil	3	1	1	500 ml	1
TA2-BPY-13,5-6.0	Subsurface soil	6	1	1	500 ml	1 .
TA2-BPY-13,6-3.0	Subsurface soil	3	1	1	500 ml	1
TA2-BPY-13,6-6.0	Subsurface soil	6	1	1	500 ml	1
TA2-BPY-13,6-6.0	Duplicate of TA2- BPY-13,6-6.0	6	1	1	500 ml	1
Total			7	7	7	7 total
			analyses	analyses	containers	containers
Lateral Sampling Grid - BPY Row 14						
TA2-BPY-14,4-3.0	Subsurface soil	3	1	1	500 ml	1
TA2-BPY-14,4-6.0	Subsurface soil	6	1	1	500 ml	1
TA2-BPY-14,5-3.0	Subsurface soil	3	1	1	500 ml	1
TA2-BPY-14,5-6.0	Subsurface soil	6	1	1	500 ml	1
TA2-BPY-14,6-3.0	Subsurface soil	3	1	1	500 ml	1
TA2-BPY-14,6-6.0	Subsurface soil	6	1	1	500 ml	1
Total			6 analyses	6 analyses	6 containers	76total containers

a - EPA Methods 6010 and 7000 for TAL metals.

NOTE: All soil samples should be preserved on ice unless otherwise noted.

NOTE: NO soil will be collected for radiological screening (i.e., for Amir) at the HEBP area.

b - USATHAMA Method for 9 HE compounds; 7-day holding time so deliver to SMO ASAP.

c - All samples will be collected in 500 ml glass jars.

d - TAL Metals and HE compound analyses can be collected in one 500 ml glass jar. Total number of containers does not include miranelli beaker samples for radiological screening.

NOTE: Equipment blanks will be collected at the end of each day or each borehole at the HEBP area. Sampling frequency can be determined in the field by the sampling team and/or geologist. These samples will be labeled as TA2-BPY-EB; a designation can be added to the BPY if it's sampled at a particular borehole (e.g., BPY-11). These samples should be collected in a 2.5 liter amber glass jar or in a l liter bottle for TAL metals and HE compounds combined.

NOTE: Collect a soil split sample (about 100 grams) at each sample location in a ziplock bag and label accordingly for future transport to Department 7584 to Nelson Capitan. Keep samples refrigerated and at the site or shed.

Table 2. Summary of analyses for soil samples to be collected from boreholes drilled in Burn Pit Y (BPY), Row 11, former HEBP area, Technical Area II (see Figure 5 for the locations of boreholes).

Borehole Locations	Sample Type or QA/QC Type	Sample Depth (in ft)	TAL Metals <sup>a</sup>	HEb	Sample Container Type <sup>c</sup>	Number of containers d
TA2-BPY-11,4-9.0	Subsurface soil	9	1	1	500 ml	1
TA2-BPY-11,4-12.0	Subsurface soil	12	1	1	500 ml	1
TA2-BPY-11,5-9.0	Subsurface soil	9	1	1	500 ml	1
TA2-BPY-11,5-12.0	Subsurface soil	12	1	1	500 ml	1
Total			4	4	4	4
			analyses		containers	containers

- a EPA Methods 6010 and 7000 for TAL metals.
- b USATHAMA Method for 9 HE compounds.
- c All samples will be collected in 500 ml glass jars.
- d One 500 ml glass jar will be submitted for both TAL Metals and HE compounds. Total number of containers does not include soil samples collected in miranelli beaker samples for radiological screening.

NOTE: All soil samples should be preserved on ice unless otherwise noted.

NOTE: NO soil will be collected for radiological screening (i.e., Amir) at the HEBP area.

NOTE: Equipment blanks will be collected at the end of each day or each borehole at the HEBP area.

Sampling frequency can be determined in the field by the sampling team and/or geologist.

These samples will be labeled as TA2-BPY-EB; a designation can be added to the BPY if it's sampled at a particular borehole (e.g., BPY-11). These samples should be collected in either a 2.5 liter amber glass jar or in a l liter bottle for TAL metals and HE compounds combined.

### Classified Waste Landfill

This location has been selected based on historical information and the results of a passive soil vapor survey (SVS) and surface geophysics. The SVS data identified two separate PCE soil vapor plumes in the Classified Waste Landfill (CWL) (Figure 3). TCE also was identified as a co-contaminant in one of the PCE soil vapor plumes.

The surface geophysics data shows an abundance of metallic debris buried at the location of the soil vapor plume. Therefore, one borehole (TA2-BH-03) will be drilled immediately north (about 10 to 20 ft) of the PCE and TCE soil vapor plume so that no buried metallic debris are encountered during drilling (Figure 4). Using the sonic coring system retrieved with drill rod, the borehole will be cored continuously and the lithology of the core will be described. Soil samples will be collected for analysis at the following depth intervals: 5, 10, 15, 20, 30, 40, 50, 75, 100, and 150 ft. As listed in detail in Tables 3a and 3b, the soil samples will be analyzed for TAL metals, HE compounds, VOCs, SVOCs, gamma spectroscopy, total uranium, and tritium. Because the passive SVS results indicate that VOCs are present in soil vapor, it is assumed that VOCs have been released to the environment in the CWL. The SVS data also indicate that 1,1,1-TCA and Freon compounds may be present in the landfill. No SVOCs were identified from the SVS investigation. However, soil samples will be collected and analyzed for SVOCs to a depth of 50 ft.

All drilling equipment and drill core material will be field-screened for VOCs with a PID or FID and for radioisotopes using alpha scintillation and G-M probes. If any potential COCs are identified above background levels, the samples will be submitted for analysis. In addition, all soil samples will be field-screened for radioisotopes and VOCs. All soil samples collected will be submitted for off-site analysis; additional samples will be collected and submitted for QA/QC (duplicates and Matrix Spike/Matrix Spike Duplicate, etc.) (Tables 3a and 3b). In addition, all samples should be preserved on ice. The drilling geologist may collect additional soil samples in more permeable zones or submit more samples for QA/QC analysis if determined necessary in the field. In addition, ground-water samples will be collected from any perched water-bearing zone(s) encountered during drilling. The samples should be collected, submitted, and analyzed for the potential COCs as listed in Tables 3a and 3b. To minimize any potential cross-contamination, all sampling

equipment should be decontaminated according to PRO 91-27 - General Equipment Decontamination (i.e., a mixture of water and Alconox soap followed by deionized water).

After the borehole has been drilled to a total depth of 150 ft, it will be backfilled to ground surface with grout. After monitor well TA2-MW-01 (Building 906) has been installed and developed with the Dresser drill rig (Phase 2), borehole TA2-BH-03 will be advanced, if time and budget permit, to first water using mud-rotary wireline punch-coring as part of Phase II drilling activities, as described later.

Table 3a. Summary of non-radiological analyses for soil samples to be collected from borehole TA2-BH-03 drilled in the vicinity of the Classified Waste Landfill, Technical Area 2.

Sample Type or QA/QC Type	Sample Depth (in ft)	VOCs <sup>a</sup>	Sample container type <sup>e</sup>	SVOCs <sup>b</sup>	HEc	TAL Metals <sup>d</sup>	Sample container type <sup>e</sup>	Total number of containers <sup>f</sup>
Subsurface soil	5	1	125 ml	1	1	1	500 ml	2
Subsurface soil	10	1	125 ml	1_	1	1	500 ml	2
MS/MSD - Include on COC	10							
Subsurface soil	15	1	125 ml	1	1	1	500 ml	2
Subsurface soil	20	1	125 ml	1	1	1	500 ml	2
Subsurface soil	30	1	125 ml	1	1	1	500 ml	2
Subsurface soil	40	1	125 ml	1	1	1	500 ml	2
Subsurface soil	50	1	125 ml	1	1	1	500 ml	2
Duplicate of sample at 50 ft	50	***			1	1	500 ml	1
Subsurface soil	<i>7</i> 5	. 1	125 ml					1
Subsurface soil	100	1	125 ml					1
Subsurface soil	150	1	125 ml					1
Total		10	10	7	8	8	8	18 total containers

- a EPA Method 8240.
- b EPA Method 8270.
- c USATHAMA Method for 9 HE compounds.
- d EPA Methods 6010 and 7000.
- e All samples will be collected in glass jars as per laboratory request (Enseco); SVOCs, HE, and TAL Metals will all be collected and submitted in a 16 oz. (500 ml) glass jar.
- f All VOC samples will be collected in a 4 oz. (125 ml) glass jar.

NOTE: A VOC and SVOC field blank should be prepared and submitted-one each for the CWL borehole.

NOTE: All soil samples should be preserved on ice unless otherwise noted.

NOTE: All soil samples should be labeled as TA2-BH-03-depth-medium (50.5-S).

NOTE: Soil will be collected in a miranelli beaker from every sample interval and submitted to Amir for radiological screening.

NOTE: Equipment blanks will be collected at the end of each day or after a particular sampling interval (e.g., 0 to 50 ft). Sampling frequency can be determined in the field by the sampling team and/or geologist. These samples will be labeled as TA2-BH-03-EB. These samples should be collected in either a 2.5 liter amber glass jar or in a l liter bottle for SVOCs, TAL metals, and HE compounds combined. A 40 ml VOA can be used for VOC samples.

# Workplan for Drilling at Technical Area 2

# Introduction

Beginning on about Tuesday, November 1, 1994 (time is approximate; assumes all drilling contracts are in place), drilling will be conducted at several locations within Technical Area II (TA-II) to collect data in support of the DOE-approved TA-II RFI Workplan and no further action (NFA) proposals. The scope of work will be divided into two phases. Phase I involves drilling 25 boreholes in TA-II ranging in depths from 30 to about 125 ft below ground level (BGL). The majority of the boreholes will be drilled to 50 ft. Phase II will involve drilling up to 2 boreholes to first water and completing them as monitor wells. This mini-Workplan only presents work and sampling and analysis (SAP) tables for Phase I. Phase II work details will be presented at a later date. All Phase I boreholes probably will be drilled with hollow stem augers and samples will be collected in split-spoon samplers lined with stainless steel liners (as done in the March 1994 drilling at Technical Area II). Phase I drilling will probably be conducted with an F-10, Mobile B-61, or a CME-75 or -95 drill rig (depending upon availability). Installation of the Phase II monitor wells will be determined after the results of the Phase I drilling, but may involve wireline coring using air or air-mist. During Phase I, continuous core will not be collected; the lithology at each borehole will be described from split-spoon samples and drill cuttings. All angled boreholes probably will be drilled with hollow stem augers, but other methods also will be evaluated.

The scope of work for Phase I drilling will involve:

# Non-Landfill Portions of TA-II

- Drilling and sampling two boreholes to 50 ft BGL at Building 904 (one at the septic leachfield and one along the high explosives [HE] catch box);
- Drilling and sampling two boreholes to 50 ft BGL at Building 907 (one at the septic leachfield and one along the HE catch box);
- Drilling and sampling three boreholes in the vicinity of Building 935 (two 50-ft boreholes, one downgradient from the former retention tank and dry well locations and one adjacent to the septic tank; and one 30-ft borehole

- east of the septic tank in a high trichloroethene [TCE] soil vapor survey [SVS] location);
- Drilling and sampling two boreholes to 50 ft BGL at Building 940 (one near the septic tank and one near the dry well);
- Drilling and sampling one borehole to 50 ft BGL west of Buildings 915 and 922 (at a high benzene-toluene-ethylbenzene-xylene [BTEX] SVS location in the vicinity of the septic tank);
- Drilling and sampling one borehole to 100 ft BGL southwest of Building
   913 (at the highest TCE SVS location in TA-II); and
- Drilling and sampling two boreholes to 50 ft BGL east of Building 919 (one in the septic leachfield and one near the seepage pit).

Thus, thirteen boreholes will be drilled and sampled in non-landfill portions of TA-II. Phase I also will involve:

## Eastern Portion of TA-II (Landfill Areas)

- Drilling and sampling four angled boreholes to a maximum of about 95 ft BGL beneath the Radioactive Waste Landfill (RWL) and one 30-ft deep borehole at the former Chemical Disposal Pit (CDP) outside of, and adjacent to, the RWL fence; and
- Drilling and sampling two angled boreholes to about 95 ft BGL beneath the Classified Waste Landfill (CWL) and five 50-ft boreholes, one at each American Car and Foundry (ACF) pit and/or cut-and-fill trench within the CWL.

Thus, six angled boreholes and 6 non-angled boreholes will be drilled and sampled in the landfill portions of TA-II (i.e., eastern portion of TA-II). A total of twenty-five boreholes will be drilled throughout TA-II as part of the Phase I drilling activities.

In addition to drilling and sampling boreholes during Phase I, surface and nearsurface soil samples also will be collected from the vicinity of Building 935. The scope of work for this sampling event will involve:

• Collecting 13 surface (0 to 0.5 ft deep) and 13 near-surface (3 to 5 ft deep) soil samples in the immediate vicinity of Building 935 (Table to be added later).

The purposes of the Phase I work are to:

- Drill and sample boreholes in the vicinity of Environmental Restoration (ER) sites in support of the DOE-approved RFI Workplan and NFA proposals;
- Determine if any potential contaminants are present in soil near the ACF pits and/or trenches and beneath part of the CWL;
- Determine concentrations of TCE in soil in the vicinity of Buildings 913 and 935; and
- Justify no further action (NFA) proposals for the 5 septic system ER sites, if appropriate.

The scope of work for Phase II will involve:

- Installing up to 3 deep boreholes within TA-II and completing them as monitor wells. One of the three boreholes drilled in the CWL last March may be advanced to the first water-bearing zone at approximately 330 ft BGL. The other two boreholes will be installed at the apexes of TA-II: one at the northern apex, possibly near Building 915, and one at the southern apex, possibly near Building 913. Both of these also will be installed in the first water-bearing zone at about 330 ft BGL; and
- Conducting geophysical logging at each potential well location.

The drilling method(s) for the Phase II monitor wells will be determined from the results of the Phase I investigation and from the availability of drill rigs.

The purposes of Phase II are to:

- Determine if ground water has been impacted by potential contaminants in soil in the vicinity of the CWL and Building 913; and
- Determine if the ground water flow direction and gradient are consistent throughout TA-II.

Geophysical logs (e.g., neutron, caliper, density, EMI) will be performed prior to installing each monitor well to confirm the location of the first water-bearing zone and evaluate the integrity of the borehole for well completion.

# **Project Personnel**

A site-specific Health and Safety Plan (HASP) has been developed and approved by DOE and Sandia Health and Safety as part of the TA-II Workplan for the planned RFI fieldwork activities at TA-II. The Technical Task Leader for the drilling activities is Rarilee Conway of Department 7582. A designated Site Safety Officer (SSO) will be on-site for all drilling activities at TA-II. A contractor technician will conduct all field screening for volatile organic compound (VOC) vapors and radiosiotopes, and monitor overall site conditions and drilling equipment. One or more designated health physics (HP) technicians will conduct field-screening for radioisotopes. The boreholes will be logged and sampled by Tom Tharp, Michael Wade, Rarilee Conway, or some other designated geologist.

The technician will conduct soil and/or ground water sampling, fill out chain-ofcustody (COC) forms, perform health and safety monitoring, obtain field sampling equipment and sample jars, deliver samples to the Sample Management Office (SMO), log lithologies, and conduct any other related field work. The technician also will provide field-screening for VOCs with a Photoionization detector (PID) or, if short-chained hydrocarbons are thought to be present, a Flame ionization detector (FID) (i.e., sample equipment, core, etc.). Tom Tharp and/or Michael Wade also will assist with sampling activities. Sandia will provide an Industrial Hygienist (IH) technician and an health physicist (HP) technician to monitor drilling and sampling equipment, soil and samples, and overall field conditions (i.e., temperature, cold stress, weather, etc.) for health and safety concerns. At the CWL and RWL, the HP will be available for the first 30 to 50 ft of drilling vertical (i.e., the primary zone of potential contamination). The HP will let the field team know when it is no longer necessary for continued radiation field-screening and/or upscaled personnel protective equipment. Potential project personnel and their phone numbers are listed in workplan HASP.

If any laboratory questions arise regarding sample containers, sample quantity, holding times, etc., the following people will be contacted from the field to reduce time and receive immediate technical advice:

Bob Friberg or a Sample Coordinator at TMA/Eberline (or Jim Lozito): 505-345-3461. Ellen LaRiviere, Quanterra (previously ENSECO): 303-421-6611.

Mike Gonzalez, Sandia SMO (848-0404).

Samples collected for VOCs, semi-volatile organic compounds (SVOCs), HE compounds, total cyanide, and/or Target Analyte List (TAL) metals will be submitted to Quanterra in Colorado unless otherwise notified. Radiological samples (total uranium, tritium, and photon emitters (gamma spectroscopy) will be submitted to TMA/Eberline in Albuquerque. In addition, soil will be collected in a plastic marinelli beaker and screened by Amir Monagheghi at the Health Physics Laboratory, Department 7715 (Radiation Protection Measurements Department) for radiological screening, but ONLY in areas where potential radioisotopes may be encountered (i.e., Building 935, CWL, and the RWL only). These samples will be collected at every interval where soil samples are collected for pre-laboratory screening purposes. Equipment rinse blanks also will be collected at various times in the field, such as after completing a borehole, and will be submitted to the analytical laboratories. The frequency of collecting equipment blanks will be determined in the field by the sampling team but will be at least once pre borehole.

The historical backgrounds for the ER sites associated with this project are available on request. The HASP provides additional information concerning potential contaminants of concern at each site. Details of the Phase I SAPs for each ER site are presented below.

# Phase I Drilling and Soil Sampling

Phase I activities involve drilling 25 boreholes from 30 to 135 ft deep at several locations within TA-II. These activities are described separately below. Because of ongoing activities at TA-II, drilling during the weekend may be necessary at the CWL and RWL. This is to avoid the potential for shrapnel and debris striking the drill rig (or Tom Tharp and Michael Wade) during TA-II testing activities. However, the drilling activities will be conducted primarily during weekdays and no other weekend field activities are planned.

Schedule for Phase I drilling activities at TA-II beginning Tuesday, November 1, 1994. Please note that the schedule may change due to potential activities at TA-II and/or any access problems.

Field Activity Dates (Estimated)	Borehole	Location
Tuesday, November 1, 1994	TA2-BH-11	Drill 50-ft borehole west of Buildings 915 and 922 - septic leachfield area; non-ER site; BTEX in soil vapor.
Wednesday, November 2	TA2-BH-08	Drill 50-ft borehole west of Building 907; septic leachfield area.
Thursday, November 3	TA2-BH-06	Drill 50-ft borehole west of Building 904; septic leachfield area.
Friday, November 4	TA2-BH-10	Drill 100-ft borehole southwest of Building 913; power line for night lighting shut-off for day; drilling at high TCE soil vapor location. Must complete this in one day.
Saturday, November 5	TA2-BH-09 (935) and TA2-BH-16 (907)	Drill 2 boreholes today - power shut off at 8 am (back on by 6 pm). Drill 30-ft borehole east of Building 935 at TCE soil vapor spot; drill 50-ft borehole north of Building 920 at Building 907 HE catch basin. Must finish both today due to power shut off in the area - weekend only.
Sunday, November 6	TA2-BH-17 and TA2-BH-18 (both at Building 940)	Drill 2 boreholes today - Both will be 50-ft deep near the Building 940 septic tank and drywell areas. Must complete these today due to power shutoff in the area - weekend only.
Monday, November 7	NO Drilling	NO Drilling - Day off.
Tuesday, November 8	TA2-BH-07	Drill 50-ft borehole along Building 904 HE Drain Trench immediately east of Buildings 914/917.
Wednesday, November 9	TA2-BH-12	Drill 50-ft borehole east of Building 919 - eastern portion of TA-II but will not interfere with testing activities.
Thursday, November 10	TA2-BH-13	Drill 50-ft borehole east of Building 919 - eastern portion of TA-II but will not interfere with testing activities.
Friday, November 11, 1994	End Phase I Drilling	Any site cleanup; drill rig decon at TA-III decontamination pad, if necessary.

Please Note: All drill rod and sampling equipment decontamination will take place at each drilling site. The drilling operation will have a mobile decontamination vehicle and augers and split-spoon samplers will be steam-cleaned at each site. The decontamination water tank will be drained into 55-gallon drums and labeled as IDW until analytical results are received for each site. All work will be performed in Level D protection, but Level C equipment will be on-hand, if required. Please refer to the TA-II site-specific workplan, sampling and analysis plan, and/or waste management plan for more details about these activities.

The estimated schedule for Phase I drilling at TA-II is as follows, and assumes that no major drill rig or health and safety issues occur during fieldwork. In addition, the schedule assumes that a minimum of one 50-ft deep borehole will be drilled and sampled each day. If the drilling contractor has at least 150 ft of hollow stem augers

available, drill rig decontamination can be performed every few days (unless decontamination is conducted at each site each day).

If time and budget permit, a third moniter well will be installed in the vicinity of the northern apex of TA-II, near Buildings 915 and 922. Assuming 10 days to well completion, the estimated dates of drilling will be from Wednesday, February 1 through about February 15 (includes decontamination and demob time).

Field activity schedules may change depending on testing activities and/or security issues at TA-II, the availability of drill rigs, and schedule conflicts during the holiday season. Field work may be completed well ahead of schedule if TA-II testing activities don't affect drilling activities. Detailed SAP tables and brief descriptions of activities planned for each site are described separately below. In addition, the following sections describe field-screening methods and other activities that apply to most of the sites in general.

All drilling equipment and drill cuttings will be field-screened for VOCs with a PID or FID and/or for radioisotopes using alpha scintillation and G-M probes. If any potential COCs are identified above background levels, samples will be collected and submitted for analysis. Samples will be submitted for off-site analysis for QA/QC (i.e., duplicates and Matrix Spike/Matrix Spike Duplicate [MS/MSD], etc.). All samples will be preserved on ice, inlouding tritium (but not other radioisotope analyses unless if it easier to do so for transporting purposes). The drilling geologist may collect additional soil samples in more permeable zones or submit more samples for QA/QC analysis if determined necessary in the field. In addition, ground water samples may be collected from any perched water-bearing zone(s) encountered during drilling, although perched ground water is not expected at the shallow depths planned for Phase I drilling. Samples will be collected, submitted, and analyzed for the potential COCs as listed in the tables in the following sections. To minimize the potential for cross-contamination, all sampling equipment will be decontaminated according to ER Operating Procedures (FOP) 94-57 (i.e., a mixture of water and Alconox soap followed by deionized water). Each borehole will be backfilled with grout after it has been drilled to the total depth.

All tritium samples will be collected in 16 oz. glass jars or plastic bottles as preferred by TMA/Eberline. Any samples collected for isotopic uranium and/or plutonium

will be collected with the tritium and submitted for all three analyses as per TMA. This will reduce sample containers, filling out COCs, and sample collection time. In addition, other samples sent to Quanterra can be combined into one liner. For example, SVOCs, HE compounds, and TAL metals analyses can all be collected into one 2-inch diameter by 6-inch long stainless steel liner and submitted as such to the lab.

In general, all soil samples will be collected in a driven split-spoon sampler (typically a 2-in. diameter) lined with stainless steel liners. The liners will then be sealed with Teflon sheets, plastic end caps, and inert duct tape. The samples will then be labeled with the appropriate I.D. (i.e., borehole number and depth) and placed on ice. Collecting samples in liners via split-spoons also was performed during TA-II drilling during March and June 1994 and at the Kauai Test Facility site in April. This is the best technical method to collect undisturbed samples, especially for VOCs and SVOCs. Although noted in the tables in the following sections, the preferred liner size and appropriate analyses for soil are as follows (as per Quanterra and TMA/Eberline):

Analysis	Minimum Stainless Steel Liner Length (inches) <sup>a</sup>	Minimum Glass Sample Jar Size
VOCs	3	
Total Cyanide	3	
TAL metals, HE compounds, and SVOCs	6	
TAL metals, HE compounds, and Total Cyanide	6	
Tritium <sup>b</sup>		250 ml
Gamma spectroscopy and total uranium		500 ml
PCBs, SVOCs, and HE compounds	6	
Tritium, isotopic uranium, and isotopic plutonium		500 ml plastic or glass jar (16 oz.)

a - The stainless steel liners are typically 3 inches or 6 inches long and 2 inches in diameter.

NOTE: Miranelli beakers will be collected for radiological screening at each interval where samples are collected for gamma spectroscopy and/or total uranium at RMMA sites only.

b - Liquid scintillation counter method.

The tritium samples and any other radiological samples can be collected in the split-spoon sampler and pushed into a jar/bottle since volatilization is not an issue or collected directly from a non-lined split-spoon sampler. In addition, most of the septic system ER sites will only require collecting VOCs and SVOCs in the first 15 to 20 ft to confirm the results of the passive SVSs. Therefore, there will be more sample material for the other analyses.

Unless contaminated soil and/or water is encountered during drilling activities, no additional soil samples will be collected during drilling. The field team has the discretion to collect additional samples at any time during these activities. The analytical results from samples collected during drilling activities will be used for waste characterization.

The following sections present the SAPs for each Phase I site and include sampling and analysis tables for all field activities.

All aqueous samples (including equipment blanks) will be sent to an off-site laboratory. For equipment blank (EB) and other aqueous samples, the following minimum quantities of water and bottle types/sizes have been requested by the analytical laboratories (Quanterra; TMA) (RCRA analytical holding times in parentheses):

TAL Metals One 500 ml poly. bottle with nitric preservative (180 days)

HE Compounds Two 1-liter amber glass bottles (7 days) SVOCs Two 1-liter amber glass bottles (7 days)

VOCs Three 40 ml VOAs (14 days)
Total Cyanide One 8 ounce poly. bottle (14 days)

Mercury One 250 ml glass bottle (preferred) with sodium hydroxide

preservative (13 days in plastic bottle; 28 days in glass bottle)

Tritium One 1-liter amber glass bottle (none)

# Building 904

Two boreholes (TA2-BH-06 and TA2-BH-07) will be drilled in the vicinity of Building 904 (see figure). Borehole TA2-BH-06 will be drilled adjacent to the septic system leachfield; borehole TA2-BH-07 will be drilled along the former HE drain

trench. Both boreholes will be drilled with a hollow stem auger drill rig and samples will be collected with a split-spoon sampler. The lithology will be described from drill cuttings and split-spoon samples.

At borehole TA2-BH-06 (leachfield), soil samples will be collected for analysis at the following depth intervals: 5, 10, 15, 20, 30, 40, and 50 ft. As listed in detail in Table 1a, the soil samples will be analyzed for SVOCs, total cyanide, high explosives, gamma spectroscopy, TAL metals, tritium, and VOCs. No VOCs or SVOCs were identified from the passive SVS investigation in the leachfield area. However, limited confirmatory sampling will be done for VOCs between 5 and 20 ft and SVOCs between 10 and 20 ft.

At borehole TA2-BH-07 (drain trench), soil samples will be collected for analysis at the following depth intervals: 5, 10, 15, 20, 30, 40, and 50 ft. As listed in detail in Table 1b, the soil samples will be analyzed for SVOCs, total cyanide, high explosives, gamma spectroscopy, TAL metals, tritium, and VOCs. Limited confirmatory sampling will be done for VOCs between 5 and 20 ft and SVOCs between 10 and 20 ft.

# **Building 907**

Two boreholes (TA2-BH-08 and TA2-BH-09) will be drilled in the vicinity of Building 907 (see attached figures). Borehole TA2-BH-08 will be drilled adjacent to the septic system leachfield; borehole TA2-BH-09 will be drilled near the HE catch box. Both boreholes will be drilled with a hollow stem auger drill rig and samples will be collected with a split-spoon sampler. The lithology will be described from drill cuttings and split-spoon samples.

At borehole TA2-BH-08, soil samples will be collected for analysis at the following depth intervals: 5, 10, 15, 20, 30, 40, and 50 ft. As listed in detail in Table 2a, the soil samples will be analyzed for SVOCs, total cyanide, high explosives, gamma spectroscopy, TAL metals, tritium, and VOCs. No VOCs or SVOCs were identified from the passive SVS investigation in the leachfield area. Limited confirmatory sampling will be done for VOCs and SVOCs between 5 and 15 ft and 10 and 20 ft, respectively.

At borehole TA2-BH-09, soil samples will be collected for analysis at the following depth intervals: 5, 10, 15, 20, 30, 40, and 50 ft. As listed in detail in Table 2b, the soil samples will be analyzed for SVOCs, total cyanide, high explosives, gamma spectroscopy, TAL metals, tritium, and VOCs.

# **Building 913**

This location has been selected based on the results of a passive SVS. The SVS investigation identified TCE in soil vapor south-southwest of Building 913. This area is **not** designated as an ER site. One borehole (TA2-BH-10) will be drilled at the location of the highest TCE soil vapor point (see figure). The borehole will be drilled with a hollow stem auger drill rig and samples will be collected with a split-spoon sampler. The lithology will be described from drill cuttings and split-spoon samples.

Soil samples will be collected for analysis at the following depth intervals: 5, 10, 15, 20, 30, 40, 50, 75, and 100 ft. As listed in detail in Table 3, the soil samples will be analyzed for SVOCs, TAL metals, tritium, and VOCs. No SVOCs were identified from the passive SVS investigation. However, soil samples will be collected and analyzed for SVOCs at 10, 15, and 20 ft. In addition, soil samples will be collected from selected depths and analyzed for VOCs by EPA Methods 8010 and 8020.

# Building 915/922

One borehole (TA2-BH-11) will be drilled in the vicinity west of Buildings 915 and 922. The borehole will be drilled adjacent to the septic system leachfield area (see attached figure). The borehole will be drilled with a hollow stem auger drill rig and samples will be collected with a split-spoon sampler. The lithology will be described from drill cuttings and split-spoon samples.

At borehole TA2-BH-11, soil samples will be collected for analysis at the following depth intervals: 5, 10, 15, 20, 30, 40, and 50 ft. As listed in detail in Table 4, the soil samples will be analyzed for SVOCs, high explosives, gamma spectroscopy, TAL metals, tritium, and VOCs. No SVOCs were identified from the passive SVS investigation in the leachfield area. However, limited confirmatory sampling will be done for SVOCs between 10 and 20 ft.

## **Building 919**

Two boreholes (TA2-BH-12 and TA2-BH-13) will be drilled in the vicinity east of Building 919. Each borehole will be drilled in the septic system leachfield area (see figure. Both boreholes will be drilled with a hollow stem auger drill rig and samples will be collected with a split-spoon sampler. The lithology will be described from drill cuttings and split-spoon samples.

At each borehole, soil samples will be collected for analysis at the following depth intervals: 5, 10, 15, 20, 30, 40, and 50 ft. As listed in detail in Table 5, the soil samples will be analyzed for high explosives, gamma spectroscopy, TAL metals, tritium, and VOCs. No VOCs or SVOCs were identified from the passive SVS investigation in the leachfield area. However, limited confirmatory sampling will be done for VOCs between 5 and 15 ft.

# **Building 935**

Three boreholes (TA2-BH-14, TA2-BH-15, and TA2-BH-16) will be drilled in the vicinity of Building 935. Borehole TA2-BH-14 will be drilled adjacent to the septic tank; borehole TA2-BH-15 will be drilled southwest of the former retention tank and dry well (see attached figure); and borehole TA2-BH-16 will be drilled southeast of Building 935 in the vicinity of high TCE concentrations in soil vapor. All three boreholes will be drilled with a hollow stem auger drill rig, two 50 ft deep and one to 30 ft deep (TA2-BH-16). Soil samples will be collected with a split-spoon sampler, and the lithology will be described from drill cuttings and split-spoon samples.

At boreholes TA2-BH-14 and TA2-BH-15 (Table 6a for both boreholes), soil samples will be collected for analysis at the following depth intervals: 5, 10, 15, 20, 30, 40, and 50 ft. As listed in detail in Table 6a, the soil samples will be analyzed for gamma spectroscopy, TAL metals, and tritium. No VOCs or SVOCs were identified from the passive SVS investigation in the leachfield area and confirmatory samples were collected during drilling activities in March 1994. These two boreholes are located within the Building 935 ER site and RMMA boundaries. Therefore, drilling will begin in Level C protection to at least 30 ft. The decision for downgrading from Level C to Level D protection will be decided by an HP.

At borehole TA2-BH-16, soil samples will be collected at the following depth intervals: 5, 10, 15, 20, and 30 (Table 6b). The samples will only be analyzed for VOCs by EPA Methods 8010 and 8020. Borehole TA2-BH-16 is located east of Building 935 and is not within the Building 935 ER site or the RMMA boundary.

# **Building 940**

Two boreholes (TA2-BH-17 and TA2-BH-18) will be drilled in the vicinity of Building 940. Borehole TA2-BH-17 will be drilled adjacent to the septic tank near the northwest side of the building. Borehole TA2-BH-18 will be drilled near the dry well southwest of the building. Both boreholes will be drilled with a hollow stem auger drill rig to 50 ft deep. Soil samples will be collected with a split-spoon sampler, and the lithology will be described from drill cuttings and split-spoon samples.

At both borehole locations, soil samples will be collected for analysis at the following depth intervals: 5, 10, 15, 20, 30, 40, and 50 ft. As listed in Table 7, the soil samples will be analyzed for HE compounds, gamma spectroscopy, TAL metals, tritium, and VOCs. No VOCs were identified from the passive SVS investigation in the leachfield area. However, limited confirmatory sampling will be done for VOCs between 5 and 15 ft.

## Radioactive Waste Landfill

Five boreholes (TA2-BH-19 through TA2-BH-23) will be drilled in the vicinity of the Radioactive Waste Landfill (RWL). Except for borehole TA2-BH-19, boreholes TA2-BH-20 through TA2-BH-23 will be angled.

Borehole TA2-BH-19 will be drilled to a depth of 30 ft in the location of a Chemical Disposal Pit (CDP) identified from historical air-photos. The CDP is located outside the northwest corner of the RWL. Although the borehole location is outside the RWL (an RMMA site), all drilling activities will be conducted as if it is an RMMA site. The borehole will be drilled with a hollow stem auger drill rig. Soil samples will be collected with a split-spoon sampler, and the lithology will be described from drill cuttings and split-spoon samples. At this borehole, soil samples will be

collected for analysis at the following depth intervals: 5, 10, 15, 20, and 30 ft. As listed in Table 8a, the soil samples will be analyzed for VOCs (confirmatory), gamma spectroscopy, TAL metals, isotopic uranium, and tritium.

The four angled boreholes will be drilled beneath trench and/or pit locations within the RWL. Each borehole however will be drilled from a minimum of 10 ft outside the RWL. Table RWL-1 shows the angles, lateral and vertical distances, and the total depth of each proposed borehole. The actual lengths of the angled boreholes range from 55 to 140 ft.

Table RWL-1. Approximate depths and angles for boreholes planned to be drilled beneath the RWL. Depths and/or angles may change depending on field conditions and sampling requirements.

Borehole Number	Angle (approximate degrees from vertical)	Lateral Distance (Approximate range in ft)	Total Length of Borehole (ft) (Approximate)	Total Depth (ft BGL) (Approximate)
TA2-BH-20	40	35	55	41
TA2-BH-21	45	40	55	40
TA2-BH-22	45	80	100	80
TA2-BH-23	45	80-100	100-140	80-95

At each of the four angled borehole locations soil samples will be collected for analysis at several depth intervals (see Tables 8b through 8e). In general, the soil samples will be analyzed for VOCs (confirmatory at some locations only), gamma spectroscopy, TAL metals, tritium, isotopic uranium, and isotopic plutonium.

## Classified Waste Landfill

Seven boreholes (TA2-BH-24 through TA2-BH-30) will be drilled in the vicinity of the Classified Waste Landfill (CWL); two of these boreholes (TA2-BH-29 and TA2-BH-30) will be angled.

Boreholes TA2-BH-24 through 28 each will be drilled 50 ft deep adjacent to four ACF pits and one ACF cut-and-fill trench. The ACF pits are reportedly 6 ft in diameter by 30 ft in depth; the cut-and-fill trench is 6-ft wide by 10-ft long by 12-ft deep. Each of these five boreholes will be drilled with a hollow stem auger drill rig to 50-ft deep. Soil samples will be collected with a split-spoon sampler, and the lithology will be

described from drill cuttings and split-spoon samples. At each of these five boreholes, soil samples will be collected for analysis at the following depth intervals: 5, 10, 15, 20, 30, 40, and 50 ft. As listed in Table 9a, the soil samples will be analyzed for HE compounds, SVOCs, isotopic uranium, gamma spectroscopy, TAL metals, tritium, PCBs, and VOCs. TCE, PCE, and BTEX were identified from the passive SVS investigations previously conducted in the CWL. However, two boreholes have already been drilled at the location of the two SVS "hot spots" and no VOCs were identified above detection limits. Limited confirmatory sampling will be done for VOCs at each of these boreholes.

The two angled boreholes will be drilled beneath trench locations within the CWL. One angled borehole (TA2-BH-30) will be drilled beneath a series of east-west oriented trenches (see Table CWL-1 below). This borehole will be drilled at an angle of 40 degrees from vertical to about 95 ft BGL (see Table CWL-1 below). The total length of the drilled borehole will be 125 ft. The other angled borehole (TA2-BH-29) will be drilled beneath a former pit and trench area (see attached Figure). This borehole will be drilled at about 40 degrees from vertical to about 60 ft BGL. The total length of the borehole will be about 75 ft.

Table CWL-1. Approximate depths and angles for boreholes planned to be drilled in the CWL. Depths and/or angles may change depending on field conditions and sampling requirements.

Borehole Number	Angle (degrees from vertical)	Lateral Distance (ft)	Total Length of Borehole (ft)	Total Depth (ft BGL)	
TA2-BH-29	40	50	75	60	
TA2-BH-30	40	95-100 ft	125	95	

At each of the two angled borehole locations, soil samples will be collected for analysis at several depth intervals (see Tables 9b and 9c). In general, the soil samples will be analyzed for VOCs, gamma spectroscopy, TAL metals, tritium, isotopic uranium, PCBs, SVOCs, and HE compounds.

# Workplan SAP Tables for Technical Area 2

The following SAP tables are for drilling and sampling activities to be conducted from October 1994 through about January 1995 at TA-2. Please note that five boreholes planned to be drilled at the CWL ACF pits have only one sampling and analysis table since the table will apply the same to all 5 boreholes (TA2-BH-24 through -28).

Table 1a. Summary of analyses for soil samples to be collected from borehole TA2-BH-06 drilled near the septic system leachfield west of Building 904, Technical Area 2.

Sample Type or QA/QC Type	Sample Depth	VOCsa	Tritium <sup>b</sup>	SVOCs <sup>c</sup>	TAL Metals <sup>d</sup>	HEe	Total Cyanide	Gamma Spec	Total # o Containe
QAQC Type	(in ft)				Wetais*		Cyannac	Spec	Containe
Subsurface soil	5	1	1				1	1	4
Subsurface soil	10	1	1	1	1	1	1	1	5
Subsurface soil	15	1	1	1	1	1	1	1	5
Subsurface soil	20		1	1	1	1	1	1	4
MS/MSD -	30								
Include on COC			<u> </u>						
Subsurface soil	30		1		1		1	1	4
Subsurface soil	40		1		1		1	1	4
Subsurface soil	50		1		1		1	1	4
Total Analyses		3	7	3	6	3	7	7	30 Total
36		<b>.</b>							Containe

- a EPA Method 8240. VOCs will be collected in 2-in. diameter by 3-in. long stainless steel liners.
- b Liquid scintillation counter method. Tritium will be collected in a split-spoon sampler and transferred into a 250 ml glass jar.
- c EPA Method 8270.
- d EPA Methods 6010 and 7000.
- e EPA Method 8330.

#### NOTE: This is NOT an RMMA site.

Note: SVOCs, High Explosives (HE), and TAL Metals samples will be collected into one 6-in. liner.

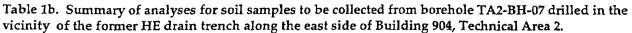
Note: A VOC and SVOC trip blank will be prepared and submitted for this borehole.

Note: All soil samples should be preserved on ice unless otherwise noted.

Note: All soil samples should be labeled as TA2-BH-06-depth.

Note: No soil will be collected from this borehole for a miranelli beaker.

Note: Equipment blanks will be collected after reaching the total depth of the borehole. The samples will be labeled as TA2-BH-06-EB. These samples should be collected in either a 2.5 liter amber glass jar or in a 1 liter bottle for SVOCs, HE, and TAL metals, and a 40 ml VOA for VOC samples.



Sample Type or QA/QC Type	Sample Depth (in ft)	VOCs <sup>a</sup>	Tritium <sup>b</sup>	SVOCs <sup>c</sup>	TAL Metals <sup>d</sup>	HEe	Total Cyanide	Gamma Spec	Total # o Containe
Subsurface soil	5	1	1	<b></b>			1	1	4
Subsurface soil	10	1	1	1	1	1	1	1	5
Subsurface soil	15	1	1	1	1	1	1	1	5
Subsurface soil	20		1	1	1	1	1	1	4
MS/MSD - Include on COC	30								
Subsurface soil	30		1		1	1	1	1	4
Subsurface soil	40		1		1	1	1	1	4
Subsurface soil	50	~	1		1	1	1	1	4.
Total Analyses 39		3	7	3	6	6	7	7	30 Total Containe

- a EPA Method 8240. VOCs will be collected in 2-in. diameter by 3-in. long stainless steel liners.
- b Liquid scintillation counter method. Tritium will be collected in a split-spoon sampler and transferred into a 250 ml glass jar.
- c EPA Method 8270.
- d EPA Methods 6010 and 7000.
- e EPA Method 8330.

Note: SVOCs, High Explosives, and TAL Metals samples will be collected into one 6-in. liner.

Note: A VOC and SVOC trip blank will be prepared and submitted for this borehole.

Note: All soil samples should be preserved on ice unless otherwise noted.

Note: All soil samples should be labeled as TA2-BH-07-depth.

Note: No soil will be collected from this borehole for a miranelli beaker.

Note: Equipment blanks will be collected after the total depth of the borehole has been drilled. The samples will be labeled as TA2-BH-07-EB. The samples should be collected in either a 2.5 liter amber glass jar or in a 1 liter bottle for SVOCs, HE, and TAL metals, and a 40 ml VOA for VOC samples.

Table 2a. Summary of analyses for soil samples to be collected from borehole TA2-BH-08 drilled near the septic system leachfield southwest of Building 907, Technical Area 2.

Sample Type or QA/QC Type	Sample Depth (in ft)	VOCs <sup>a</sup>	Tritium <sup>b</sup>	SVOC <sub>s</sub> c	TAL Metals <sup>d</sup>	HEe	Total Cyanide	Gamma Spec	Total # c Containe
Subsurface soil	5	1	1				1	1	4
Subsurface soil	10	1	1	1	1	1	1	1	5
Subsurface soil	15	1	1	1	1	1	1	1	5
Subsurface soil	20		1	1	1	1	1	1	4
MS/MSD - Include on COC	30								
Subsurface soil	30		1		1		1	1	4
Subsurface soil	40		1		1		1	1	4
Subsurface soil	50		1		1		1	1	4
Total Analyses 36		3	7	3	6	3	7	7	30 Total Containe

- a EPA Method 8240. VOCs will be collected in 2-in. diameter by 3-in. long stainless steel liners.
- b Liquid scintillation counter method. Tritium will be collected in a split-spoon sampler and transferred into a 250 ml glass jar.
- c EPA Method 8270.
- d EPA Methods 6010 and 7000.
- e EPA Method 8330.

Note: SVOCs, High Explosives, and TAL Metals samples will be collected into one 6-in. liner.

Note: A VOC and SVOC trip blank will be prepared and submitted for this borehole.

Note: All soil samples should be preserved on ice unless otherwise noted.

Note: All soil samples should be labeled as TA2-BH-08-depth.

Note: No soil will be collected from this borehole for a miranelli beaker.

Note: Equipment blanks will be collected after the total depth of the borehole has been reached. The samples will be labeled as TA2-BH-08-EB, and should be collected in either a 2.5 liter amber glass jar or in a 1 liter bottle for SVOCs, HE, and TAL metals. A 40 ml VOA will be used for VOC samples.

Table 2b. Summary of analyses for soil samples to be collected from borehole TA2-BH-09 drilled near the HE catch box along the HE drain trench south of Building 907, Technical Area 2.

Sample Type or QA/QC Type	Sample Depth (in ft)	VOCsa	Tritium <sup>b</sup>	SVOCs <sup>c</sup>	TAL Metals <sup>d</sup>	HEe	Total Cyanide	Gamma Spec	Total # o Containe
Subsurface soil	5	1	1				1	1	4
Subsurface soil	10	1	1	1	1	1	1	1	5
Subsurface soil	15	1	1	1	1	1	1	1	5
Subsurface soil	20		1	1	1	1	1	1	4
MS/MSD - Include on COC	30								
Subsurface soil	30		1		1	1	1	1	4
Subsurface soil	40		1		1	1	1	1	4
Subsurface soil	50	~	1		1	1	1	1	4
Total Analyses 39		3	7	3	6	6	7	7	30 Total Containe

- a EPA Method 8240. VOCs will be collected in 2-in. diameter by 3-in. long stainless steel liners.
- b Liquid scintillation counter method. Tritium will be collected in a split-spoon sampler and transferred into a 250 ml glass jar.
- c EPA Method 8270.
- d EPA Methods 6010 and 7000.
- e EPA Method 8330.

Note: SVOCs, High Explosives, and TAL Metals samples will be collected into one 6-in. liner.

Note: A VOC and SVOC trip blank will be prepared and submitted for this borehole.

Note: All soil samples should be preserved on ice unless otherwise noted.

Note: All soil samples should be labeled as TA2-BH-09-depth.

Note: No soil will be collected from this borehole for a miranelli beaker.

Note: Equipment blanks will be collected after the total depth of the borehole has been reached. The samples will be labeled as TA2-BH-09-EB, and should be collected in either a 2.5 liter amber glass jar or in a 1 liter bottle for SVOCs, HE, and TAL metals. A 40 ml VOA will be used for VOC samples.

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Table 3. Summary of analyses for soil samples to be collected from borehole TA2-BH-10 drilled south-southwest of Building 913, Technical Area 2.

Sample Type or QA/QC Type	Sample Depth (in ft)	VOCs <sup>a</sup>	Tritium <sup>b</sup>	SVOCs <sup>c</sup>	TAL Metals <sup>d</sup>	Total number of containers
Subsurface soil	5	1	1			2
Subsurface soil	10	1	1	1	1	3
Subsurface soil	10 <sup>e</sup>	1 <sup>e</sup>				1
Subsurface soil	15	1	1	1	1	3
Subsurface soil	20 <sup>e</sup>	1 <sup>e</sup>				1
Subsurface soil	20	1	1	1	1	3
MS/MSD - Include on COC	30					
Subsurface soil	30	1	1			2
Subsurface soil	40e	1 <sup>e</sup>				1
Subsurface soil	40	1	1			2
Subsurface soil	50	1	1			2
Subsurface soil	75	1	1			2
Subsurface soil	100 <sup>e</sup>	1e				1
Subsurface soil	100	1	1			2
Total Analyses 28	•••	13	9	3	3	25 total containers

- a EPA Methods 8010/8020. VOCs will be collected in 2-in. diameter by 3-in. long stainless steel liners.
- b Liquid scintillation counter method. Tritium will be collected in a split-spoon sampler and transferred into a 250 ml glass jar.
- c EPA Method 8270.
- d EPA Methods 6010 and 7000.
- e EPA Method 8240.
- NOTE: This is NOT an RMMA or an ER site.
- NOTE: SVOCs and TAL Metals samples will both be collected in one 6-in. liner.
- NOTE: A VOC and SVOC field blank will be prepared and submitted for this borehole.
- NOTE: All soil samples should be preserved on ice unless otherwise noted.
- NOTE: All soil samples should be labeled as TA2-BH-10-depth.
- NOTE: No soil will be collected from this borehole for a miranelli beaker.
- Note: Equipment blanks will be collected after the total depth of the borehole has been reached. The samples will be labeled as TA2-BH-10-EB, and should be collected in either a 2.5 liter amber glass jar or in a 1 liter bottle for SVOCs and TAL metals. A 40 ml VOA will be used for VOC samples.

Table 4. Summary of analyses for soil samples to be collected from borehole TA2-BH-11 drilled near the septic system leachfield southwest of Building 915/northwest of Building 922, Technical Area 2.

Sample Type or QA/QC Type	Sample Depth (in ft)	VOCsa	Tritiumb	SVOCs <sup>c</sup>	TAL Metals <sup>d</sup>	HEe	- Gamma Spec	Total # of Containers
Subsurface soil	5	1	1				1	3
Subsurface soil	10	1	1	1	1	1	1	4
Subsurface soil	15	1	1	1	1	1	1	4
Subsurface soil	20	1	1	1	1	1	1	4
MS/MSD - Include on COC	30							
Subsurface soil	30	1	1		1	1	1	4
Subsurface soil	40	1	1		1	1	1	4
Subsurface soil	50	1	1		1	1	1	4
Total Analyses 36		7	7	3	6	6	7	27 Total Containers

- a EPA Method 8240. VOCs will be collected in 2-in. diameter by 3-in. long stainless steel liners.
- b Liquid scintillation counter method. Tritium will be collected in a split-spoon sampler and transferred into a 250 ml glass jar.
- c EPA Method 8270.
- d EPA Methods 6010 and 7000.
- e EPA Method 8330.

Note: SVOCs, High Explosives, and TAL Metals samples will be collected into one 6-in. liner.

Note: A VOC and SVOC trip blank will be prepared and submitted for this borehole.

Note: All soil samples should be preserved on ice unless otherwise noted.

Note: All soil samples should be labeled as TA2-BH-11-depth.

Note: No soil will be collected from this borehole for a miranelli beaker.

Note: Equipment blanks will be collected after the total depth of the borehole has been reached. The samples will be labeled as TA2-BH-11-EB, and should be collected in either a 2.5 liter amber glass jar or in a 1 liter bottle for SVOCs, HE, and TAL metals. A 40 ml VOA will be used for VOC samples.

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Table 5. Summary of analyses for soil samples to be collected from boreholes TA2-BH-12 and TA2-BH-13 drilled in the septic system leachfield area east of Building 919, Technical Area 2. This table will be used for analyses at both boreholes.

Sample Type or QA/QC Type	Sample Depth (in ft)	VOCs <sup>a</sup>	Tritium <sup>b</sup>	TAL Metals <sup>c</sup>	HEd	Gamma Spec	Total # of Containers
Subsurface soil	5	1	1			1	4
Subsurface soil	10	1	1	1	1	1	5
Subsurface soil	15	1	1	1	1 .	1	5
Subsurface soil	20		1	1	1	1	4
MS/MSD - Include on COC	30						
Subsurface soil	30		1	1		1	4
Subsurface soil	40		1	1		1	4
Subsurface soil	50		1	1		1	4
Total Analyses 36		3	7	6	6	7	30 Total Containers

- a EPA Method 8240. VOCs will be collected in 2-in. diameter by 3-in. long stainless steel liners.
- b Liquid scintillation counter method. Tritium will be collected in a split-spoon sampler and transferred into a 250 ml glass jar.
- c EPA Methods 6010 and 7000.
- d EPA Method 8330.

Note: High Explosives (HE) and TAL Metals samples will be collected into one 6-in. liner.

Note: A VOC trip blank will be prepared and submitted for this borehole.

Note: All soil samples should be preserved on ice unless otherwise noted.

Note: All soil samples should be labeled as TA2-BH-12-depth (or TA2-BH-13-depth).

Note: No soil will be collected from this borehole for a miranelli beaker.

Note: Equipment blanks will be collected after the total depth of the borehole has been reached. The samples will be labeled as TA2-BH-12-EB (or -13-EB), and should be collected in either a 2.5 liter amber glass jar or in a 1 liter bottle for SVOCs, HE, and TAL metals. A 40 ml VOA will be

used for VOC samples.

Table 6a. Summary of analyses for soil samples to be collected from boreholes TA2-BH-14 and TA2-BH-15 drilled adjacent to the septic tank southeast of Building 935, Technical Area 2. This table will be used for analyses at both boreholes.

Sample Type or QA/QC Type	Sample Depth (in ft)	Tritium <sup>a</sup>	TAL Metals <sup>b</sup>	Gamma Spec	Total # of Containers
Subsurface soil	5	1		1	3
Subsurface soil	10	1	. 1	1	3
Subsurface soil	15	1	1	1	3
Subsurface soil	20	1	1	1	3
MS/MSD - Include on COC	30				
Subsurface soil	30	1	1	1	3
Subsurface soil	40	1	1	1	3
Subsurface soil	-50	1	1	1	3
Total Analyses 20		7	6	7	20 Total Containers

a - Liquid scintillation counter method. Tritium will be collected in a split-spoon sampler and transferred into a 250 ml glass jar.

#### NOTE: This IS an RMMA site.

Note: TAL Metals samples will be collected into one 6-in. liner.

Note: All soil samples should be preserved on ice unless otherwise noted.

Note: All soil samples should be labeled as TA2-BH-14-depth.

Note: Soil samples will be collected from this borehole at each sample location for a miranelli beaker and analyzed by Department 7715.

Note: Equipment blanks will be collected after the total depth of the borehole has been reached. The samples will be labeled as TA2-BH-14-EB, and should be collected in either a 2.5 liter amber glass jar or in a 1 liter bottle for TAL metals. A 40 ml VOA will be used for VOC samples.

b- EPA Methods 6010 and 7000.

Table 6b. Summary of analyses for soil samples to be collected from borehole TA2-BH-16 drilled in a soil vapor TCE "hot spot" east of Building 935, Technical Area 2.

Sample Type or QA/QC Type	Sample Depth (in ft)	VOCs by EPA Methods 8010 and 8020	Total # of Containers
Subsurface soil	5	1	1
Subsurface soil	10	1	1
Subsurface soil	15	1	1
Subsurface soil	20	1	1
MS/MSD - Include on COC	30		
Subsurface soil	30	1	1
Total Analyses 5		5	5 Total Containers

NOTE: This is NOT an RMMA or an ER site.

Note: Each sample will be collected into one 3-in. liner for each depth interval and the analyses labeled as 8010/8020.

Note: All soil samples should be preserved on ice unless otherwise noted. Note: A VOC field blank will be should be prepared for this borehole.

Note: All soil samples should be labeled as TA2-BH-16-depth.

Note: Soil samples will be collected from this borehole at each sample location for a miranelli beaker and analyzed by Department 7715.

Note: Equipment blanks will be collected after the total depth of the borehole has been reached. The samples will be labeled as TA2-BH-16-EB. A 40 ml VOA will be used for VOC samples.

Table 7. Summary of analyses for soil samples to be collected from boreholes TA2-BH-17 and TA2-BH-18 drilled near the septic tank on the west side of Building 940, Technical Area 2. This table will be used for both boreholes TA2-BH-17 and TA2-BH-18.

Sample Type or QA/QC Type	Sample Depth (in ft)	VOCsa	Tritium <sup>b</sup>	TAL Metals <sup>c</sup>	HEd	Gamma Spec	Total # of Containers
Subsurface soil	5	1	1			1	3
Subsurface soil	10	1	1	1	1	1	4
Subsurface soil	15	1	1	1	1	1	4
Subsurface soil	20		1	1	1	1	3
MS/MSD - Include on COC	30						
Subsurface soil	30		1	1	1	1	3
Subsurface soil	40		1	1	1	1	3
Subsurface soil	50		11	1	1	1	3
Total Analyses 29		3	7	6	6	7	23 Total Containers

- a EPA Method 8240. VOCs will be collected in 2-in. diameter by 3-in. long stainless steel liners.
- b Liquid scintillation counter method. Tritium will be collected in a split-spoon sampler and transferred into a 250 ml glass jar.
- c EPA Methods 6010 and 7000.
- d EPA Method 8330.

Note: High Explosives (HE) and TAL Metals samples will be collected into one 6-in. liner.

Note: A VOC trip blank will be prepared and submitted for this borehole.

Note: All soil samples should be preserved on ice unless otherwise noted.

Note: All soil samples should be labeled as TA2-BH-17-depth and/or TA2-BH-18-depth

Note: No soil will be collected from this borehole for a miranelli beaker.

Note: Equipment blanks will be collected after the borehole has been drilled to the total depth of about 50 ft. These samples will be labeled as TA2-BH-17-EB (or -18-EB) and should be collected in either a 2.5 liter amber glass jar or in a 1 liter bottle for TAL metals and a 40 ml VOA for VOC samples.

Table 8a. Summary of analyses for soil samples to be collected from angled borehole TA2-BH-19 drilled in the former Chemical Disposal Pit located near the Radioactive Waste Landfill, Technical Area 2.

Sample Type or QA/QC Type	Sample Depth (in ft)	VOCsa	Tritium <sup>b</sup>	Isotopic Uranium	gamma spec	TAL Metals <sup>c</sup>	Total # of Containers
Subsurface soil	5	1	1	1	1	1	4
Subsurface soil	10	1	1	1	1	1	4
Subsurface soil	15	1	1	1	1	1	4
MS/MSD - Include on COC	15						
Subsurface soil	20		1	1	1	1	3
Subsurface soil	30		11	1	1	1	3
Total Analyses 23		3	5	5	5	5	18 Total Containers

- a EPA Method 8240. VOCs will be collected in 2-in. diameter by 3-in. long stainless steel liners.
- b Liquid scintillation counter method. Tritium will be collected in a split-spoon sampler and transferred into a 500 ml glass jar or plastic bottle and analyzed with isotopic uranium.
- c EPA Methods 6010 and 7000.

NOTE: This is located outside the RWL and is not an RMMA site. However, the site will be considered as an RMMA site during this drilling event.

Note: TAL Metals samples will be collected into one 6-in. liner.

Note: All soil samples should be preserved on ice unless otherwise noted.

Note: All soil samples should be labeled as TA2-BH-19-depth.

Note: Gamma spectroscopy samples will be collected from this borehole in a miranelli beaker.

Note: Equipment blanks will be collected after the borehole has been drilled to the total depth of 30 ft.

These samples will be labeled as TA2-BH-19-EB and should be collected in either a 2.5 liter amber glass jar or in a 1 liter bottle for TAL metals and a 40 ml VOA for VOC samples.

Table 8b. Summary of analyses for soil samples to be collected from angled borehole TA2-BH-20 drilled beneath Pit 1 at the Radioactive Waste Landfill, Technical Area 2. Approximate angle is 40 degrees from vertical and 55 ft deep (41 ft BGL). Borehole will be drilled 10 ft from the RWL fence.

Sample Type or QA/QC Type	Sample Depth (in ft)	VOCs <sup>a</sup>	TAL Metals <sup>b</sup>	Tritium <sup>c</sup>	Isotopic Plutonium	Isotopic Uranium	Gamma Spec	Total # of Containers
Subsurface soil	20	1	1	1	1	1	1	4
Subsurface soil	30	1	1	1	1	1	1	4
MS/MSD - Include on COC	30							
Subsurface soil	40		1	1	1	1	1	3
Subsurface soil	50		1	1	1	1	1	_ 3
Total Analyses 22		2	4	4	4	4	4	14 Total Containers

- a EPA Method 8240. VOCs will be collected in 2-in. diameter by 3-in. long stainless steel liners.
- b EPA Methods 6010 and 7000.
- c Liquid scintillation counter method for tritium. Tritium will be collected in a split-spoon sampler and transferred into a 500 ml glass jar or plastic bottle and analyzed with isotopic uranium and plutonium.

NOTE: This IS an RMMA site (although the drill rig and sampling will be conducted outside of the RWL).

Note: TAL Metals samples will be collected into one 6-in. liner.

Note: All soil samples should be preserved on ice unless otherwise noted.

Note: All soil samples should be labeled as TA2-BH-20-depth.

Note: Tritium, isotopic uranium and isotopic plutonium all will be collected into one 500 ml plastic or glass jar.

Note: Equipment blanks will be collected after the borehole has been drilled to the total depth. These samples will be labeled as TA2-BH-20-EB and should be collected in a 2.5 liter amber glass jar or in a 1 liter bottle for TAL metals and a 40 ml VOA for VOC samples.

Table 8c. Summary of analyses for soil samples to be collected from angled borehole TA2-BH-21 drilled beneath Pit 2 at the Radioactive Waste Landfill, Technical Area 2. Approximate angle is 45 degrees from vertical and 55 ft deep (40 ft BGL). Borehole will be drilled 10 ft from the RWL fence.

Sample Type or QA/QC Type	Sample Depth (in ft)	VOCs <sup>a</sup>	TAL Metals <sup>b</sup>	Tritium <sup>C</sup>	Isotopic Plutonium	Isotopic Uranium	Gamma Spec	Total # of Containers
Subsurface soil	20	1	1	1	1	1	1	4
Subsurface soil	30	1	1	1	1	1	1	4
MS/MSD - Include on COC	30							3
Subsurface soil	40		1	1	1	1	1	
Subsurface soil	50		1	1	1	1	1	3
Total Analyses 22		2	4	4	4	4	4	14 Total Containers

- a EPA Method 8240. VOCs will be collected in 2-in. diameter by 3-in. long stainless steel liners.
- b EPA Methods 6010 and 7000.
- c Liquid scintillation counter method for tritium. Tritium will be collected in a split-spoon sampler and transferred into a 500 ml glass jar or plastic bottle and analyzed with isotopic uranium and plutonium.

#### NOTE: This IS an RMMA site.

Note: TAL Metals samples will be collected into one 6-in. liner.

Note: All soil samples should be preserved on ice unless otherwise noted.

Note: All soil samples should be labeled as TA2-BH-21-depth.

Note: Tritium, isotopic uranium, and isotopic plutonium all will be collected into one 500 ml plastic bottle or glass jar.

Note: Equipment blanks will be collected after the borehole has been drilled to the total depth. These samples will be labeled as TA2-BH-21-EB and should be collected in a 2.5 liter amber glass jar or in a 1 liter bottle for TAL metals and a 40 ml VOA for VOC samples.

Table 8d. Summary of analyses for soil samples to be collected from angled borehole TA2-BH-22 drilled beneath Trench 5 at the Radioactive Waste Landfill, Technical Area 2. Approximate angle is 45 degrees from vertical and 100 ft deep (80 ft BGL). Borehole will be drilled 10 ft from the RWL fence.

Sample Type	Sample	VOCsa	TAL	Tritium <sup>c</sup>	Isotopic	Isotopic	Gamma	Total # of
or QA/QC	Depth	VOCs*	Metalsb	Intium	Plutonium	Uraniu	Spec	Containers
Туре	(in ft)		Metais		riutomani	m	Spec	Containers
Subsurface soil	30	1	1	1	1	1	1	4
Subsurface soil	40	1	1	1	1	1	1	4
MS/MSD - Include on COC	<del>4</del> 0							
Subsurface soil	50		1	1	1	1	1	3
Subsurface soil	60		1	1	1	1	1	3
Subsurface soil	70		1	1	1	1	1	3
Subsurface soil	85		1	1	1	1	1	3
Subsurface soil	100		1	1	1	1	1	3
Total Analyses 37		2	7	7	7	7	7	23 Total Containers

- a EPA Method 8240. VOCs will be collected in 2-in. diameter by 3-in. long stainless steel liners.
- b EPA Methods 6010 and 7000.
- c Liquid scintillation counter method for tritium. Tritium will be collected in a split-spoon sampler and transferred into a 500 ml glass or plastic jar and analyzed with isotopic uranium and plutonium.

#### NOTE: This IS an RMMA site.

Note: TAL Metals samples will be collected into one 6-in. liner.

Note: All soil samples should be preserved on ice unless otherwise noted.

Note: All soil samples should be labeled as TA2-BH-22-depth.

Note: Tritium, isotopic uranium, and isotopic plutonium all will be collected into one 500 ml plastic

bottle or glass jar.

Note: Equipment blanks will be collected after the borehole has been drilled to the total depth. These samples will be labeled as TA2-BH-22-EB and should be collected in a 2.5 liter amber glass jar or in a 1 liter bottle for TAL metals and a 40 ml VOA for VOC samples.

Table 8e. Summary of analyses for soil samples to be collected from angled borehole TA2-BH-23 drilled beneath Trench 6 at the Radioactive Waste Landfill, Technical Area 2. Approximate angle is 45 degrees from vertical at a maximum of 135 ft deep and a minimum of 100 ft deep (i.e., 80 ft and 95 BGL, respectively).

Sample Type or QA/QC Type	Sample Depth (in ft)	TAL Metals <sup>a</sup>	Tritium <sup>b</sup>	Isotopic Plutonium	Isotopic Uranium	Gamma Spec	Total # of Containers
Subsurface soil	30	1	1	1	1	1	3
Subsurface soil	40	1	1	1	1	1	3
MS/MSD - Include on COC	40					***	
Subsurface soil	55	1	1	1	1	1	3.
Subsurface soil	70	1	1	1	1	1	3
Subsurface soil	85	1	1	1	1	1	3
Subsurface soil	100	1	1	1	1	1	3
Subsurface soil	120	1	1	1	1	1	3
Subsurface soil	135	1	1	1	1	1	3
Total Analyses 40		8	8	8	8	8	24 Total Containers

a - EPA Methods 6010 and 7000.

#### NOTE: This IS an RMMA site.

Note: Minimum length of borehole will be about 100 ft; maximum depth (if no auger refusal) will be 135 ft.

Note: TAL Metals samples will be collected into one 6-in. liner.

Note: All soil samples should be preserved on ice unless otherwise noted.

Note: All soil samples should be labeled as TA2-BH-23-depth.

Note: Tritium, isotopic uranium, and isotopic plutonium all will be collected into a 500 ml plastic bottle or glass jar.

Note: Equipment blanks will be collected after the borehole has been drilled to the total depth. These samples will be labeled as TA2-BH-23-EB and should be collected in a 2.5 liter amber glass jar or in a 1 liter bottle for TAL metals.

b- Liquid scintillation counter method for tritium. Tritium will be collected in a split-spoon sampler and transferred into a 500 ml glass jar or plastic bottle along with isotopic uranium and plutonium.

Table 9a. Summary of analyses for soil samples to be collected from five 50-ft deep boreholes planned to be drilled adjacent to the ACF pits in the Classified Waste Landfill, Technical Area 2. The five boreholes are TA2-BH-24, -25, -26, -27, and -28. (NOTE to SMO: this table will be applied to all five ACF boreholes; therefore, for the number of analyses and containers, multiply by 5. Also, multiply by 5 for containers/analyses for equipment and trip blanks).

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Sample Type or QA/QC Type	Sample Depth (in ft)	VOCsa	TAL Metals <sup>b</sup>	SVOCs <sup>c</sup>	HE	PCBs	Iso. U	Tritium	Gamma Spec	Total # of Containers
Subsurface soil	5	1	1	1	1	1	1	1	1	5
Subsurface soil	10	1	1	1	1	1	1	1	1	5
Subsurface soil	15	1	1	1	1	1	1	1	1	5
MS/MSD - Include on COC	15		~							
Subsurface soil	20		1	1	1	1	1	1	1	4
Subsurface soil	30		1	1	1	1	1	1	1	4
Subsurface soil	40		1	1	1	1	1	1	1	4
Subsurface soil	50		1	1	1	1	1	1	1	4
Total Analyses 52		3	7	7	7	7	7	7	7	31 Total Containers

- a EPA Method 8240. VOCs will be collected in 2-in. diameter by 3-in. long stainless steel liners.
- b EPA Methods 6010 and 7000.
- c Liquid scintillation counter method for tritium. Tritium will be collected in a split-spoon sampler and transferred into a 500 ml glass jar or plastic bottle along with isotopic uranium and plutonium.

#### NOTE: This is NOT an RMMA site.

Note: TAL Metals, SVOCs, and HE compound samples all will be collected into one 6-in. liner.

Note: PCBs will be collected into one 3-inch liner.

Note: All soil samples should be preserved on ice unless otherwise noted.

Note: Tritium and isotopic uranium will be collected into one 500 ml glass jar or plastic bottle.

Note: All soil samples should be labeled as TA2-BH-23-depth.

Note: Tritium, isotopic uranium, and isotopic plutonium all will be collected into a 500 ml plastic bottle or glass jar.

Note: Equipment blanks will be collected after the borehole has been drilled to the total depth. These samples will be labeled as TA2-BH-24-EB and should be collected in a 2.5 liter amber glass jar or in a 1 liter bottle for TAL metals and a 40 ml VOA for VOC samples (Subsequent ACF boreholes should be labeled as -25-EB; -26-EB; -27-EB; and -28-EB).

Table 9b. Summary of analyses for soil samples to be collected from angled borehole TA2-BH-29 drilled beneath pits and trenches at the Classified Waste Landfill, Technical Area 2. Approximate angle is 40 degrees from vertical and about 75 ft long (60 ft BGL).

Sample Type or QA/QC Type	Sample Depth (in ft)	VOCs <sup>a</sup>	TAL Metals <sup>b</sup>	SVOCs <sup>c</sup>	HE	PCBs	Iso. U	Tritium	Gamma Spec	Total # of Containers
Subsurface soil	20	1	1	1	1	1	1	1	1	5
Subsurface soil	30	1	1	1	1	1	1	1	1	5
Subsurface soil	40	1	1	1	1	1	1	1	1	5
MS/MSD - Include on COC	40									
Subsurface soil	50	1	1	1	1	1	1	1	1	5
Subsurface soil	60	1	1	1	1	1	1	1	1	5
Subsurface soil	70	1	1	1	1	1	1	1	1	5
Total Analyses 48		6	6	6	6	6	6	6	6	30 Total Containers

- a EPA Method 8240. VOCs will be collected in 2-in. diameter by 3-in. long stainless steel liners.
- b EPA Methods 6010 and 7000.
- c Liquid scintillation counter method for tritium. Tritium will be collected in a split-spoon sampler and transferred into a 500 ml glass jar or plastic bottle along with isotopic uranium.

NOTE: <u>Drilling and sampling will probably be conducted in Level C protection until decided otherwise</u> by the HP and SSO.

Note: TAL Metals, SVOCs, and HE compound samples all will be collected into one 6-in. liner.

Note: PCBs will be collected into one 3-inch liner.

Note: All soil samples should be preserved on ice unless otherwise noted.

Note: Tritium and isotopic uranium will be collected into one 500 ml glass jar or plastic bottle.

Note: All soil samples should be labeled as TA2-BH-29-depth.

Note: Tritium and isotopic uranium all will be collected into a 500 ml plastic bottle or glass jar.

Note: Equipment blanks will be collected after the borehole has been drilled to the total depth. These samples will be labeled as TA2-BH-29-EB and should be collected in a 2.5 liter amber glass jar or in a 1 liter bottle for TAL metals and a 40 ml VOA for VOC samples.

Table 9c. Summary of analyses for soil samples to be collected from angled borehole TA2-BH-30 drilled beneath pits and trenches at the Classified Waste Landfill, Technical Area 2. Approximate angle is 40 degrees from vertical and 125 ft long (95 ft BGL).

Sample Type or QA/QC Type	Sample Depth (in ft)	VOCs <sup>a</sup>	TAL Metals <sup>b</sup>	SVOCs <sup>¢</sup>	HE	PCBs	Iso. U	Tritium	Gamma Spec	Total # of Containers
Subsurface soil	30	1	1	1	1	1	1	1	1	5
Subsurface soil	45	1	1	1	1	1	1	1	1	5
Subsurface soil	60	1	1	1	1	1	1	1	1	5
MS/MSD - Include on COC	60									
Subsurface soil	75	1	1	1	1	1	1	1	1	. 5
Subsurface soil	90	1	1	1	1	1	1	1	1	5
Subsurface soil	115	1	1	1	1	1	1	1	1	5
Subsurface soil	125	1	1	1	1	1	1	1	1	5
Total Analyses 52		7	7	7	7	7	7	7	7	35 Total Containe:

- a EPA Method 8240. VOCs will be collected in 2-in. diameter by 3-in. long stainless steel liners.
- b EPA Methods 6010 and 7000.
- c Liquid scintillation counter method for tritium. Tritium will be collected in a split-spoon sampler and transferred into a 500 ml glass jar or plastic bottle along with isotopic uranium.

NOTE: <u>Drilling and sampling will probably be conducted in Level C protection until decided otherwise</u> by the HP and SSO.

Note: TAL Metals, SVOCs, and HE compound samples all will be collected into one 6-in. liner.

Note: PCBs will be collected into one 3-inch liner.

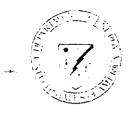
Note: All soil samples should be preserved on ice unless otherwise noted.

Note: Tritium and isotopic uranium will be collected into one 500 ml glass jar or plastic bottle.

Note: All soil samples should be labeled as TA2-BH-30-depth.

Note: Tritium and isotopic uranium all will be collected into a 500 ml plastic bottle or glass jar.

Note: Equipment blanks will be collected after the borehole has been drilled to the total depth. These samples will be labeled as TA2-BH-30-EB and should be collected in a 2.5 liter amber glass jar or in a 1 liter bottle for TAL metals and a 40 ml VOA for VOC samples.



# Department of Energy

Field Office, Albuquerque
Kirtland Area Office
P.O. Box 5400
Albuquerque, New Mexico 87115

OCT 17 1996

### CERTIFIED MAIL - RETURN RECEIPT REQUESTED

Mr. Benito Garcia, Bureau Chief New Mexico Environment Department Hazardous and Radioactive Materials Bureau 2044 Galisteo Street P.O. Box 26110 Santa Fe. NM 87505-2100

Dear Mr. Garcia:

Enclosed are two copies of the Sandia National Laboratories, New Mexico/Department of Energy (SNL/NM/DOE) response to the New Mexico Environment Department (NMED) technical comments on the 23 No Further Action (NFA) proposals submitted to NMED in June of 1995.

If you have any questions, please contact John Gould at (505) 845-6089, or Mark Jackson at (505) 845-6288.

Sincerely,

Michaeld Zamorski Acting Area Manager

### Enclosure

cc w/enclosure:

T. Trujillo, AL, ERD

W. Cox, SNL, MS 1147

N. Weber, NMED-AIP

R. Kern, NMED-AIP

D. Neleigh, EPA, Region 6 (2 copies)

cc w/o enclosure:

B. Oms, KAO-AIP

E. Krauss, SNL, MS 0141

B. Hoditschek, NMED

S. Dinwiddie, NMED

# Sandia National Laboratories Albuquerque, New Mexico October 1996

Environmental Restoration Project
Responses to NMED Technical Comments
on No Further Action Proposals
Dated June 1995

### INTRODUCTION

This document responds to comments received in a letter from the State of New Mexico Environment Department to the U.S. Department of Energy (Zamorski, July 29, 1996) documenting the review of 23 No Further Action (NFA) Proposals submitted in June 1995.

This response document is organized in numerical order by operable unit (OU) and subdivided in numerical order by site number, Each OU section provides NMED comments repeated in **bold** by comment number and by site number in the same order as provided in the call for response to comments. The DOE/SNL response is written in normal font style on a separate line under "Response". Responses to general technical comments begin on page 3 and responses to site-specific technical comments begin on page 4. Responses to general risk assessment comments begin on page 143 and responses to specific risk assessment comments begin on page 144. Additional supporting information for the site-specific comments is included as figures and tables within each comment response and as attachments to each section of this document.

SNL/NM ER Project October 1996 June 1995 NEA PROPOSES

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## RESPONSES TO NMED TECHNICAL COMMENTS ON NO FURTHER ACTION PROPOSALS DATED JUNE 1995

### GENERAL TECHNICAL COMMENTS

1. Please provide a Table of Contents so that the individual sites and their order of discussion can be more readily tracked.

<u>Response</u>: A Table of Contents is provided with each No Further Action Proposal submission sent to the regulators.

2. Information sources are listed for individual proposals within the section Sources of Supporting Information. Although the information sources might be useful for evaluation of the proposals, it is generally difficult to match the information source the referenced document. Information sources should be referenced.

<u>Response</u>: Citations in text to the references cited will be provided in future NFA proposals submissions and resubmissions.

3. The background soil sampling results should be submitted for NMED review.

Response: A Site-Wide statistical study for determining the background concentrations of metals and radionuclides in soil and water at Sandia National Laboratories/New Mexico and Kirtland Air Force Base has been recently completed and submitted to NMED in March 1996 (IT, 1996). These new background values were used to replace values provided for specific NFA proposals in this response.

4. Concerns exist over the sampling of the "septic system" solid waste management units (SWMUs). NMED believes the soil borings for drywells, seepage pits, or drain fields are inadequate. The proposal states that soil borings/samples were taken near the units (within 10 feet), but not underneath them. A sampling plan must be established to investigate underneath the seepage pits, drywells, or drain fields. Also, samples taken underneath the septic pipes/drain pipes need to be taken deeper than 3 feet.

Response: See Response to Site-Specific Technical Comment #1 below.

### 2. Site 136, OU 1303, Building 907 Septic System (TA-II)

- a. Have the septic tanks been removed?
- b. Elevated levels of barium and silver were detected in Trenches 1 and 2.
- c. Based on the results presented in Table 2, five (5) hazardous constituents (barium, beryllium, copper, nickel and zinc) were identified at concentrations in excess of the site-wide calculated UTL background concentration. The extent of contamination above background should be determined and the risk for constituents exceeding background and Screening Action Levels (SAL) should be evaluated.
- d. According to the proposal (page 5), an estimated 13 million gallons of effluent may have been discharged to the septic system. At the Liquid Waste Disposal System (LWDS), contaminants were not detected in soil boring samples, but were later detected and confirmed in groundwater 1994 Site Environmental Report (SAND95-1953)] Based upon the experience and the scenario for contaminant transport at the LWDS, it is possible that organic contaminants may not be detected in soil samples but, with the large historical fluid discharges, may have impacted the groundwater beneath TA-II.
- e. Please include the passive soil gas survey in the revised proposal.
- f. What was Sandia's logic in determining the number and the locations for the trenches?
- g. Page 5; paragraph 6: Did the vegetation continue linearly or was it only in spots? Please explain.
- h. NMED questions the value of the 1991 organic vapor survey held at waist height.
- i. Page 6; Unit Characteristics: What did the drain line consist of and how deep was the line laid? Also, please provide a more detailed map of Figure 2. In addition, please provide a more detailed schematic or map of the sanitary system and HE drain system which clearly identifies all components of the system. Please describe the construction of the seepage pit.

- j. For Tables 1 and 2: Please include the sampling results for all sampling intervals in the table, as well as the soil boring logs and the PID/FID readings for each interval.
- k. Why was the borehole TA2-BH-08 not drilled into the seepage pit? How far away was borehole TA2-BH-09 drilled from the catch box? Also, how and what is the catch box constructed of?
- 1. The most shallow borehole sample depth was 4 feet below ground level. Was there a rationale for this being the most shallow soil depth sampled?
- m. RECOMMENDATION: Based upon site concerns, including the hazardous constituent detections in soils at the site and in the perched groundwater (approximately 320 feet of depth) beneath TA-II, as well as the large historical discharges, NMED considers that NFA is not appropriate for Site 136. NMED considers that additional investigation is necessary at Site 136 and may require a RFI Workplan for this site.

Response: See Response to Site-Specific Technical Comment #1 above.

### 2. Site 136, Building 907, Septic System (TA II), OU 1303

The only chemical (non-radioactive compound) maximum concentrations that exceeded expected human health-based screening values are arsenic, beryllium and bis(2-ethylhexyl)phthalate. The reported values for arsenic and beryllium are within expected regional values. The value of 350 ppm for bis(2-ethylhexyl)phthalate is above the expected residential human health-based screening value of 46 ppm and approaching the industrial screening value of 410 ppm. Therefore, this constituent is the only one that remains a concern for this site. Additionally, the assumptions on the supporting information and data supplied in the report that applies to Site 48 would apply to Site 136.

The maximum concentrations reported for boreholes 08 and 09 (Table 2) are from different depths for the different constituents; therefore, the concentrations or exceedances of action levels cannot be compared across all constituents. Additionally, it is unclear whether the maximum concentration represents the maximum concentration across all depths for that constituent.

Response: See Response to Site-Specific Technical Comment #1 above.



U.S. Department of Energy Albuquerque Operations Office Kirtland Area Office P.O. Box 5400 Albuquerque, NM 87185-5400

UAN 2 8 2800

### CERTIFIED MAIL - RETURN RECEIPT REQUESTED

Mr. James Bearzi, Chief Hazardous and Radioactive Materials Bureau New Mexico Environment Department 2044 Galisteo Street P.O. Box 26110 Santa Fe, NM 87502-2100

Dear Mr. Bearzi:

Enclosed is one of two NMED copies of the Department of Energy and Sandia National Laboratories/New Mexico response to the NMED Notice of Deficiency (NOD), dated October 13, 1999, for Environmental Restoration sites 7, 46, 48, 50, 136, 159, 166, 227, 229, 230, 231, 233, 234, and 235. These sites were all included in the 2<sup>nd</sup> batch of No Further Action (NFA) proposals.

If you have any questions, please contact John Gould at (505) 845-6089.

Sincerely,

Michael J. Zamorski

Area Manager

Enclosure

# Sandia National Laboratories Albuquerque, New Mexico December 1999

Environmental Restoration Project Responses to NMED Notice of Deficiency No Further Action Proposals (2nd Round) Dated June 1995

### INTRODUCTION

Sandia National Laboratories/New Mexico (SNL/NM) is submitting this Notice of Deficiency (NOD) response for sites managed by the Tijeras Arroyo Operable Unit (OU) 1309 and the Technical Area (TA) II OU 1303. This response addresses Enclosures A and B comments in the October 13, 1999 NOD (NMED, 1999).

This is the second NOD response for Environmental Restoration (ER) Sites 50 and 235. Most of the following information addresses omissions in the ER Sites 50 and 235 No Further Action (NFA) Proposals (SNL/NM, 1995) and the first ER Sites 50 and 235 NOD responses (SNL/NM, 1996). This response addresses the need for reorganizing the confirmatory sampling analytical data and conducting human health and ecological risk assessments. For ER Site 50, this response also contains additional analytical data obtained during the Voluntary Corrective Measure activities recently conducted at nearby ER Site 228A (the Centrifuge Dump Site) in 1999 (SNL/NM, 1999). For ER Site 235, this response addresses the need for reorganizing the confirmatory sampling analytical data and conducting human and ecological risk assessments.

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### **Site-Specific Comments**

# RESPONSES TO NMED NOTICE OF DEFICIENCY COMMENTS ON NO FURTHER ACTION PROPOSALS ER SITES 7, 46, 48, 135, 136, 159, 165, 166, 167, 227, 229, 230, 231, 232, 233, AND 234 JUNE 1995 (2ND ROUND)

### **ENCLOSURE B**

The following discussion documents the negotiations between SNL/NM ER staff and NMED HRMB staff as requested in NMED (1999). These negotiations were finalized in a November 17, 1999 meeting.

### **OU 1303**

ER Sites 48, 135, 136, 159, 165, 166, and 167 (TA-2 Septic Systems)

Additional site characterization work proposed includes:

1. Finish compiling and provide the information requested in Stu Dindwiddie's letter to Michael Zamorski (DOE) and Joan Woodard (SNLNM) (dated December 11, 1998).

Response: The information requested in the referenced letter is listed below and is followed by the SNL/NM response.

a. Please submit maps showing the locations of boreholes with respect to seepage pits and other septic-system components for the above ER sites (48, 135, 136, 159, 165, 166, and 167).

Response: The existing site maps have been revised to reflect the best-known information on all the TA-II septic and drain system sites. The changes are based on SNL/NM Facilities Engineering drawings and Global Positioning System (GPS) mapping of visible system components. To improve the accuracy of the site maps, an excavator and GPS surveying will be used to locate system components below grade, confirm drainfield dimensions, and pinpoint effluent release locations. Planning for this work is in progress. Accurate site maps will be available in May 2000. Any further sampling at TA-II ER septic and drain system sites will be discussed with NMED HRMB staff when the maps are finalized. Note that this comment also addresses ER Sites 135 and 165, which were not incorporated in the 2nd Round of the NFA proposals. After discussions with NMED HRMB, the HE rinse-water drain from Site 48 will be investigated at the same time as co-located ER Sites 227 and 229, which are managed by Tijeras Arroyo OU 1309.

b. Please submit all analytical results of soil samples obtained from these boreholes. Data tables must include a listing of all constituents analyzed for, analytical methods, detection limits, and concentrations.

### **Site-Specific Comments**

<u>Response</u>: The requested soil analytical results for the boreholes at TA-II ER septic and drain system sites will be submitted with the revised site maps.

2. Summarize in written form, as applicable, all geologic, hydrologic, and ground-water quality data for all boreholes and ground-water monitor wells in the vicinity of TA-2.

Response: SNL/NM will summarize in written form, as applicable, all geologic, hydrologic, and groundwater quality data for all boreholes and groundwater monitor wells in the vicinity of the TA-II ER sites. This information will be presented in the Sandia North Groundwater Investigation Annual Report for FY01 or FY02.

\*



### **National Nuclear Security Administration**

Sandia Site Office P.O. Box 5400 Albuquerque, New Mexico 87185-5400



JUN 1 8 2004

### **CERTIFIED MAIL-RETURN RECEIPT REQUESTED**

Mr. John E. Kieling, Manager Permits Management Program Hazardous Waste Bureau New Mexico Environment Department 2905 Rodeo Park Rd., Building E Santa Fe, NM 87505

Dear Mr. Kieling,

On behalf of the Department of Energy (DOE) and Sandia Corporation, DOE is submitting the enclosed Solid Waste Management Unit (SWMU) Assessment Reports and Proposals for No Further Action (NFA) for Drain and Septic Systems (DSS) Sites 1010, 1028, 1083, and 1086. DOE is also submitting the Request for Supplemental Information (RSI) responses for SWMUs 48, 135, 136, 159, 165, 166, and 167; and a soil vapor summary report for Technical Area II at Sandia National Laboratories, New Mexico, EPA ID No. NM5890110518. These documents are compiled as DSS Round 5 and NFA Batch 23.

On April 29, 2004, the final Compliance Order on Consent (Consent Order) for Sandia National Laboratories was issued, replacing the HSWA Module as the sole enforceable mechanism for corrective action. The enclosed SWMU Assessment Reports/NFA Proposals and RSI responses were in the final stage of preparation when the Order was issued; thus, the enclosed documents contain language related to a NFA determination. We are requesting, consistent with the terminology in the Consent Order, an NMED determination of corrective action complete for each of these DSS sites.

This submittal includes descriptions of the site characterization work and risk assessments for DSS Sites 1010, 1028, 1083, and 1086, and SWMUs 48, 135, 136, 159, 165, 166, and 167. The risk assessments conclude that for these eleven sites: (1) there is no significant risk to human health under both the industrial and residential land-use scenarios; and (2) that there are no ecological risks associated with these sites.

Based on the information provided, DOE and Sandia are requesting a determination of corrective action complete without controls for these DSS sites.

If you have any questions, please contact John Gould at (505) 845-6089.

Sincerely,

Patty Wagner Manager

### **Enclosure**

### cc w/ enclosure:

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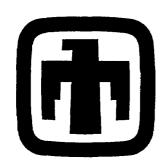
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# Sandia National Laboratories/New Mexico Environmental Restoration Project

# REQUEST FOR SUPPLEMENTAL INFORMATION RESPONSE FOR DRAIN AND SEPTIC SYSTEMS SWMU 136, BUILDING 907 SEPTIC SYSTEMS AT TECHNICAL AREA II

June 2004



United States Department of Energy Sandia Site Office

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- B DSS SWMU 136 HE Catch Box Analytical Data Summary Tables
- C DSS SWMU 136 Risk Assessment

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### **ACRONYMS AND ABBREVIATIONS**

AOP Administrative Operating Procedure

ARCH air-rotary casing hammer bgs below ground surface COC constituent of concern DSS Drain and Septic Systems

EB equipment blank

EPA U.S. Environmental Protection Agency

ER Environmental Restoration

g gram(s)

HE high explosive(s)
HI hazard index

HWB Hazardous Waste Bureau KAFB Kirtland Air Force Base

L liter(s)

MDA minimum detectable activity
MDL method detection limit

mrem millirem

NFA No Further Action

NMED New Mexico Environment Department

NOD Notice of Deficiency PCB polychlorinated biphenyl

pCi picocurie(s)

RCRA Resource Conservation and Recovery Act
RPSD Radiation Protection Sample Diagnostics
RSI Request for Supplemental Information
SNL/NM Sandia National Laboratories/New Mexico

SVOC semivolatile organic compound SWMU Solid Waste Management Unit

TA Technical Area

TAG Tijeras Arroyo Groundwater

TB trip blank

TCLP toxicity characteristic leaching procedure

TEDE total effective dose equivalent
TOP Technical Operating Procedure
VOC volatile organic compound

yr vear

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### 1.0 INTRODUCTION

### 1.1 Investigation History

In August 1994, no further action (NFA) proposals were submitted for Solid Waste Management Units (SWMUs) 135 and 165 in Technical Area (TA)-II at Sandia National Laboratories/New Mexico (SNL/NM). In July 1995, NFA proposals were also submitted for TA-II SWMUs 48, 136, 159, 166, and 167. These seven SWMUs are shown on Figure 1.1-1.

In November 1995, the New Mexico Environment Department (NMED) Hazardous Waste Bureau (HWB) responded with comments on the NFA proposals submitted for SWMUs 48, 136, 159, 166, and 167 and recommended that a Resource Conservation and Recovery Act (RCRA) Facility Investigation Work Plan, which included these SWMUs, be developed for TA-II. At that time, the SNL/NM Environmental Restoration (ER) Project decided to undertake the investigation and cleanup of these sites and others in TA-II as Voluntary Corrective Actions, and formal work plans were not submitted.

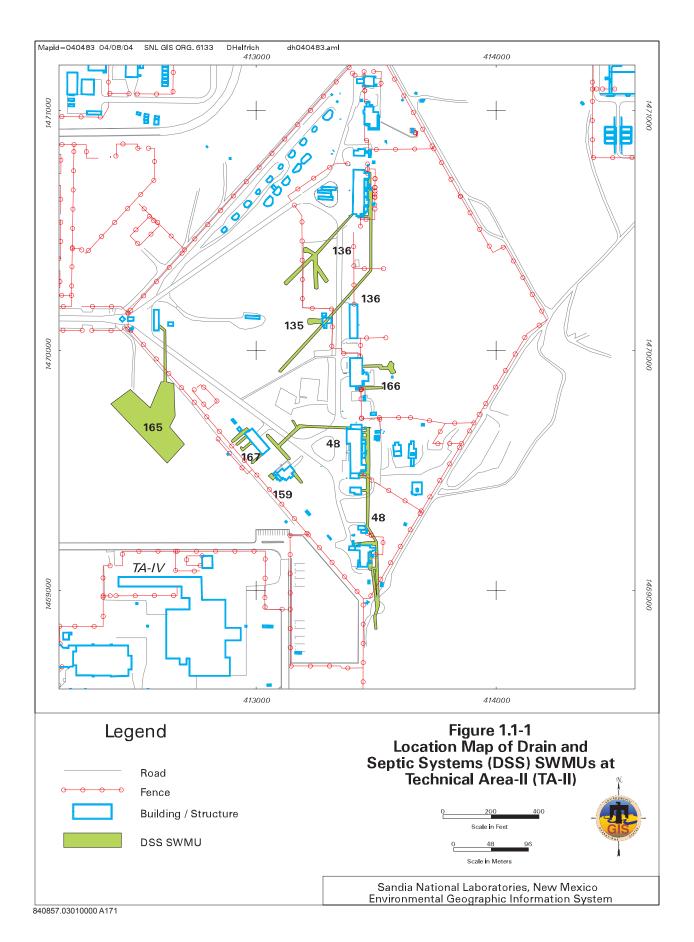
On October 13, 1999, the NMED-HWB issued a Notice of Deficiency (NOD) for these seven SWMUs. Negotiations on November 17, 1999, further defined specific procedures for sampling these seven SWMUs and transferred a requirement for groundwater reporting for these SWMUs to the ongoing Tijeras Arroyo Groundwater (TAG) Investigation. The NOD subsequently was changed by the NMED to a Request for Supplemental Information (RSI).

The requirements negotiated to fulfill the RSI for these seven TA-II SWMUs were:

- Submit revised site maps showing septic and drain system component locations (as determined by backhoe excavation).
- Submit the results for passive soil-vapor surveys and active soil-vapor monitoring wells at TA-II.
- Collect soil samples at a depth equal to the base, and 5 feet below the base, of septic tanks, seepage pits, and drain lines. Sample locations in drainfields and system outfalls were approved by HWB personnel.
- Analyze soil samples for volatile organic compounds (VOCs), semivolatile organic compounds (SVOCs), high explosive (HE) compounds, polychlorinated biphenyls (PCBs), RCRA metals, including hexavalent chromium, and total cyanide, radionuclides by gamma spectroscopy, and gross alpha/beta activity.
- Submit revised risk assessments for all seven SWMUs using all available soil data.

On January 26, 2000, the SNL/NM ER Project submitted a response to the NMED RSI, agreeing to excavations to locate system components below ground surface (bgs), confirm drainfield dimensions, pinpoint effluent release points, and investigate the SWMU 48 HE rinsewater drain line. SNL/NM also agreed to discuss additional sampling with the NMED-HWB when the maps were finalized and to submit the groundwater data requested in a subsequent TAG Investigation report.

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For tracking purposes, these seven SWMUs are included with sites listed in the SNL/NM Drain and Septic Systems (DSS) program reporting schedule. In this RSI response, they will be referred to as the "Drain and Septic Systems SWMUs at TA-II."

# 1.2 Additional Investigation Information

Although not specifically required as part of the RSI, this report presents additional information for several TA-II SWMUs as follows:

- In May 2003, soil-vapor monitoring wells were installed at SWMUs 159 and 165 as part of a separate site-wide DSS investigation. Additional details and sampling results for these wells are presented in the soil-vapor sampling chapter of this RSI response.
- Residual material in catch (settling) boxes for HE compound particulates located on HE rinse-water drain lines at SWMUs 48 and 136 was sampled as part of the site characterization process. The results are presented in the SWMU 48 and SWMU 136 chapters of this RSI response.

### 1.3 Report Organization

This RSI response presents the required information as follows:

- The soil-vapor survey information is presented as a whole and is not discussed on a site-by-site basis.
- Because NFA proposals were previously submitted for these SWMUs, only a brief
  description and history for each site is presented. Each SWMU is discussed in a
  separate report. The soil sampling analytical results and risk assessments for
  each site are presented in separate annexes for each SWMU.

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### 2.0 SOIL SAMPLING AT TA-II

## 2.1 Soil Sampling Methodology

Soil samples were collected at the TA-II DSS SWMUs using a variety of methods. Some shallow soil samples were collected from trenches excavated with a backhoe. For deep borehole sampling, either auger or air-rotary casing hammer (ARCH) drill rigs were used to drill down to the top of the desired sample interval. A drive sampler (split-spoon or thin-wall tube sampler) lined with stainless steel or brass sleeves was then mechanically advanced into the undisturbed soil below the drilled depth. For shallow soil sampling, a Geoprobe™ sample tube system with an inner butyl acetate liner was used through hollow-stem augers. The length of the recovered interval varied with the length of the sampling system, ranging from 2 feet using a split-spoon-type sampler, to up to 4 feet using a Geoprobe™ system. Following retrieval from the borehole, the sample for VOC analysis was collected by immediately capping and sealing either one of the metal liners from the split-spoon sampler or a cut portion of the butyl acetate liner from the Geoprobe™ sampler.

For the non-VOC analyses, the soil remaining in the sample sleeves or liner was emptied into a decontaminated mixing bowl, and aliquots of soil were transferred into appropriate sample containers for analysis. On occasion, the amount of soil recovered in the first sampling run was insufficient for sample volume requirements. In this case, additional sampling runs were completed until an adequate soil volume was recovered. Soil recovered from these additional runs was emptied into the mixing bowl and blended with the soil already collected. Aliquots of the blended soil were then transferred into sample containers and submitted for analysis.

All samples were documented and handled in accordance with applicable SNL/NM operating procedures and transported to on- and off-site laboratories for analysis.

### 2.2 Soil Sampling Events for DSS SWMUs at TA-II

In August and September 1992, 10 boreholes were drilled and sampled in the SWMU 165 drainfield. Samples were collected and analyzed for VOCs, SVOCs, HE compounds, metals, radionuclides by gamma spectroscopy, gross alpha/beta activity, and tritium. In November 1992, the groundwater monitoring well TA2-SW1-320 was installed in the shallow aquifer beneath the SWMU 165 drainfield, and soil samples collected from the borehole during drilling were analyzed for VOCs, SVOCs, PCBs, HE compounds, metals, cyanide, radionuclides by gamma spectroscopy, gross alpha/beta activity, and tritium.

In October and November 1993, trenches were excavated across septic and other drain system drain lines at SWMUs 48, 136, 165, and 166. At each trench-drain line intersection, samples were collected at three depths: the surface (0 to 0.5 feet bgs), at the top of the piping, and immediately below the piping. Samples were analyzed for VOCs, SVOCs, HE compounds, metals, radionuclides by gamma spectroscopy, and tritium.

From March to December 1994, 18 boreholes to depths of at least 50 feet were drilled throughout TA-II. The locations were chosen to be in and around the anomalies identified by the passive soil-vapor surveys, and also near the septic tanks, drain lines, and catch boxes that may have had releases. Fourteen borehole locations were near or within the seven SWMUs

addressed in this RSI response. The borehole locations are shown on the appropriate sample location maps for each SWMU. The SWMU 135 borehole, TA2-BH-01, was completed as groundwater monitoring well TA2-W-01. All borehole soil samples were analyzed for VOCs, SVOCs, HE compounds, metals, radionuclides by gamma spectroscopy, and tritium. Some samples were also analyzed for cyanide and gross alpha/beta activity.

In August 1995, soil samples were collected from borings drilled next to the septic tanks at SWMUs, 48, 136, 159, 165, 166, and 167 using a Geoprobe™ sampling system. Samples were collected starting at the approximate depth of the septic tank bottom and analyzed for VOCs, SVOCs, HE compounds, metals, including hexavalent chromium, and total cyanide, and radionuclides by gamma spectroscopy.

In August and October 2000, additional soil sampling was conducted at the seven TA-II SWMUs to fulfill the RSI requirements. Borehole soil samples were collected at depths starting at the base, and 5 feet below the base, of septic tanks, seepage pits, drywells, and septic drainfield drain lines. Sample locations in drainfields and system outfalls were approved by NMED-HWB personnel. The samples were analyzed for VOCs, SVOC, PCBs, HE compounds, RCRA metals, total cyanide, and radionuclides by gamma spectroscopy.

# 2.2.1 Soil Sampling Events at DSS SWMU 136

Soil samples were collected from nine boreholes, five trenches, and the HE catch box at DSS SWMU 136. The septic and HE waste-water drain line trench samples were collected in 1993 by filling sample containers with soil collected with a backhoe at three depths for each sample location. The deepest sample was collected at a depth equal to the base of the drain line. Boreholes adjacent to the septic tanks, septic system, and HE waste-water drain lines were drilled in 1994 and 1995 using a hollow-stem auger. Samples were collected using a hollow-stem auger and a 2-foot-long, split-spoon-type drive sampler. Samples were collected next to septic tanks at depths equal to, and below, the units using a Geoprobe<sup>TM</sup> sampling system inside hollow-stem augers.

Samples beneath the seepage pit and in the drainfields were collected in 2000 with a Geoprobe<sup>TM</sup> from two 3-foot-long sampling intervals at each boring location. Drainfield sampling intervals started at 10 and 15 feet bgs in the one drainfield lateral boring, and 8 and 13 feet bgs in the other two drainfield lateral borings. Sample intervals started at 13 and 18 feet bgs in the single boring through, and beneath, the seepage pit. Samples in the HE waste-water outfall ditch were collected near the surface (0.75 feet bgs) and at 2 and 7 feet bgs at the point where the HE drain line discharged to the outfall ditch, as well as at 4 and 9 feet bgs at two locations along the outfall ditch. Sample containers were filled with soil collected with a backhoe. The HE catch box was sampled at three locations along the bottom of the unit, at approximately 8 feet bgs.

The soil samples in 2000 and 2001 were collected in accordance with the procedures developed for, and described in, the Operable Unit 1295 Sampling and Analysis Plan (SNL/NM October 1999) and subsequent "Field Implementation Plan, Characterization of Non-Environmental Restoration Drain and Septic Systems" (SNL/NM November 2001) approved by the NMED. The 1994 and 1995 sampling activities were conducted using similar procedures.

### 3.0 DSS SWMU 136: BUILDING 907 SEPTIC SYSTEMS

### 3.1 Site Description

Building 907, located in the northern half of TA-II (Figure 3.1-1), was constructed about 1948 and used for a variety of purposes. The septic system was connected to the City of Albuquerque sanitary sewer system in 1992, and the building was demolished in October 2002. Additional information on the operational history for Building 907 can be found on the SNL/NM ER Project web page (SNL/NM January 2003) and in the original NFA proposal (SNL/NM June 1995). The Building 907 septic and HE drain line systems comprising SWMU 136 are shown on Figures 3.1-2 and 3.1-3. A summary of the drain systems investigated at Building 907 is presented in Table 3.1-1.

### 3.2 DSS SWMU 136 Soil Sampling Results and Conclusions

Soil sampling was conducted at DSS SWMU 136 as described in Section 2.2. Figures 3.1-2 and 3.1-3 show the soil sampling locations at DSS SWMU 136. The analytical data summary tables are presented in Annex A. Because there were several sampling events at this site, the results are grouped by general area or location in the analytical tables. The HE catch box investigation is discussed in Section 3.3.

### **VOCs**

VOC analytical results for soil samples collected from the DSS SWMU 136 boreholes and trenches are summarized in Table A-1, and method detection limits (MDLs) for the VOC soil analyses are presented in Table A-2. Seven VOCs were detected in the soil samples collected at this site. Three additional VOCs were detected only in equipment blanks (EBs) or trip blanks (TBs). Most of the VOCs detected are common laboratory contaminants and may not indicate soil contamination at this site.

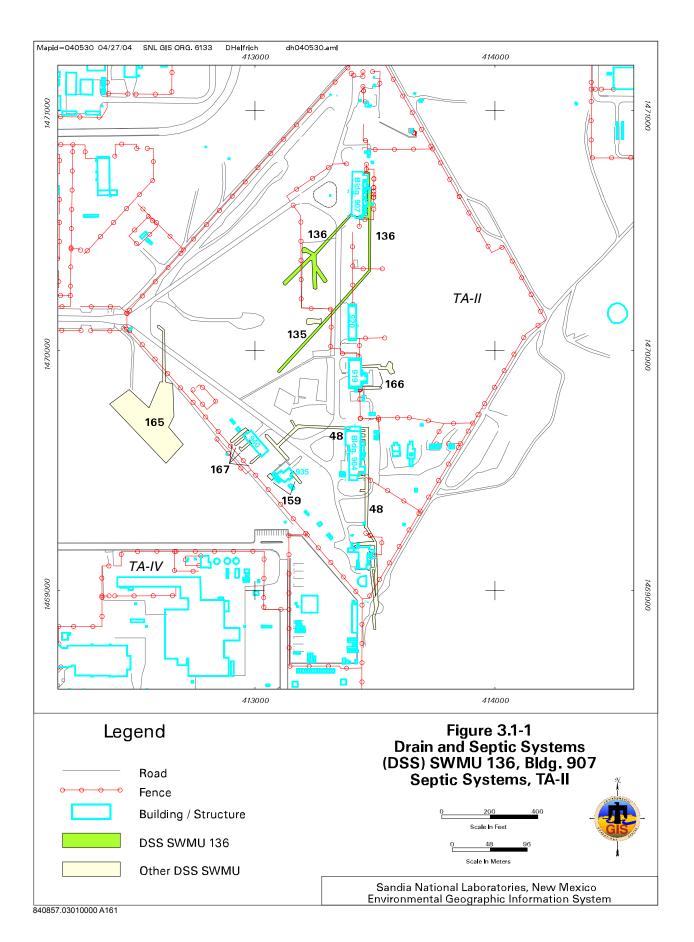
### **SVOCs**

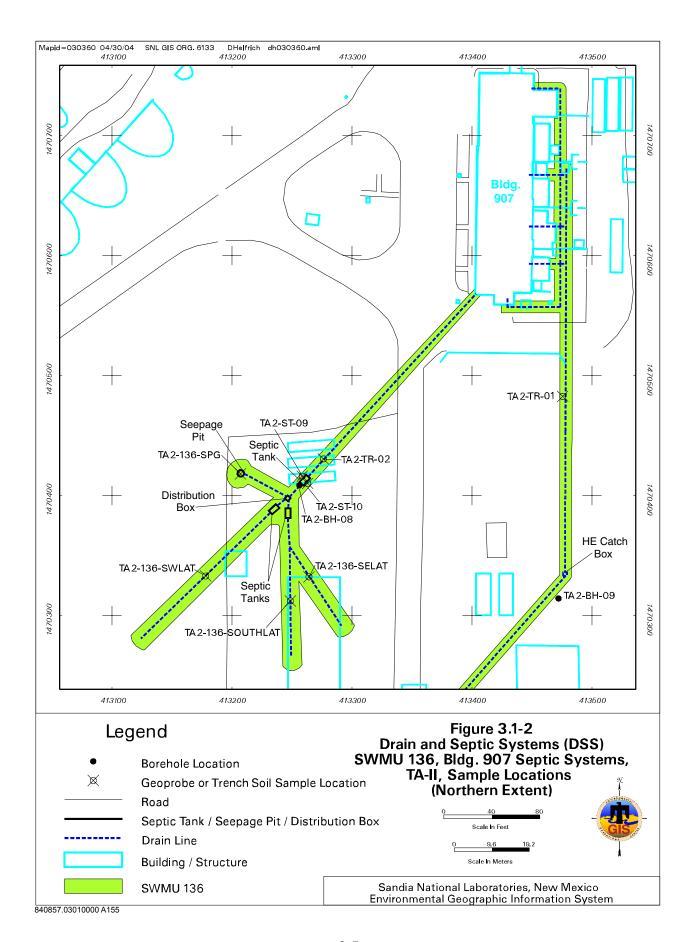
SVOC analytical results for the soil samples collected from the DSS SWMU 136 boreholes and trenches are summarized in Table A-3, and MDLs for the SVOC soil analyses are presented in Table A-4. A total of 12 SVOCs were detected in the soil samples. Eight SVOCs were detected in the 16.5-foot-bgs sample from borehole TA2-BH-08. Five SVOCs were detected in the 13-foot-bgs primary sample from the seepage pit borehole, TA2-136-SPG, but none were detected in the duplicate sample from the same interval.

### **PCBs**

PCB analytical results for the soil samples collected from the DSS SWMU 136 boreholes are summarized in Table A-5, and MDLs for the PCB analyses are presented in Table A-6. No PCBs were detected in any soil sample collected. Aroclor-1260 was detected in the EB for these samples.

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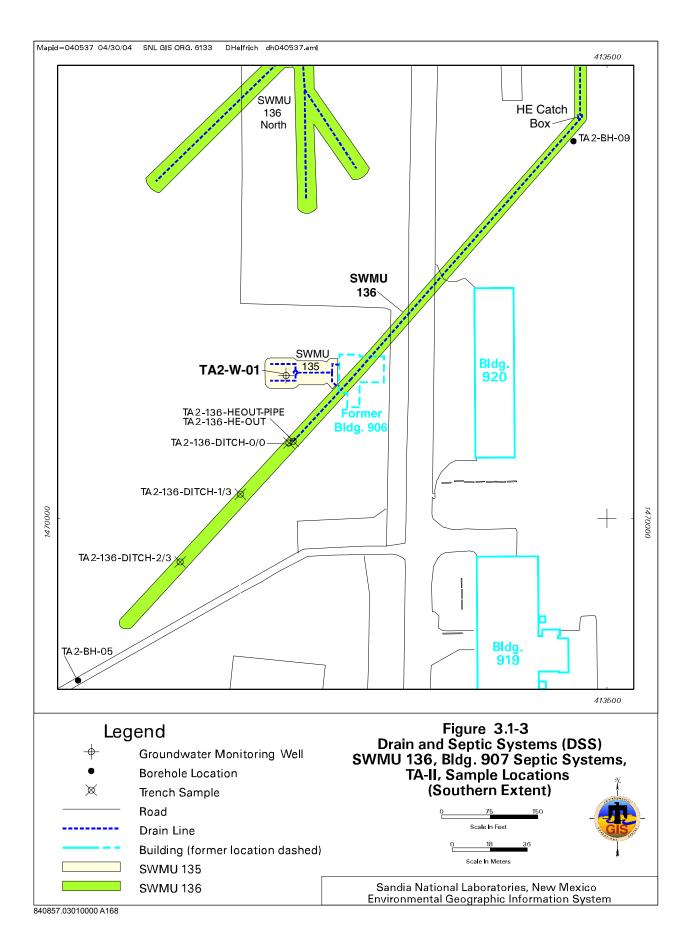


Table 3.1-1 Summary of DSS SWMU 136, Building 907 Septic Systems

	Systems		
Building 907	Operational	Ruilding 907 Drain Systems Descriptions	System Status
Dialiti Systems	1040 1000	The contine water consisted of a 450-nallon sentir tank	The Building 907 Septic System was
l-septic system	1940-1992	two 1000-gallon septic tanks located downstream of the first, a	connected to the City of Albuquerque
1-HE waste-		5-foot-diameter by 13-foot-deep seepage pit, and approximately	sanitary sewer system in 1992.
water drain		400 feet of leachfield drain piping (Figure 3.1-2).	:
svstem			The NMED inspected the septic system in
,		The HE waste-water drain system drained Building 907 assembly	late 1995. The northern (first) tank was
		bays and passed through a catch box to collect HE particulates	pumped out in February 1996, and the
		by mechanical filtration of the waste water. The catch box is	other two tanks were found to be dry at
		approximately 200 feet south of the building. The HE waste-	that time. Eight 55-gallon drums of septic
		water drain system extended an additional 675 feet to the	tank waste were generated.
		southwest, passed under Building 906, and discharged into a	
		200-foot-long, open ditch west of Building 919 (Figure 3.1-3). HE	The septic tank concrete was sampled
		residues would not have been flushed down the floor drains after	and confirmed to be clean in early 1996.
		the mid-1950s when weapons assembly operations were	The septic tank waste was characterized
		discontinued. The catch box reportedly partially collapsed in	and managed according to SNL/NM
		1991 or 1992	policy. The system was subsequently
			abandoned in place.
			The HE catch box was sampled in
			December 2001 and was abandoned in
			place by backfilling on June 23, 2003.

= Drain and Septic Systems.
= High explosive(s).
= New Mexico Environment Department.
= Sandia National Laboratories/New Mexico.
= Solid Waste Management Unit. DSS HE NMED SNL/NM SWMU

### **HE Compounds**

The HE compound analytical results for the soil samples collected from the DSS SWMU 136 boreholes and trenches are summarized in Table A-7, and MDLs for the HE compound soil analyses are presented in Table A-8. One HE compound was detected in the 6.67-foot-bgs sample collected in borehole TA2-ST-10, adjacent to the first septic tank.

### Metals, Including Hexavalent Chromium, and Cyanide

Metals, including hexavalent chromium, and cyanide analytical results for the soil samples collected from the DSS SWMU 136 boreholes and trenches are summarized in Table A-9, and MDLs for the metals and cyanide soil analyses are presented in Table A-10. Eleven metals were detected in these samples at concentrations above their respective NMED-approved background concentrations. The elevated detections were as follows:

- Lead and silver were detected in the drain line trench samples at concentrations above their respective NMED-approved background concentrations.
- Arsenic, barium, beryllium, chromium, cobalt, copper, and vanadium were
  detected in the borehole samples near the drain lines or septic tanks at
  concentrations above their respective NMED-approved background
  concentrations.
- Cadmium, chromium, lead, mercury, and silver were detected in the HE drain line outfall ditch samples at concentrations above their respective NMED-approved background concentrations.
- Arsenic, barium, chromium, and mercury were detected in the drainfield borehole samples at concentrations above their respective NMED-approved background concentrations.
- No elevated metals concentrations were detected in the seepage pit borehole samples.

Hexavalent chromium was not detected in the two samples that were analyzed for this metal.

Cyanide was not detected in any sample where an analysis for it was performed.

Six metals were detected at trace concentrations in the EBs associated with sampling of boreholes BH-05 and BH-08 in March and November 1994. Two metals were detected at trace concentrations in the EB associated with the August 2000 borehole sampling activities.

### Radionuclides

Analytical results for the gamma spectroscopy analysis of the soil samples collected from the DSS SWMU 136 boreholes and trenches are summarized in Table A-11. Three radioisotopes were detected at elevated activities in these samples. The results are as follows:

- Cesium-137 was detected in two HE drain line trench samples and one of the HE drain line outfall ditch samples at activities above the NMED-approved background.
- Thorium-232 was not detected in any sample above the NMED-approved background activity.
- Uranium-235 was detected in two of the HE drain line outfall ditch samples at activities above the NMED-approved background.
- Uranium-238 was not detected in any sample above the NMED-approved background activity.

However, although not detected, the minimum detectable activities (MDAs) for some uranium-235 and uranium-238 analyses exceed their respective background activities in several site soil samples because the standard gamma spectroscopy count time for soil samples (6,000 seconds) was not sufficient to reach the NMED-approved background activities established for SNL/NM soils (Dinwiddie September 1997). Even though the MDAs may be slightly elevated, they are still very low, and the risk assessment outcome for the site is not significantly impacted by their use.

### **Tritium**

Tritium analytical results for the soil samples collected from the DSS SWMU 136 boreholes and trenches are summarized in Table A-12. Tritium was detected in five of the borehole BH-09 samples at activities above the SNL/NM-established tritium background of 420 picocuries (pCi)/liter (L) (0.021 pCi/gram [g]) for soil (Tharp February 1999). Tritium MDAs exceed the SNL/NM-established background for soil in six samples from borehole BH-08. In contrast to the drain line trench samples, the tritium samples from boreholes BH-08 and BH-09 were analyzed by liquid scintillation counting, and the activity was reported in pCi/g instead of pCi/L. For comparison with the more common laboratory distillation values, these activities can be converted from pCi/g to pCi/L using the assumption of 5 percent soil moisture and a soil density of 1 g/cubic centimeter. Although the calculated MDA exceeds the SNL/NM-established background value for some analyses, these results still indicate no significant levels of tritium are present in the soil at the site.

A low tritium activity was measured in the EB associated with the sampling of borehole BH-05 in March 1994, but not in the EBs associated with the borehole BH-08 or BH-09 sampling in November 1994.

### 3.2.1 Soil Sampling Quality Assurance/Quality Control Samples and Data Validation

Quality assurance/quality control samples were collected according to the ER Project guidelines and operating procedures in effect at the time of sampling. These included duplicate, EB, and TB samples. EB samples were analyzed for the same analytical suite as the associated soil samples. TB samples, for VOCs only, were included with soil sample shipments sent to laboratories for VOC analyses. The analytical results for the EB samples appear only on the data tables for the site where they were collected. However, the results would have been used

in the data validation process for all the samples analyzed at that time. EB and TB results are discussed with the associated analytical results in Section 3.2.

As shown in the data summary tables in Annex A, to assess the precision and repeatability of sampling and analytical procedures, duplicate soil samples (designated "DU") were collected and analyzed for VOCs, SVOCs, PCBs, HE compounds, metals, cyanide, radionuclides by gamma spectroscopy, and tritium. The results were comparable for the primary and duplicate sample analyses and are summarized as follows:

- Similar concentrations of two of the three VOCs detected were in both the primary and duplicate 30.2-foot-interval samples from borehole BH-05. The analyte 2-butanone was detected only in the primary sample but not in the duplicate. Acetone was detected in both the primary and duplicate 13-foot samples from the seepage pit borehole, while 2-butanone was detected only in the duplicate sample (Table A-1).
- The SVOC bis(2-ethylhexyl) phthalate was detected at similar concentrations in both the primary and duplicate 30-foot sample from borehole BH-05. Five SVOCs were detected in the 13-foot-interval primary sample but none were detected in the duplicate from the seepage pit borehole (Table A-3).
- No PCBs were detected in either the 13-foot-interval primary or duplicate samples from the seepage pit borehole (Table A-5).
- No HE compounds were detected in either the 30-foot-interval primary or duplicate samples from borehole BH-05 or in the 13-foot-interval primary or duplicate samples in the seepage pit borehole (Table A-7).
- Metals concentrations were similar in both the 30-foot-interval primary and duplicate samples from borehole BH-05 and in the 13-foot-interval primary and duplicate samples in the seepage pit borehole (Table A-9).
- Gamma spectroscopy activities were similar in both the 30-foot-interval primary and duplicate samples from borehole BH-05. Uranium-235 was detected in the 13-foot-interval primary sample, but not in the duplicate sample from the seepage pit borehole (Table A-11).
- Low but somewhat dissimilar tritium activities were measured in both the 30-foot-interval primary and duplicate samples from borehole BH-05 (Table A-12).

All laboratory data were reviewed and verified/validated according to "Verification and Validation of Chemical and Radiochemical Data Technical Operating Procedure (TOP) 94-03," Rev. 0 (SNL/NM July 1994) or SNL/NM ER Project "Data Validation Procedure for Chemical and Radiochemical Data," in Administrative Operating Procedure (AOP) 00-03 (SNL/NM December 1999). In addition, SNL/NM Department 7713 (Radiation Protection Sample Diagnostics [RPSD] Laboratory) reviewed all on-site gamma spectroscopy results according to "Laboratory Data Review Guidelines," Procedure No. RPSD-02-11, Issue No. 2 (SNL/NM July 1996). The data are acceptable for use in this RSI response.

### 3.3 DSS SWMU 136 HE Catch Box Investigation

The DSS SWMU 136 HE catch box (Figures 3.1-2 and 3.1-3) had partially collapsed in 1991 or 1992. The material inside the unit was sampled in December 2001. Three samples of material from the base of the unit were collected and analyzed for VOCs, SVOCs, HE compounds, toxicity characteristic leaching procedure (TCLP) metals, radionuclides by gamma spectroscopy, and tritium. Low levels of VOCs, SVOCs, metals, and radionuclides including tritium were detected. No HE compounds were detected. The HE catch box analytical data are presented separately in Annex B.

During a meeting with NMED on June 3, 2003, the DSS SWMU 136 HE catch box sampling data were reviewed, and additional investigation requirements were decided (Bowles 2003). The requirements were:

- 1. Abandon the HE catch box in place by backfilling to grade with native soil and/or clean fill. Removal of the material in the unit was not required by the NMED.
- 2. Tabulate and evaluate all data associated with the site HE drain lines and catch boxes into the risk assessment.
- 3. Request an NFA determination based upon the completed site characterization work and the risk assessment results.

The HE catch box was subsequently backfilled with native soil on June 23, 2003. All site soil sampling data, including the HE catch box samples, were incorporated into the associated DSS SWMU 136 risk assessment (Annex C). This RSI response constitutes a request for NFA.

### 3.4 Site Sampling Data Gaps

Analytical data from the site assessment were sufficient for characterizing the nature and extent of possible constituent of concern (COC) releases. There are no further data gaps regarding characterization of DSS SWMU 136.

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### 4.0 CONCEPTUAL SITE MODEL

The conceptual site model for DSS SWMU 136, the Building 907 Septic Systems, is based upon the COCs identified in the soil samples collected from trenches across drain lines, from boreholes drilled near drain lines, septic tanks, and beneath the drainfield and seepage pit, and from the HE catch box at this site. This section summarizes the nature and extent of contamination and the environmental fate of the COCs.

### 4.1 Nature and Extent of Contamination

Potential COCs at DSS SWMU 136 are VOCs, SVOCs, PCBs, HE compounds, metals, including hexavalent chromium, cyanide, and radionuclides. Seven VOCs, twelve SVOCs, and one HE compound were detected in the soil samples. There were no PCBs, hexavalent chromium or cyanide detected in any of the soil samples. Eleven metals were detected at concentrations above the approved maximum background concentrations for SNL/NM North Area Supergroup soils (Dinwiddie September 1997) or above the nonquantified background concentrations. When a metal concentration exceeded its maximum background screening value, had MDLs above background, or had no quantified background value, it was considered further in the risk assessment process. Cesium-137, uranium-235, and tritium were detected at activities above the corresponding background activities. For some of the uranium-235, uranium-238, and tritium analyses, the MDAs exceed the corresponding background activities.

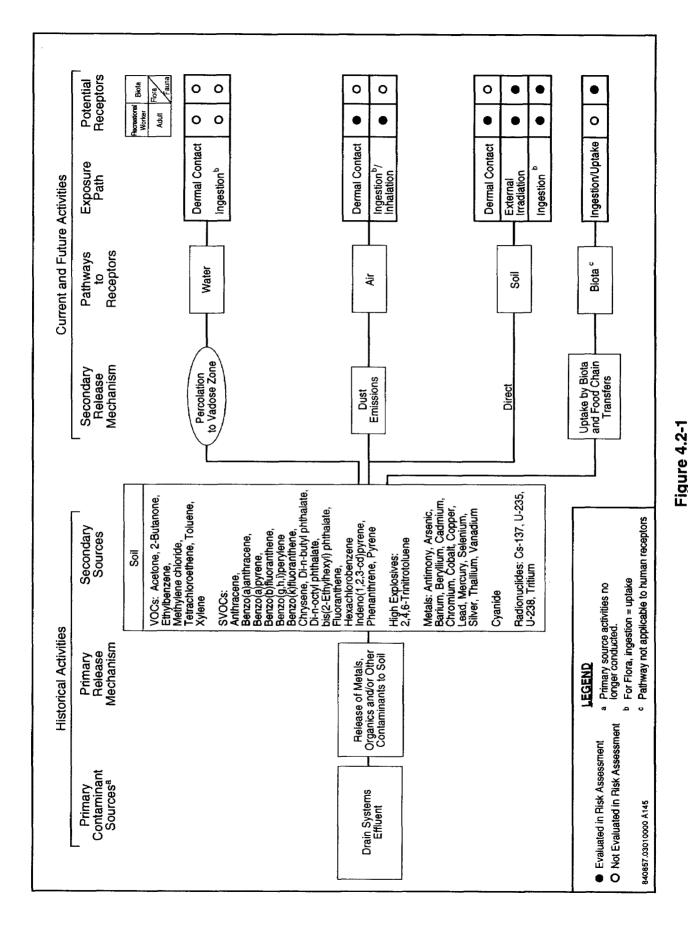
For the HE catch box samples, one VOC, and ten SVOCs were detected. No HE compounds were detected. Cesium-137 and tritium were detected at activities above the corresponding background activities. For one of the uranium-235 analyses, the MDA exceeded the corresponding background activity. Because the metals were analyzed by the TCLP method (for waste characterization purposes), the results do not represent total metal concentrations and were not evaluated in the risk assessment.

### 4.2 Environmental Fate

Potential COCs may have been released into the vadose zone via aqueous effluent discharged from the HE drain line outfall ditch and the septic system drainfield and seepage pit. Possible secondary release mechanisms include the uptake of COCs that may have been released into the soil beneath the HE drain line outfall ditch, drainfield, and seepage pit (Figure 4.2-1).

Two water-bearing zones, a shallow groundwater system and the regional aquifer, underlie DSS SWMU 136. The depth to the shallow groundwater system is approximately 300 feet bgs. The shallow groundwater system is not used as a water supply. The depth to the regional aquifer is approximately 545 feet bgs. Both the City of Albuquerque and Kirtland Air Force Base (KAFB) utilize the regional aquifer as a water supply. Groundwater flow in the shallow groundwater system is to the southeast, while regional groundwater flow is predominantly to the north-northwest in this portion of KAFB. The nearest downgradient water-supply wells are KAFB-1 and KAFB-4, which are approximately 1.3 and 1.1 miles northwest and southwest of the site, respectively. The depth to the shallow and regional aquifers at the site (approximately

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Conceptual Site Model Flow Diagram for DSS SWMU 136, Building 907 Septic Systems

300 and 545 feet bgs) most likely precludes migration of potential COCs into the groundwater system. The potential pathways to receptors include soil ingestion, dermal contact, and inhalation, which could occur as a result of receptor exposure to contaminated subsurface soil at the site. No intake routes through plant, meat, or milk ingestion are considered appropriate for either the industrial or residential land-use scenarios. Annex C provides additional discussion on the fate and transport of COCs at DSS SWMU 136.

Table 4.2-1 summarizes the potential COCs for DSS SWMU 136. All potential COCs were retained in the conceptual model and evaluated in both the human health and ecological risk assessments. The current and future land use for DSS SWMU 136 is industrial (DOE et al. September 1995).

The potential human receptors at the site are considered to be an industrial worker and resident. The exposure routes for the receptors are dermal contact and ingestion/inhalation; however, these are realistic possibilities only if contaminated soil is excavated at the site. The major exposure route modeled in the human health risk assessment is soil ingestion for the COCs. The inhalation pathway is included because of the potential to inhale dust and volatiles. The dermal pathway is included because of the potential for receptors to be exposed to the contaminated soil.

Potential biota receptors include flora and fauna at the site. Major exposure routes for biota include direct soil ingestion, ingesting COCs through food chain transfers, and direct contact with COCs in the soil. Annex C provides additional discussion of the exposure routes and receptors at DSS SWMU 136.

### 4.3 Site Assessment

Site assessment at DSS SWMU 136 included risk assessments for both human health and ecological risk. This section briefly summarizes the site assessment results, and Annex C discusses the risk assessment performed for DSS SWMU 136 in more detail.

### 4.3.1 Summary

The site assessment concluded that DSS SWMU 136 poses no significant threat to human health under either the industrial or residential land-use scenarios. Ecological risks are expected to be very low.

### 4.3.2 Risk Assessments

Risk assessments were performed for both human health and ecological risk at DSS SWMU 136. This section summarizes the results.

Table 4.2-1
Summary of Potential COCs in Soil for DSS SWMU 136, Building 907 Septic Systems (Includes Data from the HE Catch Box Sampling)

		COCs Detected or with	Maximum Background	Maximum		Number of Samples Where COCs Detected with Concentrations
	Number	Concentrations Greater than Background or	Area	Concentration	Average	Background or
COC Type	Samples <sup>a</sup>	Nonquantified Background	Supergroup <sup>5</sup> (mg/kg)	(All Samples) (mg/kg)	Concentration (mg/kg)	Nonquantified Background <sup>e</sup>
VOCs	31	Acetone	NA	0.047	0.0141	24
	32	2-Butanone	NA	0.044 J	0.0037	5
	39	Ethylbenzene	NA	0.0012 J	00000	-
	39	Methylene Chloride	NA	0.008	0.0413	12
	39	Tetrachoroethene	۷N	0.0026 J	0.0050	3
	39	Toluene	NA	0.0046 J	0.0050	11
	39	Xylene	NA	0.0018 J	0.0054	2
SVOCs	36	Anthracene	NA	0.0249 J	0.0877	3
	36	Benzo(a)anthracene	NA	0.048 J	0.0821	-
	36	Benzo(a)pyrene	NA	0.041 J	0.0830	2
	36	Benzo(b)fluoranthene	ΝA	0.100 J	0.0877	က
	36	Benzo(g,h,i)perylene	NA	0.044 J	0.1067	
	36	Benzo(k)fluoranthene	AN	0.062 J	0.0991	2
	36	Chrysene	AN	0.071 J	0.0879	4
	36	Di-n-butyl phthalate	NA	0.086 J	0.0886	4
	36	Di-n-octyl phthalate	ΑN	0.036 J	2060.0	1
	36	bis(2-Ethylhexyl)	ΥN	0.350	0.1242	7
		phthalate				
	36	Fluoranthene	AN	0.0523 J	0.0878	4
	36	Hexachlorobenzene	ΝA	0.0066 J	0.0876	1
	36	Ideno(1,2,3-cd)pyrene	ΝA	0.0370 J	0.0953	1
	36	Phenanthrene	ΑN	0.0251 J	0.0871	3
	36	Pyrene	ΨN	0.0748 J	0.1129	2

Refer to footnotes at end of table.

Table 4.2-1 (Continued)
Summary of Potential COCs in Soil for DSS SWMU 136, Building 907 Septic Systems (Includes Data from the HE Catch Box Sampling)

							Number of Samples
				Maximum			Where COCs Detected with
			COC Cotosto Cotosto	Background			Concentrations
			Concentrations Greater	Limit/North	Maximum		Greater than
		Number	than Dookaround or	Area	Concentration	Average	Background or
		of Soil	Noncliantified	Supergroup <sup>b</sup>	(All Samples)	Concentration <sup>d</sup>	Nonquantified
	COC Type	Samplesa	Background	(mg/kg)	(mg/kg)	(mg/kg)	Backgrounde
PCBs		14	None	NA	NA	NA	None
HF Compounds	ď	41	2.4.6-Trinitrotoluene	NA	L 0970.0	0.1778	
Metals		25	Antimony	3.9	ND (12)	3.36	None
		41	Arsenic	4.4	ND (50)	3.88	3
		41	Barium	200	944	149.8	5
		25	Beryllium	9.0	ND (3.4)	0.423	•
		41	Cadmium	6.0	ND (10)	0.62	
		39	Chromium	12.8	20.2	7.68	သ
		25	Cobalt	7.1	10.2	4.73	-
		25	Copper	17	ND (20)	8.63	•
		41	Lead	11.2	58.4	6.62	3
		41	Mercury	ØN	1.13	0.068	3
		41	Selenium	Ø	(05) QN	1.40	None
		41	Silver	Ø.	6.09	2.12	3
		25	Thallium	Ø.	ND (200)	8.45	None
		25	Vanadium	33	33.2	20.71	Ļ
Hexavalent Chromium	romium	2	None	ØN	NA	NA	None
Cyanide		27	Cvanide	QN ON	NA	NA	None
Radioniclides	Gamma Spectroscopy	32	Cesium-137	0.084	0.22	NC	9
(pCi/a)		26	Uranium-235	0.18	ND (0.348)	NC	10
(n: ) L	-	26	Uranium-238	1.3	ND (1.96)	NC	7
	Triting	30	Tritium	0.0219	10.4	NC	41

Refer to footnotes at end of table.

### Table 4.2-1 (Concluded)

### Summary of Potential COCs in Soil for DSS SWMU 136, Building 907 Septic Systems (Includes Data from the HE Catch Box Sampling)

Number of soil samples includes duplicates, splits and HE catch box samples.

Dinwiddie September 1997.

<sup>4</sup>Average concentration includes all samples except blanks. The average is calculated as the sum of detected amounts and one-half of the MDLs he maximum MDL or MDA above background or nonquantified background. For other COCs, the value listed is the maximum amount detected. •Maximum concentration for metals and cyanide, or maximum activity for radionuclides, is the greater of either the maximum amount detected or

or nondetect results, divided by the number of samples.

See appropriate data table for sample locations.

An average MDA is not calculated because of the variability in instrument counting error and the number of reported nondetect activities for gamma spectroscopy or tritium analyses.

Tharp February 1999. 0.021 pCi/g = 420 pCi/L, assuming a soil density of 1 gram/cubic centimeter and a 5 percent soil moisture.

= Constituent of concern. ပ္ပင္ပ

= Drain and Septic Systems. DSS HE

= High explosive(s)

Analytical result was qualified as an estimated value.

Minimum detectable activity. MDA

= Milligram(s) per kilogram. = Method detection limit. MDL

= Not applicable.

= Not detected above the MDL or MDA, shown in parentheses. Not calculated

= Nonquantified background value.

= Polychlorinated biphenyl. = Picocurie(s) per gram.

Semivolatile organic compound. = Picocurie(s) per liter. mg/kg NA NC NC ND () ND () PCB PCB PCI/L SVOC SWMU

= Solid Waste Management Unit. Volatile organic compound.

### 4.3.2.1 Human Health

DSS SWMU 136 has been recommended for an industrial land-use scenario (DOE et al. September 1995). Because acetone, 2-butanone, ethylbenzene, methylene chloride, tetrachloroethene, toluene, xylene, anthracene, benzo(a)anthracene, benzo(a)pyrene, benzo(b)fluoranthene, benzo(g,h,i)perylene, benzo(k)fluoranthene, chrysene, di-n-butyl phthalate, di-n-octyl phthalate, bis(2-ethylhexyl) phthalate, fluoranthene, hexachlorobenzene, ideno(1,2,3-cd)pyrene, phenanthrene, pyrene, 2,4,6-trinitrotoluene, antimony, arsenic, barium, beryllium, cadmium, chromium, cobalt, copper, lead, mercury, selenium, silver, thallium, vanadium, cyanide, cesium-137, uranium-235, uranium-238, and tritium are present above background or have nonquantified background levels, it was necessary to perform a human health risk assessment analysis for the site, which included these COCs. Annex C provides a complete discussion of the risk assessment process, results, and uncertainties. The risk assessment process provides a quantitative evaluation of the potential adverse human health effects from constituents in the site's soil by calculating the hazard index (HI) and excess cancer risk for both industrial and residential land-use scenarios.

The HI calculated for the COCs at DSS SWMU 136 is 0.11 for the industrial land-use scenario, which is less than the numerical standard of 1.0 suggested by risk assessment guidance (EPA 1989). The incremental HI risk, determined by subtracting risk associated with background from potential nonradiological COC risk (without rounding), is 0.07. The excess cancer risk for DSS SWMU 136 COCs is 4E-6 for an industrial land-use scenario. NMED guidance states that cumulative excess lifetime cancer risk must be less than 1E-5 (Bearzi January 2001); thus the excess cancer risk for this site is below the suggested acceptable risk value. The incremental excess cancer risk is 1.57E-6. Both the incremental HI and excess cancer risk are below NMED guidelines.

The HI calculated for the COCs at DSS SWMU 136 is 1.36 for the residential land-use scenario, which is greater than the numerical standard of 1.0 suggested by risk assessment guidance (EPA 1989). The incremental HI risk, determined by subtracting risk associated with background from potential nonradiological COC risk (without rounding), is 0.89. The excess cancer risk for DSS SWMU 136 COCs is 2E-5 for a residential land-use scenario. NMED guidance states that cumulative excess lifetime cancer risk must be less than 1E-5 (Bearzi January 2001); thus the excess cancer risk for this site is above the suggested acceptable risk value. The incremental excess cancer risk is 5.95E-6. Both the incremental HI and the excess cancer risk are below NMED guidelines.

Although both the HI and excess cancer risk are slightly above the NMED guideline for the residential land-use scenario, maximum concentrations were used in the risk calculation. Because the site has been adequately characterized, average concentrations are more representative of actual site conditions. Using the 95% upper confidence limit of the average concentrations for the main contributors to excess cancer risk reduces the total and incremental HI to 0.76 and 0.49, respectively. The total and incremental estimated excess cancer risk are reduced to 2E-6 and 1.83E-6, respectively. Thus, by using realistic concentrations in the risk calculations that more accurately depict actual site conditions, the total and incremental risk calculations are below NMED guidelines.

For the radiological COCs, four of the constituents (cesium-137, uranium-235, uranium-238, and tritium) had detected values greater than the corresponding background values. The incremental total effective dose equivalent (TEDE) and corresponding estimated cancer risk from radiological COCs are much lower than the U.S. Environmental Protection Agency (EPA)

guidance values; the estimated TEDE is 1.2E-1 millirem (mrem)/year (yr) for the industrial landuse scenario. This value is much lower than the EPA's numerical guidance of 15 mrem/yr (EPA 1997a). The corresponding incremental estimated cancer risk value is 1.2E-6 for the industrial land-use scenario. Furthermore, the incremental TEDE for the residential land-use scenario that results from a complete loss of institutional controls is 3.1E-1 mrem/yr with an associated risk of 3.5E-6. The guideline for this scenario is 75 mrem/yr (SNL/NM February 1998). Therefore, DSS SWMU 136 is eligible for unrestricted radiological release.

The incremental nonradiological and radiological carcinogenic risks are tabulated and summed in Table 4.3.2-1.

Table 4.3.2-1
Summation of Incremental Radiological and Nonradiological Risks from DSS SWMU 136, Building 907 Septic Systems Carcinogens

Scenario	Nonradiological Risk	Radiological Risk	Total Risk
Industrial	1.57E-6	1.2E-6	2.8E-6
Residential	1.83E-6	3.5E-6	5.3E-6

DSS = Drain and Septic Systems. SWMU = Solid Waste Management Unit.

Uncertainties associated with the calculations are considered small relative to the conservatism of the risk assessment analysis. Therefore, it is concluded that this site poses insignificant risk to human health under both the industrial and residential land-use scenarios

### 4.3.2.2 Ecological

An ecological assessment that corresponds with the procedures in the EPA's Ecological Risk Assessment Guidance for Superfund (EPA 1997b) also was performed as set forth by the NMED Risk-Based Decision Tree in the "RPMP [RCRA Permits Management Program] Document Requirement Guide" (NMED March 1998). An early step in the evaluation compared COC concentrations and identified potentially bioaccumulative constituents (see Annex C, Sections IV, VII.2, and VII.3). This methodology also required developing a site conceptual model and a food web model, as well as selecting ecological receptors, as presented in "Predictive Ecological Risk Assessment Methodology, Environmental Restoration Program, Sandia National Laboratories, New Mexico" (IT July 1998). The risk assessment also includes the estimation of exposure and ecological risk.

Table 18 of Annex C presents the results of the ecological risk assessment. Ecological risks associated with DSS SWMU 136 were estimated through a risk assessment that incorporated site-specific information when available. Initial predictions of potential risk to plants from exposure to several metals were based upon maximum measured soil concentrations, highly conservative plant toxicity benchmarks, and assumptions of high bioavailability. Actual risk to this receptor is expected to be low based upon more realistic exposure assumptions. Predictions of potential risk to omnivorous and insectivorous deer mice from exposures to mercury and thallium were also attributed to conservative exposure assumptions. For the burrowing owl, the initial prediction of risk for exposure to mercury is attributed to the assumption of 100-percent area use by this receptor. A more realistic assumption of area use

for this receptor resulted in a hazard quotient of only 0.21. The very small size of this site (0.46 acres) also limits the potential for significant risk to ecological receptors at this site, particularly at the population or community levels. Based upon this final analysis, the potential for ecological risks associated with DSS SWMU 136 is expected to be low.

### 4.4 Baseline Risk Assessments

This section discusses the baseline risk assessments for human health and ecological risk.

### 4.4.1 Human Health

Because the results of the human health risk assessment summarized in Section 4.3.2.1 indicate that DSS SWMU 136 poses insignificant risk to human health under both the industrial and residential land-use scenarios, a baseline human health risk assessment is not required for this site.

### 4.4.2 Ecological

Because the results of the ecological risk assessment summarized in Section 4.3.2.2 indicate that ecological risks at DSS SWMU 136 are expected to be low, a baseline ecological risk assessment is not required for the site.

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### 5.0 NO FURTHER ACTION PROPOSAL

### 5.1 Rationale

Based upon field investigation data and the human health and ecological risk assessment analyses, an NFA decision is recommended for DSS SWMU 136 for the following reasons:

- The soil has been sampled for all potential COCs.
- No COCs are present in the soil at levels considered hazardous to human health for either an industrial or residential land-use scenario.
- None of the COCs warrant ecological concern after conservative exposure assumptions are analyzed.

### 5.2 Criterion

Based upon the evidence provided in Section 5.1, DSS SWMU 136 is proposed for an NFA decision according to Criterion 5, which states, "the SWMU/AOC [area of concern] has been characterized or remediated in accordance with current applicable state or federal regulations, and the available data indicate that contaminants pose an acceptable level of risk under current and projected future land use" (NMED March 1998).

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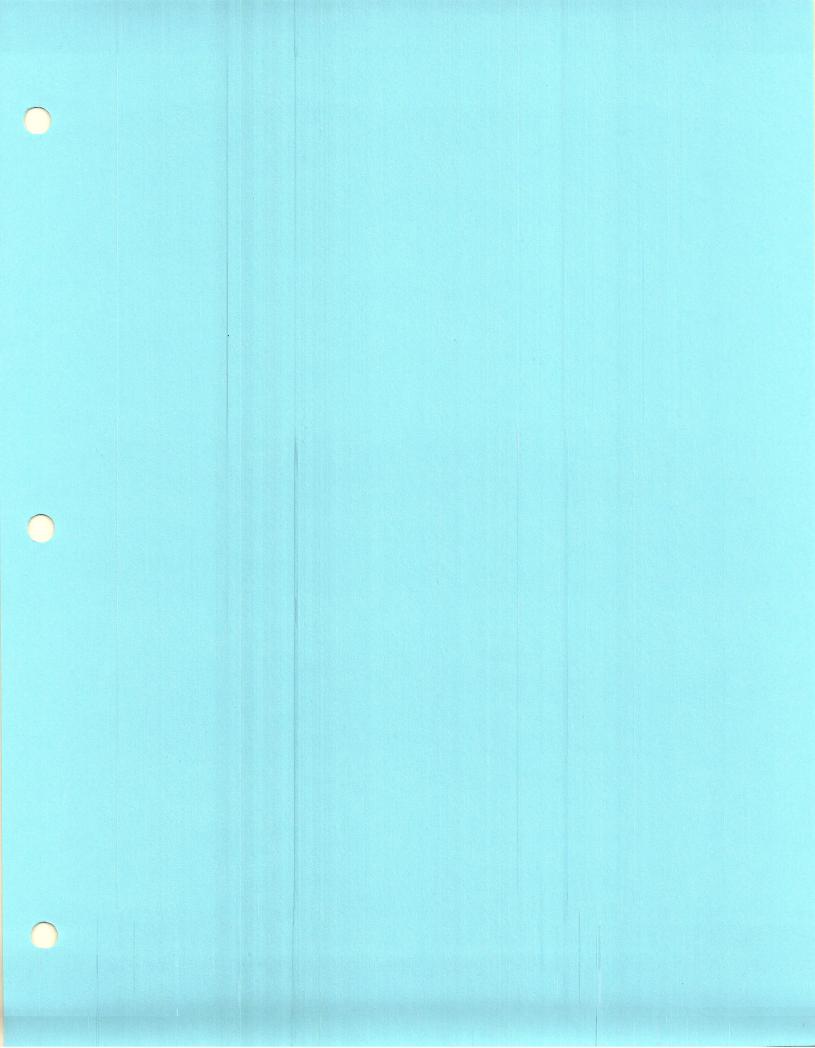
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USGS, see U.S. Geological Survey.



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603366	000000	603075	603075	2094	2044	508417	508417	Quality Ass	603366	003300	600000	935500	832503	603366	603366	Drainfield b	603366	603366	603366	603366	603366	603075	000073	ECOUNT IN	DE design line	- 1	600006	- 12	Seenane nit	04202	2092	2094	2004	2004	2044	2044	2004	500417	200417	508417	200417	508417	DOTE NOT NOT	Boroholo ca	508001	500001	508001	None	None	Drain line tru	Number	Record		
TA2-136-SPT-TB	TA2-128-SET-ER	TA2-136-HEOUT-PIPE-TB75	TA2-136-HE-OUT-TB-2.0	TA2-BH-09 (EB)	TA2-BH-08 (EB)	TA2-BH-05 (TB)	TA2-BH-05 (EB)	Quality Assurance/Quality Control Samples (pg/L)	1A2-136-SWLAT-3-13	TAG 136 CWI AT C 13	TA9-136-SWI AT-S-B	TA2-136-SOLITHI AT-S-13	TA2-136-SOUTHLAT-S-8	TA2-136-SELAT-S-15	603366 TA2-136-SELAT-S-10	orehole samples	TA2-136-DITCH-2/3-S-9	TA2-136-DITCH-2/3-S-4	TA2-136-DITCH-1/3-S-9	TA2-136-DITCH-1/3-S-4	TA2-136-DITCH-0/0-S-7	TA2-136-HEOUT-S-2.0 <sup>C</sup>	TA2-136-HEOUT-PIPE-S/5	ECONTE OCCIONA SERVICE CONTENTE CONTE CONTENTE CONTENTE CONTENTE CONTENTE CONTENTE CONTENTE CONTENTE C	nation samples	TA2-136-SPG-S-18	TA3-136-SPG-DH-13	TA9-136-SPG-S-13	϶Ϯ	TA2-ST-10-6 67 (On-site laboratory)	TA2-ST-09-6 67 (On-site laboratory)	TA2-BH-09-15-5	TA9-RH-09-10-5	TA2-BH-09-5 75	TA9_RH_08_16_25	TA9-RH-08-10 75	TA9 BH-08-5 95	TA2 BH 05 (DI I)	TA2 BU 05	TA2 BH 05	100 BL 05	TA2-BH-05	Borellote salliples lied didni inico di soprio con	males near drain lines or septic tanks	TA2-TB-02	TA2-TB-02	TA2-TR-02	TA2_TB.01	TA2-TR-01	Drain line trench samples	Ci Cambio io	EB Sample ID	Sample Attributes	
NA	NA	NA	NA	NA	NA	NA.	2 3	NA		13	8	13	8	15	10		9	4	9	4	7	2.0	3	0.75		18	13	13		6.67	6.67	15.5	10.5	5.75	16.25	10.75	5.25	30.2	30.2	20.5	15.5	11.5	6.5		5	3.5	0.5	6	4	0		Sample Depth (#)		
8-23-00	8-23-00	3-8-00	3-8-00	11-6-94	11-2-94	3-10-94	3 10-04	3-10-94		8-22-00	8-22-00	8-22-00	8-22-00	8-22-00	8-22-00		8-22-00	8-22-00	8-21-00	8-21-00	8-21-00	0.00	3 9 00	3-8-00		8-22-00	8-22-00	8-22-00		8-29-95	8-29-95	11-6-94	11-6-94	11-6-94	11-2-94	11-2-94	11-2-94	3-10-94	3-10-94	3-10-94	3-10-94	3-10-94	3-10-94		11-1-93	11-1-93	11-1-93	10-28-93	10-28-93	10-28-93		Sample Date		
ת	20	ND (0.224 J)	ND (0.224 J)	NI NI	25	2000	36.1/10	10		18 J (20)	B	ND (20 U)	16 J (20	18 J (20	16 J (20)		18 J (20)	I	12 J (20	02) r 61	02) L 61	1.41	ND (2 42)	ND (2.42)		14 J (20)	17 J (20)	14 J (20)		11 J (20)	ND (5)	NR.	NR	NR	NR	NR	NR	16	13	13	22	14	13		28	47	5.1 J (10)	12	12	6.1 J (10)		Acetone		
ND (2.6)	0.38 J (1)	ND (0.085)	ND (0.085)	ND (0)	ND (S)	ND (3)		ND (5)			ND (2.6)	ND (2.6)	ND (2.6)	ND (2.6)	ND (2.6)		ND (2.6)		ND (2.6)	ND (2.0)	ND (2.0)	1000	ND (0 145)	ND (0.145 J)				ND (2.6)	l	ND (5)	ND (5)	ND (500)	ND (500)	ND (500)	ND (500)	ND (500)	ND (500)	ND (5)	ND (5)	ND (5)	ND (5)		ND (5)			ND (5)	ND (5)	ND (5)	ND (5)	ND (5)		Bromoform		
		ND (1.18)	ND (1.16)	ND (1 10)	N. C.	NB	ND (10)	ND (10)		ND (2.5)	ND (2.5)	ND (2.5)	ND (2.5)	(C.2)	11000		ND (2.5)	ND (2.5)	ND (2.5)	ND (2.3)	ND (2.5)	ND (3 E)	ND (1.76)	ND (1.76)		ND (2.5)	33 J	ND (2.5)		ND (5)	ND (5)	NR	NR	NR	NR.	S	NR	ND (10)	5.2 J (10)	4.6 J (10)	4.4 J (10)	ND (10)	ND (10)		ND (10)	ND (10)	ND (10)	ND (10)	ND (10)	ND (10)		2-Butanone		
ND (1.3)	0.34 J (1)	(GOO'O) CIN	ND (0.000)	ND (O DBQ)	ND (1)	ND (1)	ND (5)	ND (5)		ND (1.3)	ND (1.3)	ND (1.3)	ND (1.3)	ND (1.3)	NO (1.3)	ND (4.3)	ND (1.3)	ND (1.0)	ND (1.3)	ND (1-2)	ND (1.3)	ND (1.3)	ND (0.111)	ND (0.111 J)		ND (1.3)	ND (1.3)	ND (1.3)		ND (1)	ND (1)	ND (100)	ND (100)	ND (100)	ND (100)	ND (100)	ND (100)	ND (5)	ND (5)	ND (5)	ND (5)	ND (5)	ND (5)		ND (5)	NO (a)	ND (5)	ND (5)	ND (5)	ND (5)		Dibromochloromethane		
	ND (0.21)		ND (0.07)	ND (0.07)	ND (1)	ND (1)	ND (5)	ND (5)		ND (1.1)	ND (1.1)	NC (1.1)	ND (1.1)	ND (1.1)	ND (1 1)	ND (1.1)	140 (1.1)	ND (1.1)	ND (1 1)	ND (1-1)	ND (1.1)	ND (1 1)	ND (0.19)	ND (0.23)		NO (1.1)	20(1.1)	20(3.5)	NO (1 1)	ND (I)	ND (I)	ND (100)	ND (100)	ND (100)	ND (100)	ND (100)	ND (100)	ND (5)	(c) (d)	ND (S)	ND (O)	NU (5)	ND (5)	ND (7)	ND (5)	NO (5)	ND (S)	ND (S)	ND (6)	ND (5)		1,2-Dichloropropane		VOCs (EPA Method 8000 <sup>a</sup> ) (μg/kg)
ND (1.6)		ND (0.35)	ND (0.051)	ND (0.051)	ND (0.5)	ND (0.5)	ND (5)	ND (5)		ND (1.6)	NO (1.0)	ND (1.0)	ND (1.6)	ND (1.6)	NO (1 6)	ND (1.6)		ND (1 6)	ND (1.6)	ND (1.6)	ND (1.6)	ND (1.6)	ND (0.212)	ND (0.212)	ND (0 040)	NO (1.0)	ND (1.0)	ND (1.6)	ND (1 8)	NO (1)	20 (3)	ND (1)	ND (50)	ND (50)	ND (50)	ND (50)	ND (50)	ND (5)	NO (5)	ND (S)	NO (5)	ND (S)	1000	NID (E)	(c) ON	ND (5)	١.	121/5	ND (S)	ND (5)		Ethylbenzene		Ͻ <sup>a</sup> ) (μ <b>g/kg</b> )
	ם:	R	ND (2.59 U)	ND (3.34 U)	ND (5)	ND (5)	ND (5)	NU (5)		140 (3)	ND (5)	ND (5)	ND (S)	ND (5)	ND (5)	ND (5)		ND (5)	ND (0.971)	ND (1.71 O)	ND /1 71 IN	130 (3)	ND (5)	ND (S)	ND (5)	10 (1)	ND (1)	ND (1)	ND (500)	ND (500)	ND (500)	ND (500)	ND (500)	1	50	47.16	43 1 (5)	30.00	19 1 (5)	4.1(5)		7 8	88		7 4	5.3	2	Methylene chloride						
NO (1:0)	ND (1 5)	ND (0.57)	ND (0.385)	ND (0.385)	ND (0.5)	ND (0.5)	ND (5)	ND (S)	• (5)	100 (1.0)	ND (1.5)	ND (1.5)	ND (1.5)	ND (1.5)	ND (1.5)	ND (1.5)		ND (1.5)	ND (0.582)	110 (0:002)	ND (0 582)		2 J (5)	ND (1.5)	ND (1.5)		ND (1)	ND (1)	ND (50)	ND (50)	ND (50)	ND (50)	ND (50)	ND (50)		ļ		ND (5)		1		ND (5)	2.6 J (5)		ND (5)	ND (5)	ND (E)	Tetrachloroethene						
100	ND (1 1)	ND (0.51 U)	ND (0.262)	ND (0.262)	ND (0.5)	ND (0.5)	(c) UN	ND (5)	ND (E)		ND (1.1)	ND (1.1)	ND (1.1)	ND (1.1)	ND (1.1)	2 J (5)		) ND (1.1)	1.5 J (5)	ND (1.1)	1.8 J (5)	ND (1.1)	ND (0.259)	NI (0.000)	ND (0.259)			ND (1 1)	ND (1.1)		8	ND (1)	ND (50)	ND (50)	ND (50)	ND (50)	ND (50)	ND (50)	ND (5)	ND (5)	ND (5)	2.3 J (5)	28.1 (5	2.2 J (5		ND	(5) 4.6 J (5)		ND (5)	ND (5)	NO (8)	Toluene		
	ND (3.4)	ND (0.61)	ND (0.437)	ND (0.437)	ND (0.5)	ND (0.5)	NO (3)	20 (3)	20 (8)		ND (3.4)	ND (3.4)	ND (3.4)	ND (3.4)		ND (3.4)				ND (3.4)	ND (3.4)	ND (3.4)	ND (0.66)	ND (0.69)	ND (0.68)		ND (3.4)	ND (3.4)	ND (3.4)		ND (2)	ND (2)	ND (50)	ND (50)	ND (50)	ND (50)	ND (50)	ND (50)	ND (5)	ND (5)		ND (5)				ND (			ND (5)	ND (5)	NO (5)	Xylene		

Note: Values in **bold** represent detected analytes. <sup>a</sup>EPA November 1986.

bAnalysis request/chain-of-custody record.

CSame location as sample TA2-136-DITCH-0/0-S-7.

BH = Borehole.

DITCH = Sample in HE drain line outfall ditch.

DSS = Drain and Septic Systems.

DU = Duplicate sample.

EB = Equipment blank.

EPA = U.S. Environmental Protection Agency.

ER = Environmental Restoration

## Table A-1 (Concluded) Summary of DSS SWMU 136, Building 907 Septic Systems Confirmatory Soil Sampling, VOC Analytical Results October 1993–August 2000 (On- and Off-Site Laboratories)

Table A-2 Summary of DSS SWMU 136, Building 907 Septic Systems Confirmatory Soil Sampling, VOC Analytical MDLs October 1993–August 2000 (On- and Off-Site Laboratories)

	EPA Method 8000a
	Detection Limits
Analyte	(μg/kg)
Acetone	2.42-10
Benzene	0.235-50
Bromodichloromethane	0.194-100
Bromoform	0.145-500
Bromomethane	0.478-500
2-Butanone	1.76–10
Carbon disulfide	0.988-5
Carbon tetrachloride	0.144-50
Chlorobenzene	0.206–200
Chloroethane	0.286-500
2-Chloroethyl vinyl ether	1.8
Chloroform	0.204-50
Chloromethane	0.192-500
Dibromochloromethane	0.111-100
1,2-Dibromoethane	200
1,2-Dichlorobenzene	50
1,3-Dichlorobenzene	50
1,4-Dichlorobenzene	50
1,1-Dichloroethane	0.231-50
1,2-Dichloroethane	0.17-100
1,1-Dichloroethene	0.262-50
cis-1,2-Dichloroethene	0.327-0.7
trans-1,2-Dichloroethene	0.232-50
1,2-Dichloroethene	5
1,2-Dichloropropane	0.19–100
cis-1,3-Dichloropropene	0.216–200
trans-1,3-Dichloropropene	0.163-100
Ethylbenzene	0.212-50
2-Hexanone	1.33–10
Methylene chloride	0.571-500
4-Methyl-2-pentanone	1.17–10
Styrene	0.198–5
1,1,2,2-Tetrachloroethane	0.195–100
Tetrachloroethene	0.316-50
Toluene	0.259-50
1,1,1-Trichloroethane	0.157–50
1,1,2-Trichloroethane	0.177–100
Trichloroethene	0.439-50
1,1,2-Trichloro-1,2,2-trifluoroethane	100

Refer to footnotes at end of table.

### Table A-2 (Concluded) Summary of DSS SWMU 136, Building 907 Septic Systems Confirmatory Soil Sampling, VOC Analytical MDLs October 1993–August 2000 (On- and Off-Site Laboratories)

	EPA Method 8000 <sup>a</sup> Detection Limits
Analyte	(μg/kg)
Vinyl acetate	1.17–10
Vinyl chloride	0.255–100
Xylene	0.68–50

Note: Because of the long time period covering sample collection at this site, MDL ranges are presented. MDLs were not routinely reported, or were reported as ranges, by the laboratories for analyses performed in the early- to mid-1990s.

<sup>a</sup>EPA November 1986.

DSS = Drain and Septic Systems.

EPA = U.S. Environmental Protection Agency.

MDL = Method detection limit.
μg/kg = Microgram(s) per kilogram.
SWMU = Solid Waste Management Unit.
VOC = Volatile organic compound.

Refer to footnotes at end of table.

Table A-3
Summary of DSS SWMU 136, Building 907 Septic Systems
Confirmatory Soil Sampling, SVOC Analytical Results
October 1993–August 2000
(Off-Site Laboratories)

603366	2094	2044	508417	Quality As	603366	603366	603366	603366	603366	603366	Drainfield	603366	603366	603366	603366	603366	603075	603075	HE drain I	603366	603366	603366	Seepage	2094	2094	2094	2044	2044	2044	508417	508417	508417	508417	508417	508417	Boreholes	508001	508001	508001	None	None	None	Drain line	Record Number <sup>b</sup>	
TA2-136-SPT-EB	TA2-BH-09 (EB)	TA2-BH-08 (EB)	TA2-BH-05 (EB)	Quality Assurance/Quality Control Samples (μg/L)	TA2-136-SWLAT-S-13	TA2-136-SWLAT-S-8	TA2-136-SOUTHLAT-S-13	TA2-136-SOUTHLAT-S-8	TA2-136-SELAT-S-15	TA2-136-SELAT-S-10	Drainfield borehole samples	TA2-136-DITCH-2/3-S-9	TA2-136-DITCH-2/3-S-4	TA2-136-DITCH-1/3-S-9	TA2-136-DITCH-1/3-S-4	TA2-136-DITCH-0/0-S-7	TA2-136-HE-OUT-S-2.0 <sup>c</sup>	TA2-136-HEOUT-PIPE-S75c	HE drain line outfall ditch samples	TA2-136-SPG-S-18	TA2-136-SPG-DU-13	TA2-136-SPG-S-13	Seepage pit borehole samples	TA2-BH-09-20.0	TA2-BH-09-16.0	TA2-BH-09-10.0	TA2-BH-08-21.0	TA2-BH-08-16.5	TA2-BH-08-11.0	TA2-BH-05 (DU)	TA2-BH-05	TA2-BH-05	TA2-BH-05	TA2-BH-05		₃ t	TA2-TB-02	TA9-TB-09	TA2-TB-02	TA2-TB-01	TA2-TR-01	TA2-TR-01	Drain line trench samples	ER Sample ID	Sample Attributes
NA	N A	AN	NA	es (μg/L)	13	8	13	8	15	10		9	4	9	4	7	2.0	0.75		18	13	13		20	16	10	21	16.5	7	38	30	21.2	14.6	11	6	-	5	3 7 6	0.5	5	4	0		Sample	
8-23-00	11-6-94	11-2-94	3-10-94		8-22-00	8-22-00	8-22-00	8-22-00	8-22-00	8-22-00		8-22-00	8-22-00	8-21-00	8-21-00	8-21-00	3-8-00	3-8-00		8-22-00	8-22-00	8-22-00		11-6-94	11-6-94	11-6-94	11-2-94	11-2-94	11-2-94	3-10-94	3-10-94	3-10-94	3-10-94	3-10-94	3-10-94		11-1-93	11-1-93	11-1-93	10-28-93	10-28-93	10-28-93	Ì	Sample Date	
ND (0.58)	ND (10)	ND (10)	ND (10)		ND (13)	ND (13)	ND (13)	ND (13)	ND (13)	ND (13)		ND (13)	R	ND (5.99)		ND (13)	ND (13)	ND (13)		ND (330)	ND (330)	ND (330)	ND (330)	48 J (330)	ND (330)	ND (330)	ND (330)	ND (330)	ND (330)	ND (330)	ND (330)	1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	ND (330)	ND (330)	ND (330)	ND (330)	ND (330)	ND (330)		Benzo(a)anthracene					
ND (0.6)	ND (10)	ND (10)	ND (10)		ND (15)	ND (15)	ND (15)	ND (15)	ND (15 J)	ND (15)		ND (15)	ND (15 J)	ND (15)	ND (15)	ND (15)	730	ND (5.66)		ND (15)	ND (15)	41 J (330)		ND (330)	ND (330)	ND (330)	N		ND (330)	ND (330)	ND (330)	ND (330)	ND (330)	ND (330)	ND (330)		ND (330)	ND (330)	ND (330)	ND (330)	ND (330)	ND (330)		Benzo(a)pyrene	
ND (0.88)	ND (10)	ND (10)	ND (10)		ND (17)	ND (17)	ND (17)	ND (17)	ND (17 J)	ND (17)		ND (17)	ND (17 J)	ND (17)	ND (17)	ND (17)	R	ND (8.99)		ND (17)	ND	41 J (330)		ND (330)	ND (330)	ND (330)	ND (330)			ND (330)	ND (330)	ND (330)	ND (330)	ND (330)	ND (330)		ND (330)	ND (330)	ND (330)	ND (330)	ND (330)	ND (330)		Benzo(b)fluoranthene	
ND (0.95)	ND (10)	ND (10)	ND (10)		ND (140)	ND (140)	ND (140)	ND (140)	ND (140 J)	ND (140)		ND (140)	ND (140 J)	ND (140)	ND (140)	ND (140)	R	ND (8.99)		ND (140)	ND (140)	ND (140)		ND (330)	ND (330)	ND	ND		ND (330)	ND (330)	ND (330)	ND (330)	ND (330)	ND (330)	ND (330)		ND (330)	ND (330)	ND (330)	ND (330)	ND (330)	ND (330)		Benzo(g,h,i)perylene	
ND (0.74)	ND (10)	ND (10)	ND (10)		ND (33)	ND (33)	ND (33)	ND (33)	ND (33 J)	ND (33)		ND (33)	ND (33 J)	ND (33)	ND (33)	ND (33)	В	ND (8.99)		ND (33)	ND (33)	62 J (330)		ND (330)	ND (330)	ND (330)	ND (330)	ND (330)	ND (330)		ND (330)	ND (330)	ND (330)	ND (330)	ND (330)	ND (330)		Benzo(k)fluoranthene							
ND (0.42)	ND (10)	ND (10)	ND (10)		ND (15)	ND (15)	ND (15)	ND (15)	ND (15)	ND (15)		ND (15)	R	ND (6.33)		ND (15)	ND (15)	36 J (330)		ND (330)	ND (330)	ND (330)	ND (330)	71 J (330)	ND (330)	ND (330)	ND (330)	ND (330)	ND (330)	ND (330)	ND (330)		ND (330)	ND (330)	ND (330)	ND (330)	ND (330)	ND (330)		Chrysene	SVOCs (EPA Me				
ND (1.3)	ND (10)	ND (10)	ND (10)		ND (25)	50 J (330)	ND (25)	ND (25)	ND (25)	ND (25)		ND (25)	R	ND (14)		ND (25)	ND (25)	ND (25)		ND (330)	ND (330)	ND (330)	ND (330)	ND (330)	ND (330)		ND (330)	ND (330)	86 J (330)	ND (330)	ND (330)	ND (330)		Di-n-butyl phthalate	SVOCs (EPA Method 8270 <sup>a</sup> ) (µg/kg)										
ND (1.3)	ND (10)	ND (10)	ND (10)		ND (27)	ND (27)	ND (27)	ND (27)	(L /2) ON	ND (27)		ND (27)	ND (27 J)	ND (27)	ND (27)	ND (27)	æ	ND (8.99)		ND (27)	ND (27)	36 J (330)		ND (330)	ND (330)	ND (330)	ND (330)	ND (330)	ND (330)		ND (330)	ND (330)	ND (330)	ND (330)	ND (330)	ND (330)		Di-n-octyl phthalate							
ND (3.5 U)	ND (10)	1 J (10)	2.8 J (10)	2011	ND (170 U)	ND (52 U)	ND (190 U)	ND (41 U)	ND (1100)	ND (70 U)		(0 0/1) UN	ND (350 U)	ND (140 U)	ND (110 U)	ND (140 U)	B	ND (19.6)	<b>i</b>	ND (82 U)	ND (36)			ND (330)	290 J (330)	280 J (330)	ND (330)	350	290 J (330)	99 J (330)		ND (330)	ND (330)	ND (330)	ND (330)	ND (330)			bis(2-Ethylhexyl) phthalate						
ND (0.6)			ND (10)		ND (25)	ND (25)	ND (25)	ND (25)	ND (25)	ND (25)		ND (25)	סכ	ND (5)		ND (25)	ND (25)	ND (25)		ND (330)	ND (330)	ND (330)	ND (330)	45 J (330)	ND (330)	ND (330)			ľ		ND (330)		ND (330)	ND (330)	ND (330)	ND (330)	ND (330)	ND (330)	, ij	Fluoranthene					
ND (0.61)	ND (10)	ND (10)	ND (10)	NID (40)	ND (82)	ND (82)	ND (82)	ND (82)	ND (82 J)	ND (82)		ND (82)	ND (82 J)	ND (82)	ND (82)	ND (82)	D	ND (8.99)		ND (82)	ND (82)	ND (82)		ND (330)	ND (330)	ND (330)	ND (330)		N	ND (330)	ND (330)	ND (330)	ND (330)	ND (330)	ND (330)		ND (330)	ND (330)	ND (330)	ND (330)	ND (330)	ND (330)	ND (200)	Indeno(1,2,3-cd)pyrene	
ND (0./1)	ND (10)	NO (10)	ND (10)	ND (10)	ND (160)	ND (160)	ND (100)	ND (160)	ND (160)	ND (160)	1100	ND (100)	ND (160)	ND (160)	ND (160)	ND (160)	I	ND (8.00)	5	ND (160)	ND (160)	ND (160)		ND (330)	ND (330)	ND (330)	ND (330)	) L (320)	ND (330)	ND (330)	ND (330)	ND (330)	ND (330)	ND (330)	ND (330)		ND (330)	ND (330)	ND (330)	ND (330)	ND (330)	ND (330)	(VCC) CIN	Pyrene	

### Table A-3 (Concluded) Summary of DSS SWMU 136, Building 907 Septic Systems Confirmatory Soil Sampling, SVOC Analytical Results October 1993–August 2000 (Off-Site Laboratories)

```
Note: Values in bodd represent detected analytes.

a PA November 1986.

banalysis request/hatin-of-custody record.

Same location as sample TA2-136-DITCH-0/0-S-7.

BH

STICH

Benchold as propie in HE drain line outfall ditch.

DITCH

E Sample in ME drain line outfall ditch.

ESS

E Dain and Septic Systems.

DUI

E Equipment blank.

EPA

E LUS. Emironmental Protection Agency.

ERA

EPA

ELIVIT = Sample in HE drain line outfall ditch.

HE OUT

E Sample in HE drain line outfall ditch.

HE OUT

E High = subjointered in the outfall ditch.

HE OUT

E High = subjointered in the outfall ditch.

HE US

Analytical result was qualified as an estimated value.

MDL

Analytical result was qualified as an estimated value.

MDL

Microgram(s) per lite;

ANA

NO ()

ENd = Not applicable.

NO ()

E Not described above the MDL shown in parentheses.

SUUTHLAT

Southern lateral.

SOUTHLAT

Southwestern lateral.

SWMU

Solf waste Management Unit.

TA

The Tencha Area.

Tran = Tench - Southwestern lateral.

SWMU

E Analytical result was qualified as not detected.
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Table A-4
Summary of DSS SWMU 136, Building 907 Septic Systems
Confirmatory Soil Sampling, SVOC Analytical MDLs
October 1993–August 2000
(Off-Site Laboratories)

	EPA Method 8270 <sup>a</sup>
	Detection Limit
Analyte	(μg/kg)
Acenaphthene	4–3300
Acenaphthylene	3.66–330
Anthracene	4.66–330
Benzo(a)anthracene	5.99–330
Benzo(a)pyrene	2330
Benzo(b)fluoranthene	2.33–330
Benzo(g,h,i)perylene	5–330
Benzo(k)fluoranthene	5–330
Benzoic acid	1600
Benzyl alcohol	330
4-Bromophenyl phenyl ether	4.66–330
Butylbenzyl phthalate	12–330
Carbazole	5–330
4-Chlorobenzenamine	54–330
bis(2-Chloroethoxy)methane	5.99-330
bis(2-Chloroethyl)ether	6.66–330
bis-Chloroisopropyl ether	5.99-330
4-Chloro-3-methylphenol	19.6–6700
2-Chloronaphthalene	3.66-330
2-Chlorophenol	5–6700
4-Chlorophenyl phenyl ether	3.33–330
Chrysene	6.33–330
o-Cresol	7.66–330
Dibenz[a,h]anthracene	2.67-330
Dibenzofuran	2.66-330
1,2-Dichlorobenzene	4.33–330
1,3-Dichlorobenzene	3.33-330
1,4-Dichlorobenzene	5.99-3300
3,3'-Dichlorobenzidine	28-660
2,4-Dichlorophenol	7.99-330
Diethylphthalate	6.33-330
2,4-Dimethylphenol	6.99-330
Dimethylphthalate	11.7–330
Di-n-butyl phthalate	14-330
Dinitro-o-cresol	16–1600
2,4-Dinitrophenol	15–1600
2,4-Dinitrotoluene	5–3300
2,6-Dinitrotoluene	3–330
Di-n-octyl phthalate	8.99–330
Diphenylamine	7–84.9
bis(2-Ethylhexyl) phthalate	7–330

Refer to footnotes at end of table.

# Table A-4 (Concluded) Summary of DSS SWMU 136, Building 907 Septic Systems Confirmatory Soil Sampling, SVOC Analytical MDLs October 1993–August 2000 (Off-Site Laboratories)

	EPA Method 8270 <sup>a</sup> Detection Limit
Analyte	(μg/kg)
Fluoranthene	3.33–330
Fluorene	3–330
Hexachlorobenzene	4.66–330
Hexachlorobutadiene	6.66–330
Hexachlorocyclopentadiene	2.33–330
Hexachloroethane	4.33–330
Indeno(1,2,3-cd)pyrene	6.67–330
Isophorone	2.33–330
2-Methylnaphthalene	4–330
4-Methylphenol	5.66–330
Naphthalene	3.33–330
2-Nitroaniline	56–1600
3-Nitroaniline	37–1600
4-Nitroaniline	52–1600
Nitrobenzene	11–330
2-Nitrophenol	3.66–330
4-Nitrophenol	21–6700
n-Nitrosodiphenylamine	30–330
n-Nitrosodipropylamine	6.66–3300
Pentachlorophenol	616700
Phenanthrene	4–330
Phenol	3.66-6700
Pyrene	8.66-3300
1,2,4-Trichlorobenzene	4.66-3300
2,4,5-Trichlorophenol	24.3-1600
2,4,6-Trichlorophenol	5.33–330

Note: Because of the long time period covering sample collection at this site, MDL ranges are presented. MDLs were not routinely reported, or were reported as ranges, by the laboratories for analyses performed in the early- to mid-1990s.

<sup>a</sup>EPA November 1986.

DSS = Drain and Septic Systems.

EPA = U.S. Environmental Protection Agency.

MDL = Method detection limit.
 μg/kg = Microgram(s) per kilogram.
 SWMU = Solid Waste Management Unit.
 SVOC = Semivolatile organic compound.

# Table A-5 Summary of DSS SWMU 136, Building 907 Septic Systems Confirmatory Soil Sampling, PCB Analytical Results August 2000 (Off-Site Laboratory)

				PCBs	
				(EPA Method 8082a	)
	Sample Attribut	es		(μg/kg)	•
Record		Sample			
Numberb	ER Sample ID	Depth (ft)	Sample Date	Aroclor-1260	
	pit borehole samples				
603366	TA2-136-SPG-S-13	13	8-22-00	ND (31 J) H	
603366	TA2-136-SPG-DU-13	13	8-22-00	ND (31 J) H	
603366	TA2-136-SPG-S-18	18	8-22-00	ND (31 J) H	
HE drain l	ine outfall ditch samples			(0.0)	
603366	TA2-136-DITCH-0/0-S-7	7	8-21-00	ND (31 J) H	
603366	TA2-136-DITCH-1/3-S-4	4	8-21-00	ND (31 J) H	
603366	TA2-136-DITCH-1/3-S-9	9	8-21-00	ND (31 J) H	
603366	TA2-136-DITCH-2/3-S-4	4	8-22-00	ND (31 J) H	
603366	TA2-136-DITCH-2/3-S-9	9	8-22-00	ND (31 J) H	
Drainfield	borehole samples			115 (01 0) 11	
603366	TA2-136-SELAT-S-10	10	8-22-00	ND (31 J) H	
603366	TA2-136-SELAT-S-15	15	8-22-00	ND (31 J) H	
603366	TA2-136-SOUTHLAT-S-8	8	8-22-00	ND (31 J) H	
603366	TA2-136-SOUTHLAT-S-13	13	8-22-00	ND (31 J) H	
603366	TA2-136-SWLAT-S-8	8	8-22-00	ND (31 J) H	
603366	TA2-136-SWLAT-S-13	13	8-22-00	ND (31 J) H	
Quality As	surance/Quality Control Sample		3 22 00	140 (010)11	
603366	TA2-136-SPT-EB	I NA I	8-22-00	<u> </u>	3 J

Note: Values in **bold** represent detected analytes.

<sup>a</sup>EPA November 1986.

<sup>b</sup> Analysis	s request/chain-of-custody record.		
DITCH	<ul> <li>Sample in HE drain line outfall ditch.</li> </ul>	μg/kg μg/L	= Microgram(s) per kilogram. = Microgram(s) per liter.
DSS DU	<ul><li>= Drain and Septic Systems.</li><li>= Duplicate sample.</li></ul>	NA	= Not applicable.
EB	= Equipment blank.	ND()	= Not detected above the MDL, shown in parentheses.
EPA	<ul> <li>U.S. Environmental Protection</li> <li>Agency.</li> </ul>	PCB S	<ul><li>Polychlorinated biphenyl.</li></ul>
ER	= Environmental Restoration.	SELAT	<ul><li>Soil sample.</li><li>Southeastern lateral.</li></ul>
ft	= Foot (feet).	SOUTHLAT	= Southern lateral.
H	= The holding time was exceeded for	SPG	= Seepage pit.
HE	the associated sample analysis. = High explosive(s).	SPT SWLAT	= Septic tanks project. = Southwestern lateral.
ID	= Identification.	SWMU	= Solid Waste Management
J	<ul> <li>Analytical result was qualified as</li> </ul>		Unit.
MDL	an estimated value. = Method detection limit.	TA	= Technical Area.

# Table A-6 Summary of DSS SWMU 136, Building 907 Septic Systems Confirmatory Soil Sampling, PCB Analytical MDLs August 2000 (Off-Site Laboratory)

	EPA Method 8082 <sup>a</sup> Detection Limit
Analyte	(μg/kg)
Aroclor-1016	30
Aroclor-1221	30
Aroclor-1232	30
Aroclor-1242	30
Aroclor-1248	30
Aroclor-1254	31
Aroclor-1260	31

# <sup>a</sup>EPA November 1986.

DSS = Drain and Septic Systems.

EPA = U.S. Environmental Protection Agency.

MDL = Method detection limit.

μg/kg = Microgram(s) per kilogram.

PCB = Polychlorinated biphenyl.

SWMU = Solid Waste Management Unit.

Table A-7
Summary of DSS SWMU 136, Building 907 Septic Systems
Confirmatory Soil Sampling, HE Compound Analytical Results
October 1993–August 2000
(On- and Off-Site Laboratories)

				HE
				(EPA Method 8330a)
	Sample Att	<del></del>	···	(mg/kg)
Record		Sample		
Number <sup>b</sup>	ER Sample ID	Depth (ft)	Sample Date	2,4,6-Trinitrotoluene
	rench samples			
None	TA2-TR-01	0	10-28-93	ND (1)
None	TA2-TR-01	4	10-28-93	ND (1)
None	TA2-TR-01	6	10-28-93	ND (1)
508001	TA2-TR-02	0.5	11-1-93	ND (1)
508001	TA2-TR-02	3.5	11-1-93	ND (1)
508001	TA2-TR-02	5	11-1-93	ND (1)
	amples near drain lines or septi			
508417	TA2-BH-05	6	3-10-94	ND (1)
508417	TA2-BH-05	11	3-10-94	ND (1)
508417	TA2-BH-05	14.6	3-10-94	ND (1)
508417	TA2-BH-05	21.2	3-10-94	ND (1)
508417	TA2-BH-05	30	3-10-94	ND (1)
508417	TA2-BH-05 (DU)	30	3-10-94	ND (1)
2044	TA2-BH-08-11.0	11	11-2-94	ND (0.25)
2044	TA2-BH-08-16.5	16.5	11-2-94	ND (0.25)
2044	TA2-BH-08-21.0	21	11-2-94	ND (0.25)
2094	TA2-BH-09-10.0	10	11-6-94	ND (0.25)
2094	TA2-BH-09-16.0	16	11-6-94	ND (0.25)
2094	TA2-BH-09-20.0	20	11-6-94	ND (0.25)
2094	TA2-BH-09-28.0	28	11-6-94	ND (0.25)
2094	TA2-BH-09-38.5	38.5	11-6-94	ND (0.25)
04283	TA2-ST-09-6.67 (On-site	6.67	8-29-95	ND (0.076)
	laboratory)			(2,2,5)
04283	TA2-ST-10-6.67 (On-site	6.67	8-29-95	0.076 J (1
	laboratory)			•
	borehole samples			
603366	TA2-136-SPG-S-13	13	8-22-00	ND (0.020)
603366	TA2-136-SPG-DU-13	13	8-22-00	ND (0.020)
603366	TA2-136-SPG-S-18	18	8-22-00	ND (0.020)
HE drain lin	e outfall ditch samples			
603075	TA2-136-HEOUT-PIPE-S75°	0.75	3-8-00	ND (0.0141)
603075	TA2-136-HE-OUT-S-2.0 <sup>c</sup>	2.0	3-8-00	ND (0.0141)
603366	TA2-136-DITCH-0/0-S-7	7	8-21-00	ND (0.020)
603366	TA2-136-DITCH-1/3-S-4	4	8-21-00	ND (0.020)
603366	TA2-136-DITCH-1/3-S-9	9	8-21-00	ND (0.020)
603366	TA2-136-DITCH-2/3-S-4	4	8-22-00	ND (0.020)
603366	TA2-136-DITCH-2/3-S-9	9	8-22-00	ND (0.020)

Refer to footnotes at end of table.

# Table A-7 (Concluded)

# Summary of DSS SWMU 136, Building 907 Septic Systems Confirmatory Soil Sampling, HE Compound Analytical Results

October 1993-August 2000 (On- and Off-Site Laboratories)

Record	Sample Attr	ributes		HE (EPA Method 8330 <sup>a</sup> ) (mg/kg)
Number <sup>b</sup>	ED Commit ID	Sample		
	ER Sample ID porehole samples	Depth (ft)	Sample Date	2,4,6-Trinitrotoluene
			_	
603366	TA2-136-SELAT-S-10	10	8-22-00	ND (0.020)
603366	TA2-136-SELAT-S-15	15	8-22-00	ND (0.020)
603366	TA2-136-SOUTHLAT-S-8	8	8-22-00	ND (0.020)
603366	TA2-136-SOUTHLAT-S-13	13	8-22-00	ND (0.020)
603366	TA2-136-SWLAT-S-8	8	8-22-00	ND (0.020)
603366	TA2-136-SWLAT-S-13	13	8-22-00	
Quality Ass	urance/Quality Control Samples	(uo/L)	0 22-00	ND (0.020)
508417	TA2-BH-05 (EB)	NA NA	3-10-94	ND (0)
2044	TA2-BH-08 (EB)	NA NA	11-2-94	ND (2)
2094	TA2-BH-09 (EB)	NA NA	11-6-94	ND (0.11)
603366	TA2-136-SPT-EB	NA NA	8-23-00	ND (0.11) ND (0.11)

Note: Values in **bold** represent detected analytes.

HEOUT-PIPE = Sample in HE drain line outfall ditch.

ID = Identification.

J() = The reported value is greater than or equal to the MDL but is less than the practical quantitation limit, shown in parentheses.

MDL = Method detection limit. μg/L = Microgram(s) per liter. = Milligram(s) per kilogram. mg/kg NA = Not applicable.

ND()

= Not detected above the MDL, shown in parentheses.

= Soil sample. **SELAT** = Southeastern lateral. SOUTHLAT = Southern lateral. SPG = Seepage pit. SPT = Septic tanks project. ST = Septic tank. **SWLAT** = Southwestern lateral.

SWMU = Solid Waste Management Unit.

TA = Technical Area

TR = Trench.

<sup>&</sup>lt;sup>a</sup>EPA November 1986.

<sup>&</sup>lt;sup>b</sup>Analysis request/chain-of-custody record.

<sup>&</sup>lt;sup>c</sup>Same location as sample TA2-136-DITCH-0/0-S-7.

BH = Borehole.

DITCH = Sample in HE drain line outfall ditch.

DSS = Drain and Septic Systems.

DU = Duplicate sample. EΒ = Equipment blank.

**EPA** = U.S. Environmental Protection Agency.

<sup>=</sup> Environmental Restoration ER

ft = Foot (feet).

**HE-OUT** = Sample in HE drain line outfall ditch.

<sup>=</sup> High explosive(s). HE

# Table A-8 Summary of DSS SWMU 136, Building 907 Septic Systems Confirmatory Soil Sampling, HE Compound Analytical MDLs October 1993–August 2000 (On- and Off-Site Laboratories)

	EPA Method 8330 <sup>a</sup>
	Detection Limit
Analyte	(mg/kg)
2-Amino-4,6-dinitrotoluene	0.0134-0.25
4-Amino-2,6-dinitrotoluene	0.0101-0.25
1,3-Dinitrobenzene	0.0134–5
2,4-Dinitrotoluene	0.012–5
2,6-Dinitrotoluene	0.0157–5
HMX	0.0168–5
Nitrobenzene	0.014–5
2-Nitrotoluene	0.0152-0.25
3-Nitrotoluene	0.0116-0.25
4-Nitrotoluene	0.0116-0.25
PETN	0.124-0.5
RDX	0.0125–5
Tetryl	0.0155–5
1,3,5-Trinitrobenzene	0.0119–5
2,4,6-Trinitrotoluene	0.0141–5

Note: Because of the long time period covering sample collection at this site, MDL ranges are presented. MDLs were not routinely reported, or were reported as ranges, by the laboratories for analyses performed in the early- to mid-1990s. aEPA November 1986.

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DSS = Drain and Septic Systems.

EPA = U.S. Environmental Protection Agency.

HE = High Explosive(s).

HMX = Octahydro-1,3,5,7-tetranitro-1,3,5,7-tetrazocine.

MDL = Method detection limit.mg/kg = Milligram(s) per kilogram.PETN = Pentaerythritol tetranitrate.

RDX = Hexahydro-1,3,5-trinitro-1,3,5-triazine.

SWMU = Solid Waste Management Unit.

Tetryl = Methyl-2,4,6-trinitrophenylnitramine.

AL/6-04/WP/SNL04:r5517.doc

Table A-9
Summary of DSS SWMU 136, Building 907 Septic Systems
Confirmatory Soil Sampling, Metals, Including Hexavalent Chromium, and Cyanide Analytical Results
October 1993–August 2000
(On- and Off-Site Laboratories)

Backgroun	603366	603366	603366	603366	603366	603075	603075	HE drain lin	603366	603366	603366	Seepage pi	04282		04282	2094	2094	2094	2094	2094	2094	2094	2094	2044	2044	2044	2044	2044	2044	2044	2044	2044	2044	508417	508417	508417	508417	508417	Borehole sa	508001	508001	None	None	None	Drain line tr	Numberb		
Background Concentration—North Area Supergroupe	TA2-136-DITCH-2/3-S-9	TA2-136-DITCH-2/3-S-4	TA2-136-DITCH-1/3-S-9	TA2-136-DITCH-1/3-S-4	TA2-136-DITCH-0/0-S-7	TA2-136-HE-OUT-S-2.0 <sup>d</sup>	TA2-136-HEOUT-PIPE-S75 <sup>th</sup>	HE drain line outfall ditch samples	TA2-136-SPG-S-18	TA2-136-SPG-DU-13	603366 TA2-136-SPG-S-13	t borehole samples	(On-site laboratory)	(On-site laboratory)	TA2-ST-09-6.67	TA2-BH-09-38 5	TA2-BH-09-28.0	TA2-BH-09-20.75	TA2-BH-09-20.0	TA9-RH-09-16 0	TA2 BH 00-15 75	TA2 BH 00 10 75	TA2-BH-U9-5.5	1A2-BH-08-50.0	TA2-BH-08-41.0	TA2-BH-08-31.0	TA2-BH-08-30.75	TA2-BH-08-21.00	TA2-BH-08-20.75	TA2-BH-08-16-50	TA2-BH-08-11.0	TA2-BH-08-10.5	TA2-BH-08-5.0	TA2-BH-05 (DU)	TA2-BH-05	TA2-BH-05	TA2-BH-05	TA2-BH-05	Borehole samples near drain lines or septic tanks	TA2-113-02	TA2-TR-02	TA2-TR-07	TA2-TR-01	TA2-TR-01	Drain line trench samples	ER Sample ID	Sample Attributes	
supergroup <sup>e</sup>	9	4	9	4	7	2.0	0.75		18	13	13		6.67		6.67	38.5	28	20.75	28	16	15.75	10.75	3.0	<u></u>	41	31	30.75	21	20.75	16.5	5 =	10.5	5	30	30	14.6	11	6	ic tanks	5 6	3.5	0.5	6 4	• 0	,	Depth (ft)		
	8-21-00	8-21-00	8-21-00	8-21-00	8-21-00	3-8-00	3-8-00		8-22-00	8-22-00	8-22-00		8-29-95	200	8-29-95	11-6-94	11-6-94	11-6-94	11-6-94	11-6-94	11-6-94	11-6-94	11-6-94	11 6 04	11-2-94	11-2-94	11-2-94	11-2-94	11-2-94	11-2-94	11-2-94	11-2-94	11-2-94	3-10-94	3-10-94	3-10-94	3-10-94	3-10-94		11-1-93	11-1-93	11-1-93	10-28-93	10-28-93	2000	Date	Cample	
3.9	NR	NR	NR	NR	RN	NR	N.	5	NR	NR	NR		NO (10)	25 (20)	ND (10)	ND (6)	ND (6)	NR R	ND (6)	ND (6)	NR :	NR (	ND (12)	ND (c)	ND (6)	ND (6)	NR	ND (6)	NR	ND (12)	NR O		NE NE	ND (6)	ND (6)	ND (6)	ND (6)	2.1 J (6)		ND (6)	ND (6)	ND (6)	ND (6)	ND (8)	5 6	Antimony		
4.4	2.1	4	2	3	2.9	3.62	3.54		1.2	2.2	2.1		ייני	20 (60	ND (50)	2.3	3	NR	2.3	=	N.	N.P.	5.7	200	0.96.0 (1)	3.5	NR NR	3.1	NR	2.8	NR.	NR	NR	3.8	3.4	1.4	2.1	2		2.9	3.4	2.1	17	) - J	10	Arsenic		
200	93.2	169	101	162	140	118	132	33	58.5	62.9	61.5			38	310	83.2	169	RN	116	31.2	N.	NR NR	944	NR G	16.8	37.6	NA	159	NA	174	NN S		NR.	150	149	97.9	72.8	130		96.6	101	80.5	74.4	104	77.9	Barium		
0.8	NR P	Z Z	Z	<b>1</b>	N.	NR	3	Š	ZI	S N	Z Z			200	0 ND (3.4)	0.35	0.46	NR	0.34	ND (0.2)	NR.	_	z	N. S.	024	0 1	NR	0.48	NR	0.32 J (0.4)	NH S	2 2	S S	ND (0.2)	0.25	0.00	ND (0.2)	ND (0.2)		0.45	0.41	0.26	0.26	0.26	0.36	Beryllium		
0.9	ND (0.03)	0.185 J (0.476)	00	70	ND (0.03)	ND (0.03)	ND (0.03)			ND (10		ND (0.5)	ND (0.5)	NR	ND (0.5)	ND (0.5)	NR	NH NH	ND (1)	NR	ND (0.5)	NO (0.0)	NI	ND (0.5)	NR	ND (1)	NR	N N	N N	0.71	ND (0.5)	ND (0.5)	ND (0.5)	ND (0.5)		ND (0.5)	ND (0.5)	ND (0.5)	ND (0.5)	ND (0.5)	0.68	Cadmium						
9.71	7.7	6./	9.5	2 3	10.4	9.67	19.7		5.8	50	5.6			2	NE NE	7.3	9.5	N.R.	7.6	6.3	N.R.			Z i	42	4 15.5	Z	8.3		5.1	NR	4 9	i K	8.4	6.9	9.3	6.7	9.3		6.5	6.2	6	4.3	3.9	6.3	Chromium		
2	3 3	5 3	5 5	3 3	S	¥.		N N	2	2 3	2 2	,		ND (0.38)	ND (0.38)	N N	N.	N.	NR R	NB	N.R	NR	NR R	N.	S :			S S	NR NR	NR	NR.	2 3	3 3	i ≅	NR	NA S	S Z	5 5		NR	NR	NR R	NR.	\$ :	NR NR	(VI)	Chromium	
/.1	71 1	5 3	200	2 3	5 3	S S		NR I	CINI	2 2	2 3	Š	-	ND (10)	ND (10)	4.7	5.5	NR	4.6	3.2	NR	NR	2.4	NR R	2.8	3	3	5 6	NR R	3.2	NR.	42	ž Z	5.2	5.7	6.1	60.3	4.00		4.9	5	4.7	3.5	3.8	3.6	Cobalt		
-	\$   <del>3</del>	5 5	2 3	5 3		; <u>₹</u>	3	NR I	145	20 3	2 2	Š		ND (20)	b) ND (20)		11.3	N.	7.7	4.9	NR	ND ND	2.7 J (4)	NR	3.9	6.4		8.6	NR NR	4.3	¥.	3.4		10.3	9.3	10.8	2.5	30		11.4	11.6	9.7	12.2	7.3	9.6	Copper		Metals (EPA
74	NO (0.091)	ND (0.091)	ND (0.091)	ND (0.091	ND (0.091)	N N			140 (0.00)	ND (0.001)	ND (0.091)	10000		NA NA	NR R	Z	ND (0.5)	ND (0.5)	Z R	N <sub>R</sub>	ND (0.5)	ND (0.5)	NR	ND (0.5)	ND (0.5)	$\neg$	NA CO.O.	NE	ND (0.5)	NR.	ND (0.5)	NA	ND (0.5)	S N	NR	Z :	2 3		5	NR	NA NA	N <sub>B</sub>	NR.	N <sub>E</sub>	S.	Cyanide		Methods 600
11.4	4.50	$\dagger$	$\dagger$	44.		9.4		58.4	r		39.5	1		ND (10)	ND (10)	4.8	5.8	Z	4.3	2.2	N.R	NR.	4.2	NR	5.6	2.4	8 2	2 0.0	NR.	3.4	NR.	3.8	2 2	200	3.6	5	2.6	37	27	7.7	10.2	4.3	4.3	4.4	14.5	Lead		Metals (EPA Methods 6000/7000/9010A/908 <sup>a</sup> ) (mg/kg)
2	831			200				NR		Z .	N .	NO		120	140	214	284	Z Z	226	148	NR	NR	54.6	NR.	134	140	548	NI SUP	2	133	NR	=	20 3	228	268	215	188	177	177	225	206	193	171		.5	Manganese		V908 <sup>a</sup> ) (mg/k
2	0.01	NO (0.0)	ND (0.017)	ND (0.01)	ND (0.017)	0.181		_	1000	ND (0.017)	ND (0.017)	ND (0 017		ND (0.06)	ND (0.06)	ND (0.1)	100	Z Z	ND (0.1	ND (0.1	NR R	N.R.	ND (0.1)	NR	ND (0.1	ND (0.1	ND (0.1	ND (o.		ND (0.1	NR	ND (0.1	5	ND (0.1	ND (0.1	ND (0.1)	ND (0.1	ND (0.1)	2000	ND (0.1)	ND (0.1	ND (0.	ND (0.1)	ND (0.1)	ND (0.1)	e Mercury		g)
	1	NR.						1.13 NR	ļ		1	ND	-	7.2 J (15)	ND (4)	╞		1				N.R	4	NR R		3		NA C	1	6			S :	ND 7.9		8	4	ω	62		6.7		3.5	1) 4.5		Nickel		
		ND (0.24)	ND (0.24	ND (0.24)	ND (0.24)	NO (0, 146)	(0.467)	0.465 J	}	ND (0.24)	ND (0.24)	ND (0.24		ND (50)	(OC) UN	ND (0.3)	ND (0.5)	NO 00	ND (0.0)	ND (0.5)	NR.	N.R.	10	NR	ND (0.5)	ND (0.5)	ND (0.5)	NR O	2000	0.48 J (0.5)	NE NE	ND (0.5)	S .	NE C	NB (3	ND (2)	0.19 J (1)	0.24 J (1)	011.1(1)	NO	ND (1)	ND (1)	NB O	ND (1)	ND (1)	Selenium		
1	<u> </u>	+	ND (0	ND (0	ND (0.14)	200				) ND (0.14)	ND (0	ND (0.14)		50) ND (10)		4	+	$\dagger$	+	$\dagger$	+	Z	(1) ND			ND (1)	-	N .	200	0 2	NA	7	NA.	NA	(1) 0.93 J (1)	0.79 J	0.72 J	2	081 J (1)		0.52 J (1)			1) ND (1)		<u>u</u>		
	_	7	1		ND :	<del>                                     </del>		60.9 NR		NR		NR	-	10) ND (200)	10)		- 1	ءِ		(1) ND (1)			(2) ND (2)				_		3	ND (1)	7	<b>ID (1)</b> 0.61 J (1)	NR R	+	ND (0.5)	╁	) ND (1)	<del>- 1</del>	┥.				ļ		l	1 1	1	
	2.3	NR NR	NR R	NR	NR :	NS E	20	NR		NA	NR	NR	-	00) NR		3	+	$\dagger$	$\dagger$	$\dagger$	+	T			-			+	+		N R		NR	+	20	2.2		+	$\dashv$	-	+		+	S S		-    -		-
	33	NR	NR	N <sub>R</sub>	S .	NE S	N.D.	NR		NR	NR	N.R	_	ND (10)		ND (10)	17.6	24	ND IS	23 -		N N	5	I NI	11.9	12	3	NR.	30.5	NI C	NA NA	23.9	NR R	NR R	24.4	30.5			31.7	5	24	20.7	16.6	16.8	16.6			
	76	NR	NR	NR	NR.	NF :	Z D	NR		S	NR	NR R		)) 19 J (38)		z	26.3	37.9	NR	27.6	18.4	2 2	ND 13.4	NH	17.3	17.5	<b>33.2</b> 58.8	NR	39.3	NR G	19 NH	17.8	N N	N.R.	28.3	30.8	17.9	17.7	19.5	F0.0	26.8	27	ος α	17.5	34.1	1		

Refer to footnotes at end of table.

# Table A-9 (Concluded) Summary of DSS SWMU 136, Building 907 Septic Systems Confirmatory Soil Sampling, Metals, Including Hexavalent Chromium, and Cyanide Analytical Results October 1993–August 2000 (On- and Off-Site Laboratories)

	Sample Attributes											Metals (EPA N	ethods 6000	Metals (EPA Methods 6000/7000/9010A/908-)	(mg/kg)								
Record		Sample	Sample							Chromium	:	)					<u>}</u>	Colori.	Cilvio	The limit	l ranium	Vanadium	Zinc
Numberb	ER Sample ID	Depth (ft)	Date	Antimony	Arsenic	Barium	Beryllium	Cadmium	Chromium	(V)	Cobalt	Copper	Cyanide	Lead	Manganese	Mercury	Nicker	Ocicilian	CHACL	⊦	Cignitian	* ci. ci. ci.	1
Drainfield bo	Drainfield borehole samples														,	15 62 64	,	200	200		NO	Ž.	200
603366	TA2-136-SELAT-S-10	10	8-22-00	NR	2.6	109	NR	ND (0.03)	8.3	NR NR	NR.	L	ND (0.091)	4.8 J	N.	D (0.017)	Z	ND (0.24)		5 3	2 2	Ď	20 2
603366	TA2-136-SELAT-S-15	15	8-22-00	NR	2.6	162	NR R	ND (0.03)	4.6	NR	NR.	NR.	ND (0.091)	3,	NH.	(0.01/)	Z	ND (0.24)	ND (0.14)	2	5 3	5 3	5 3
603366	TA2-136-SOUTH  AT-S-8	30	8-22-00	NH NH	2.9	92.9	NR	ND (0.03)	8.7	NR NR	NR	NR	ND (0.091)	4.5 J	NR NR	ND (0.017)	NR	ND (0.24)	ND (0.14)	N.	Z,	i E	Z
603366	TA2-136-SOLITH AT-S-13	3	8-22-00	S	2.2	288	N N	ND (0.03)	5.4	NR	Z Z	NR	ND (0.091)	3.2 J	NH NH	ND (0.017)	NR.	ND (0.24)	ND (0.14)	NE	N.	Z	Z
603366	TA2-136-SWLAT-S-8	8	8-22-00	NR	2.3	97.3	N.	ND (0.03)	20.2	NR	NR R	NR	ND (0.091)	10.4 J	S	0.16	NH	ND (0.24)	ND (0.14)	Z	i H	5	3
603366	TA2-136-SWLAT-S-13	13	8-22-00	NR	6	598	NR R	ND (0.03)	6.3	NA	NR	N.	ND (0.091)	4.5 J	N.H.	ND (0.017)	Z	ND (0.24)	ND (0.14)	3	2 3	3 3	75
Background	Background Concentration—North Area Supergroup <sup>e</sup>	upergroup <sup>e</sup>		3.9	4.4	200	0.8	0.9	12.8	NO	7.1	17	NO.	11.2	831	6.1	25.4	^	2	\$1.1	2.3	5	,,
Quality Assu	Quality Assurance/Quality Control Samples (mg/L)	s (mg/L)									→	:	5		200	0 0000		_			1 50000 0	ND (0 01)	0.59
508417	TA2-BH-05 (EB)	N A	3-10-94	ND (0.06)		ND (0.01)	ND (0.002) ND (0.005)	ND (0.005)	ND (0.01)	N R	ND (0.01)	ND (0.02)	3	ND (0.005)	ND (0.01)	ND (0.01) ND (0.0002) ND (0.04)		ND (0.000)	(0.01)	(0.000)		(0.01)	0.000
2044	TA2-BH-08 (EB)	NA	11-2-94	ND (0.06)	ND (0.01)	ND (0.01)	ND (0.002)	ND (0.005)	0.0049 J (0.01)	NR.	ND (0.01)	0.018 J (0.02)	ND (0.01)	ND (0.003)	ND (0.01)	ND (0.0002)	ND (0.04)	0.0046 J (0.005)	ND (0.01)	ND (0.01)	NR NR	ND (0.01)	0.018 J (0.02)
2094	TA2-BH-09 (EB)	N <sub>P</sub>	11-6-94	ND (0.06)	ND (0.01)	ND (0.01)	ND (0.002)	ND (0.005)	ND (0.01)	N.	ND (0.01)	ND (0.02)	ND (0.01)	ND (0.003)	ND (0.01)	ND (0.0002) ND (0.04)		ND (0.005)	ND (0.01)	ND (0.01)	N.	ND (0.01)	0.0023 J (0.02)
603366	TA2-136-SPT-EB	NA.	8-23-00	NR R	ND (0.0014)	0.0033 J (0.2)	NF.	ND (0.0003)	0.0064	NR R	N <sub>R</sub>	N <sub>B</sub>	ND (1.6)	ND (0.0019)	N.S.	ND (0.0001)	NR	ND (0.0024)	ND (0.0014 J)	NE NE	SE SE		N.R

Note: Values in bold exceed background concentrations.

<sup>a</sup>EPA November 1986.

<sup>b</sup>Analysis request/chain-of-custody record. <sup>C</sup>Sample depth shown in brackets.

<sup>e</sup>Dinwiddie September 1997. <sup>d</sup>Same location as sample TA2-136-DITCH-0/0-S-7.

USGS 1994.

= Borehole.
= Sample in HE drain line outfall ditch.
= Drain and Septic Systems.
= Duplicate sample.
= Equipment blank.
= U.S. Environmental Protection Agency.
= Environmental Restoration
= Foot (feet).
= High explosive(s).

= Publ (very).

= High explosive(s).

= Sample in HE drain line outfall ditch.

= Identification.

= Analytical result was qualified as an estimated value.

= The reported value is greater than or equal to the MDL but is less than the practical quantitation limit, shown in parentheses.

= Net reported detection limit.

= Milligram(s) per kilogram.

= Not iligram(s) per kilogram.

= Not detected above the MDL, shown in parentheses.

= Not detected but the MDL, shown in parentheses, equals or exceeds the background concentration.

= Not quantified.

= Not quantified.

= Not exported.

= Southern lateral.

= Southern lateral.

= Southern lateral.

= Septic tanks project.

= Septic tanks project.

= Septic tanks project.

= Septic tanks project.

= Southwastern lateral.

HE HE HE HE-OUT HEOUT-PIPE ID J () MNDL mg/kg mg/L NA ND () 
# Table A-10

# Summary of DSS SWMU 136, Building 907 Septic Systems Confirmatory Soil Sampling, Metals, Including Hexavalent Chromium, and Cyanide Analytical MDLs October 1993–August 2000 (On- and Off-Site Laboratories)

	EPA Method 6000/7000/9010A/908 <sup>a</sup>
	Detection Limits
Analyte	(mg/kg)
Antimony	6– 50
Arsenic	0.131–3
Barium	0.0465–200
Beryllium	0.2–5
Cadmium	0.03–5
Chromium	0.0645–20
Chromium (VI)	0.054
Cobalt	1–50
Copper	2–25
Cyanide	0.091-0.5
Lead	0.099-3
Manganese	150
Mercury	0.0152-0.5
Nickel	4–50
Selenium	0.146-3
Silver	0.062-5
Thallium	0.5–3
Vanadium	1–50
Zinc	2–50

Note: Because of the long time period covering sample collection at this site, MDL ranges are presented. MDLs were not routinely reported, or were reported as ranges, by the laboratories for analyses performed in the early- to mid-1990s.

<sup>a</sup>EPA November 1986.

DSS = Drain and Septic Systems.

EPA = U.S. Environmental Protection Agency.

MDL = Method detection limit.mg/kg = Milligram(s) per kilogram.SWMU = Solid Waste Management Unit.

Table A-11
Summary of DSS SWMU 136, Building 907 Septic Systems
Confirmatory Soil Sampling, Gamma Spectroscopy Analytical Results
October 1993–August 2000
(On- and Off-Site Laboratories)

							Othyita / CDA	Activity (EDA 900 Od) (ACI/a)			
	Sample Attributes					`	ייין ובועוו	(8/10/1/ 0:008			000
Record		Sample	Sample	Cesium-137	1-137	Thorium-232	m-232	Uranium-235	m-235	Uranium-238	m-238
Numberb	ER Sample ID	Depth (ft)	Date	Result	Error	Result	Error <sup>c</sup>	Result	Error <sup>c</sup>	Result	Error
Drain line	Drain line trench samples										
None	TA2-TR-01 (Off-site laboratory)	0	10-28-93	0.22	0.083	0.81	0.3	ND (0.062)	1	ND (1.5)	:
None	TA2-TR-01 (Off-site laboratory)	4	10-28-93	ND (0.053)		0.94	0.3	0.036	0.084	(1.5) ND	:
None	TA2-TR-01 (Off-site laboratory)	9	10-28-93	ND (0.048)	;	1.1	0.31	0.034	0.071	ND (1.4)	
508002	TA2-TR-02 (Off-site laboratory)	0.5	11-1-93	0.21	0.073	0.88	0.26	0.072	0.093	ND (1.5)	:
508002	TA2-TR-02 (Off-site laboratory)	3.5	11-1-93	ND (0.049)	;	1	0.36	ND (0.062)	:	ND (1.3)	:
508002	+-	5	11-1-93	ND (0.051)	;	1	0.27	0.085	0.07	ND (1.4)	1
Borehole	Borehole samples near drain lines or septic tanks	tanks					İ				
508418	TA2-BH-05 (Off-site laboratory)	5	3-10-94	ND (0.056)	:	1.1	0.33	NR	:	RN	:
508418	TA2-BH-05 (Off-site laboratory)	10	3-10-94	ND (0.047)		1	0.28	NR.	:	NR	:
508418	TA2-BH-05 (Off-site laboratory)	14.5	3-10-94	ND (0.055)	:	0.94	0.33	an En	:	NR	
508418	TA2-BH-05 (Off-site laboratory)	20	3-10-94	ND (0.057)	;	1.1	0.33	NR		NR	
508418	TA2-BH-05 (Off-site laboratory)	30	3-10-94	ND (0.052)	;	0.98	0.33	RN		NR	:
508418	TA2-BH-05 (DU) (Off-site	30	3-10-94	ND (0.057)	1	6.0	0.33	RN	:	a E	;
	laboratory)										
04284	TA2-ST-09-6.67	6.67	8-29-95	ND (0.0529)	;	0.0535	0.192	ND (0.348)		ND (1.96)	:
Seepade	Seepage pit borehole samples										
603365	TA2-136-SPT-S-13	13	8-22-00	ND (0.049)	1	0.81	0.476	0.145	0.219	ND (0.705)	-
603365	TA2-136-SPT-DU-13	13	8-22-00	ND (0.0471)	:	0.921	0.542	ND (0.248)	-	ND (0.671)	:
603365	TA2-136-SPT-S-18	18	8-22-00	ND (0.0333)	:	0.851	0.439	0.107	0.166	ND (0.521)	;
HE drain	HE drain line outfall ditch samples										
603074	TA2-136-HEOUT-PIPE-S75 <sup>d</sup>	0.75	3-8-00	ND (0.0348)	;	0.836	0.504	0.273	0.19	0.993	0.828
603074	TA2-136-HE-OUT-S-2.0 <sup>d</sup>	2.0	3-8-00	ND (0.0378)		0.896	0.497	0.163	0.146	0.952	0.541
603365	TA2-136-DITCH-0/0-S-7	7	8-21-00	0.0276	0.0184	0.903	0.426	0.124		ND (0.66)	:
603365	TA2-136-DITCH-1/3-S-4	4	8-21-00	ND (0.0355)	1	0.906	0.504	0.242	0.178	ND (0.544)	:
603365	TA2-136-DITCH-1/3-S-9	6	8-21-00	ND (0.0251)	1	0.638	0.36	ND (0.179)	;	ND (0.629)	;
603365	TA2-136-DITCH-2/3-S-4	4	8-22-00	0.0948	0.0396	0.92	0.477	ND (0.223)	:	ND (0.615)	1
603365	TA2-136-DITCH-2/3-S-9	6	8-22-00	ND (0.0251)	1	0.717	0.375	ND (0.183)	:	ND (0.637)	
Backgrou	Background activity-North Area Supergroupe	edn e		0.084	Ϋ́	1.54	NA	0.18	NA A	1.3	ΑΝ

Refer to footnotes at end of table.

Confirmatory Soil Sampling, Gamma Spectroscopy Analytical Results Summary of DSS SWMU 136, Building 907 Septic Systems (On- and Off-Site Laboratories) October 1993-August 2000 Table A-11 (Concluded)

							ctivity (EPA	Activity (EPA 900.0 <sup>a</sup> ) (pCi/g)			
	Sample Attributes					1	000	1 Iranium-235	2.235	Uranium-238	-238
		0	olda	Cesium-137	n-13/	1 norman - 252	11-636	Ola IId	200		
Diocold		Carrylle The At	Carl Fig.	+linood	n rord	Besilt	Frror	Result	<b>E</b> rror <sup>c</sup>	Result	Error
Number	ER Sample ID	Deptin (iii)	Dale	ווחפפע	5	1000					
<b>Drainfield</b> b	Drainfield borehole samples					1100	0.630	VICO 0/ 014		(10 824)	;
10000	C+ C +4 -10 CC+ C+1	ç	8-21-00	R-21-00   ND (0.0437)	;	0.844	0.4/0	ND (0.221)		120:01	
603365	603365 IAZ-136-SELAI-5-10	2 9		V-7-00 00 014		0.458	0.249	0.0801	0.129	ND (0.437)	;
603365	603365 TA2-136-SELAT-S-13	ည	8-22-00	ND 0.0277				197 101		(PET 0) CIN	:
20000	A TA LITTI CO GC, CAT	α	8-22-00	ND (0.0322)	1	0.808	0.404	ND (0/ 1/0)		201	
603365	603365 IAZ-130-50017LA1-3-0	,		(00000)		0.803	0.352	ND (0.184)	;	ND (0.494)	:
603365	ENRIGHT TAP-136-SOUTHLAT-S-13	<u></u>	8-22-00	ND (0.0329)	;	20.5	3	+ 33,	0,00	(33.0) (14.	
3	0 0 10 00 00 00	a	8-22-00	(9750 O) UN	:	ND (0.147)		201.0	0.109	(CC-C)	
603365	IA2-136-SWLAI-5-8	٥	20-0	7 3 3 3 3	00100	0800	0.247	ND (0.184)	;	0.703	0.468
ROSSEE	AN3385 TA2.136-SWI AT-S-13	<del>1</del> 3	8-22-00	0.0346	0.0403	00.0	2				VIV
202000	0.0000000000000000000000000000000000000	1		0.084	Ϋ́	1.54	¥	0.18	Y Y		Į.
Backgroun	Background activity—North Area Supergroup	b.		2000							
	(Inclinity Complete Complete (nCi/ml)	(nCi/m)									
Cuality Ass	surance/quainty control samples	2	, 0	(000 0) 214		az	1	N.	:	œ	•
508418	508418 TA2-BH-05 (EB)	NA	3-10-94	3-10-94 ND (0.020)	:	117		(30 + 0)		ND (0.358)	:
10000	TAN 496 COT ER	ΝA	8-23-00	8-23-00   ND (0.0202)	;	ND (0.124)		ND (0.130)		7,555,7	
- 602500	77-120-02-74										

Note: Values in bold exceed background soil activity.

PEPA November 1986

OAnalysis request/chain-of-custody record.

<sup>c</sup>Two standard deviations around the mean detected activity.

dSame location as sample TA2-136-DITCH-0/0-S-7.

<sup>e</sup>Dinwiddie September 1997.

= Borehole. BH DITCH DSS

= Sample in HE drain line outfall ditch. = Drain and Septic Systems. = Duplicate sample.

Equipment blank.
U.S. Environmental Protection Agency.
Environmental Restoration.
Foot (feet). EPA

High explosive(s).Sample in HE drain line outfall ditch. = !dentification. HE H**E**OUT-PIPE

 Not detected above the MDA, shown in parentheses.
 Not detected but the MDA (shown in parentheses) exceeds background activity. = Minimum detectable activity. Not applicable.

= Error not calculated for nondetect results. = Technical Area. = Trench.

= Solid Waste Management Unit.

= Southwestern lateral. = Septic tanks project.

= Septic tank.

SELAT SOUTHLAT SPT ST SWLAT SWMU

= Picocurie(s) per gram. = Picocurie(s) per milliliter.

pCi/g pCi/mL S

Soil sample.Southeastern lateral.Southern lateral.

# Table A-12 Summary of DSS SWMU 136, Building 907 Septic Systems Confirmatory Soil Sampling, Tritium Analytical Results October 1993–March 2000 (Off-Site Laboratories)

	Sample Attribute	s		Activity (EPA 906.	0a) (pCi/L)
Record		Sample	Sample		
Number <sup>b</sup>	ER sample ID	Depth (ft)	Date	Result	Error <sup>c</sup>
Drain line	trench samples		<del></del>		
None	TA2-TR-01	0	10-28-93	65	160
None	TA2-TR-01	4	10-28-93	120	160
None	TA2-TR-01	6	10-28-93	15	150
508002	TA2-TR-02	0.5	11-1-93	360	180
508002	TA2-TR-02	3.5	11-1-93	ND (290)	
508002	TA2-TR-02	5	11-1-93	260	170
Borehole	samples near drain lines o	r septic tan	ks		
508418	TA2-BH-05	5	3-10-94	370	160
508418	TA2-BH-05	10	3-10-94	110	180
508418	TA2-BH-05	14.5	3-10-94	350	160
508418	TA2-BH-05	20	3-10-94	220	150
508418	TA2-BH-05	30	3-10-94	35	140
508418	TA2-BH-05 (DU)	30	3-10-94	220	140
2046	TA2-BH-08-5.75	5.75	11-2-94	ND (0.023) pCi/g	
2046	TA2-BH-08-10.25	10.25	11-2-94	ND (0.034) pCi/g	••
2046	TA2-BH-08-15.75	15.75	11-2-94	ND (0.02) pCi/g	
2046	TA2-BH-08-20.5	20.5	11-2-94	ND (0.024) pCi/g	
2046	TA2-BH-08-30.5	30.5	11-2-94	ND (0.047) pCi/g	
2046	TA2-BH-08-40.5	40.5	11-2-94	ND (0.025) pCi/g	
2046	TA2-BH-08-50.5	50.5	11-2-94	ND (0.028) pCi/g	
2092	TA2-BH-09-5.25	5.25	11-6-94	0.031 pCi/g	0.03
2092	TA2-BH-09-11.0	11	11-6-94	0.059 pCi/g	0.036
2092	TA2-BH-09-15.25	15.25	11-6-94	0.01 pCi/g	0.024
2092	TA2-BH-09-20.5	20.5	11-6-94	0.055 pCi/g	0.031
2092	TA2-BH-09-30.0	30	11-6-94	0.047 pCi/g	0.026
2092	TA2-BH-09-39.0	39	11-6-94	0.025 pCi/g	0.027
2092	TA2-BH-09-39.0	39	11-6-94	0.012 pCi/g	0.026
HE drain	line outfall ditch sample				
603075	TA2-136-HE-OUT-S-2.0	2_	3-8-00	187	219
	nd Activityd			420 <sup>e</sup>	
Quality A	ssurance/Quality Control S	amples (pC			
	TA2-BH-05 (EB)	NA	3-10-94	1.8	NR
2046	TA2-BH-08 (EB)	NA	11-2-94	ND (180)	
2092	TA2-BH-09 (EB)	NA	11-6-94	ND (180)	

Note: Values in **bold** exceeded background soil activity.

<sup>&</sup>lt;sup>a</sup>EPA November 1986.

<sup>&</sup>lt;sup>b</sup>Analysis request/chain-of-custody record.

<sup>&</sup>lt;sup>c</sup>Two standard deviations around the mean detected activity.

<sup>&</sup>lt;sup>d</sup>Tharp February 1999.

<sup>°420</sup> pCi/L = 0.021 pCi/g, assuming a soil density of 1 gram/cubic centimeter and 5 percent soil moisture.

BH = Borehole

DITCH = Sample in HE drain line outfall ditch.

# Table A-12 (Concluded)

# Summary of DSS SWMU 136, Building 907 Septic Systems Confirmatory Soil Sampling, Tritium Analytical Results October 1993–March 2000

(Off-Site Laboratories)

DSS = Drain and Septic Systems.

DU = Duplicate sample. EB = Equipment blank.

EPA = U.S. Environmental Protection Agency.

ER = Environmental Restoration.

ft = Foot (feet). HE = High explosive(s).

HE-OUT = HE drain line outfall. Same location as TA2-136-DITCH-0/0 sample.

ID = Identification.

MDA = Minimum detectable activity.

NA = Not applicable.

ND () = Not detected above the MDA, shown in parentheses.

ND () = Not detected, but the MDA shown in parentheses, exceeds background activity.

NR = Not Reported.

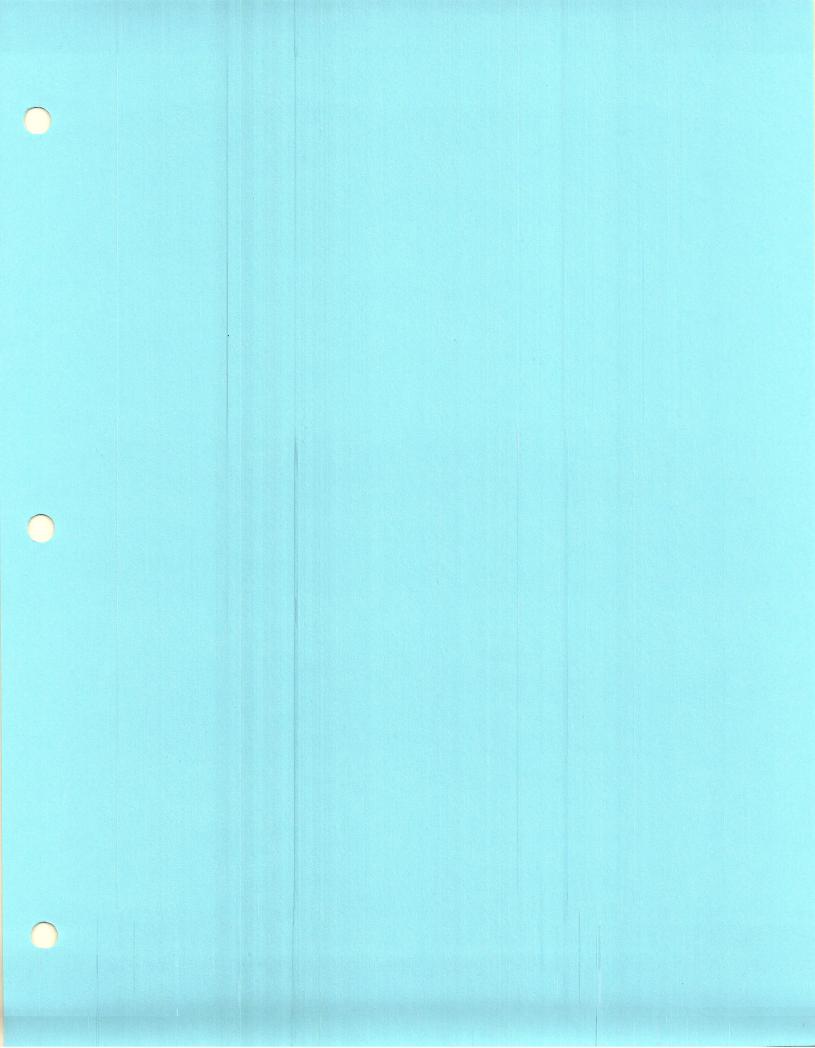
pCi/g = Picocurie(s) per gram. pCi/L = Picocurie(s) per liter.

SWMU = Solid Waste Management Unit.

TA = Technical Area.

TR = Trench.

-- = Error not calculated for nondetect results.



ANNEX B
DSS SWMU 136
HE Catch Box Analytical Data Summary Tables

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# Table B-1

# Summary of DSS SWMU 136, Building 907 Septic Systems, HE Catch Box Confirmatory Sampling, VOC Analytical Results December 2001

(Off-Site Laboratory)

	Sample Attributes	VOCs (EPA Method 8260ª) (μg/kg)
Record Number <sup>b</sup>	ER Sample ID	Toluene
605146	TA2-136-SEEP-AGE-001-S	0.353 J (1.04)
605146	TA2-136-SEEP-AGE-002-S	0.857 J (1.04)
605146	TA2-136-SEEP-AGE-003-S	0.408 J (1.02)

Note: Values in **bold** represent detected analytes.

<sup>a</sup>EPA November 1986.

<sup>b</sup>Analysis request/chain-of-custody record.

DSS = Drain and Septic Systems.

EPA = U.S. Environmental Protection Agency.

ER = Environmental Restoration.

ΗE = High explosive(s) = Identification. ΙD

= The reported value is greater than or equal to the MDL but is less than the J()

practical quantitation limit, shown in parentheses.

= Microgram(s) per kilogram. μg/kg = Method detection limit. MDL

= Soil sample.

SEEP-AGE = Sample from base of catch box. SWMU = Solid Waste Management Unit.

TA = Technical Area.

VOC = Volatile organic compound.

# Table B-2

# Summary of DSS SWMU 136, Building 907 Septic Systems, HE Catch Box Confirmatory Sampling, VOC Analytical MDLs December 2001

(Off-Site Laboratory)

	EPA Method 8260a
	Detection Limit
Analyta	
Analyte Acetone	(μg/kg)
Benzene	3.45–3.59 0.441–0.459
Bromodichloromethane	
	0.48-0.5
Bromoform	0.48-0.5
Bromomethane	0.49-0.51
2-Butanone	3.67–3.82
Carbon disulfide	2.31–2.41
Carbon tetrachloride	0.48-0.5
Chlorobenzene	0.402-0.418
Chloroethane	0.794–0.827
Chloroform	0.51-0.531
Chloromethane	0.363-0.378
Dibromochloromethane	0.490.51
1,1-Dichloroethane	0.461–0.48
1,2-Dichloroethane	0.422-0.439
1,1-Dichloroethene	0.49-0.51
cis-1,2-Dichloroethene	0.461-0.48
trans-1,2-Dichloroethene	0.52-0.541
1,2-Dichloropropane	0.471-0.49
cis-1,3-Dichloropropene	0.422-0.439
trans-1,3-Dichloropropene	0.245-0.255
Ethylbenzene	0.373-0.388
2-Hexanone	3.7–3.85
Methylene chloride	1.32-1.38
4-methyl-2-Pentanone	3.95-4.11
Styrene	0.382-0.398
1,1,2,2-Tetrachloroethane	0.892-0.929
Tetrachloroethene	0.373-0.388
Toluene	0.333-0.347
1,1,1-Trichloroethane	0.52-0.541
1,1,2-Trichloroethane	0.529-0.551
Trichloroethene	0.441-0.459
Vinyl acetate	1.75–1.82
Vinyl chloride	0.549-0.571
Xylene	0.382-0.398

# <sup>a</sup>EPA November 1986.

DSS = Drain and Septic Systems.

EPA = U.S. Environmental Protection Agency.

HE = High explosive(s).

MDL = Method detection limit.

μg/kg = Microgram(s) per kilogram.

SWMU = Solid Waste Management Unit.

VOC = Volatile organic compound.

Summary of DSS SWMU 136, Building 907 Septic Systems, HE Catch Box Confirmatory Sampling, SVOC Analytical Results (Off-Site Laboratory) December 2001 Table B-3

	_		_	12	~	_	4
		bis(2-Ethylhexyl) phthalate	(F. 7) CIN	ŀ	135 J (333	40.6 J (333	
		Pyrene	ND /8 67 .II	20.00	74.8 J	(I. 78.8) (IN	75 .5.5
		Phenanthrene	10 2 1 /22 2	่า	25.1 J (33.3)	15 4 .1 (33 3)	22.20
ug/kg)		-lexachlorobenzene	2	_	ND (4.67 J)	C EC   /333	0.50 0 1000
$\overline{}$		=)uoranthene	20.02	52.3 0	48.5 J	0001 000	32.3 0 (33.3)
SVOCs (EPA Method 8270a)		əlslarliq İvind-n-iC	١.	44,8 J (333)	23.5 J (333)	1 200	ND (20.7 J)
ÖΛS		уилузеле	.ار	9.56 J (33.3)	57.1	0000	ND (6.33 J)
		ienzo(k)fluoranthene	3	ND (5.5)	139 1 (333)	3	ND (5 J)
		enzo(b)fluoranthene	8	ND (2.33 J)	69	95.10	ND (2.33 J)
		ифгаселе	¥	24.9 J (33.3)	7 70 1 (93 3)	1.16 0 33.3	5.92 J (33.3)
	Sample Attributes	•	EH Sample ID	TA2-136-SEEP.AGE-001-S		1 A2-136-SEEF-AGE-002-5	605146 TA2-136-SEEP-AGE-003-S
		Record	Number	80511B	0	605146	605146

Note: Values in bold represent detected analytes.

<sup>a</sup>EPA November 1986.

Danalysis request/chain-of-custody record.

DSS = Drain and Septic Systems.

EPA = U.S. Environmental Protection Agency.

ER = Environmental Restoration

HE = High explosive(s).

Analytical result was qualified as an estimated value.
 The reported value is greater than or equal to the MDL but is less than the practical quantitation limit, shown in parentheses.

= Method detection limit. Ç₹ 04

= Microgram(s) per kilogram.
 = Not detected above the MDL, shown in parentheses.

ug/kg ND ()

= Sample from base of catch box. = Soil sample. SEEP-AGE

 Solid Waste Management Unit.
 Semivolatile organic compound.
 Technical Area. SWMU

# Table B-4 Summary of DSS SWMU 136, Building 907 Septic Systems, HE Catch Box Confirmatory Sampling, SVOC Analytical MDLs December 2001 (Off-Site Laboratory)

	EDA Mothod B0703
	EPA Method 8270 <sup>a</sup> Detection Limits
Anolito	
Analyte	(μg/kg) 4–1300
Acenaphthene	
Acenaphthylene	3.66-1300
Anthracene	4.66–1300
Benzo(a)anthracene	5.99–1300
Benzo(a)pyrene	5.66–1300
Benzo(b)fluoranthene	8.99-1300
Benzo(g,h,i)perylene	8.99–1300
Benzo(k)fluoranthene	8.99-1300
Benzoic acid	1600-6400
Benzyl alcohol	330–1300
4-Bromophenyl phenyl ether	4.66–1300
Butylbenzyl phthalate	12–1300
4-Chlorobenzenamine	54–1300
bis(2-Chloroethoxy)methane	5.99-1300
bis(2-Chloroethyl)ether	6.66-1300
bis-Chloroisopropyl ether	5.99-1300
4-Chloro-3-methylphenol	19.6–1300
2-Chloronaphthalene	3.66-1300
2-Chlorophenol	5-1300
4-Chlorophenyl phenyl ether	3.33-1300
Chrysene	6.33-1300
o-Cresol	7.66–1300
Dibenz[a,h]anthracene	4.66-1300
Dibenzofuran	2.66-1300
1,2-Dichlorobenzene	4.33-1300
1,3-Dichlorobenzene	3.33-1300
1,4-Dichlorobenzene	5.99-1300
3,3'-Dichlorobenzidine	28–2600
2,4-Dichlorophenol	7.99–1300
Diethylphthalate	6.33-1300
2,4-Dimethylphenol	6.99-1300
Dimethylphthalate	27.3–1300
Di-n-butyl phthalate	14–1300
Dinitro-o-cresol	33.3-6400
2,4-Dinitrophenol	15.7–6400
2,4-Dinitrotoluene	5–1300
2,6-Dinitrotoluene	3–1300
Di-n-octyl phthalate	8.99–1300
Diphenyl amine	15.7–84.9
bis(2-Ethylhexyl) phthalate	19.6–1300
Fluoranthene	5–1300
Fluorene	3-1300
Hexachlorobenzene	4.66–1300
T TO ACCUMENTAGE TO ACCUMENT	7.00-1000

Refer to footnotes at end of table.

# Table B-4 (Concluded) Summary of DSS SWMU 136, Building 907 Septic Systems, HE Catch Box Confirmatory Sampling, SVOC Analytical MDLs December 2001

(Off-Site Laboratory)

	EPA Method 8270 <sup>a</sup> Detection Limits
Analyte	(μg/kg)
Hexachlorobutadiene	6.66–1300
Hexachlorocyclopentadiene	2.33-1300
Hexachloroethane	4.33–1300
Indeno(1,2,3-cd)pyrene	8.99–1300
Isophorone	2.33–1300
2-Methylnaphthalene	4–1300
4-Methylphenol	5.66-1300
Naphthalene	3.33-1300
2-Nitroaniline	56-6400
3-Nitroaniline	37-6400
4-Nitroaniline	52-6400
Nitrobenzene	36.6–1300
2-Nitrophenol	3.66-1300
4-Nitrophenol	79–6400
n-Nitrosodiphenylamine	30–1300
n-Nitrosodipropylamine	6.66-1300
Pentachlorophenol	65–6400
Phenanthrene	4–1300
Phenol	3.66-1300
Pyrene	8.66-1300
1,2,4-Trichlorobenzene	4.66-1300
2,4,5-Trichlorophenol	24.3–6400
2,4,6-Trichlorophenol	5.331300

# <sup>a</sup>EPA November 1986.

DSS = Drain and Septic Systems.

EPA = U.S. Environmental Protection Agency.

HE = High explosive(s).

MDL = Method detection limit.

μg/kg = Microgram(s) per kilogram.

SWMU = Solid Waste Management Unit.

SVOC = Semivolatile organic compound.

# Table B-5

# Summary of DSS SWMU 136, Building 907 Septic Systems, HE Catch Box Confirmatory Sampling, HE Compound Analytical Results December 2001

# (On-Site Laboratory)

	Sample Attributes	HE
Record		(EPA Method 8330)a
Number <sup>b</sup>	ER Sample ID	(μg/kg)
605146	TA2-136-SEEP-AGE-001-S	ND
605146	TA2-136-SEEP-AGE-002-S	ND
605146	TA2-136-SEEP-AGE-003-S	ND

# <sup>a</sup>EPA November 1986.

<sup>b</sup>Analysis request/chain-of-custody record.

DSS

= Drain and Septic Systems.

EPA

= U.S. Environmental Protection Agency.

ER

= Environmental Restoration

HE

= High explosive(s).

ID

= Identification.

μg/kg

= Microgram(s) per kilogram.

ND

Not detected.Soil sample.

S

SEEP-AGE = Sample from base of catch box.

SWMU

= Solid Waste Management Unit.

TA

= Technical Area.

# Table B-6 Summary of DSS SWMU 136, Building 907 Septic Systems, HE Catch Box Confirmatory Sampling, HE Analytical MDLs December 2001 (On-Site Laboratory)

	EPA Method 8330 <sup>a</sup> Detection Limits
Analyte	(μg/kg)
2-Amino-4,6-dinitrotoluene	13.4
4-Amino-2,6-dinitrotoluene	10.1
2,4-Dinitrotoluene	12
2,6-Dinitrotoluene	15.7
HMX	16.8
Nitrobenzene	14
1,3-Dinitrobenzene	13.4
2-Nitrotoluene	15.2
3-Nitrotoluene	11.6
4-Nitrotoluene	11.6
RDX	12.5
Tetryl	15.5
1,3,5-Trinitrobenzene	11.9
2,4,6-Trinitrotoluene	14.1

# <sup>a</sup>EPA November 1986.

DSS = Drain and Septic Systems.

EPA = U.S. Environmental Protection Agency.

HE = High explosive(s).

HMX = Octahydro-1,3,5,7-tetranitro-1,3,5,7-tetrazocine.

MDL = Method detection limit. μg/kg = Microgram(s) per kilogram.

RDX = Hexahydro-1,3,5-trinitro-1,3,5-triazine.

SWMU = Solid Waste Management Unit.

Tetryl = Methyl-2,4,6-trinitrophenylnitramine.

Summary of DSS SWMU 136, Building 907 Septic Systems, HE Catch Box Confirmatory Sampling, RCRA Metals TCLP Analytical Results Table B-7

(On-Site Laboratory) December 2001

	Sample Attributes			RCRA M	RCRA Metals (EPA Methods 6000/7000a) (mg/L	ethods 6000/7	'000 <sup>a</sup> ) (mg/L)		
Record Number <sup>b</sup>	ER Sample ID	Arsenic	Barium	Cadmium	Chromium	Lead	Mercury	Selenium	Silver
605146	605146 TA2-136-SEEP-AGE-001-S ND (0.0333 J)	(L 8880.0) ON	0.372	0.269	ND (0.00691)	ND (0.0172)	0.269 ND (0.00691) ND (0.0172) ND (0.00073) ND (0.0492)	ND (0.0492)	ND (0.00448)
605146	605146 TA2-136-SEEP-AGE-002-S ND (0.0333 J)	ND (0.0333 J)	0.519	0.153	ND (0.00691)	ND (0.0172)	0.153 ND (0.00691) ND (0.0172) ND (0.00073) ND (0.0492)	ND (0.0492)	0.00587 J (0.05)
605146	605146 TA2-136-SEEP-AGE-003-S ND (0.0333 J)	(C 8883 (C) (C)	77.0	0.77 0.0073 J (0.05) ND (0.00691) ND (0.0172) ND (0.00073) ND (0.0492)	ND (0.00691)	ND (0.0172)	ND (0.00073)	ND (0.0492)	ND (0.00448)

Note: Values in bold represent detected analytes.

Note: Data presented is from TCLP analysis, no MDL table is provided.

<sup>a</sup>EPA November 1986.

= Drain and Septic Systems.

U.S. Environmental Protection Agency.

= Environmental Restoration. bAnalysis request/chain-of-custody record.

DSS = Drain and Septic Systems.

EPA = U.S. Environmental Protec

ER = Environmental Restoration

HE = High explosive(s).

Identification.The reported value is greater than or equal to the MDL but is less than the practical quantitation limit, shown in parentheses.

M M

= Method detection limit.

= Milligram(s) per liter.

= Not detected above the MDL shown in parentheses. Resource Conservation and Recovery Act.

mg/L ND () RCRA SEEP-AGE

Sample from base of catch box. = Solid Waste Management Unit. SWMU

= Toxicity Characteristic Leaching Procedure. = Technical Area. TA TCLP

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Summary of DSS SWMU 136, Building 907 Septic Systems, HE Catch Box Confirmatory Sampling, Gamma Spectroscopy Analytical Results December 2001 Table B-8

(On-Site Laboratory)

	O Attributes					Activity (EPA 900.0°)(POV9)	100	Ti i con I	238
	Sample Aminomes		407	Thorns	Thorium-232	Cranur	n-235	Olaniani-230	2003-11
7		Cesin	Cesium-15/				L	#1.70	O CY
חוממש	<u>.</u>	41.10	Free	Besult	Error	orc Result Error	Frror	nesau	
Crackmin	ER Sample ID	Hesaul	2	1000	707	720 0/ 011		ND (0.845)	;
1001		0 462		0.829	0.401	ND (0.207)	:	ואם ומים	
17.1 A.S.	TA9-136-SEEF-AGE-001-0	9	١		0000	0 440	105	(0.779) ND (0.779)	;
2		C 4 2 0		0.838	0.388	211.0	0.13	1	
7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7	TA9-136-0FFP-A(3F-002-0	20.10				1070	0000	ND (0 844)	,
2		9640		0000	0.42/	791.0	0.500	2.2.2.2.1	
F115	EDE145 TA2.136-SEEP-AGE-003-S	0.130	1			0,1	V.V		42
2		7000	ΔN	1.54	Ž	0.18	Y	5:-	
Karolli	Rankaround Activity—North Area Supergroup	0.00	2						
2									

Note: Values in bold exceeded background soil activity.

aEPA November 1986.

bAnalysis request/chain-of-custody record.

cTwo standard deviations around the mean detected activity.

<sup>d</sup>Dinwiddie September 1997.

= U.S. Environmental Protection Agency. = Drain and Septic Systems.

= Environmental Restoration. = High explosive(s).

= Identification.

= Minimum detectable activity. Not applicable.

MDA

= Not detected but the MDA shown in parentheses, exceeds background activity. ■ Not detected above the MDA, shown in parentheses.

2 2 2

= Picocurie(s) per gram.

pCi/g

= Sample from base of catch box. Soil sample. SEEP-AGE

= Solid Waste Management Unit.

SWMU

= Technical Area.

= Error not calculated for nondetect results.

EPA 缶里

# Table B-9

# Summary of DSS SWMU 136, Building 907 Septic Systems, HE Catch Box Confirmatory Sampling, Tritium Analytical Results November 2001

# (On-Site Laboratory)

Sample Attributes		Activity (EPA 906.0a) (pCi/g)	
Record Number <sup>b</sup>	ER Sample ID	Result	Errorc
605145	TA2-136-SEEP-AGE-001-S	3.48	10.1
605145	TA2-136-SEEP-AGE-002-S	2.46	8.68
605145	TA2-136-SEEP-AGE-003-S	10.4	11.4
Backgrou	nd Activity <sup>d</sup>	0.021e	

Note: Values in bold exceeded background soil activity.

e0.021 pCi/g = 420 pCi/L, assuming a soil density of 1 gram/cubic centimeter and 5 percent soil moisture.

DSS = Drain and Septic Systems.

EPA = U.S. Environmental Protection Agency.

ER = Environmental Restoration.

HE = High explosive(s).
ID = Identification.

pCi/g = Picocurie(s) per gram. pCi/L = Picocurie(s) per liter.

S = Soil sample.

SEEP-AGE = Sample from base of catch box. SWMU = Solid Waste Management Unit.

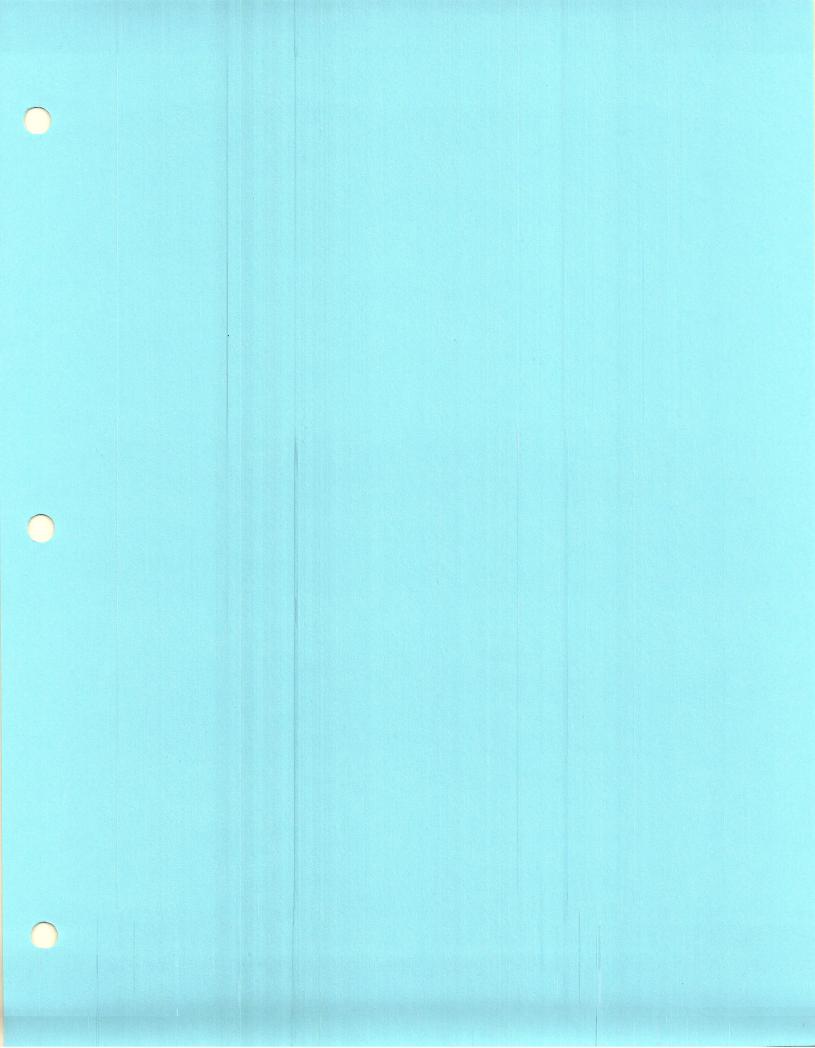
TA = Technical Area.

<sup>&</sup>lt;sup>a</sup>EPA November 1986.

<sup>&</sup>lt;sup>b</sup>Analysis request/chain-of-custody record.

<sup>&</sup>lt;sup>c</sup>Two standard deviations around the mean detected activity.

dTharp February 1999.



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# **DSS SWMU 136: RISK ASSESSMENT REPORT**

# I. Site Description and History

Drain and Septic Systems (DSS) Solid Waste Management Unit (SWMU) 136, the Building 907 Septic Systems, at Sandia National Laboratories/New Mexico (SNL/NM), is located in Technical Area (TA)-II on federally owned land controlled by Kirtland Air Force Base (KAFB) and permitted to the U.S. Department of Energy (DOE). Building 907 had a septic system and a high-explosive (HE) compound waste-water drain system. The septic system, located on the west side of Building 907, consisted of a 450-gallon septic tank, two 1,000-gallon septic tanks located downstream of the first, a 5-foot-diameter by 13-foot-deep seepage pit, and approximately 400 feet of leachfield drain piping. The HE waste-water system drained Building 907 assembly bays and passed through a catch box to collect HE particulates by mechanical filtration of the waste water. The catch box is approximately 200 feet south of the building. The HE waste-water drain system extended an additional 675 feet to the southwest, passing under Building 906, and discharged into a 200-foot-long, open ditch west of Building 919. HE residues would not have been flushed down the floor drains after the mid-1950s when weapons assembly operations were discontinued. Available information indicates that Building 907 was constructed in 1948 (SNL/NM March 2003), and it is assumed that the septic and HE waste-water drain system were also constructed at that time. In 1992, the septic system discharges were routed to the City of Albuquerque sanitary sewer system. The old septic system line was disconnected and capped, and the system was abandoned in place concurrent with this change. The catch box reportedly partially collapsed in 1991 or 1992. The building was demolished in October 2002.

Environmental concern about DSS SWMU 136 is based upon the potential for the release of constituents of concern (COCs) in effluent discharged to the environment via the septic system and HE waste-water drain line at this site. Because operational records were not available, the investigation was planned to be consistent with other DSS site investigations and to sample for the COCs most commonly found at similar facilities.

The ground surface in the vicinity of the site is essentially flat or slopes slightly to the west. The closest major drainage is Tijeras Arroyo, located approximately 3,000 feet east of the site. No springs or perennial surface-water bodies are located within 2 miles of the site. Average annual rainfall in the SNL/NM and KAFB area, as measured at Albuquerque International Sunport, is 8.1 inches (NOAA 1990). Surface-water runoff in the vicinity of the site is minor because the surface slope is flat or inclines to the west. Infiltration of precipitation is almost nonexistent as virtually all of the moisture subsequently undergoes evapotranspiration. The estimates of evapotranspiration for the KAFB area range from 95 to 99 percent of the annual rainfall (SNL/NM March 1996). Most of the area immediately surrounding DSS SWMU 136 is unpaved with some native vegetation, and no storm sewers are used to direct surface water away from the site.

DSS SWMU 136 lies at an average elevation of approximately 5,409 feet above mean sea level. The groundwater beneath the site occurs in unconfined conditions in essentially unconsolidated silts, sands, and gravels. Two water-bearing zones, a shallow groundwater system and the regional aquifer, underlie DSS SWMU 136. The depth to the shallow groundwater system is approximately 300 feet below ground surface (bgs). The shallow

groundwater system is not used for water supply purposes. The depth to the regional aquifer is approximately 545 feet bgs (SNL/NM May 2003). Both the City of Albuquerque and KAFB utilize the regional aquifer as a water supply. Groundwater flow in the shallow groundwater system is to the southeast, while regional groundwater flow is predominantly to the north-northwest in this portion of KAFB. The nearest downgradient water-supply wells are southwest and northwest of the site and include KAFB-1 and KAFB-4, which are approximately 1.2 and 1.1 miles away, respectively.

# II. Data Quality Objectives

Between 1992 and 1994, borehole drilling, monitoring well installation, and sampling in trenches were performed in accordance with the DOE-approved "Interim RCRA [Resource Conservation and Recovery Act] Facility Investigation [RFI] Workplan for Technical Area 2 " (SNL/NM 1991). Beginning in late 1994, borehole drilling and sampling were performed in accordance with the Quality Assurance Project Plan (QAPjP) for the RFI for TA-II (SNL/NM August 1994).

The sampling events completed in 2000 and 2001 were conducted in accordance with the Data Quality Objectives (DQOs) presented in the "Sampling and Analysis Plan [SAP] for Characterizing and Assessing Potential Releases to the Environment From Septic and Other Miscellaneous Drain Systems at Sandia National Laboratories/New Mexico" (SNL/NM October 1999). Negotiations held on November 17, 1999, with the New Mexico Environment Department (NMED) Hazardous Waste Bureau (HWB) defined specific procedures for soil sampling at the seven DSS SWMUs located in TA-II and transferred a requirement for groundwater reporting for these SWMUs to the ongoing Tijeras Arroyo groundwater investigation. The DQOs outlined the quality assurance (QA)/quality control (QC) requirements necessary for producing defensible analytical data suitable for risk assessment purposes. The sampling conducted at this site was designed to:

- Determine whether hazardous waste or hazardous constituents were released at the site.
- Characterize the nature and extent of any releases.
- Provide analytical data of sufficient quality to support risk assessments.

Table 1 summarizes the rationale for determining the sampling locations at this site. The source of potential COCs at DSS SWMU 136 was effluent discharged to the environment from the septic system and along the HE waste-water drain line at this site.

The soil samples were collected at five areas across DSS SWMU 136. The septic and HE waste-water drain line trench samples were collected in 1993 by filling sample containers with soil collected with a backhoe at three depths for each sample location. The deepest sample was collected at a depth equal to the base of the drain line. Boreholes adjacent to the septic tanks, septic system, and HE waste-water drain lines were drilled in 1994 and 1995 using a hollow-stem auger. Samples were collected using a hollow-stem auger and a 2-foot-long, split-spoon-type drive sampler. Samples were collected next to septic tanks at depths equal to, and below, the units using a Geoprobe™ sampling system inside hollow-stem augers.

Table 1
Summary of Sampling Performed to Meet DQOs

DSS SWMU 136 Sampling Areas	Potential COC Source	Number of Sampling Locations	Sample Density (samples/acre)	Sampling Location Rationale
Soil adjacent to, and beneath, the septic tank, septic drain lines, and HE waste-water drain lines	Effluent discharged to the environment from the drain lines and septic tank	7	NA	Evaluate potential COC releases to the environment from effluent discharged from the septic tank and drain line systems.
Soil beneath the septic system seepage pit	Effluent discharged to the environment from the seepage pit	1	NA	Evaluate potential COC releases to the environment from effluent discharged from the seepage pit.
Soil beneath the septic system drainfield	Effluent discharged to the environment from the drainfield	3	NA	Evaluate potential COC releases to the environment from effluent discharged from the drainfield.
Soil beneath the HE drain line outfall ditch	Effluent discharged to the environment from the outfall ditch	3	NA	Evaluate potential COC releases to the environment from effluent discharged from the outfall ditch.
Material in the HE catch box	Effluent discharged to the environment from the HE catch box	3	NA	Evaluate potential COC releases to the environment from effluent discharged from the HE catch box.

COC = Constituent of concern.

DQO = Data Quality Objective.

DSS = Drain and Septic Systems.

HE = High explosive(s). NA = Not applicable.

SWMU = Solid Waste Management Unit.

Samples beneath the seepage pit and in the drainfields were collected in 2000 with a Geoprobe™ from two 3-foot-long sampling intervals at each boring location. Drainfield sampling intervals started at 10 and 15 feet bgs in the one drainfield lateral boring, and 8 and 13 feet bgs in the other two drainfield lateral borings. Sample intervals started at 13 and 18 feet bgs in the single boring through, and beneath, the seepage pit. Samples in the HE waste-water outfall ditch were collected near the surface (0.75 feet bgs) and at 2 and 7 feet bgs at the point where the HE drain line discharged to the outfall ditch, as well as at 4 and 9 feet bgs at two locations along the outfall ditch. Sample containers were filled with soil collected with a backhoe. The HE catch box was sampled at three locations along the bottom of the unit, at approximately 8 feet bgs. The soil samples in 2000 and 2001 were collected in accordance with the procedures developed for, and described in, the Operable Unit (OU) 1295 SAP (SNL/NM October 1999) and subsequent "Field Implementation Plan [FIP], Characterization of Non-Environmental Restoration Drain and Septic Systems" (SNL/NM November 2001) approved by the NMED. The 1994 and 1995 sampling activities were conducted using similar procedures. Table 2 summarizes the types of confirmatory and QA/QC samples collected at the site and the laboratories that performed the analyses.

The DSS SWMU 136 soil samples were analyzed for volatile organic compounds (VOCs), semivolatile organic compounds (SVOCs), HE compounds, polychlorinated biphenyls (PCBs), metals, hexavalent chromium, cyanide, radionuclides by gamma spectroscopy, and H-3. The samples were analyzed by off-site laboratories (Enseco, Inc. [ENS], General Engineering Laboratories, Inc. [GEL], Severn Trent Laboratories [STL], and Thermo Analytical Inc./Eberline Laboratories [TMA]) and at the on-site SNL/NM Environmental Restoration (ER) Chemistry Laboratory (ERCL) and Radiation Protection Sample Diagnostics (RPSD) Laboratory. Table 3 summarizes the analytical methods and data quality requirements based upon the subsequently developed OU 1295 SAP and FIP.

The QA/QC samples were collected during the sampling effort according to the ER Project QAPjP. The QA/QC samples consisted of four trip blanks and four equipment blanks (EBs) for VOCs, four EBs for SVOCs, HE compounds, and metals, three EBs for cyanide and H-3, two EBs for gamma spectroscopy, and one EB for PCBs. With the exception of hexavalent chromium, at least one field duplicate sample was collected for all analyses. No significant QA/QC problems were identified in the QA/QC samples.

All of the DSS SWMU 136 soil sample results were verified/validated by SNL/NM. The off-site laboratory results from ENS, GEL, STL, and TMA were reviewed according to either SNL/NM ER Project "Verification and Validation of Chemical and Radiochemical Data," Technical Operating Procedure (TOP) 94-03 (SNL/NM July 1994) or earlier ER Project Administrative Operating Procedures. The gamma spectroscopy data from the RPSD Laboratory were reviewed according to "Laboratory Data Review Guidelines," Procedure No. RPSD-02-11, Issue No. 2 (SNL/NM July 1996) or an earlier procedure. The reviews confirmed that the analytical data are defensible and therefore acceptable for use in this response to the Request for Supplemental Information (RSI). Therefore, the DQOs have been fulfilled.

Number of Confirmatory Soil and QA/QC Samples Collected at DSS SWMU 136 (Includes samples collected from the HE Catch Box) Table 2

								Gamma	
						Hexavalent		Spectroscopy	
Sample Type	VOCs	SVOCs	PCBs	里	Metals <sup>a</sup>	Chromium	Cyanide	Radionuclides	H-3
Confirmatory	37	35	13	39	48	2	26	30	29
Dinlicates	0	2	-	2	2	0	1	2	1
FRe and TReb	1 00	4	-	4	4	0	3	2	3
Total Samples	47	41	7.	45	54	2	30	34	33
	ENS.	ENS. GEL.	STL	ENS, ERCL,	ENS, ERCL,	ERCL	ENS, STL	GEL, RPSD, TMA	GEL, RPSD,
>	ERCL, GEL,	STL		GEL, STL	GEL, STL				T MA
	SIL								

<sup>a</sup>Numbers do not include the TCLP metals analyses for the HE catch box samples.

bTBs for VOCs only.

Drain and Septic Systems. = Equipment blank.

= Enseco, Inc., Arvada Colorado. = Environmental Restoration Chemistry Laboratory.

General Engineering Laboratories, Inc.
High explosive(s).
Polychlorinated biphenyl.
Quality assurance.

= Quality control.

= Radiation Protection Sample Diagnostics Laboratory. = Semivolatile organic compound. = Severn Trent Laboratories. EB ENS ERCL GEL HE PCB QA QC QC RPSD STL SVOC

= Toxicity Characteristic Leaching Procedure. = Solid Waste Management Unit. SWMU

= Thermo Analytical Inc./Eberline. = Volatile organic compound.

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Summary of Data Quality Requirements for DSS SWMU 136 (Includes samples collected from the HE Catch Box) Table 3

Analytical Methoda	Data Quality Level	ENS	ERCL	GEL	RPSD	STL	TMA
VOCs EPA Method 8000	Defensible	17	2	5	None	13	None
SVOCs EPA Method 8270	Defensible	17	None	5	None	13	None
PCBs EPA Method 8082	Defensible	None	None	None	None	13	None
HE Compounds EPA Method 8330	Defensible	19	ည	2	None	13	None
Metals EPA Methods 6000/7000/908	Defensible	31	2	2	None	13	None
Hexavalent Chromium EPA Method 7196A	Defensible	None	2	None	None	None	None
Total Cyanide EPA Method 9010A	Defensible	13	None	None	None	13	None
Gamma Spectroscopy Radionuclides EPA Method 901.1	Defensible	None	None	2	17	None	11
H-3 EPA Method 906.0	Defensible	None	None	τ-	က	None	25

Note: The number of samples does not include QA/QC samples, such as duplicates, trip blanks, and equipment blanks, or the TCLP metals analyses for the HE catch box samples.

<sup>a</sup>EPA November 1986.

Drain and Septic Systems.

= Enseco Inc., Arvada, Colorado. DSS ENS EPA ERCL

U.S. Environmental Protection Agency.

= Environmental Restoration Chemistry Laboratory.

General Engineering Laboratories, Inc. GEL

 Polychlorinated biphenyl. = High explosive(s).

Quality assurance. PCB QA

Volatile organic compound.

= Thermo Analytical Inc./Eberline Laboratories. TMA VOC

= Toxicity Characteristic Leaching Procedure.

TCLP

 Semivolatile organic compound. ⇒ Solid Waste Management Unit.

Severn Trent Laboratories.

RPSD STL SVOC SWMU

Quality control.

Radiation Protection Sample Diagnostics Laboratory.

# III. Determination of Nature, Rate, and Extent of Contamination

#### III.1 Introduction

The determination of the nature, migration rate, and extent of contamination at DSS SWMU 136 is based upon an initial conceptual model validated with confirmatory sampling at the site. The initial conceptual model was developed from archival site research, site inspections, soil sampling, and passive soil-vapor sampling. The DQOs contained in the RFI Workplan (SNL/NM 1991) and the SAP (SNL/NM October 1999) as well as negotiations with the NMED-HWB held on November 17, 1999, identified the sample locations, sample density, sample depth, and analytical requirements for the August 2000 sampling. The sample data were subsequently used to develop the final conceptual model for DSS SWMU 136, which is presented in Section 4.0 of the associated RSI response. The quality of the data specifically used to determine the nature, migration rate, and extent of contamination is described in the following sections.

#### III.2 Nature of Contamination

Both the nature of contamination and the potential for the degradation of COCs at DSS SWMU 136 were evaluated using laboratory analyses of the soil samples. The analytical requirements included analyses for VOCs, SVOCs, HE compounds, PCBs, metals, hexavalent chromium, cyanide, radionuclides by gamma spectroscopy, and H-3 activities. The analytes and methods listed in Tables 2 and 3 are appropriate to characterize the COCs and potential degradation products at DSS SWMU 136.

# III.3 Rate of Contaminant Migration

The septic system at DSS SWMU 136 was deactivated in the early 1990s when Building 907 was connected to an extension of the City of Albuquerque sanitary sewer system. The HE waste-water drain system was not used for disposing of HE residues after the mid-1950s. The migration rate of COCs that may have been introduced into the subsurface via the septic system at this site was therefore dependent upon the volume of aqueous effluent discharged to the environment from this system when it was operational. Any migration of COCs from this site after use of the septic system was discontinued has been predominantly dependent upon precipitation. However, it is highly unlikely that sufficient precipitation has fallen on the site to reach the depth at which COCs may have been discharged to the subsurface from this system. Analytical data generated from the soil sampling conducted at the site are adequate to characterize the rate of COC migration at DSS SWMU 136.

#### III.4 Extent of Contamination

Subsurface soil samples were collected from five trenches excavated with a backhoe across drain lines, the HE waste-water outfall ditch, and boreholes drilled at nine locations beneath the effluent release points and areas (drain lines, septic tank, seepage pit, drainfield) at the site to

assess whether releases of effluent from the septic system caused any environmental contamination. Three samples were also collected inside the HE catch box.

The DSS SWMU 136 soil samples were collected at sampling depths at which effluent discharged from drain lines, septic tank, seepage pit, drainfield lateral lines, and the HE wastewater outfall ditch would have entered the subsurface environment at the site. This sampling procedure was required by NMED regulators, and similar sampling has been used at numerous DSS-type sites at SNL/NM. The soil samples are considered to be representative of the soil potentially contaminated with the COCs at this site and are sufficient to determine the vertical extent, if any, of COCs.

# IV. Comparison of COCs to Background Screening Levels

Site history and characterization activities are used to identify potential COCs. The DSS SWMU 136 RSI response describes the identification of COCs and the sampling that was conducted in order to determine the concentration levels of those COCs across the site. Generally, COCs evaluated in this risk assessment include all detected organic compounds and all inorganic and radiological COCs for which samples were analyzed. When the detection limit of an organic compound is too high (i.e., could possibly cause an adverse effect to human health or the environment), the compound is retained. Nondetected organic compounds not included in this assessment were determined to have detection limits low enough to ensure protection of human health and the environment. In order to provide conservatism in this risk assessment, the calculation uses only the maximum concentration value of each COC found for the entire site. The SNL/NM maximum background concentration (Dinwiddie September 1997) was selected to provide the background screen listed in Tables 4 through 7.

By agreement with the NMED, two metals samples analyzed by the on-site laboratory are not included in the risk assessment due to high method detection limits (MDLs) (Pavletich May 2003). The justification being that sufficient data was collected to adequately characterize the site, and the risk assessment would not be negatively impacted by exclusion of these samples.

Nonradiological inorganic constituents that are essential nutrients, such as iron, magnesium, calcium, potassium, and sodium, are not included in this risk assessment (EPA 1989). Both radiological and nonradiological COCs are evaluated. The nonradiological COCs evaluated include inorganic and organic compounds.

Tables 4 and 5 list the nonradiological COCs for the human health and ecological risk assessments at DSS SWMU 136, respectively. Tables 6 and 7 list the radiological COCs for the human health and ecological risk assessments, respectively. All tables show the associated SNL/NM maximum background screening values (Dinwiddie September 1997). Section VI.4 discusses the results presented in Tables 4 and 6; Sections VII.2 and VII.3 discuss Tables 5 and 7.

# V. Fate and Transport

The primary releases of COCs at DSS SWMU 136 were to the subsurface soil resulting from the discharge of waste water from the Building 907 Septic Systems. Wind, water, and biota are natural mechanisms of COC transport from the primary release point; however, because the

Comparison to the Associated SNL/NM Background Screening Value, BCF, and Log Kow Nonradiological COCs for Human Health Risk Assessment at DSS SWMU 136 with Table 4

Ö	Maximum Concentration (All Samples)	SNL/NM Background Concentration (mg/kg) <sup>a</sup>	Is Maximum COC Concentration Less Than or Equal to the Applicable SNL/NM Background Screening Value?	BCF (maximum aquatic)	Log K <sub>ow</sub> (for organic COCs)	Bioaccumulator? <sup>b</sup> (BCF>40, Log K <sub>ow</sub> >4)
Inorganic						
Antimony	ь9	3.9	No	16,000€	ì	Yes
Arsenic	9	4.4	No	441	,	Yes
Barium	944	200	No	1709	ļ	Yes
Beryllium	1.2	0.80	No	19f	ļ	ON
Cadmium	7.88	6.0	No	64f		Yes
Chromium, total	20.2	12.8	No	16'		No
Cobalt	10.2	7.1	No	10,000 <sup>h</sup>	-	Yes
Copper	17.2	17	No	6f	,	oN N
Cyanide	0.25 <sup>d</sup>	NC	Unknown	NC	1	Unknown
Lead	58.4	11.2	No	49 <sup>f</sup>	ţ	Yes
Manganese	548	831	Yes	100,000 <sup>h</sup>	1	Yes
Mercury	1.13	<0.1	No	$5,500^{f}$	•	Yes
Nickel	18.4	25.4	Yes	47'	,	Yes
Selenium	19	₹	No	800e	,	Yes
Silver	6.09	₹	No	0.5f		No
Thallium	19	<1.1	No	119'	,	Yes
Uranium	2.2	2.3	Yes	209	,	ON.
Vanadium	33.2	33	No	3,0009	1	Yes
Zinc	58.8	92	Yes	471	,	Yes

Refer to footnotes at end of table.

Comparison to the Associated SNL/NM Background Screening Value, BCF, and Log Kow Nonradiological COCs for Human Health Risk Assessment at DSS SWMU 136 with Table 4 (Continued)

***************************************						
<b>303</b>	Maximum Concentration (All Samples) (mg/kg)	SNL/NM Background Concentration (mg/kg) <sup>a</sup>	Is Maximum COC Concentration Less Than or Equal to the Applicable SNL/NM Background Screening Value?	BCF (maximum aquatic)	Log K <sub>ow</sub> (for organic COCs)	Bioaccumulator? <sup>b</sup> (BCF>40, Log K <sub>ow</sub> >4)
Organic						
Acetone	0.047	NA	NA	i69.0	-0.24	No
Anthracene	0.0249 J	AN	NA	917k	4.45 <sup>k</sup>	Yes
Benzo(a)anthracene	0.048 J	ΝΑ	NA	10,000 <sup>k</sup>	5.61 <sup>k</sup>	Yes
Benzo(a)pyrene	0.041 J	AN	AN	3,000	6.04	Yes
Benzo(b)fluoranthene	0.1 J	NA	NA	_	6.124 <sup>k</sup>	Yes
Benzo(g,h,i)perylene	0.044 J	NA	NA	58,884 <sup>k</sup>	6.58 <sup>k</sup>	Yes
Benzo(k)fluoranthene	0.062 J	NA	NA	93,325 <sup>k</sup>	6.84 <sup>k</sup>	Yes
2-Butanone	0.044 J	NA	NA	1)	0.29i	No
Chrysene	0.071 J	NA	NA	18,000k	5.91k	Yes
Di-n-butyl phthalate	0.086 J	NA	NA	6,761	4.61 <sup>k</sup>	Yes
Di-n-octyl phthalate	0.036 J	NA	AN	9,334 <sup>k</sup>	5.22 <sup>k</sup>	Yes
Ethylbenzene	0.0012 J	NA	NA	15.5	3.15	No
bis(2-Ethylhexyl) phthalate	0.35	NA	NA	851	7.6 <sup>k</sup>	Yes
Fluoranthene	0.052	NA	NA	12,302 <sup>k</sup>	4.90k	Yes
Hexachlorobenzene	0.0066 J	AN	NA	31,622	5.31	Yes
Indeno(1,2,3-cd)pyrene	0.037 J	NA	AN	59,407k	6.58 <sup>k</sup>	Yes
Methylene chloride	0.008	NA	AN	5j	1.25	No
Phenanthrene	0.025 J	NA	NA	23,800f	4.63	Yes
Pyrene	0.075 J	NA	NA	36,300f	5.32 <sup>k</sup>	Yes
Tetrachloroethene	0.0026 J	NA	NA	49j	2.67 <sup>k</sup>	Yes
Toluene	0.0046 J	AN	AN	10.7	2.69f	No

Refer to footnotes at end of table.

Comparison to the Associated SNL/NM Background Screening Value, BCF, and Log K<sub>ow</sub> Nonradiological COCs for Human Health Risk Assessment at DSS SWMU 136 with Table 4 (Concluded)

_						
<b>303</b>	Maximum Concentration (All Samples) (mg/kg)	SNL/NM Background Concentration (mg/kg)ª	Is Maximum COC Concentration Less Than or Equal to the Applicable SNL/NM Background	BCF (maximum	Log K <sub>ow</sub> (for organic	Bioaccumulator? <sup>b</sup> (BCF>40, Log K >4)
0 1 0 Tail it at 1				adagie)	(\$200	· Mo
z,4,o-1 rinitrotoluene	0.076 J	¥	٧Z	453m	1.6k	λογ
Vylene	- 0700				•	2
Aylerie	0.0018 J	AN	¥×	23.4)	٠- بر	No
					2	

Note: **Bold** indicates the COCs that exceed the background screening values and/or are bioaccumulators.

<sup>a</sup>Dinwiddie September 1997, North Area Supergroup.

bNMED March 1998.

cln agreement with NMED, two metals samples evaluated by the on-site laboratory were not included in the risk assessment due to high MDLs.

dConcentration is one-half the maximum MDL.

Callahan et al. 1979.

Yanicak March 1997.

Nanderploeg et al. 1975. 9Neumann 1976.

**USGS 1994** 

Howard 1990.

<sup>k</sup>Micromedex 1998.

Howard 1989.

 Bioconcentration factor. Talmage and Opresko 1995. BCF

= Drain and Septic Systems. = Constituent of concern. 2000 DSS

Estimated concentration.

 Octanol-water partition coefficient. = Logarithm (base 10).

Milligram(s) per kilogram. = Method detection limit.

= Not applicable.

= New Mexico Environment Department. = Not calculated. NMED

 Sandia National Laboratories/New Mexico. Solid Waste Management Unit. SNL/NM SWMU

= U.S. Geological Survey. USGS

= Information not available.

Comparison to the Associated SNL/NM Background Screening Value, BCF, and Log Kow Nonradiological COCs for Ecological Risk Assessment at DSS SWMU 136 with Table 5

3.9 Yes 16,000 <sup>d</sup> 4.4 Yes 16,000 <sup>d</sup> 6.80 Yes 170 <sup>f</sup> 0.80 Yes 170 <sup>f</sup> 0.80 Yes 170 <sup>f</sup> 0.80 Yes 170 <sup>f</sup> 12.8 No 64 <sup>e</sup> 7.1 Yes 6 <sup>e</sup> 7.1 Yes 6 <sup>e</sup> 7.1 Yes 6 <sup>e</sup> 7.1 Yes 6 <sup>e</sup> 7.1 No 49 <sup>e</sup> 831 <sup>h</sup> Yes 100,000 <sup>g</sup> 47 <sup>e</sup> 41 Unknown 800 <sup>d</sup> 41 Unknown 119 <sup>e</sup> 2.3 Yes 3,000 <sup>f</sup> 76 Yes 3,000 <sup>f</sup> 76 Yes 3,000 <sup>f</sup> NA NA NA 6,761 <sup>i</sup> NA NA NA 6,761 <sup>i</sup> NA NA NA 15.5i		Maximum Concentration (Samples ≤ 5 ft	SNL/NM Background Concentration	Is Maximum COC Concentration Less Than or Equal to the Applicable SNL/NM Background	BCF (Maximum	Log K <sub>ow</sub> (for Organic	Bioaccumulator? <sup>b</sup> (BCF>40, Log K >4)
3°         3.9         Yes         16,000 <sup>d</sup> 4         4.4         Yes         44e           169         200         Yes         47e           0.45         0.80         Yes         170f           0.45         0.80         Yes         19e           7.88         0.9         No         64e           15.7         12.8         No         16e           0.25°         NC         Unknown         NC           58.4         11.2         No         49e           6.7         25.4         Yes         47e           6.7         25.4         Yes         47e           60.9         <1	-	(Sugar) (CS		i Dana Billion Dana	Adams)	(8000	/ MO_ 621
4         4.4         Yes         44e           169         200         Yes         170f           0.45         0.80         Yes         170f           7.88         0.9         No         64e           7.1         Yes         10,000s           5         7.1         Yes         6e           11.6         17         Yes         6e           0.25c         NC         Unknown         A9e           58.4         11.2         No         49e           58.4         11.2         No         49e           6.7         25.4         Yes         47e           6.7         25.4         Yes         47e           60.9         <1		3c	3.9	Yes	16,000 <sup>d</sup>		Yes
169         200         Yes         170f           0.45         0.80         Yes         19e           7.88         0.9         No         64e           15.7         12.8         No         16e           15.7         12.8         No         16e           11.6         17         Yes         6e           0.25c         NC         Unknown         NC           58.4         11.2         No         49e           225         831h         Yes         100,0009           6.7         25.4         Yes         47e           60.9         <1		4	4.4	Yes	44e	-	Yes
0.45         0.80         Yes         19e           7.88         0.9         No         64e           15.7         12.8         No         16e           5         7.1         Yes         6e           11.6         17         Yes         6e           0.25c         NC         Unknown         NC           58.4         11.2         No         49e           58.4         11.2         No         49e           58.4         11.2         No         49e           58.4         11.2         No         49e           60.25c         NO         No         5,500e           6.7         25.4         Yes         47e           60.9         <1		169	200	Yes	170	ı	Yes
7.88         0.9         No         64e           15.7         12.8         No         16e           5         7.1         Yes         10,0009           11.6         17         Yes         6e           0.25c         NC         Unknown         NC           58.4         11.2         No         49e           58.4         11.2         No         49e           58.4         11.2         No         49e           60.25c         NO         47e         47e           60.9         <1		0.45	0.80	Yes	19e	1	No
15.7         12.8         No         16e           5         7.1         Yes         10,0009           11.6         17         Yes         6e           0.25c         NC         Unknown         NC           58.4         11.2         No         49e           225         831h         Yes         100,0009           1.13         <0.1		7.88	0.0	No	64e		Yes
5         7.1         Yes         10,0009           11.6         17         Yes         6°           0.25°         NC         Unknown         NC           58.4         11.2         No         49°           225         831°         Yes         100,0009           6.7         25.4         Yes         47°           6.7         25.4         Yes         47°           60.9         <1		15.7	12.8	No	16e		No
11.6       17       Yes       6e         0.25°       NC       Unknown       NC         58.4       11.2       No       49°         58.4       11.2       No       49°         225       831°       Yes       100,0009         1.13       <0.1		5	7.1	Yes	10,0009		Yes
0.25¢         NC         Unknown         NC           58.4         11.2         No         49e           225         831h         Yes         100,0009           1.13         <0.1	_	11.6	17	Yes	99		No
58.4         11.2         No         49e           225         831h         Yes         100,0009           1.13         <0.1		0.25℃	NC	Unknown	SC	1	Unknown
225         831h         Yes         100,0009           1.13         <0.1         No         5,500e           6.7         25.4         Yes         47e           0.465 J         <1         Unknown         800d           60.9         <1         Unknown         119e           0.5c         <1.1         Unknown         119e           2.3         Yes         20f           24         33         Yes         3,000f           37         76         Yes         47e           0.047         NA         NA         6,761i           0.086 J         NA         NA         15.5i		58.4	11.2	No	49e	-	Yes
1.13         <0.1         No         5,500e           6.7         25.4         Yes         47e           0.465 J         <1		225	831h	Yes	100,0009	1	Yes
6.7         25.4         Yes         47e           0.465 J         <1		1.13	<0.1	No	5,500e	an-	Yes
0.465 J         <1         Unknown         800 <sup>d</sup> 60.9         <1		6.7	25.4	Yes	47e	-	Yes
60.9         <1         No         0.5e           0.5°         <1.1		0.465 J	√	Unknown	800d	1	Yes
0.5e         <1.1         Unknown         119e           1.7         2.3         Yes         20f           24         33         Yes         3,000f           37         76         Yes         47e           0.047         NA         NA         0.69i           0.086J         NA         6,761i           0.0012J         NA         NA         15.5i		6.09	<b>^</b>	No	0.5e	1	S.
1.7         2.3         Yes         20f           24         33         Yes         3,000f           37         76         Yes         47e           0.047         NA         NA         0.69f           0.086 J         NA         6,761i           0.0012 J         NA         NA         15.5i		$0.5^{c}$	<1.1	Unknown	119e	1	Yes
24         33         Yes         3,000f           37         76         Yes         47e           0.047         NA         NA         0.69i           0.086 J         NA         6,761i           0.0012 J         NA         15.5i		1.7	2.3	Yes	201	1	9
37         76         Yes         47e           0.047         NA         NA         0.69i           0.086 J         NA         6,761i           0.0012 J         NA         15.5i		24	33	Yes	3,000	1	Yes
0.047         NA         NA         0.69i           0.086 J         NA         6,761i           0.0012 J         NA         15.5i		37	92	Yes	47e	-	Yes
0.047         NA         NA         0.69i           0.086 J         NA         NA         6,761i           0.0012 J         NA         15.5i							
0.086 J NA NA 6,761 i 0.0012 J NA NA 15.5 i		0.047	AN	AN	0.69	-0.24 <sup>i</sup>	No
NA NA 15.5i	e)	0.086 J	NA	AN	6,761	4.61k	Yes
		0.0012 J	NA	NA	15.5i	3.15	No

Refer to footnotes at end of table.

Comparison to the Associated SNL/NM Background Screening Value, BCF, and Log Kow Nonradiological COCs for Ecological Risk Assessment at DSS SWMU 136 with Table 5 (Concluded)

202	Maximum Concentration (Samples ≤ 5 ft bgs) (mg/kg)	SNL/NM Background Concentration (mg/kg) <sup>a</sup>	Is Maximum COC Concentration Less Than or Equal to the Applicable SNL/NM Background Screening Value?	BCF (Maximum Aquatic)	Log K <sub>ow</sub> (for Organic COCs)	Bioaccumulator? <sup>⊳</sup> (BCF>40, Log K <sub>ow</sub> >4)
Methylene chloride	0.008	NA	ΨN	5	1.25	No
Tetrachloroethene	0.0026 J	ΑN	ΑN	49	2.67 <sup>k</sup>	Yes
Toluene	0.0046 J	AN	ΨN	10.7e	2.69e	No
Xylene	0.0018 J	NA	ΨN	23.4	1.5k	N <sub>O</sub>

Note: Bold indicates the COCs that exceed the background screening values and/or are bioaccumulators.

<sup>a</sup>Dinwiddie September 1997, North Area Supergroup.

NMED March 1998,

Parameter was not detected. Concentration is one-half the detection limit.

dCallahan et al. 1979.

Yanicak March 1997.

Neumann 1976.

<sup>3</sup>Vanderploeg, et al. 1975.

**"USGS 1994** 

Howard 1990.

Howard 1989.

\*Micromedex 1998.

= Bioconcentration factor. = Below ground surface. BCF pgs

= Constituent of concern. 200

= Drain and Septic Systems. = Foot (feet). DSS

Estimated concentration.

 Octanol-water partition coefficient. = Logarithm (base 10).

= Solid Waste Management Unit. = U.S. Geological Survey. SWMU USGS

 Sandia National Laboratories/New Mexico. = New Mexico Environment Department.

SNL/NM

NMED

= Milligram(s) per kilogram.

= Not applicable. = Not calculated.

= Information not available.

Radiological COCs for Human Health Risk Assessment at DSS SWMU 136 with Comparison to the Associated SNL/NM Background Screening Value and BCF Table 6

C	Maximum Activity (All Samples)	SNL/NM Background Activity	Is Maximum COC Activity Less Than or Equal to the Applicable SNL/NM Background Screening Value?	BCF (Maximum Aquatic)	ls COC a Bioaccumulator?≎ (BCF >40)
Cs-137	0.22	0.084	S.	3,000d	Yes
H-3	10.4	0.021e	S.	٧N	No
Th-232		1.54	Yes	3,000f	Yes
U-235	ND (0.35)	0.18	S.	€000	Yes
U-238	ND (1.96)	1.3	No	900€	Yes

Note: **Bold** indicates COCs that exceed background screening values and/or are bloaccumulators.

aValue listed is the greater of either the maximum detection or the highest MDA.

Dinwiddie September 1997, North Area Supergroup.

°NMED March 1998.

dWhicker and Schultz 1982.

Tharp February 1999. 420 pCi/L = 0.021 pCi/g, assuming a soil density of 1 gram/cubic centimeter and 5 percent soil moisture.

Baker and Soldat 1992.

BCF = Bioconcentration factor.

COC = Constituent of concern.

DSS = Drain and Septic Systems.

MDA = Minimum detectable activity

Not applicable.

ND() = Not detected, but the MDA (shown in parentheses) exceeds background.

NMED = New Mexico Environment Department

g = Picocurie(s) per gram.

SNL/NM = Sandia National Laboratories/New Mexico.

SWMU = Solid Waste Management Unit.

Comparison to the Associated SNL/NM Background Screening Value and BCF Radiological COCs for Ecological Risk Assessment at DSS SWMU 136 with Table 7

ls COC a Bioaccumulator?≎ (BCF >40)	Yes	No	Yes	
BCF Bic	3,000	AN	3,000	- COO
Is Maximum COC Activity Less Than or Equal to the Applicable SNL/NM Background Screening Value?	No	Yes	Yes	ON A
SNL/NM Background Activity (pCi/g) <sup>b</sup>	0.084	0.021e	1.54	0.18
Maximum Activity (Samples ≤ 5 ft bgs) (pCi/g)³	0.22	0.002	0.94	0.273
000	Cs-137	H-3	Th-232	11-235

Note: Bold indicates COCs that exceed background screening values and/or are bioaccumulators.

aValue listed is the greater of either the maximum detection or the highest MDA.

<sup>b</sup>Dinwiddie September 1997, North Area Supergroup.

°NMED March 1998.

<sup>d</sup>Whicker and Schultz 1982.

Tharp February 1999. 420 pCi/L = 0.021 pCi/g, assuming a soil density of 1 gram/cubic centimeter and 5 percent soil moisture.

Baker and Soldat 1992.

Bioconcentration factor. BCF

= Below ground surface. bgs

= Drain and Septic Systems. = Constituent of concern. DSS

= Foot (feet).

Minimum detectable activity. = Not applicable. MDA

¥

= Not detected, but the MDA (shown in parentheses) exceeds background. CON

= New Mexico Environment Department. NMED

= Picocurie(s) per gram.

 Sandia National Laboratories/New Mexico. SNL/NM

Solid Waste Management Unit. SWMU

discharge was to subsurface soil, none of these are considered to be of potential significance as transport mechanisms at this site. Because the septic and HE drain systems are no longer active, additional infiltration of water is not expected. Infiltration of precipitation is essentially nonexistent at DSS SWMU 136, as virtually all of the moisture either drains away from the site or evaporates. Because the regional aquifer at this site is approximately 545 feet bgs, the potential for COCs to reach groundwater through the unsaturated zone above the water table is extremely low.

The COCs at DSS SWMU 136 include both inorganic and organic constituents. The inorganic COCs include both radiological and nonradiological analytes. With the exception of cyanide, the inorganic COCs are elemental in form and are not considered to be degradable. Transformations of these inorganic constituents could include changes in valence (oxidation/reduction reactions) or incorporation into organic forms (e.g., the conversion of selenite or selenate from soil to seleno-amino acids in plants). Cyanide can be metabolized by soil biota. Radiological COCs will undergo decay to stable isotopes or radioactive daughter elements. However, because of the long half-lives of the radiological COCs, the aridity of the environment at this site, and the lack of potential contact with biota, none of these mechanisms are expected to result in significant losses or transformations of the inorganic COCs.

The organic COCs at DSS SWMU 136 include both VOCs and SVOCs. Organic COCs may be degraded through photolysis, hydrolysis, and biotransformation. Photolysis requires light and therefore takes place in the air, at the ground surface, or in surface water. Hydrolysis includes chemical transformations in water and may occur in the soil solution. Biotransformation (i.e., transformation caused by plants, animals, and microorganisms) may occur; however, biological activity may be limited by the arid environment at this site. Because of the depth of the COCs in the soil, the loss of VOCs through volatilization is expected to be minimal.

Table 8 summarizes the fate and transport processes that can occur at DSS SWMU 136. The COCs at this site include both radiological and nonradiological inorganic analytes as well as organic analytes. Wind, surface water, and biota are considered to be of low significance as potential transport mechanisms at this site. Significant leaching into the subsurface soil is unlikely, and leaching into the regional groundwater at this site is highly unlikely. The potential for transformation of COCs is low, and loss through decay of the radiological COCs is insignificant because of the long half-lives.

Table 8
Summary of Fate and Transport at DSS SWMU 136

Transport and Fate Mechanism	Existence at Site	Significance
Wind	Yes	Low
Surface runoff	Yes	Low
Migration to regional groundwater	No	None
Food chain uptake	Yes	Low
Transformation/degradation	Yes	Low to moderate

DSS = Drain and Septic Systems. SWMU = Solid Waste Management Unit.

#### VI. Human Health Risk Assessment

#### VI.1 Introduction

The human health risk assessment of this site includes a number of steps that culminate in a quantitative evaluation of the potential adverse human health effects caused by constituents located at the site. The steps to be discussed include the following:

<ul> <li>Step 1. Site data are described that provide information on the potential COCs, as well as the relevant physical characteristics and properties of the site.</li> <li>Step 2. Potential pathways are identified by which a representative population might be exposed to the COCs.</li> <li>Step 3. The potential intake of these COCs by the representative population is calculated using a tiered approach. The first component of the tiered approach is a screening procedure that compares the maximum concentration of the COC to an SNL/NM maximum background screening value. COCs that are not eliminated during the first screening procedure are carried forward in the risk assessment process.</li> <li>Step 4. Toxicological parameters are identified and referenced for COCs that were not eliminated during the screening procedure.</li> <li>Step 5. Potential toxicity effects (specified as a hazard index [HI]) and estimated excess cancer risks are calculated for nonradiological COCs and background. For radiological COCs, the incremental total effective dose equivalent (TEDE) and incremental estimated cancer risk are calculated by subtracting applicable background concentrations directly from maximum on-site contaminant values. This background subtraction applies only when a radiological COC occurs as contamination and exists as a natural background radionuclide.</li> <li>Step 6. These values are compared with guidelines established by the U.S. Environmental Protection Agency (EPA), NMED, and the DOE to determine whether further evaluation and potential site cleanup are required. Nonradiological COC risk values also are compared to background risk so that an incremental risk can be calculated.</li> <li>Step 7. Uncertainties of the above steps are addressed.</li> </ul>		
the COCs.  Step 3. The potential intake of these COCs by the representative population is calculated using a tiered approach. The first component of the tiered approach is a screening procedure that compares the maximum concentration of the COC to an SNL/NM maximum background screening value. COCs that are not eliminated during the first screening procedure are carried forward in the risk assessment process.  Step 4. Toxicological parameters are identified and referenced for COCs that were not eliminated during the screening procedure.  Step 5. Potential toxicity effects (specified as a hazard index [Hi]) and estimated excess cancer risks are calculated for nonradiological COCs and background. For radiological COCs, the incremental total effective dose equivalent (TEDE) and incremental estimated cancer risk are calculated by subtracting applicable background concentrations directly from maximum on-site contaminant values. This background subtraction applies only when a radiological COC occurs as contamination and exists as a natural background radionuclide.  Step 6. These values are compared with guidelines established by the U.S. Environmental Protection Agency (EPA), NMED, and the DOE to determine whether further evaluation and potential site cleanup are required. Nonradiological COC risk values also are compared to background risk so that an incremental risk can be calculated.	Step 1.	
tiered approach. The first component of the tiered approach is a screening procedure that compares the maximum concentration of the COC to an SNL/NM maximum background screening value. COCs that are not eliminated during the first screening procedure are carried forward in the risk assessment process.  Step 4. Toxicological parameters are identified and referenced for COCs that were not eliminated during the screening procedure.  Step 5. Potential toxicity effects (specified as a hazard index [HI]) and estimated excess cancer risks are calculated for nonradiological COCs and background. For radiological COCs, the incremental total effective dose equivalent (TEDE) and incremental estimated cancer risk are calculated by subtracting applicable background concentrations directly from maximum on-site contaminant values. This background subtraction applies only when a radiological COC occurs as contamination and exists as a natural background radionuclide.  Step 6. These values are compared with guidelines established by the U.S. Environmental Protection Agency (EPA), NMED, and the DOE to determine whether further evaluation and potential site cleanup are required. Nonradiological COC risk values also are compared to background risk so that an incremental risk can be calculated.	Step 2.	
during the screening procedure.  Step 5. Potential toxicity effects (specified as a hazard index [Hi]) and estimated excess cancer risks are calculated for nonradiological COCs and background. For radiological COCs, the incremental total effective dose equivalent (TEDE) and incremental estimated cancer risk are calculated by subtracting applicable background concentrations directly from maximum on-site contaminant values. This background subtraction applies only when a radiological COC occurs as contamination and exists as a natural background radionuclide.  Step 6. These values are compared with guidelines established by the U.S. Environmental Protection Agency (EPA), NMED, and the DOE to determine whether further evaluation and potential site cleanup are required. Nonradiological COC risk values also are compared to background risk so that an incremental risk can be calculated.	Step 3.	tiered approach. The first component of the tiered approach is a screening procedure that compares the maximum concentration of the COC to an SNL/NM maximum background screening value. COCs that are not eliminated during the first screening procedure are
risks are calculated for nonradiological COCs and background. For radiological COCs, the incremental total effective dose equivalent (TEDE) and incremental estimated cancer risk are calculated by subtracting applicable background concentrations directly from maximum on-site contaminant values. This background subtraction applies only when a radiological COC occurs as contamination and exists as a natural background radionuclide.  Step 6. These values are compared with guidelines established by the U.S. Environmental Protection Agency (EPA), NMED, and the DOE to determine whether further evaluation and potential site cleanup are required. Nonradiological COC risk values also are compared to background risk so that an incremental risk can be calculated.	Step 4.	
Protection Agency (EPA), NMED, and the DOE to determine whether further evaluation and potential site cleanup are required. Nonradiological COC risk values also are compared to background risk so that an incremental risk can be calculated.	Step 5.	risks are calculated for nonradiological COCs and background. For radiological COCs, the incremental total effective dose equivalent (TEDE) and incremental estimated cancer risk are calculated by subtracting applicable background concentrations directly from maximum on-site contaminant values. This background subtraction applies only when a radiological COC occurs as contamination and exists as a natural background
	Step 6.	Protection Agency (EPA), NMED, and the DOE to determine whether further evaluation and potential site cleanup are required. Nonradiological COC risk values also are
	Step 7.	

# VI.2 Step 1. Site Data

Section I of this risk assessment provides the site description and history for DSS SWMU 136. Section II presents a comparison of results to DQOs. Section III discusses the nature, rate, and extent of contamination.

By agreement with the NMED, two metals samples analyzed by the on-site laboratory are not included in the risk assessment due to high MDLs (Pavletich May 2003). The justification being that sufficient data was collected to adequately characterize the site, and the risk assessment would not be negatively impacted by exclusion of these samples.

# VI.3 Step 2. Pathway Identification

DSS SWMU 136 has been designated with a future land-use scenario of industrial (DOE et al. September 1995) (see Appendix 1 for default exposure pathways and parameters). However, the residential land-use scenario is also considered in the pathway analysis. Because of the location and characteristics of the potential contaminants, the primary pathway for human exposure is considered to be soil ingestion for the nonradiological COCs and direct gamma

exposure for the radiological COCs. The inhalation pathway for both nonradiological and radiological COCs is included because the potential exists to inhale dust. Soil ingestion is included for the radiological COCs as well. The dermal pathway is included for the nonradiological COCs because of the potential for the receptor to be exposed to contaminated soil. No water pathways to the groundwater are considered; depth to the regional aquifer at DSS SWMU 136 is approximately 545 feet bgs. No intake routes through plant, meat, or milk ingestion are considered appropriate for either the industrial or residential land-use scenarios. Figure 1 shows the conceptual model flow diagram for DSS SWMU 136.

### **Pathway Identification**

Nonradiological Constituents	Radiological Constituents
Soil ingestion	Soil ingestion
Inhalation (dust)	Inhalation (dust)
Dermal contact	Direct gamma

# VI.4 Step 3. Background Screening Procedure

This section discusses Step 3, the background screening procedure, which compares the maximum COC concentration to the background screening level. The methodology and results are described in the following sections.

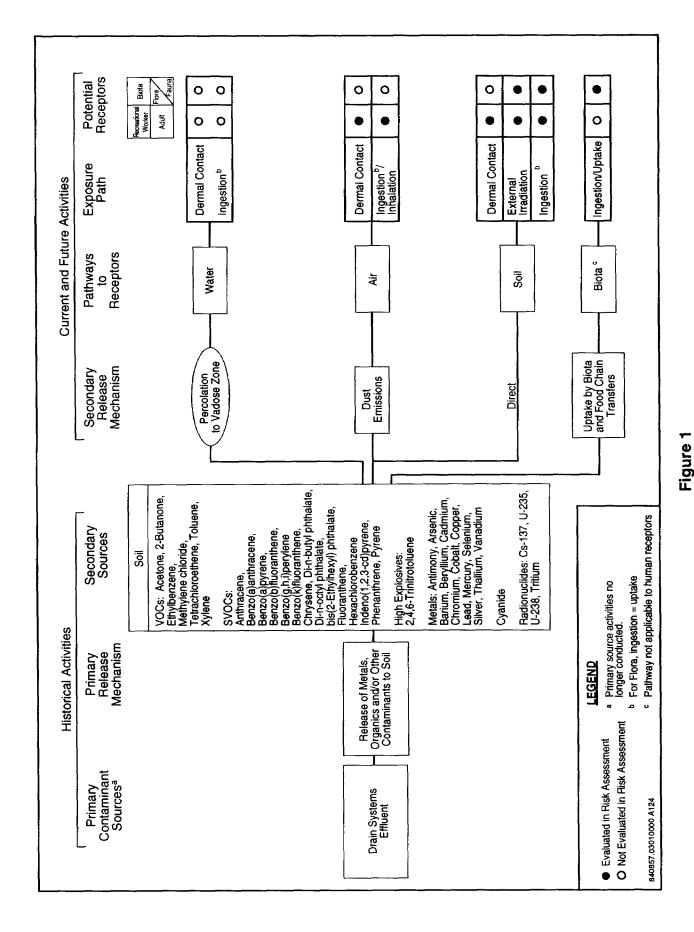
# VI.4.1 Methodology

Maximum concentrations of nonradiological COCs were compared to the approved SNL/NM maximum screening levels for this area. The SNL/NM maximum background concentration was selected to provide the background screen in Table 4 and used to calculate risk attributable to background in Section VI.6.2. Only the COCs that were detected above the corresponding SNL/NM maximum background screening levels or did not have either a quantifiable or calculated background screening level are considered in further risk assessment analyses.

For the radiological COCs that exceed the SNL/NM background screening levels, background values are subtracted from the individual maximum radionuclide concentrations. Those that do not exceed these background levels are not carried any further in the risk assessment. This approach is consistent with DOE Order 5400.5, "Radiation Protection of the Public and the Environment" (DOE 1993). Radiological COCs that do not have background screening values and were detected above the analytical minimum detectable activity (MDA) are carried through the risk assessment at the maximum levels. The resultant radiological COCs remaining after this step are referred to as background-adjusted radiological COCs.

#### VI.4.2 Results

Tables 4 and 6 show the DSS SWMU 136 maximum COC concentrations that were compared to the SNL/NM maximum background values (Dinwiddie September 1997) for the human health risk assessment. For the nonradiological COCs, 14 constituents were measured at concentrations greater than the corresponding background screening values. One constituent



Conceptual Site Model Flow Diagram for DSS SWMU 136, Building 907 Septic Systems

does not have a quantified background screening concentration; therefore, it is unknown whether this COC exceeds background. Twenty-three nonradiological COCs are organic compounds that do not have corresponding background screening values.

The maximum concentration value for lead is 58.4 milligrams (mg)/kilogram (kg). The EPA intentionally does not provide any human health toxicological data on lead; therefore, no risk parameter values could be calculated. However, the NMED guidance for lead screening concentrations for construction and industrial land-use scenarios are 750 and 1,500 mg/kg, respectively (Olson and Moats March 2000). The EPA screening guidance value for a residential land-use scenario is 400 mg/kg (Laws July 1994). The maximum concentration value for lead at this site is lower than all the screening values; therefore, lead is eliminated from further consideration in the human health risk assessment.

For the radiological COCs, four constituents (Cs-137, H-3, U-235, and U-238) had measured activity or MDA values greater than the background screening levels. The greater of either the maximum detection or the highest MDA is conservatively used in the risk assessment.

# VI.5 Step 4. Identification of Toxicological Parameters

Tables 9 (nonradiological) and 10 (radiological) list the COCs retained in the risk assessment and provide the values for the available toxicological information. The toxicological values for the nonradiological COCs presented in Table 9 were obtained from the Integrated Risk Information System (IRIS) (EPA 2003), the Health Effects Assessment Summary Tables (HEAST) (EPA 1997a), the Technical Background Document for Development of Soil Screening Levels (NMED December 2000), Risk Assessment Information System (ORNL 2003), and the EPA Regions 6 and 9 electronic databases (EPA 2002a, EPA 2002b). Dose conversion factors (DCFs) used in determining the excess TEDE values for radiological COCs for the individual pathways are the default values provided in the RESRAD computer code (Yu et al. 1993a) as developed in the following documents:

- DCFs for ingestion and inhalation were taken from "Federal Guidance Report No. 11, Limiting Values of Radionuclide Intake and Air Concentration and Dose Conversion Factors for Inhalation, Submersion, and Ingestion" (EPA 1988).
- DCFs for surface contamination of the site were taken from DOE/EH-0070, "External Dose-Rate Conversion Factors for Calculation of Dose to the Public" (DOE 1988).
- DCFs for volume contamination (exposure to contamination deeper than the immediate surface of the site) were calculated using the methods discussed in "Dose-Rate Conversion Factors for External Exposure to Photon Emitters in Soil" (Kocher 1983) and in ANL/EAIS-8, "Data Collection Handbook to Support Modeling the Impacts of Radioactive Material in Soil" (Yu et al. 1993b).

Table 9
Toxicological Parameter Values for DSS SWMU 136 Nonradiological COCs

	$RfD_{O}$		RfD <sub>inh</sub>		SF <sub>o</sub>	SF <sub>inh</sub>	Cancer	í
200	(mg/kg-d)	Confidence	(mg/kg-d)	Confidencea	(mg/kg-d) <sup>-1</sup>	(mg/kg-d) <sup>-1</sup>	Class	ABS
Inorganic								
Antimony	4E-4c		1	I	ı	ı	-	0.01 <sup>d</sup>
Arsenic	3E-4°	∑	1	1	1.5E+0°	1.5E+1º	A	$0.03^{d}$
Barium	7E-2°	Σ	1.4E-4e	1	1	1	۵	0.01₫
Beryllium	2E-3°	L to M	5.7E-6°	Σ	ŀ	8.4E+0°	B1	0.01⁴
Cadmium	5E-4°	T	5.7E-5f	1	-	6.3E+0°	B1	0.001⁴
Chromium, total	1.5E+0°		1	ŀ		ł	D	0.01 <sup>d</sup>
Cobalt	2E-29	ı	5.7E-69	1	_	9.8E+0f	_	0.01
Copper	3.7E-2f	-	ı		1	ı	۵	0.01 <sup>d</sup>
Cyanide	2E-2°	Σ	1	1		1	۵	0.1d
Mercury	3E-4e	1	8.6E-5°	Σ		-	D	0.01 <sup>d</sup>
Selenium	5E-3°	T	1	1		1	D	0.01
Silver	5E-3°		1	-	-	-	۵	0.01
Thallium	6.6E-59	ŀ	ı	1	1	-	-	0.01
Vanadium	7E-3e		-	1	-	1	1	0.01 <sup>d</sup>
Organic								
Acetone	1E-1º	1	1E-1f	ı	1	-	۵	0.01h
Anthracene	3E-1º		3E-1e	1	-	-	۵	0.134
Benzo(a)anthracene	1	ı	ı	1	7.3E-1f	3.1E-1f	B2	0.13⁴
Benzo(a)pyrene	1	ı	ı	1	7.3E+0°	3.1E+0f	B2	0.13 <sup>d</sup>
Benzo(b)fluoranthene	1	I	ı	-	7.3E-1f	3.1E-1f	B2	0.13⁴
Benzo(g,h,i)perylene	ı	1	1	1	7.3E+0f	3.1E+0f	B2	0.13 <sup>d</sup>
Benzo(k)fluoranthene	1	1	1	-	7.3E-2f	3.1E-2 <sup>f</sup>	B2	0.13 <sup>d</sup>
2-Butanone	6E-1°		2.9E-1°		ı	1	D	0.19
bis(2-Ethylhexyl) phthalate	2E-2f	1	2E-2f	ı	1.4E-2f	1.4E-2f	1	0.01 <sup>h</sup>
Chrysene	ı	1	1	1	7.3E-3f	3.1E-3f	B2	0.134
Di-n-butyl phthalate	1E-1º	_	1E-1 <sup>f</sup>	1	1	1	۵	0.14
Di-n-octyl phthalate	2E-2e	-	2E-2f	_	-	l	1	0.19
Ethylbenzene	1E-1º		2.9E-1c	Γ	3.85E-39	3.85E-39	ı	0.1 <sup>d</sup>
Fluoranthene	4F-2c		4F-2f	-	-	ı	_	0.134

Refer to footnotes at end of table.

# Toxicological Parameter Values for DSS SWMU 136 Nonradiological COCs Table 9 (Concluded)

	RfDo		RfDinh		SFo	SFinh	Cancer	
200	(mg/kg-d)	Confidencea	(mg/kg-d)	Confidencea	(mg/kg-d) <sup>-1</sup>	(mg/kg-d) <sup>-1</sup>	Class	ABS
Hexachlorobenzene	8E-4°	Σ	8E-4 <sup>f</sup>	1	1.6E+0c	1.6E+0°	B2	0.19
Indeno(1.2.3-cd)pyrene	ı	1		1	7.3E-1f	3.1E-1f	B2	0.13 <sup>d</sup>
Methylene chloride	6F-2c	Σ	8.6E-1e	1	7.5E-3°	1.6E-3°	B2	0.19
Phenanthrane	3F-1c		3E-1f	1	-	1	۵	0.13 <sup>d</sup>
Dyrana	3F-2c		3E-2f	1	1	****	_ O	0.1
Tetrachloroethene	1F-20	Σ	1.1E-1f	1	5.2E-2f	1.2E-2f	1	0.1
Tolliene	2F-10	2	1.1E-1º	Σ	1		D	0.14
2.4.6-Trinitrotoluene	5E-4c	Σ	5E-4 <sup>f</sup>	1	3E-2°	3E-2f	ပ	0.1
Xylene	2E+0°	Σ	2E-1f	1	-	1	۵	0.14

<sup>a</sup>Confidence associated with IRIS (EPA 2003) database values. Confidence: L = low, M = medium, H = high. PEPA weight-of-evidence classification system for carcinogenicity (EPA 1989) taken from IRIS (EPA 2003):

A = Human carcinogen.
 B1 = Probable human carcinogen. Limited human data are available.
 B2 = Probable human carcinogen. Sufficient evidence in animals and inadequate or no evidence in humans.
 C = Possible human carcinogen.
 D = Not classifiable as to human carcinogenicity.

Toxicological parameter values from IRIS electronic database (EPA 2003).

<sup>d</sup>Toxicological parameter values from NMED (December 2000). Toxicological parameter values from HEAST (EPA 1997a)

Toxicological parameter values from EPA Region 6 electronic database (EPA 2002a)

<sup>a</sup>Toxicological parameter values from EPA Region 9 electronic database (EPA 2002b)

Toxicological parameter values from Risk Assessment Information System (ORNL 2003).

Toxicological parameter values for phenanthrene could not be found. Anthracene was used as a surrogate.

= New Mexico Environment Department.

RfDinh Gastrointestinal absorption coefficient. = Constituent of concern. 000 ABS

Inhalation chronic reference dose.

Oral chronic reference dose

Inhalation slope factor. = Oral slope factor.

SFinh  $RfD_o$ 

Solid Waste Management Unit.

SWMU

= Information not available.

 Drain and Septic Systems. DSS

= Health Effects Assessment Summary Tables. U.S. Environmental Protection Agency. EPA

= Integrated Risk Information System. HEAST SES

= Milligram(s) per kilogram-day. mg/kg-d

Per milligram per kílogram-day. (mg/kg-d)<sup>-1</sup>

Table 10
Radiological Toxicological Parameter Values for
DSS SWMU 136 COCs Obtained from RESRAD Risk Coefficients <sup>a</sup>

сос	SF <sub>o</sub> (1/pCi)	SF <sub>inh</sub> (1/pCi)	SF <sub>ev</sub> (g/pCi-yr)	Cancer Class <sup>b</sup>
Cs-137	3.20E-11	1.90E-11	2.10E-06	Α
H-3	7.20E-14	9.60E-14	0	A
U-235	4.70E-11	1.30E-08	2.70E-07	A
U-238	6.20E-11	1.20E-08	6.60E-08	A

<sup>&</sup>lt;sup>a</sup>Yu et al. 1993a.

<sup>b</sup>EPA weight-of-evidence classification system for carcinogenicity (EPA 1989): A = Human carcinogen for high dose and high dose rate (i.e., greater than 50 rem per year). For low-level environmental exposures, the carcinogenic effect has not been observed and documented.

1/pCi = One per picocurie.

COC = Constituent of concern.

DSS = Drain and Septic Systems.

EPA = U.S. Environmental Protection Agency.

g/pCi-yr = Gram(s) per picocurie-year.

SF<sub>ev</sub> = External volume exposure slope factor.

SF<sub>inh</sub> = Inhalation slope factor.
 SF<sub>o</sub> = Oral (ingestion) slope factor.
 SWMU = Solid Waste Management Unit.

# VI.6 Step 5. Exposure Assessment and Risk Characterization

Section VI.6.1 describes the exposure assessment for this risk assessment. Section VI.6.2 provides the risk characterization, including the HI and excess cancer risk for both the potential nonradiological COCs and associated background for industrial and residential land-use scenarios. The incremental TEDE and incremental estimated cancer risk are provided for the background-adjusted radiological COCs for both industrial and residential land-use scenarios.

#### VI.6.1 Exposure Assessment

Appendix 1 provides the equations and parameter input values used to calculate intake values and subsequent HI and excess cancer risk values for the individual exposure pathways. The appendix shows parameters for both industrial and residential land-use scenarios. The equations for nonradiological COCs are based upon the Risk Assessment Guidance for Superfund (RAGS) (EPA 1989). Parameters are based upon information from the RAGS (EPA 1989), the Technical Background Document for Development of Soil Screening Levels (NMED December 2000), as well as other EPA and NMED guidance documents. The parameters reflect the reasonable maximum exposure (RME) approach advocated by the RAGS (EPA 1989). For radiological COCs, the coded equations provided in RESRAD computer code are used to estimate the incremental TEDE and cancer risk for individual exposure pathways. Further discussion of this process is provided in the "Manual for Implementing Residual Radioactive Material Guidelines Using RESRAD" (Yu et al. 1993a).

Although the designated land-use scenario for this site is industrial, risk and TEDE values for a residential land-use scenario are also presented.

### VI.6.2 Risk Characterization

Table 11 shows an HI of 0.11 for the DSS SWMU 136 nonradiological COCs and an estimated excess cancer risk of 4E-6 for the designated industrial land-use scenario. The numbers presented include exposure from soil ingestion, dermal contact, and dust and volatile inhalation for nonradiological COCs. Table 12 shows that for the DSS SWMU 136 associated background constituents, the HI is 0.04 with an excess cancer risk of 3E-6 for the designated industrial land-use scenario.

For the radiological COCs, contribution from the direct gamma exposure pathway is included. For the industrial land-use scenario, a TEDE was calculated that results in an incremental TEDE of 1.2E-1 millirem (mrem)/year (yr). In accordance with EPA guidance found in Office of Solid Waste and Emergency Response (OSWER) Directive No. 9200.4-18 (EPA 1997b), an incremental TEDE of 15 mrem/yr is used for the probable land-use scenario (industrial in this case); the calculated dose value for DSS SWMU 136 for the industrial land use is well below this guideline. The estimated excess cancer risk is 1.2E-6.

For the nonradiological COCs under the residential land-use scenario, the HI is 1.36 and the estimated excess cancer risk is 2E-5 (Table 11). The numbers in the table include exposure from soil ingestion, dermal contact, and dust and volatile inhalation. Although the EPA (1991) generally recommends that inhalation not be included in a residential land-use scenario, this pathway is included because of the potential for soil in Albuquerque, New Mexico, to be eroded and for dust to be present in predominantly residential areas. Because of the nature of the local soil, other exposure pathways are not considered (see Appendix 1). Table 12 shows that for the DSS SWMU 136 associated background constituents, the HI is 0.47 with an estimated excess cancer risk of 1E-5.

For the radiological COCs, the incremental TEDE for the residential land-use scenario is 3.1E-1 mrem/yr. The guideline being used is an excess TEDE of 75 mrem/yr (SNL/NM February 1998) for a complete loss of institutional controls (residential land use in this case); the calculated dose value for DSS SWMU 136 for the residential land-use scenario is well below this guideline. Consequently, DSS SWMU 136 is eligible for unrestricted radiological release as the residential land-use scenario resulted in an incremental TEDE of less than 75 mrem/yr to the on-site receptor. The estimated excess cancer risk is 3.5E-6. The excess cancer risk from the nonradiological and radiological COCs should be summed to provide risk estimates for persons exposed to both types of carcinogenic contaminants, as noted in OSWER Directive No. 9200.4-18, "Establishment of Cleanup Levels for CERCLA [Comprehensive Environmental Response, Compensation, and Liability Act] Sites with Radioactive Contamination" (EPA 1997b). This summation is tabulated in Section VI.9, "Summary."

Table 11
Risk Assessment Values for DSS SWMU 136 Nonradiological COCs

	Maximum		l Land-Use nario <sup>a</sup>		al Land-Use nario <sup>a</sup>
	Concentration	Hazard	Cancer	Hazard	Cancer
COC	(mg/kg)	Index	Risk	Index	Risk
Inorganic					
Antimony	6 <sup>b</sup>	0.02	-	0.20	_
Arsenic	6	0.02	4E-6	0.28	2E-5
Barium	944	0.02	_	0.18	
Beryllium	1.2	0.00	5E-10	0.01	1E-9
Cadmium	7.88	0.02	3E-9	0.20	5E-9
Chromium, totalc	20.2	0.00	_	0.00	_
Cobalt	10.2	0.00	5E-9	0.01	1E-8
Copper	17.2	0.00		0.01	_
Cyanide	0.25 <sup>b</sup>	0.00	_	0.00	-
Mercury	1.13	0.00	-	0.05	_
Selenium	1 <sup>b</sup>	0.00	_	0.00	_
Silver	60.9	0.01	_	0.16	-
Thallium	1b	0.02	_	0.20	_
Vanadium	33.2	0.00		0.06	_
Organic			L	1	
Acetone	0.047	0.00	_	0.00	
Anthracene	0.0249 J	0.00	_	0.00	
Benzo(a) anthracene	0.048 J	0.00	2E-8	0.00	8E-8
Benzo(a)pyrene	0.041 J	0.00	2E-7	0.00	7E-7
Benzo(b)fluoranthene	0.1 J	0.00	5E-8	0.00	2E-7
Benzo(g,h,i)perylene	0.044 J	0.00	2E-7	0.00	7E-7
Benzo(k)fluoranthene	0.062 J	0.00	3E-9	0.00	1E-8
2-Butanone	0.044 J	0.00	-	0.00	_
Chrysene	0.071 J	0.00	3E-10	0.00	1E-9
Di-n-butyl phthalate	0.086 J	0.00	-	0.00	
Di-n-octyl phthalate	0.036 J	0.00	_	0.00	_
Ethylbenzene	0.0012 J	0.00	1E-9	0.00	2E-9
bis(2-Ethylhexyl) phthalate	0.35	0.00	2E-9	0.00	8E-9
Fluoranthene	0.052	0.00	-	0.00	
Hexachlorobenzene	0.0066 J	0.00	6E-9	0.00	2E-8
Indeno (1,2,3-cd) pyrene	0.037 J	0.00	2E-8	0.00	6E-8
Methylene chloride	0.008	0.00	5E-8	0.00	1E-7
Phenanthrene	0.025 J	0.00	_	0.00	_
Pyrene	0.075 J	0.00	_	0.00	
Tetrachloroethene	0.0026 J	0.00	8E-10	0.00	2E-9
Toluene	0.0046 J	0.00	-	0.00	
2,4,6-Trinitrotoluene	0.076 J	0.00	1E-9	0.00	5E-9
Xylene	0.0018 J	0.00	_	0.00	
Total		0.11	4E-6	1.36	2E-5

<sup>&</sup>lt;sup>a</sup>EPA 1989.

<sup>&</sup>lt;sup>b</sup>Maximum concentration is one-half of the detection limit.

<sup>&</sup>lt;sup>c</sup>Chromium total was considered to be chromium VI, (most conservative).

# Table 11 (Concluded) Risk Assessment Values for DSS SWMU 136 Nonradiological COCs

COC = Constituent of concern.

DSS = Drain and Septic Systems.

EPA = U.S. Environmental Protection Agency.

J = Estimated Concentration.
mg/kg = Milligram(s) per kilogram.
SWMU = Solid Waste Management Unit.
Information not available.

Table 12
Risk Assessment Values for DSS SWMU 136 Nonradiological Background Constituents

	Background		l Land-Use nario <sup>b</sup>		al Land-Use nario <sup>b</sup>
	Concentration <sup>a</sup>	Hazard	Cancer	Hazard	Cancer
COC	(mg/kg)	Index	Risk	Index	Risk
Antimony	3.9	0.01	_	0.13	-
Arsenic	4.4	0.02	3E-6	0.20	1E-5
Barium	200	0.00	_	0.04	_
Beryllium	0.80	0.00	3E-10	0.01	7E-10
Cadmium	0.9	0.00	2E-10	0.02	6E-10
Chromium, totalc	12.8	0.00	_	0.00	_
Cobalt	7.1	0.00	4E-9	0.01	8E-9
Copper	17	0.00	-	0.01	-
Cyanide	NC	_	-	_	_
Mercury	<0.1	_	_	_	_
Selenium	<1	-		-	<del>-</del>
Silver	<1	_	-		_
Thallium	<1.1	_	-	-	
Vanadium	33	0.00		0.06	-
To	otal	0.04	3E-6	0.47	1E-5

<sup>&</sup>lt;sup>a</sup>Dinwiddie September 1997, North Area Supergroup.

Chromium, total is considered to be chromium III.

COC = Constituent of concern.

DSS = Drain and Septic Systems.

EPA = U.S. Environmental Protection Agency.

mg/kg = Milligram(s) per kilogram.

SWMU = Solid Waste Management Unit.

= Information not available.

# VI.7 Step 6. Comparison of Risk Values to Numerical Guidelines

The human health risk assessment analysis evaluates the potential for adverse health effects for both the industrial (the designated land-use scenario for this site) and residential land-use scenarios.

For the nonradiological COCs under the industrial land-use scenario, the HI is 0.11 (less than the numerical guideline of 1 suggested in the RAGS [EPA 1989]). The excess cancer risk is estimated at 4E-6. NMED guidance states that cumulative excess lifetime cancer risk must be less than 1E-5 (Bearzi January 2001); thus the excess cancer risk for this site is below the suggested acceptable risk value. This assessment also determines risks considering background concentrations of the potential nonradiological COCs for both the industrial and residential land-use scenarios. Assuming the industrial land-use scenario, for the nonradiological COCs the HI is 0.04 and the estimated excess cancer risk is 3E-6. The incremental risk is determined by subtracting risk associated with background from potential COC risk. These numbers are not rounded before the difference is determined and, therefore, may appear to be inconsistent with numbers presented in tables and within the text. For conservatism, the background constituents that do not have quantified background screening

<sup>&</sup>lt;sup>b</sup>EPA 1989.

concentrations are assumed to have a hazard quotient (HQ) of 0.00. The incremental HI is 0.07 and the estimated incremental cancer risk is 1.57E-6 for the industrial land-use scenario. These incremental risk calculations indicate insignificant risk to human health from nonradiological COCs considering an industrial land-use scenario.

For the radiological COCs under the industrial land-use scenario, the incremental TEDE is 1.2E-1 mrem/yr, which is significantly lower than EPA's numerical guideline of 15 mrem/yr. The incremental estimated excess cancer risk 1.2E-6.

The calculated HI for the nonradiological COCs under the residential land-use scenario is 1.36, which is above the numerical guidance. The excess cancer risk is estimated to be 2E-5. NMED guidance states that cumulative excess lifetime cancer risk must be less than 1E-5 (Bearzi January 2001); thus the excess cancer risk for this site is also above the suggested acceptable risk value. The HI for associated background for the residential land-use scenario is 0.47; the estimated excess cancer risk is 1E-5. The incremental HI is 0.89 and the estimated incremental cancer risk is 5.95E-6 for the residential land-use scenario. The incremental excess cancer risk calculation is below guidelines from the nonradiological COCs considering a residential land-use scenario.

The incremental TEDE for a residential land-use scenario from the radiological components is 3.1E-1 mrem/yr, which is significantly lower than the numerical guideline of 75 mrem/yr suggested in the SNL/NM "RESRAD Input Parameter Assumptions and Justification" (SNL/NM February 1998). The estimated excess cancer risk is 3.5E-6.

# VI.8 Step 7. Uncertainty Discussion

The determination of the nature, rate, and extent of contamination at DSS SWMU 136 is based upon an initial conceptual model that was validated with sampling conducted at the site. The sampling was implemented in accordance with the RFI Workplan (SNL/NM 1991), the SAP (SNL/NM October 1999), and negotiations with the NMED-HWB. The DQOs contained in these two documents are appropriate for use in risk assessments. The data from soil samples collected at effluent release points are representative of potential COC releases to the site. The analytical requirements and results satisfy the DQOs, and data quality was verified/validated in accordance with SNL/NM procedures. Therefore, there is no uncertainty associated with the data quality used to perform the risk assessment at DSS SWMU 136.

Because of the location, history, and future land use, there is low uncertainty in the land-use scenario and the potentially affected populations that were considered in performing the risk assessment analysis. Based upon the COCs found in near-surface soil and the location and physical characteristics of the site, there is low uncertainty in the exposure pathways relevant to the analysis.

An RME approach is used to calculate the risk assessment values. Specifically, the parameter values in the calculations are conservative and calculated intakes may be overestimated. Maximum measured values of COC concentrations are used to provide conservative results.

Table 9 shows the uncertainties (confidence levels) in nonradiological toxicological parameter values. There is a mixture of estimated values and values from the IRIS (EPA 2003), HEAST (EPA 1997a), and the Technical Background Document for Development of Soil Screening

Levels (NMED December 2000). Where values are not provided, information is not available from the HEAST (EPA 1997a), IRIS (EPA 2003), Technical Background Document for Development of Soil Screening Levels (NMED December 2000), the Risk Assessment Information System (ORNL 2003) or the EPA regions (EPA 2002a, EPA 2002b, EPA 2002c). Because of the conservative nature of the RME approach, uncertainties in toxicological values are not expected to change the conclusion from the risk assessment analysis.

Risk assessment values for the nonradiological COCs are within the acceptable range for human health under the industrial land-use scenario compared to established numerical guidance.

Although both the HI and estimated excess cancer risk are slightly above the NMED guidelines for the residential land-use scenario, maximum concentrations were used in the risk calculation. Because the site has been adequately characterized, average concentrations are more representative of actual site conditions. Using the 95% upper confidence limit (UCL) of the average concentrations (Appendix 2) for the main contributors to excess cancer risk (antimony at 4.0 mg/kg, arsenic at 3.2 mg/kg [below background and eliminated as a COC], barium at 400 mg/kg, cadmium at 1.6 mg/kg, and thallium at 0.70 mg/kg) reduces the total and incremental HI to 0.76 and 0.49, respectively. The total and incremental estimated excess cancer risk are reduced to 2E-6 and 1.83E-6, respectively. Thus, by using realistic concentrations in the risk calculations that more accurately depict actual site conditions, the total and incremental risk calculations are below NMED guidelines.

For the radiological COCs, the conclusion of the risk assessment is that potential effects on human health for both the industrial and residential land-use scenarios are within guidelines and represent only a small fraction of the estimated 360 mrem/yr received by the average U.S. population (NCRP 1987).

The overall uncertainty in all of the steps in the risk assessment process is not considered to be significant with respect to the conclusion reached.

#### VI.9 Summary

DSS SWMU 136 contains identified COCs consisting of some inorganic, organic, and radiological compounds. Because of the location of the site, the designated industrial land-use scenario, and the nature of contamination, potential exposure pathways identified for this site include soil ingestion, dermal contact, and dust and volatile inhalation for chemical COCs and soil ingestion, dust inhalation, and direct gamma exposure for radionuclides. The same exposure pathways are applied to the residential land-use scenario.

Using conservative assumptions and an RME approach to risk assessment, calculations for the nonradiological COCs show that for the industrial land-use scenario the HI (0.11) is significantly lower than the accepted numerical guidance from the EPA. The estimated excess cancer risk is 4E-6. Thus, excess cancer risk is also below the acceptable risk value provided by the NMED for an industrial land-use scenario (Bearzi January 2001). The incremental HI is 0.07 and the incremental excess cancer risk is 1.57E-6 for the industrial land-use scenario. The incremental risk calculations indicate insignificant risk to human health for the industrial land-use scenario.

Using conservative assumptions and an RME approach to risk assessment, calculations for the nonradiological COCs show that for the residential land-use scenario the HI (1.36) is above the accepted numerical guidance from the EPA. The estimated excess cancer risk is 2E-5. Thus, excess cancer risk is also above the acceptable risk value provided by the NMED for a residential land-use scenario (Bearzi January 2001). The incremental HI is 0.89, and the incremental excess cancer risk is 5.95E-6 for the residential land-use scenario. The incremental risk calculations indicate insignificant risk to human health for the residential land-use scenario.

Although both the HI and estimated excess cancer risk are slightly above the NMED guidelines for the residential land-use scenario, maximum concentrations were used in the risk calculation. Because the site has been adequately characterized, average concentrations are more representative of actual site conditions. Using the 95% UCL of the average concentrations for the main contributors to excess cancer risk (antimony at 4.0 mg/kg, arsenic at 3.2 mg/kg [below background and eliminated as a COC], barium at 400 mg/kg, cadmium at 1.6 mg/kg, and thallium at 0.70 mg/kg) (Appendix 2) reduces the total and incremental HI to 0.76 and 0.49, respectively. The total and incremental estimated excess cancer risk are reduced to 2E-6 and 1.83E-6, respectively. Thus, by using realistic concentrations in the risk calculations that more accurately depict actual site conditions, the total and incremental risk calculations are below NMED guidelines.

The incremental TEDE and corresponding estimated cancer risk from the radiological COCs are much lower than EPA guidance values. The estimated TEDE is 1.2E-1 mrem/yr for the industrial land-use scenario, which is much lower than the EPA's numerical guidance of 15 mrem/yr (EPA 1997b). The corresponding incremental estimated cancer risk value is 1.2E-6 for the industrial land-use scenario. Furthermore, the incremental TEDE for the residential land-use scenario that results from a complete loss of institutional control is 3.1E-1 mrem/yr with an associated risk of 3.5E-6. The guideline for this scenario is 75 mrem/yr (SNL/NM February 1998). Therefore, DSS SWMU 136 is eligible for unrestricted radiological release.

The summation of the incremental nonradiological and radiological carcinogenic risks is tabulated in Table 13.

Table 13
Summation of Incremental Radiological and Nonradiological Risks from DSS SWMU 136

Scenario	Nonradiological Risk	Radiological Risk	Total Risk
Industrial	1.57E-6	1.2E-6	2.8E-6
Residential	1.83E-6	3.5E-6	5.3E-6

DSS = Drain and Septic Systems. SWMU = Solid Waste Management Unit.

Uncertainties associated with the calculations are considered small relative to the conservatism of the risk assessment analysis. Therefore, it is concluded that this site poses insignificant risk to human health under both the industrial and residential land-use scenarios.

# VII. Ecological Risk Assessment

#### VII.1 Introduction

This section addresses the ecological risks associated with exposure to constituents of potential ecological concern (COPECs) in the soil at DSS SWMU 136. A component of the NMED Risk-Based Decision Tree (NMED March 1998) is to conduct an ecological assessment that corresponds with that presented in EPA's Ecological RAGS (EPA 1997c). The current methodology is tiered and contains an initial scoping assessment followed by a more detailed risk assessment. Initial components of NMED's decision tree (a discussion of DQOs, data assessment, and evaluations of bioaccumulation as well as fate and transport potential) are addressed in previous sections of this report. Following the completion of the scoping assessment, a determination is made as to whether a more detailed examination of potential ecological risk is necessary. If deemed necessary, the scoping assessment proceeds to a risk assessment whereby a more quantitative estimate of ecological risk is conducted. Although this assessment incorporates conservatisms in the estimation of ecological risks, ecological relevance and professional judgment also are used as recommended by the EPA (1998) to ensure that predicted exposures of selected ecological receptors reflect those reasonably expected to occur at the site.

# VII.2 Scoping Assessment

The scoping assessment focuses primarily on the likelihood of exposure of biota at, or adjacent to, the site to constituents associated with site activities. Included in this section are an evaluation of existing data and a comparison of maximum detected concentrations to background concentrations, examination of bioaccumulation potential, and fate and transport potential. A scoping risk-management decision (Section VII.2.4) summarizes the scoping results and assesses the need for further examination of potential ecological impacts.

#### VII.2.1 Data Assessment

As indicated in Section IV (Tables 5 and 7), inorganic constituents in the soil within the 0- to 5-foot depth interval that exceed background concentrations or that do not have a quantified background concentration include the following:

- Cadmium
- Chromium (total)
- Cyanide
- Lead
- Mercury
- Selenium
- Silver
- Thallium
- Cs-137
- U-235
- U-238

Organic analytes detected in the soil include the following:

- Acetone
- · Di-n-butyl phthalate
- Ethylbenzene
- · Methylene chloride
- Tetrachloroethene
- Toluene
- Xylene

#### VII.2.2 Bioaccumulation

Among the COPECs listed in Section VII.2.1, the following are considered to have bioaccumulation potential in aquatic environments (Section IV, Tables 5 and 7):

- Cadmium
- Lead
- Mercury
- Selenium
- Thallium
- Cs-137
- U-235
- U-238
- Di-n-butyl phthalate
- Tetrachloroethene

However, as directed by the NMED (March 1998), bioaccumulation for inorganic constituents is assessed exclusively based upon maximum reported bioconcentration factors (BCFs) for aquatic species. Because only aquatic BCFs are used to evaluate the bioaccumulation potential for metals, bioaccumulation in terrestrial species is likely to be overpredicted.

# VII.2.3 Fate and Transport Potential

The potential for the COPECs to migrate from the source of contamination to other media or biota is discussed in Section V. As noted in Table 8 (Section V), wind is expected to be of low significance as a transport mechanism for COPECs at this site, and surface-water runoff is potentially of low significance. Migration to groundwater is not anticipated. Food chain uptake is expected to be of low significance. Degradation (decay) and transformation for the inorganic COPECs and radionuclides are expected to be of low significance. Of the organic COPECs, acetone, ethylbenzene, methylene chloride, tetrachloroethene, toluene, and xylenes are VOCs that may be lost to the atmosphere if near the soil surface.

# VII.2.4 Scoping Risk-Management Decision

Based upon information gathered through the scoping assessment, it is concluded that complete ecological pathways may be associated with DSS SWMU 136 and that COPECs also exist at the site. As a consequence, a risk assessment is deemed necessary to predict the potential level of ecological risk associated with the site.

#### VII.3 Risk Assessment

As concluded in Section VII.2.4, both complete ecological pathways and COPECs are associated with DSS SWMU 136. The risk assessment performed for the site involves a quantitative estimate of current ecological risks using exposure models in association with exposure parameters and toxicity information obtained from the literature. The estimation of potential ecological risks is conservative to ensure that ecological risks are not underpredicted.

Components within the risk assessment include the following:

- Problem Formulation—sets the stage for the evaluation of potential exposure and risk.
- Exposure Estimation—provides a quantitative estimate of potential exposure.
- Ecological Effects Evaluation—presents benchmarks used to gauge the toxicity of COPECs to specific receptors.
- Risk Characterization—characterizes the ecological risk associated with exposure of the receptors to environmental media at the site.
- Uncertainty Assessment—discusses uncertainties associated with the estimation of exposure and risk.
- Risk Interpretation—evaluates ecological risk in terms of HQs and ecological significance.
- Risk Assessment Scientific/Management Decision Point—presents the decision to risk managers based upon the results of the ecological risk assessment.

#### VII.3.1 Problem Formulation

Problem formulation is the initial stage of the ecological risk assessment that provides the introduction to the risk evaluation process. Components that are addressed in this section include a discussion of ecological pathways and the ecological setting, identification of COPECs, and selection of ecological receptors. The conceptual model, ecological food webs, and ecological endpoints (other components commonly addressed in a risk assessment) are presented in "Predictive Ecological Risk Assessment Methodology, Environmental Restoration Program, Sandia National Laboratories, New Mexico" (IT July 1998) and are not duplicated here.

# VII.3.1.1 Ecological Pathways and Setting

DSS SWMU 136 is approximately 0.46 acres in size and is located in an area dominated by grassland habitat. The site is unpaved and open to use by wildlife. No threatened or endangered species are known to occur at this site (IT February 1995), and no surface-water bodies, seeps, or springs are associated with the site.

Complete ecological pathways may exist at this site through the exposure of plants and wildlife to COPECs in surface soil. It is assumed that direct uptake of COPECs from soil is the major route of exposure for plants and that exposure of plants to wind-blown soil is minor. Exposure modeling for the wildlife receptors is limited to the food and soil ingestion pathways and external radiation. Because of the lack of surface water at this site, exposure to COPECs through the ingestion of surface water is considered insignificant. Inhalation and dermal contact are also considered insignificant pathways with respect to ingestion (Sample and Suter 1994). Groundwater is not expected to be affected by COCs at this site.

#### VII.3.1.2 COPECs

The Building 907 drain systems were the primary sources of COPECs at DSS SWMU 136. Inorganic and organic COPECs identified for this site are listed in Section VII.2.1. The inorganic COPECs include both radiological and nonradiological analytes. The inorganic analytes were screened against background concentrations and those that exceed the approved SNL/NM background screening levels (Dinwiddie September 1997) for the area or that do not have a quantified background screening value are considered to be COPECs. Nonradiological inorganic constituents that are essential nutrients, such as iron, magnesium, calcium, potassium, and sodium, are not included in this risk assessment as set forth by the EPA (1989). All organic analytes detected within the upper 5 feet of soil are considered to be COPECs for the site. In order to provide conservatism, this ecological risk assessment is based upon the maximum soil concentrations of the COPECs measured in the upper 5 feet of soil at this site. Tables 5 and 7 present the maximum concentrations for the COPECs.

#### VII.3.1.3 Ecological Receptors

A nonspecific perennial plant has been selected as the receptor to represent plant species at the site (IT July 1998). Vascular plants are the principal primary producers at the site and are key to the diversity and productivity of the wildlife community associated with the site. The deer mouse (*Peromyscus maniculatus*) and the burrowing owl (*Speotyto cunicularia*) are used to represent wildlife use. Because of its opportunistic food habits, the deer mouse is used to represent a mammalian herbivore, omnivore, and insectivore. The burrowing owl represents a top predator at this site. The burrowing owl is present at SNL/NM and is designated a species of management concern by the U.S. Fish and Wildlife Service in Region 2, which includes the state of New Mexico (USFWS September 1995).

#### VII.3.2 Exposure Estimation

For the nonradiological COPECs, direct uptake from the soil is considered the only significant route of exposure for terrestrial plants. Exposure modeling for the wildlife receptors is limited to

food and soil ingestion pathways. Inhalation and dermal contact are considered insignificant pathways with respect to ingestion (Sample and Suter 1994). Drinking water is also considered an insignificant pathway because of the lack of surface water at this site. The deer mouse is modeled under three dietary regimes: as an herbivore (100 percent of its diet as plant material), as an omnivore (50 percent of its diet as plants and 50 percent as soil invertebrates), and as an insectivore (100 percent of its diet as soil invertebrates). The burrowing owl is modeled as a strict predator on small mammals (100 percent of its diet as deer mice). Because the exposure in the burrowing owl from a diet consisting of equal parts of herbivorous, omnivorous, and insectivorous mice would be equivalent to the exposure consisting of only omnivorous mice, the diet of the burrowing owl is modeled with intake of omnivorous mice only. Both species are modeled with soil ingestion comprising 2 percent of the total dietary intake. Table 14 presents the species-specific factors used in modeling exposures in the wildlife receptors. Justification for use of the factors presented in this table is described in the ecological risk assessment methodology document (IT July 1998).

Although home range is also included in this table, exposures for this risk assessment are modeled using an area use factor of 1.0, implying that all food items and soil ingested come from the site being investigated. The maximum COPEC concentrations measured in surface soil samples are used to conservatively estimate potential exposures and risks to plants and wildlife at this site.

For the radiological dose-rate calculations, the deer mouse is modeled as an herbivore (100 percent of its diet as plants), and the burrowing owl is modeled as a strict predator on small mammals (100 percent of its diet as deer mice). Both are modeled with soil ingestion comprising 2 percent of the total dietary intake. Receptors are exposed to radiation both internally and externally from Cs-137, U-235, and U-238. Internal and external dose rates to the deer mouse and the burrowing owl are approximated using modified dose-rate models from the DOE (1995) as presented in the ecological risk assessment methodology document for the SNL/NM ER Project (IT July 1998). Radionuclide-dependent data for the dose-rate calculations were obtained from Baker and Soldat (1992). The external dose-rate model examines the totalbody dose rate to a receptor residing in soil exposed to radionuclides. The soil surrounding the receptor is assumed to be an infinite medium uniformly contaminated with gamma-emitting radionuclides. The external dose-rate model is the same for both the deer mouse and the burrowing owl. The internal total-body dose-rate model assumes that a fraction of the radionuclide concentration ingested by a receptor is absorbed by the body and concentrated at the center of a spherical body shape. This provides for a conservative estimate of absorbed dose. This concentrated radiation source at the center of the body of the receptor is assumed to be a "point" source. Radiation emitted from this point source is absorbed by the body tissues to contribute to the absorbed dose. Alpha and beta emitters are assumed to transfer 100 percent of their energy to the receptor as they pass through tissues. Gamma-emitting radionuclides transfer only a fraction of their energy to the tissues because gamma rays interact less with matter than do beta or alpha emitters. The external and internal dose-rate results are summed to calculate a total dose rate from exposure to Cs-137, U-235, and U-238 in soil.

Table 15 provides the transfer factors used in modeling the concentrations of COPECs through the food chain. Table 16 presents maximum concentrations in soil and derived concentrations in tissues of the various food chain elements that are used to model dietary exposures for each of the wildlife receptors.

Exposure Factors for Ecological Receptors at DSS SWMU 136 Table 14

Receptor Species (				Food Intake		
	, con () con ()	Trophic	Body Weight	Rate (kg/dav) <sup>b</sup>	Dietary Composition <sup>c</sup>	Home Kange (acres)
Deer Mouse	Mammalia/	Herbivore	2.39E-2d	3.72E-3	Plants: 100%	2.7E-1e
(Peromyscus	Rodentia				(+ Soil at 2% of intake)	
maniculatus)			1		2001	2 7E_18
Deer Mouse	Mammalia/	Omnivore	2.39E-2d	3.72E-3	Plants: 50%	7.7 -
	Codentia				Invertebrates: 50%	
(Feromyscus					(+ Soil at 2% of intake)	
maniculatus)					,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,	2 7E 1e
Door Mouse	Mammalia/	Insectivore	2.39E-2d	3.72E-3	Invertebrates: 100%	7.7
מפו אוסמס	1		-		(+ Soil at 2% of intake)	
(Peromyscus	Kodentia				(	
maniculatus)						
Burrowing owl	Aves/	Carnivore	1.55E-1f	1.73E-2	Rodents: 100%	3.5=+19
	Ctrigiformos				(+ Soil at 2% of intake)	

<sup>a</sup>Body weights are in kg wet weight.

bFood intake rates are estimated from the allometric equations presented in Nagy (1987). Units are kg dry weight per day. Dietary compositions are generalized for modeling purposes. Default soil intake value of 2% of food intake.

dSilva and Downing 1995.

eEPA 1993, based upon the average home range measured in semiarid shrubland in Idaho.

Dunning 1993.

= Drain and Septic Systems. 9Haug et al. 1993. DSS

= U.S. Environmental Protection Agency. = Kilogram(s). = Solid Waste Management Unit.

EPA kg SWMU

Table 15
Transfer Factors Used in Exposure Models for COPECs at DSS SWMU 136

	Soil-to-Plant	Soil-to-Invertebrate	Food-to-Muscle
COPEC	Transfer Factor	Transfer Factor	Transfer Factor
Inorganic			
Cadmium	5.5E-1 <sup>a</sup>	6.0E-1b	5.5E-4 <sup>a</sup>
Chromium (total)	4.0E-2 <sup>c</sup>	1.3E-1 <sup>d</sup>	3.0E-2 <sup>c</sup>
Cyanide	0.0E+0 <sup>e</sup>	0.0E+0e	0.0E+0e
Lead	9.0E-2°	4.0E-2 <sup>b</sup>	8.0E-4°
Mercury	1.0E+0°	1.0E+0 <sup>f</sup>	2.5E-1 <sup>a</sup>
Selenium	5.0E-1°	1.0E+0 <sup>f</sup>	1.0E-1°
Silver	1.0E+0 <sup>c</sup>	2.5E-1 <sup>b</sup>	5.0E-3°
Thallium	4.0E-3 <sup>a</sup>	1.0E+0f	4.0E-2ª
Organic <sup>9</sup>			
Acetone	5.3E+1	1.3E+1	1.0E-8
Di-n-butyl phthalate	8.4E-2	2.2E+1	1.1E-3
Ethylbenzene	5.9E-1	1.9E+1	3.3E-5
Methylene chloride	7.3E+0	1.5E+1	3.6E-7
Tetrachloroethene	1.1E+0	1.8E+1	1.1E-5
Toluene	1.0E+0	1.8E+1	1.3E-5
Xylene	5.5E-1	1.9E+1	3.7E-5

<sup>&</sup>lt;sup>a</sup>Baes et al. 1984.

<sup>g</sup>Soil-to-plant and food-to-muscle transfer factors from equations developed in Travis and Arms (1988). Soil-to-invertebrate transfer factors from equations developed in Connell and Markwell (1990). All three equations based upon relationship of the transfer factor to the Log K<sub>ow</sub> value of compound.

COPEC = Constituent of potential ecological concern.

DSS = Drain and Septic Systems.

K<sub>ow</sub> = Octanol-water partition coefficient.

Log = Logarithm (base 10).

NCRP = National Council on Radiation Protection and Measurements.

SWMU = Solid Waste Management Unit.

bStafford et al. 1991.

<sup>&</sup>lt;sup>c</sup>NCRP January 1989.

<sup>&</sup>lt;sup>d</sup>Ma 1982.

<sup>&</sup>lt;sup>e</sup>No data found for food chain transfers of cyanide; however, because of its high metabolic activity, cyanide is assumed not to transfer in the food chain.

fDefault value.

Table 16
Media Concentrations<sup>a</sup> for COPECs at DSS SWMU 136

	Soil	Plant	Soil	Deer Mouse
COPEC	(maximum) <sup>a</sup>	Foliage <sup>b</sup>	Invertebrate <sup>b</sup>	Tissues <sup>c</sup>
Inorganic				
Cadmium	7.9E+0	4.3E+0	4.7E+0	8.1E-3
Chromium (total)	1.6E+1	6.3E-1	2.0E+0	1.5E-1
Cyanide	2.5E-1 <sup>d</sup>	0.0E+0	0.0E+0	0.0E+0
Lead	5.8E+1	5.3E+0	2.3E+0	1.2E-2
Mercury	1.1E+0	1.1E+0	1.1E+0	9.0E-1
Selenium	4.7E-1	2.3E-1	4.7E-1	1.1E-1
Silver	6.1E+1	6.1E+1	1.5E+1	6.1E-1
Thallium	5.0E-1 <sup>d</sup>	2.0E-3	5.0E-1	3.3E-2
Organic				
Acetone	4.7E-2	2.5E+0	6.0E-1	5.1E-8
Di-n-butyl phthalate	8.6E-2 <sup>e</sup>	7.2E-3	1.9E+0	3.2E-3
Ethylbenzene	1.2E-3 <sup>e</sup>	7.0E-4	2.3E-2	1.2E-6
Methylene chloride	8.0E-3	5.9E-2	1.2E-1	1.0E-7
Tetrachloroethene	2.6E-3e	2.9E-3	4.7E-2	8.2E-7
Toluene	4.6E-3 <sup>e</sup>	4.6E-3	8.3E-2	1.8E-6
Xylene	1.8E-3 <sup>e</sup>	9.9E-4	3.4E-2	2.1E-6

<sup>&</sup>lt;sup>a</sup>In milligrams per kilogram. All biotic media are based upon dry weight of the media. Soil concentration measurements are assumed to have been based upon dry weight. Values have been rounded to two significant digits after calculation.

COPEC = Constituent of potential ecological concern.

DSS = Drain and Septic Systems.

SWMU = Solid Waste Management Unit.

<sup>&</sup>lt;sup>b</sup>Product of the soil concentration and the corresponding transfer factor.

<sup>&</sup>lt;sup>c</sup>Based upon the deer mouse with an omnivorous diet. Product of the average concentration ingested in food and soil times the food-to-muscle transfer factor times a wet weight-dry weight conversion factor of 3.125 (EPA 1993).

<sup>&</sup>lt;sup>d</sup>Maximum concentration of parameter is one-half of the detection limit.

eEstimated value.

# VII.3.3 Ecological Effects Evaluation

Table 17 shows benchmark toxicity values for the plant and wildlife receptors. For plants, the benchmark soil concentrations are based upon the lowest-observed-adverse-effect level (LOAEL). For wildlife, the toxicity benchmarks are based upon the no-observed-adverse-effect level (NOAEL) for chronic oral exposure in a taxonomically similar test species. Sufficient toxicity information was not available to estimate the LOAELs or NOAELs for some COPECs.

The benchmark used for exposure of terrestrial receptors to radiation was 0.1 rad/day. This value has been recommended by the International Atomic Energy Agency (IAEA 1992) for the protection of terrestrial populations. Because plants and insects are less sensitive to radiation than vertebrates (Whicker and Schultz 1982), the dose of 0.1 rad/day should also protect other groups within the terrestrial habitat of DSS SWMU 136.

#### VII.3.4 Risk Characterization

Maximum concentrations in soil and estimated dietary exposures are compared to plant and wildlife benchmark values, respectively. Table 18 presents the results of these comparisons. HQs are used to quantify the comparison with benchmarks for plant and wildlife exposure.

HQs for plants exceed unity for cadmium, total chromium, lead, mercury, and silver. For the omnivorous and insectivorous deer mice, HQs exceed unity for thallium. When mercury is assumed to be entirely in organic form, the HQs for both the deer mice (all dietary regimes) and the burrowing owl exceed unity; however, the values do not exceed unity when the mercury is assumed to be in inorganic form. Because of a lack of sufficient toxicity information, HQs for plants could not be determined for cyanide and five of the seven organic COPECs. Similarly for the burrowing owl, HQs could not be determined for cyanide, silver, thallium, and all of the organic COPECs except di-n-butyl phthalate. As directed by the NMED, HIs are calculated for each of the receptors (the HI is the sum of chemical-specific HQs for all pathways for a given receptor). All receptors have total HI values greater than unity, with a maximum HI of 55 for plants.

Tables 19 and 20 summarize the internal and external dose-rate model results for Cs-137, U-235, and U-238 for the deer mouse and burrowing owl, respectively. The total radiation dose rate to the deer mouse is predicted to be 2.7E-4 rad/day and that for the burrowing owl is 2.5E-4 rad/day. The dose rates for the deer mouse and the burrowing owl are less than the benchmark of 0.1 rad/day.

#### VII.3.5 Uncertainty Assessment

Many uncertainties are associated with the characterization of ecological risks at DSS SWMU 136. These uncertainties result from assumptions used in calculating risk that could overestimate or underestimate true risk presented at the site. For this risk assessment, assumptions are made that are more likely to overestimate exposures and risk rather than to underestimate them. These conservative assumptions are used to be more protective of the ecological resources potentially affected by the site. Conservatisms incorporated into this risk assessment include the use of maximum analyte concentrations measured in soil samples to

Toxicity Benchmarks for Ecological Receptors at DSS SWMU 136 Table 17

COPEC Inorganic Cadmium		Mam	Mammalian NOAELs	S.		Avian NOAELs	
COPEC Inorganic Cadmium			Test	Deer			Burrowing
Inorganic Cadmium	Plant Benchmark <sup>a,b</sup>	Mammalian Test Species <sup>c,d</sup>	Species NOAEL d.e	Mouse NOAELe.f	Avian Test Species <sup>d</sup>	Test Species	OWI PONIE
Cadmium							
	3	rath	1.0	1.9	mallard	1.45	1.45
Chromium (total)	-	rat	2,737	5,354	black duck	1.0	1.0
Cyanide	1	rati	68.7	126	1	-	1
Lead	99	rat	8.0	15.7	American	3.85	3.85
Morono, (orocoro)	000	1		000	Kestrel		
Mercury (Organic)	0.3	rat	0.03	0.00	mailard	0.0064	0.0064
Mercury (inorganic)	0.3	mouse	13.2	14.0	Japanese quail	0.45	0.45
Selenium	1	rat	0.2	0.391	screech owl	0.44	0.44
Silver	2	rat	17.8i	34.8		ı	1
Thallium	-	ratk	0.0074	0.015	-	1	1
Organic							
Acetone	1	rat	10	19.6			
Di-n-butyl phthalate	200	mouse	550	582	ringed dove	0.11	0.11
Ethylbenzene	_	rat	291	569	1	ı	
Methylene chloride		rat	5.85	11.4	ı	i	
Tetrachloroethene	_	mouse	1.4	1.5	I	1	ı
Toluene	200	mouse	26	27.5	ŀ	1	1
Xylene	I	mouse	2.1	2.22	-	ŀ	1

aln mg/kg soil dry weight.

<sup>b</sup>Efroymson et al. 1997.

<sup>o</sup>Body weights (in kg) for the NOAEL conversion are as follows: lab mouse, 0.030; lab rat, 0.350, (except where noted). <sup>d</sup>Sample et al. (1996), except where noted.

eln mg/kg body weight per day.

fassed upon NOAEL conversion methodology presented in Sample et al. (1996), using a deer mouse body weight of 0.0239 kg and a mammalian scaling factor of 0.25.

The avian scaling factor of 0.0 was used, making the NOAEL <sup>3</sup>Based upon NOAEL conversion methodology presented in Sample et al. (1996). independent of body weight.

# Toxicity Benchmarks for Ecological Receptors at DSS SWMU 136 Table 17 (Concluded)

Body weight: 0.303 kg.

Based upon a rat lowest-observed-adverse-effect level of 89 mg/kg/d (EPA 2003) and an uncertainty factor of 0.2.

= Constituent of potential ecological concern. \*Body weight: 0.365 kg.
'EPA 2003.
COPEC = Constituent of DSS = Drain and Sep EPA = U.S. Environm kg = Kilogram(s).
mg = Milligram(s).
mg/kg/d = Milligram(s) PNOAEL = No observed (s) SWMU = Solid Waste N

= U.S. Environmental Protection Agency. = Drain and Septic Systems.

= Milligram(s) per kilogram per day. = No observed adverse effect level. = Solid Waste Management Unit.

= Insufficient toxicity data.

HQs for Ecological Receptors at DSS SWMU 136 Table 18

		Deer Mouse HQ	Deer Mouse HQ	Deer Mouse HQ	Burrowing Owl
COPEC	Plant HQ <sup>a</sup>	(Herbivorous) <sup>a</sup>	(Omnivorous) <sup>a</sup>	(Insectivorous) <sup>a</sup>	HQª
Inorganic					
Cadmium	2.6E+0	3.7E-1	3.9E-1	4.0E-1	1.3E-2
Chromium (total)	1.6E+1	2.7E-5	4.8E-5	6.8E-5	5.2E-2
Cyanide	1	6.2E-6	6.2E-6	6.2E-6	
Lead	1.2E+0	6.4E-2	4.9E-2	3.5E-2	3.4E-2
Mercury (organic)	3.8E+0	2.9E+0	2.9E+0	2.9E+0	1.6E+1
Mercury (inorganic)	3.8E+0	1.3E-2	1.3E-2	1.3E-2	2.3E-1
Selenium	4.7E-1	9.6E-2	1.4E-1	1.9E-1	3.1E-2
Silver	3.0E+1	2.8E-1	1.8E-1	7.3E-2	1
Thallium	5.0E-1	1.3E-1	2.8E+0	5.4E+0	1
Organic					
Acetone	1	2.0E-2	1.2E-2	4.8E-3	1
Di-n-butyl phthalate	4.3E-4	2.4E-6	2.6E-4	5.1E-4	5.0E-3
Ethylbenzene	I	2.0E-7	3.2E-6	6.2E-6	
Methylene chloride		8.0E-4	1.2E-3	1.7E-3	-
Tetrachloroethene	ļ	3.1E-4	2.6E-3	4.9E-3	1
Toluene	2.3E-5	2.6E-5	2.5E-4	4.7E-4	1
Xylene	1	7.2E-5	1.2E-3	2.4E-3	•
q.H	5.5E+1	3.8E+0	6.4E+0	9.0E+0	1.6E+1

<sup>a</sup>**Bold** values indicate the HQ or HI exceeds unity.

<sup>b</sup>The HI is the sum of individual HQs.

Constituent of potential ecological concern.Drain and Septic Systems.Hazard index. COPEC

DSS HI HQ SWMU

Hazard quotient.Solid Waste Management Unit.Insufficient toxicity data available for risk estimation purposes.

Table 19
Total Dose Rates for Deer Mice
Exposed to Radionuclides at DSS SWMU 136

Radionuclide	Maximum Concentration (pCi/g)	Total Dose (rad/day)
Cs-137	0.095	1.7E-5
U-235	0.273	7.4E-6
U-238	ND (1.5)	2.4E-4
Total Dose		2.7E-4

<sup>a</sup>External dose from H-3 assumed to be insignificant.

DSS = Drain and Septic Systems.

MDA = Minimum detectable activity.

ND ( ) = Not detected, but the MDA (shown in parentheses)

exceeds background.

pCi/g = Picocurie(s) per gram.

SWMU = Solid waste management unit.

Table 20
Total Dose Rates for Burrowing Owls
Exposed to Radionuclides at DSS SWMU 136

Radionuclide	Maximum Concentration (pCi/g)	Total Dose (rad/day)
Cs-137	0.095	1.5E-5
U-235	0.273	5.7E-6
U-238	ND (1.5)	2.3E-4
Total [	Total Dose	

<sup>a</sup>External dose from H-3 assumed to be insignificant.

DSS = Drain and Septic Systems.

MDA = Minimum detectable activity.

ND ( ) = Not detected, but the MDA (shown in parentheses)

exceeds background.

pCi/g = Picocurie(s) per gram.

SWMU = Solid waste management unit.

evaluate risk, the use of wildlife toxicity benchmarks based upon NOAEL values, and the incorporation of strict herbivorous and strict insectivorous diets for predicting the extreme HQ values for the deer mouse. Each of these uncertainties, which are consistent among each of the SWMU-specific ecological risk assessments, is discussed in greater detail in the uncertainty section of the ecological risk assessment methodology document for the SNL/NM ER Project (IT July 1998).

Uncertainties associated with the estimation of risk to ecological receptors following exposure to Cs-137, U-235, and U-238 are primarily related to those inherent in the radionuclide-specific data. Radionuclide-dependent data are measured values that have their associated errors.

The dose-rate models used for these calculations are based upon conservative estimates of receptor shape, radiation absorption by body tissues, and intake parameters. The goal is to provide a realistic but conservative estimate of a receptor's internal and external exposure to radionuclides in soil.

Although the HQs for thallium exceed unity for both the omnivorous and insectivorous deer mice (2.8 and 5.4, respectively), thallium was not detected in any of the soil samples collected from the 0- to 5-foot depth interval (Table 5). For this reason, the exposure point concentration (0.5 mg/kg) for thallium is based upon one-half the detection limit for this analyte. Although the potential for risk to these two receptors from exposure to thallium cannot be ruled out, the conservative estimates of that risk, based upon this exposure point concentration, are low. Therefore, it is considered unlikely that actual concentrations of thallium at this site are sufficient to pose a risk to ecological receptors.

The assumption of an area use factor of 1.0 is a source of uncertainty for the burrowing owl at this site. Because DSS SWMU 136 is approximately 0.46 of an acre in size and the home range of the burrowing owl is 35 acres, an area use factor of approximately 0.013 would be justified for this receptor. This is sufficient to reduce the burrowing owl HQ for organic mercury from 16 to 0.21.

For the deer mice and burrowing owl, HQs for mercury only exceed unity when the values are based upon the highly conservative assumption that all mercury at this site is in organic form. Because the exposure point concentration (1.13 mg/kg) is measured as total mercury, this likely represents a mixture of both organic and inorganic forms of mercury, probably dominated by mercury in inorganic form. When the mercury at the site is assumed to be in inorganic form, the HQs for the deer mice and burrowing owl are 0.013 and 0.23, respectively, indicating no risk to these receptors.

In the estimation of ecological risk, background concentrations are included as a component of maximum on-site concentrations. Conservatisms in the modeling of exposure and risk can result in the prediction of risk to ecological receptors when exposed at background concentrations. In the case of total chromium, the background screening value is 12.8 mg/kg. which accounts for 82 percent of the estimated maximum chromium exposure at this site; therefore, 82 percent of the HQs for total chromium are shown in Table 18. However, because the plant toxicity benchmark for chromium is only 1.0 mg/kg, the HQ for plants from exposure to background concentrations of chromium would be predicted to be 12.8. The plant toxicity benchmark for this metal is based upon chromium VI, which may be more toxic to plants than the more common chromium III. The majority of the total chromium measured at DSS SWMU 136 is expected to be chromium III. For this reason, it is uncertain whether the calculated HQ for total chromium accurately predicts the potential risk to plants. Further, this benchmark is conservatively based upon laboratory tests using soil amendments with a highly available form of chromium (K<sub>2</sub>Cr<sub>2</sub>O<sub>7</sub>) (Efroymson et al. 1997). It is likely that only a small fraction the chromium in the soil at DSS SWMU 136 is in a form that is highly available for plant uptake. For these reasons, the plant toxicity benchmark for total chromium probably overestimates risk to plants to a significant degree.

A further source of uncertainty associated with the prediction of ecological risks at this site is the use of the maximum measured concentrations to evaluate exposure and risk. This results in a conservative exposure scenario that does not necessarily reflect actual site conditions. To evaluate the potential effect on risk predictions by using maximum concentrations as exposure

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point concentrations, 95% UCLs of the mean soil concentrations were calculated (Appendix 2) for cadmium (6.3 mg/kg), total chromium (11.0 mg/kg), lead (28.4 mg/kg), and mercury (0.44 mg/kg). The 95% UCL for chromium is less than its background screening level (12.8 mg/kg), indicating that average chromium exposures at this site are within background levels. The 95% UCL for lead is less than its plant toxicity benchmark (50 mg/kg), indicating that on average, this COPEC does not pose a risk to ecological receptors at this site. Further, exposures to plants at the 95% UCL concentrations for cadmium and mercury reduce the HQs to 2.1 and 1.5, respectively, indicating low average risk to this receptor from these two COPECs.

In the case of silver, the calculated 95% UCL (Appendix 2) exceeds the maximum measured concentration. This is because the data are highly skewed by two outlier detections of this COPECs. Of the nine samples analyzed for silver, all but two were nondetections (the nondetections were less than 1.0 mg/kg, which coincides with the background screening level). Of the two detections, the maximum concentration (60.9 mg/kg) is 15 times higher than the other (3.96 mg/kg). The plant HQ for the second highest measured concentration is 2.0. All plant HQs for the nondetections would be less than 0.5. Therefore, the potential for risk to plants at this site, based upon the maximum value, is likely to overestimate actual risk, which is likely to be low.

Based upon this uncertainty analysis, the potential for ecological risks at DSS SWMU 136 is expected to be low. HQs greater than unity were predicted; however, closer examination of the exposure assumptions reveal an overestimation of risk primarily attributed to conservative toxicity benchmarks; the use of maximum concentrations, maximum bioavailability, and maximum area use to estimate exposure; and the contribution of background risk.

# VII.3.6 Risk Interpretation

Ecological risks associated with DSS SWMU 136 are estimated through a risk assessment that incorporates site-specific information when available. Initial predictions of potential risk to plants from exposure to several metals are based upon maximum measured soil concentrations, highly conservative plant toxicity benchmarks, and assumptions of high bioavailability. Actual risk to this receptor is expected to be low based upon more realistic exposure assumptions. Predictions of potential risk to omnivorous and insectivorous deer mice from exposures to mercury and thallium also can be attributed to conservative exposure assumptions. For the burrowing owl, the initial prediction of risk from exposure to mercury is due to the assumption of 100-percent area use by this receptor. A more realistic assumption of area use for this receptor results in an HQ of only 0.21. The very small size of this site (0.46 acre) also limits the potential for significant risk to ecological receptors at this site, particularly at the population or community levels. Based upon this final analysis, the potential for ecological risks associated with DSS SWMU 136 is expected to be low.

# VII.3.7 Risk Assessment Scientific/Management Decision Point

After potential ecological risks associated with the site have been assessed, a decision is made regarding whether the site should be recommended for no further action (NFA) or whether additional data should be collected to assess actual ecological risk at the site more thoroughly.

With respect to this site, ecological risks are predicted to be low. The scientific/management decision is to recommend this site for NFA.

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# APPENDIX 1 EXPOSURE PATHWAY DISCUSSION FOR CHEMICAL AND RADIONUCLIDE CONTAMINATION

# Introduction

Sandia National Laboratories/New Mexico (SNL/NM) uses a default set of exposure routes and associated default parameter values developed for each future land-use designation being considered for SNL/NM Environmental Restoration (ER) Project sites. This default set of exposure scenarios and parameter values are invoked for risk assessments unless site-specific information suggests other parameter values. Because many SNL/NM solid waste management units (SWMUs) have similar types of contamination and physical settings, SNL/NM believes that the risk assessment analyses at these sites can be similar. A default set of exposure scenarios and parameter values facilitates the risk assessments and subsequent review.

The default exposure routes and parameter values used are those that SNL/NM views as resulting in a Reasonable Maximum Exposure (RME) value. Subject to comments and recommendations by the U.S. Environmental Protection Agency (EPA) Region VI and New Mexico Environment Department (NMED), SNL/NM will use these default exposure routes and parameter values in future risk assessments.

At SNL/NM, all SWMUs exist within the boundaries of the Kirtland Air Force Base. Approximately 240 potential waste and release sites have been identified where hazardous. radiological, or mixed materials may have been released to the environment. Evaluation and characterization activities have occurred at all of these sites to varying degrees. Among other documents, the SNL/NM ER draft Environmental Assessment (DOE 1996) presents a summary of the hydrogeology of the sites and the biological resources present. When evaluating potential human health risk the current or reasonably foreseeable land use negotiated and approved for the specific SWMU/AOC, aggregate, or watershed will be used. The following references generally document these land uses: Workbook: Future Use Management Area 2 (DOE et al. September 1995); Workbook: Future Use Management Area 1 (DOE et al. October 1995); Workbook: Future Use Management Areas 3, 4, 5, and 6 (DOE and USAF January 1996); Workbook: Future Use Management Area 7 (DOE and USAF March 1996). At this time, all SNL/NM SWMUs have been tentatively designated for either industrial or recreational future land use. The NMED has also requested that risk calculations be performed based upon a residential land-use scenario. Therefore, all three land-use scenarios will be addressed in this document.

The SNL/NM ER Project has screened the potential exposure routes and identified default parameter values to be used for calculating potential intake and subsequent hazard index (HI), excess cancer risk and dose values. The EPA (EPA 1989) provides a summary of exposure routes that could potentially be of significance at a specific waste site. These potential exposure routes consist of:

- Ingestion of contaminated drinking water
- · Ingestion of contaminated soil

- · Ingestion of contaminated fish and shellfish
- Ingestion of contaminated fruits and vegetables
- Ingestion of contaminated meat, eggs, and dairy products
- Ingestion of contaminated surface water while swimming
- Dermal contact with chemicals in water
- · Dermal contact with chemicals in soil
- Inhalation of airborne compounds (vapor phase or particulate)
- External exposure to penetrating radiation (immersion in contaminated air; immersion in contaminated water; and exposure from ground surfaces with photon-emitting radionuclides)

Based upon the location of the SNL/NM SWMUs and the characteristics of the surface and subsurface at the sites, we have evaluated these potential exposure routes for different landuse scenarios to determine which should be considered in risk assessment analyses (the last exposure route is pertinent to radionuclides only). At SNL/NM SWMUs, there is currently no consumption of fish, shellfish, fruits, vegetables, meat, eggs, or dairy products that originate on site. Additionally, no potential for swimming in surface water is present due to the high-desert environmental conditions. As documented in the RESRAD computer code manual (ANL 1993), risks resulting from immersion in contaminated air or water are not significant compared to risks from other radiation exposure routes.

For the industrial and recreational land-use scenarios, SNL/NM ER has, therefore, excluded the following four potential exposure routes from further risk assessment evaluations at any SNL/NM SWMU:

- Ingestion of contaminated fish and shellfish
- Ingestion of contaminated fruits and vegetables
- Ingestion of contaminated meat, eggs, and dairy products
- Ingestion of contaminated surface water while swimming
- · Dermal contact with chemicals in water

That part of the exposure pathway for radionuclides related to immersion in contaminated air or water is also eliminated.

Based upon this evaluation, for future risk assessments the exposure routes that will be considered are shown in Table 1.

Table 1
Exposure Pathways Considered for Various Land-Use Scenarios

Industrial	Recreational	Residential
Ingestion of contaminated drinking water	Ingestion of contaminated drinking water	Ingestion of contaminated drinking water
Ingestion of contaminated soil	Ingestion of contaminated soil	Ingestion of contaminated soil
Inhalation of airborne compounds (vapor phase or particulate)	Inhalation of airborne compounds (vapor phase or particulate)	Inhalation of airborne compounds (vapor phase or particulate)
Dermal contact (nonradiological constituents only) soil only	Dermal contact (nonradiological constituents only) soil only	Dermal contact (nonradiological constituents only) soil only
External exposure to penetrating radiation from ground surfaces	External exposure to penetrating radiation from ground surfaces	External exposure to penetrating radiation from ground surfaces

# Equations and Default Parameter Values for Identified Exposure Routes

In general, SNL/NM expects that ingestion of compounds in drinking water and soil will be the more significant exposure routes for chemicals; external exposure to radiation may also be significant for radionuclides. All of the above routes will, however, be considered for their appropriate land-use scenarios. The general equation for calculating potential intakes via these routes is shown below. The equations are taken from "Assessing Human Health Risks Posed by Chemicals: Screening-Level Risk Assessment" (NMED March 2000) and "Technical Background Document for Development of Soil Screening Levels" (NMED December 2000). Equations from both documents are based upon the "Risk Assessment Guidance for Superfund" (RAGS): Volume 1 (EPA 1989, 1991). These general equations also apply to calculating potential intakes for radionuclides. A more in-depth discussion of the equations used in performing radiological pathway analyses with the RESRAD code may be found in the RESRAD Manual (ANL 1993). RESRAD is the only code designated by the U.S. Department of Energy (DOE) in DOE Order 5400.5 for the evaluation of radioactively contaminated sites (DOE 1993). The Nuclear Regulatory Commission (NRC) has approved the use of RESRAD for dose evaluation by licensees involved in decommissioning, NRC staff evaluation of waste disposal requests, and dose evaluation of sites being reviewed by NRC staff. EPA Science Advisory Board reviewed the RESRAD model. EPA used RESRAD in their rulemaking on radiation site cleanup regulations. RESRAD code has been verified, undergone several benchmarking analyses, and been included in the International Atomic Energy Agency's VAMP and BIOMOVS Il projects to compare environmental transport models.

Also shown are the default values SNL/NM ER will use in RME risk assessment calculations for industrial, recreational, and residential land-use scenarios, based upon EPA and other governmental agency guidance. The pathways and values for chemical contaminants are discussed first, followed by those for radionuclide contaminants. RESRAD input parameters that are left as the default values provided with the code are not discussed. Further information relating to these parameters may be found in the RESRAD Manual (ANL 1993) or by directly accessing the RESRAD websites at: http://web.ead.anl.gov/resrad/home2/ or http://web.ead.anl.gov/resrad/documents/.

# Generic Equation for Calculation of Risk Parameter Values

The equation used to calculate the risk parameter values (i.e., hazard quotients/HI, excess cancer risk, or radiation total effective dose equivalent [TEDE] [dose]) is similar for all exposure pathways and is given by:

Risk (or Dose) = Intake x Toxicity Effect (either carcinogenic, noncarcinogenic, or radiological)

= 
$$C \times (CR \times EFD/BW/AT) \times Toxicity Effect$$
 (1)

where:

C = contaminant concentration (site specific)

CR = contact rate for the exposure pathway

EFD= exposure frequency and duration

BW = body weight of average exposure individual

AT = time over which exposure is averaged.

For nonradiological constituents of concern (COCs), the total risk/dose (either cancer risk or HI) is the sum of the risks/doses for all of the site-specific exposure pathways and contaminants. For radionuclides, the calculated radiation exposure, expressed as TEDE is compared directly to the exposure guidelines of 15 millirem per year (mrem/year) for industrial and recreational future use and 75 mrem/year for the unlikely event that institutional control of the site is lost and the site is used for residential purposes (EPA 1997).

The evaluation of the carcinogenic health hazard produces a quantitative estimate for excess cancer risk resulting from the COCs present at the site. This estimate is evaluated for determination of further action by comparison of the quantitative estimate with the potentially acceptable risk of 1E-5 for nonradiological carcinogens. The evaluation of the noncarcinogenic health hazard produces a quantitative estimate (i.e., the HI) for the toxicity resulting from the COCs present at the site. This estimate is evaluated for determination of further action by comparison of this quantitative estimate with the EPA standard HI of unity (1). The evaluation of the health hazard from radioactive compounds produces a quantitative estimate of doses resulting from the COCs present at the site. This estimated dose is used to calculate an assumed risk. However, this calculated risk is presented for illustration purposes only, not to determine compliance with regulations.

The specific equations used for the individual exposure pathways can be found in RAGS (EPA 1989) and are outlined below. The RESRAD Manual (ANL 1993) describes similar equations for the calculation of radiological exposures.

#### Soil Ingestion

A receptor can ingest soil or dust directly by working in the contaminated soil. Indirect ingestion can occur from sources such as unwashed hands introducing contaminated soil to food that is then eaten. An estimate of intake from ingesting soil will be calculated as follows:

$$I_s = \frac{C_s * IR * CF * EF * ED}{BW * AT}$$

#### where:

I<sub>s</sub> = Intake of contaminant from 55.....g = Chemical concentration in soil (mg/kg) IR = Ingestion rate (mg soil/day) = Intake of contaminant from soil ingestion (milligrams [mg]/kilogram [kg]-day)

CF = Conversion factor (1E-6 kg/mg)

EF = Exposure frequency (days/year)

ED = Exposure duration (years)

BW = Body weight (kg)

AT = Averaging time (period over which exposure is averaged) (days)

It should be noted that it is conservatively assumed that the receptor only ingests soil from the contaminated source.

# Soil Inhalation

A receptor can inhale soil or dust directly by working in the contaminated soil. An estimate of intake from inhaling soil will be calculated as follows (EPA August 1997):

$$I_{s} = \frac{C_{s} * IR * EF * ED * \left(\frac{1}{VF} \text{ or } \frac{1}{PEF}\right)}{RW * AT}$$

where:

= Intake of contaminant from soil inhalation (mg/kg-day)

I<sub>s</sub> = Intake of contaminant non-collection C<sub>s</sub> = Chemical concentration in soil (mg/kg) IR = Inhalation rate (cubic meters [m³]/day)

EF = Exposure frequency (days/year)

ED = Exposure duration (years)

VF = soil-to-air volatilization factor (m<sup>3</sup>/kg)

PEF = particulate emission factor (m<sup>3</sup>/kg)

BW = Body weight (kg)

AT = Averaging time (period over which exposure is averaged) (days)

#### Soil Dermal Contact

$$D_a = \frac{C_s * CF * SA * AF * ABS * EF * ED}{BW * AT}$$

where:

 $D_a$  = Absorbed dose (mg/kg-day)  $C_s$  = Chemical concentration in soil (mg/kg) CF = Conversion factor (1E-6 kg/mg)

SA = Skin surface area available for contact (cm<sup>2</sup>/event)

AF = Soil to skin adherence factor (mg/cm<sup>2</sup>)

ABS = Absorption factor (unitless)

EF = Exposure frequency (events/year)

ED = Exposure duration (years)

BW = Body weight (kg)

AT = Averaging time (period over which exposure is averaged) (days)

#### **Groundwater Ingestion**

A receptor can ingest water by drinking it or through using household water for cooking. An estimate of intake from ingesting water will be calculated as follows (EPA August 1997):

$$I_{w} = \frac{C_{w} * IR * EF * ED}{BW * AT}$$

where:

 $I_w$  = Intake of contaminant from water ingestion (mg/kg/day)  $C_w$  = Chemical concentration in water (mg/liter [L])

IR = Ingestion rate (L/day)

EF = Exposure frequency (days/year)

ED = Exposure duration (years)

BW = Body weight (kg)

AT = Averaging time (period over which exposure is averaged) (days)

# **Groundwater Inhalation**

The amount of a constituent taken into the body via exposure to volatilization from showering or other household water uses will be evaluated using the concentration of the constituent in the water source (EPA 1991 and 1992). An estimate of intake from volatile inhalation from groundwater will be calculated as follows (EPA 1991):

$$I_{w} = \frac{C_{w} * K * IR_{i} * EF * ED}{BW * AT}$$

where:

= Intake of volatile in water from inhalation (mg/kg/day)

IR; = Inhalation rate (m³/day)

EF = Exposure frequency (days/year)

ED = Exposure duration (years)

BW = Body weight (kg)

AT = Averaging time (period over which exposure is averaged—days)

For volatile compounds, volatilization from groundwater can be an important exposure pathway from showering and other household uses of groundwater. This exposure pathway will only be evaluated for organic chemicals with a Henry's Law constant greater than 1x10-5 and with a molecular weight of 200 grams/mole or less (EPA 1991).

Tables 2 and 3 show the default parameter values suggested for use by SNL/NM at SWMUs. based upon the selected land-use scenarios for nonradiological and radiological COCs,

respectively. References are given at the end of the table indicating the source for the chosen parameter values. SNL/NM uses default values that are consistent with both regulatory guidance and the RME approach. Therefore, the values chosen will, in general, provide a conservative estimate of the actual risk parameter. These parameter values are suggested for use for the various exposure pathways, based upon the assumption that a particular site has no unusual characteristics that contradict the default assumptions. For sites for which the assumptions are not valid, the parameter values will be modified and documented.

# Summary

SNL/NM will use the described default exposure routes and parameter values in risk assessments at sites that have an industrial, recreational, or residential future land-use scenario. There are no current residential land-use designations at SNL/NM ER sites, but NMED has requested this scenario to be considered to provide perspective of the risk under the more restrictive land-use scenario. For sites designated as industrial or recreational land use, SNL/NM will provide risk parameter values based upon a residential land-use scenario to indicate the effects of data uncertainty on risk value calculations or in order to potentially mitigate the need for institutional controls or restrictions on SNL/NM ER sites. The parameter values are based upon EPA guidance and supplemented by information from other government sources. If these exposure routes and parameters are acceptable, SNL/NM will use them in risk assessments for all sites where the assumptions are consistent with site-specific conditions. All deviations will be documented.

Table 2
Default Nonradiological Exposure Parameter Values for Various Land-Use Scenarios

Parameter	Industrial	Recreational	Residential
General Exposure Parameters		•	
		8.7 (4 hr/wk for	
Exposure Frequency (day/yr)	250 <sup>a,b</sup>	52 wk/yr) <sup>a,b</sup>	350 <sup>a,b</sup>
Exposure Duration (yr)	25 <sup>a,b,c</sup>	30 <sup>a,b,c</sup>	30 <sup>a,b,c</sup>
	70 <sup>a,b,c</sup>	70 Adulta,b,c	70 Adult <sup>a,b,c</sup>
Body Weight (kg)		15 Child <sup>a,b,c</sup>	15 Child <sup>a,b,c</sup>
Averaging Time (days)			
for Carcinogenic Compounds	25,550 <sup>a,b</sup>	25,550 <sup>a,b</sup>	25,550 <sup>a,b</sup>
(= 70 yr x 365 day/yr) for Noncarcinogenic Compounds (= ED x 365 day/yr)	9,125 <sup>a,b</sup>	10,950 <sup>a,b</sup>	10,950 <sup>a,b</sup>
Soil Ingestion Pathway	· · · · · · · · · · · · · · · · · · ·	<u> </u>	
Ingestion Rate (mg/day)	100 <sup>a,b</sup>	200 Child <sup>a,b</sup>	200 Childa,b
( ) ,		100 Adult <sup>a,b</sup>	100 Adult <sup>a,b</sup>
Inhalation Pathway			
		15 Child <sup>a</sup>	10 Child <sup>a</sup>
Inhalation Rate (m³/day)	20 <sup>a,b</sup>	30 Adulta	20 Adult <sup>a</sup>
Volatilization Factor (m³/kg)	Chemical Specific	Chemical Specific	Chemical Specific
Particulate Emission Factor (m <sup>3</sup> /kg)	1.36E9 <sup>a</sup>	1.36E9 <sup>a</sup>	1.36E9 <sup>a</sup>
Water Ingestion Pathway			
	2.4 <sup>a</sup>	2.4 <sup>a</sup>	2.4 <sup>a</sup>
Ingestion Rate (liter/day)			
Dermal Pathway			
		0.2 Child <sup>a</sup>	0.2 Childa
Skin Adherence Factor (mg/cm²)	0.2a	0.07 Adulta	0.07 Adult <sup>a</sup>
Exposed Surface Area for Soil/Dust		2,800 Childa	2,800 Childa
(cm²/day)	3,300a	5,700 Adulta	5,700 Adult <sup>a</sup>
Skin Adsorption Factor	Chemical Specific	Chemical Specific	Chemical Specific

<sup>&</sup>lt;sup>a</sup>Technical Background Document for Development of Soil Screening Levels (NMED 2000).

ED = Exposure duration.

EPA = U.S. Environmental Protection Agency.

hr = Hour(s).

kg = Kilogram(s).

m = Meter(s).

mg = Milligram(s).

NA = Not available.

wk = Week(s).

yr = Year(s).

<sup>&</sup>lt;sup>b</sup>Risk Assessment Guidance for Superfund, Vol. 1, Part B (EPA 1991).

<sup>°</sup>Exposure Factors Handbook (EPA August 1997).

Table 3
Default Radiological Exposure Parameter Values for Various Land-Use Scenarios

Parameter	Industrial	Recreational	Residential
General Exposure Parameters			
	8 hr/day for		
Exposure Frequency	250 day/yr	4 hr/wk for 52 wk/yr	365 day/yr
Exposure Duration (yr)	25 <sup>a,b</sup>	30 <sup>a,b</sup>	30 <sup>a,b</sup>
Body Weight (kg)	70 Adult <sup>a,b</sup>	70 Adult <sup>a,b</sup>	70 Adult <sup>a,b</sup>
Soil Ingestion Pathway			
Ingestion Rate	100 mg/day <sup>c</sup>	100 mg/day <sup>c</sup>	100 mg/day <sup>c</sup>
Averaging Time (days) (= 30 yr x 365 day/yr)	10,950 <sup>d</sup>	10,950 <sup>d</sup>	10,950 <sup>d</sup>
Inhalation Pathway			
Inhalation Rate (m³/yr)	7,300 <sup>d,e</sup>	10,950 <sup>e</sup>	7,300 <sup>d,e</sup>
Mass Loading for Inhalation g/m <sup>3</sup>	1.36E-5 <sup>d</sup>	1.36E-5 <sup>d</sup>	1.36E-5 <sup>d</sup>
Food Ingestion Pathway			
Ingestion Rate, Leafy Vegetables			
(kg/yr)	NA	NA	16.5 <sup>c</sup>
Ingestion Rate, Fruits, Non-Leafy			
Vegetables & Grain (kg/yr)	NA	NA	101.8 <sup>b</sup>
Fraction Ingested	NA	NA	0.25 <sup>b,d</sup>

<sup>&</sup>lt;sup>a</sup>Risk Assessment Guidance for Superfund, Vol. 1, Part B (EPA 1991).

EPA = U.S. Environmental Protection Agency.

g = Gram(s)

hr = Hour(s).

kg = Kilogram(s).

m = Meter(s).

mg = Milligram(s).

NA = Not applicable.

wk = Week(s).

yr = Year(s).

<sup>&</sup>lt;sup>b</sup>Exposure Factors Handbook (EPA August 1997).

<sup>°</sup>EPA Region VI guidance (EPA 1996).

dFor radionuclides, RESRAD (ANL 1993).

eSNL/NM (February 1998).

# References

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DOE and USAF, see U.S. Department of Energy and U.S. Air Force.

EPA, see U.S. Environmental Protection Agency.

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- U.S. Environmental Protection Agency (EPA), 1997. (OSWER No. 9200.4-18) Establishment of Cleanup Levels for CERCLA Sites with Radioactive Contamination, U.S. EPA Office of Radiation and Indoor Air, Washington D.C, August 1997.

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# APPENDIX 2 CALCULATION OF THE UPPER CONFIDENCE LIMITS OF MEAN CONCENTRATIONS

For conservatism, Sandia National Laboratories/New Mexico uses the maximum concentration of the constituents of concern (COCs) for initial risk calculation. If the maximum concentrations produce risk above New Mexico Environment Department (NMED) guidelines, conservatism with this approach is evaluated and, if appropriate, a more realistic approach is applied. When the site has been adequately characterized, an estimate of the mean concentration of the COCs is more representative of actual site conditions. The NMED has proposed the use of the 95% upper confidence limit (UCL) of the mean to represent average concentrations at a site (NMED December 2000). The 95% UCL is calculated according to NMED guidance (Tharp June 2002) using the U.S. Environmental Protection Agency ProUCL program (EPA April 2002). Attached are the outputs from that program and the calculated UCLs used in the risk analysis.

# References

EPA, see U.S. Environmental Protection Agency.

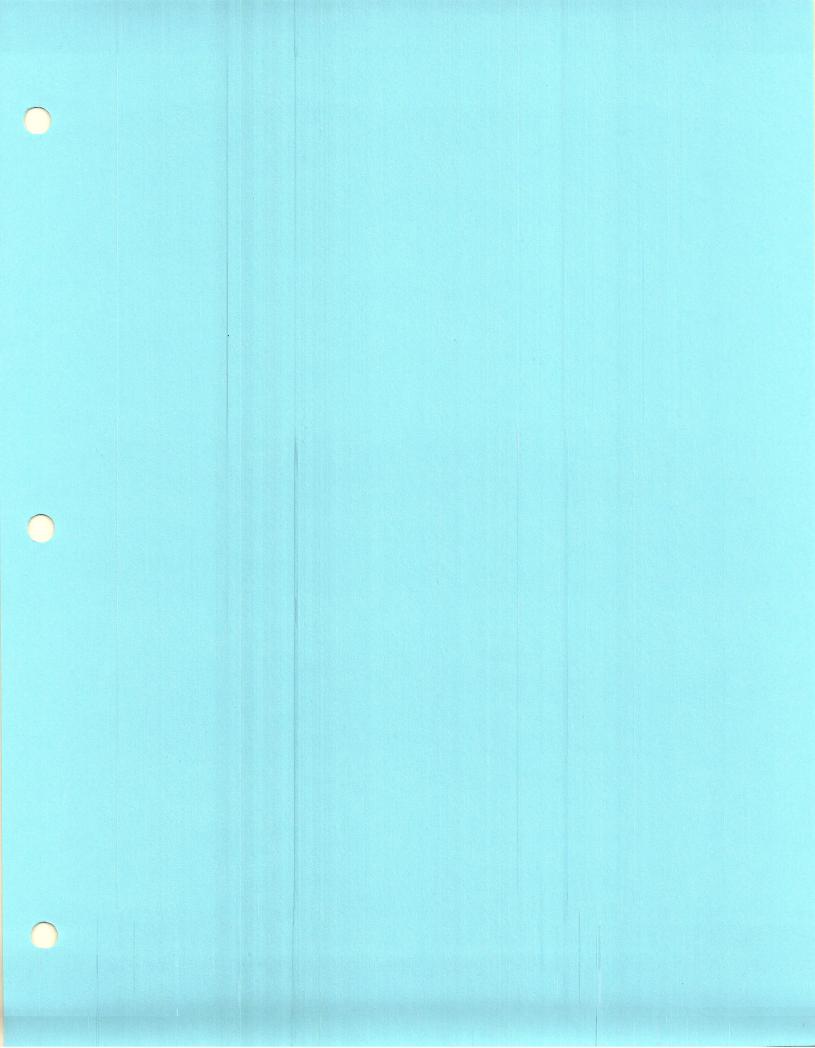
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NMED, see New Mexico Environment Department.

Tharp, T. (Weston Solutions, Inc.), June 2002. *Personal communication* with K. Olsen (Hazardous Waste Bureau, New Mexico Environment Department). June 12, 2002.

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**ATTACHMENT** 

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SWMU 136 Human Health		
Summary Statistics for	Antimony	
Number of Samples	23	
Minimum	2.1	
Maximum	6	
Mean	3.221739	
Median	3	
Standard Deviation	0.896436	
Variance	0.803597	
Coefficient of Variation	0.278246	
Skewness	2.851382	
	1 = 1 = 1 = 1	
Shapiro-Wilk Test Statisitic	0.461552	
Shapiro-Wilk 5% Critical Value	0.914	
Data not Lognormal at 5% Signit	ficance Level	
Data not Normal: Try Non-param	netric UCL	
95 % UCL (Assuming No	ormal Data)	~
Student's-t	3.542707	
	1	
95 % UCL (Adjusted for	Skewness)	
Adjusted-CLT	3.647943	-
Modified-t	3.56123	
	<b>-</b>	
95 % Non-parametric UC	CL C	
CLT	3.529195	
Jackknife	3.542707	
Standard Bootstrap	3.520877	
Bootstrap-t	4.467871	
Chebyshev (Mean, Std)	4.036503	

SWMU 136 Human Health			
Summary Statistics for	Arsenic	Summary Statistics for	In(Arsenic)
Number of Samples	39	Minimum	-0.04082
Minimum	0.96	Maximum	1.791759
Maximum	6	Mean	0.954062
Mean	2.803077	Standard Deviation	0.40486
Median	2.6	Variance	0.163912
Standard Deviation	1.113378		
Variance	1.239611	Shapiro-Wilk Test Statisitic	0.971368
Coefficient of Variation	0.397199	Shapiro-Wilk 5% Critical Value	0.939
Skewness	0.960525	Data are Lognormal at 5% Signif	ficance Level
95 % UCL (Assuming	Normal Data)	Estimates Assuming Lognormal	Distribution
Student's-t	3.103654	MLE Mean	2.817973
		MLE Standard Deviation	1.189272
95 % UCL (Adjusted	for Skewness)	MLE Coefficient of Variation	0.422031
Adjusted-CLT	3.125627	MLE Skewness	1.341262
Modified-t	3.108224	MLE Median	2.596234
		MLE 80% Quantile	3.655265
95 % Non-parametric	UCL	MLE 90% Quantile	4.368008
CLT	3.096327	MLE 95% Quantile	5.053377
Jackknife	3.103654	MLE 99% Quantile	6.657649
Standard Bootstrap	3.095943		
Bootstrap-t	3.145494	MVU Estimate of Median	2.590783
Chebyshev (Mean, Std)	3.580195	MVU Estimate of Mean	2.811611
		MVU Estimate of Std. Dev.	1.176467
		MVU Estimate of SE of Mean	0.188068
		UCL Assuming Lognormal Dis	tribution
		95% H-UCL	3.180339
		95% Chebyshev (MVUE) UCL	3.631382
		99% Chebyshev (MVUE) UCL	4.682867
		Recommended UCL to use:	
		Student's-t or H-UCL	

SWMU 136 Human Health	
Summary Statistics for	Barium
Number of Samples	39
Minimum	16.8
Maximum	944
Mean	144.3744
Median	101
Standard Deviation	160.66
Variance	25811.62
Coefficient of Variation	1.112801
Skewness	4.000599
Shapiro-Wilk Test Statisitic	0.915257
Shapiro-Wilk 5% Critical Value	0.939
Data not Lognormal at 5% Signification	
Data not Normal: Try Non-parame	etric UCL
99 % UCL (Assuming No	
Student's-t	206.8521
99 % UCL (Adjusted for S	Skewness)
Adjusted-CLT	236.6992
Modified-t	209.5989
99 % Non-parametric UC	L
CLT	204.2224
Jackknife	206.8521
Standard Bootstrap	204.0166
Bootstrap-t	319.0228
Chebyshev (Mean, Std)	400.3466

SWMU 136	Human Health	
Summary S	tatistics for	Cadmium
Number of		39
Minimum		0.015
Maximum		7.88
Mean		0.395256
Median		0.25
Standard D	eviation	1.242799
Variance		1.54455
Coefficient of Variation		3.144286
Skewness		6.047805
Shapiro-Wilk Test Statisitic		0.772481
	k 5% Critical Value	0.939
Data not Lognormal at 5% Signif		
Data not Normal: Try Non-param		etric UCL
	% UCL (Assuming No	ormal Data)
Student's-t		0.798125
	% UCL (Adjusted for	
Adjusted-Cl	_T	1.064203
Modified-t		0.830246
	% Non-parametric U0	
CLT		0.785303
Jackknife		0.798125
Standard Bo	ootstrap	0.772959
Bootstrap-t		2.432334
Chebyshev	(Mean, Std)	1.638056

SWMU 136 Human Health	
OVVIVIO 130 Fiditian Fleatin	·
Summary Statistics for	Mercury
Number of Samples	39
Minimum	0.0085
Maximum	1.13
Mean	0.070038
Median	0.05
Standard Deviation	0.177882
Variance	0.031642
Coefficient of Variation	2.53977
Skewness	5.863381
Shapiro-Wilk Test Statisitic	0.753697
Shapiro-Wilk 5% Critical Value	0.939
Data not Lognormal at 5% Signifi	icance Level
Data not Normal: Try Non-param	etric UCL
95 % UCL (Assuming No	rmal Data)
Student's-t	0.118061
95 % UCL (Adjusted for \$	Skewness)
Adjusted-CLT	0.145466
Modified-t	0.122518
95 % Non-parametric UC	EL
CLT	0.11689
Jackknife	0.118061
Standard Bootstrap	0.116448
Bootstrap-t	0.28065
Chebyshev (Mean, Std)	0.194197

SWMU 136 Human Health		
SVIVIO 130 Human Health	<u> </u>	
Summary Statistics for	Thallium	
Number of Samples	23	
Minimum	0.05	
Maximum	1	<del></del>
Mean	0.491304	
Median	0.5	
Standard Deviation	0.226059	
Variance	0.051103	
Coefficient of Variation	0.460121	
Skewness	0.106846	
	•	
Shapiro-Wilk Test Statisitic	0.693756	""
Shapiro-Wilk 5% Critical Value	0.914	
Data not Lognormal at 5% Signifi	cance Leve	
Data not Normal: Try Non-param	etric UCL	
95 % UCL (Assuming No	rmal Data)	
Student's-t	0.572245	
95 % UCL (Adjusted for S	Skewness)	
Adjusted-CLT	0.569959	
Modified-t	0.57242	
95 % Non-parametric UC		
CLT	0.568837	
Jackknife	0.572245	
Standard Bootstrap	0.566495	
Bootstrap-t	0.569387	
Chebyshev (Mean, Std)	0.696768	

SWMU 136 Ecological			
Summary Statistics for	Cadmium	Summary Statistics for	In(Cadmium)
Number of Samples	9	Minimum	-4.19971
Minimum	0.015	Maximum	2.064328
Maximum	7.88	Mean	-1.55037
Mean	1.086111	Standard Deviation	1.890337
Median	0.25	Variance	3.573375
Standard Deviation	2.555013		
Variance	6.528092	Shapiro-Wilk Test Statisitic	0.869297
Coefficient of Variation	2.352442	Shapiro-Wilk 5% Critical Value	0.829
Skewness	2.967649	Data are Lognormal at 5% Significance Level	
95 % LICI (Assumin	g Normal Data) Estimates Assuming Lognormal Distribut		Dietzibution
95 % UCL (Assuming Normal Data) Student's-t 2.669834		MLE Mean	1.266577
Ottudent 3-1	2.003034	MLE Standard Deviation	7.454177
95 % UCL (Adjusted	for Skewness)	MLE Coefficient of Variation	5.885291
Adjusted-CLT	3.387195	MLE Skewness	221.5027
Modified-t	2.810249	MLE Median	0.21217
		MLE 80% Quantile	1.048086
95 % Non-parametric UCL		MLE 90% Quantile	2.407802
CLT	2.486985	MLE 95% Quantile	4.755185
Jackknife	2.669834	MLE 99% Quantile	17.22833
Standard Bootstrap	2.394459		
Bootstrap-t	22.9891	MVU Estimate of Median	0.173251
Chebyshev (Mean, Std)	4.798459	MVU Estimate of Mean	0.858952
		MVU Estimate of Std. Dev.	1.892579
99 % Non-parametri	c UCL	MVU Estimate of SE of Mean	0.544778
Chebyshev (Mean, Std)	9.560131		
		UCL Assuming Lognormal Dis	tribution
		95% H-UCL	51.84426
		95% Chebyshev (MVUE) UCL	3.233583
		99% Chebyshev (MVUE) UCL	6.279422
		Recommended UCL to use:	
		99 % Chebyshev (MVUE) UCL	

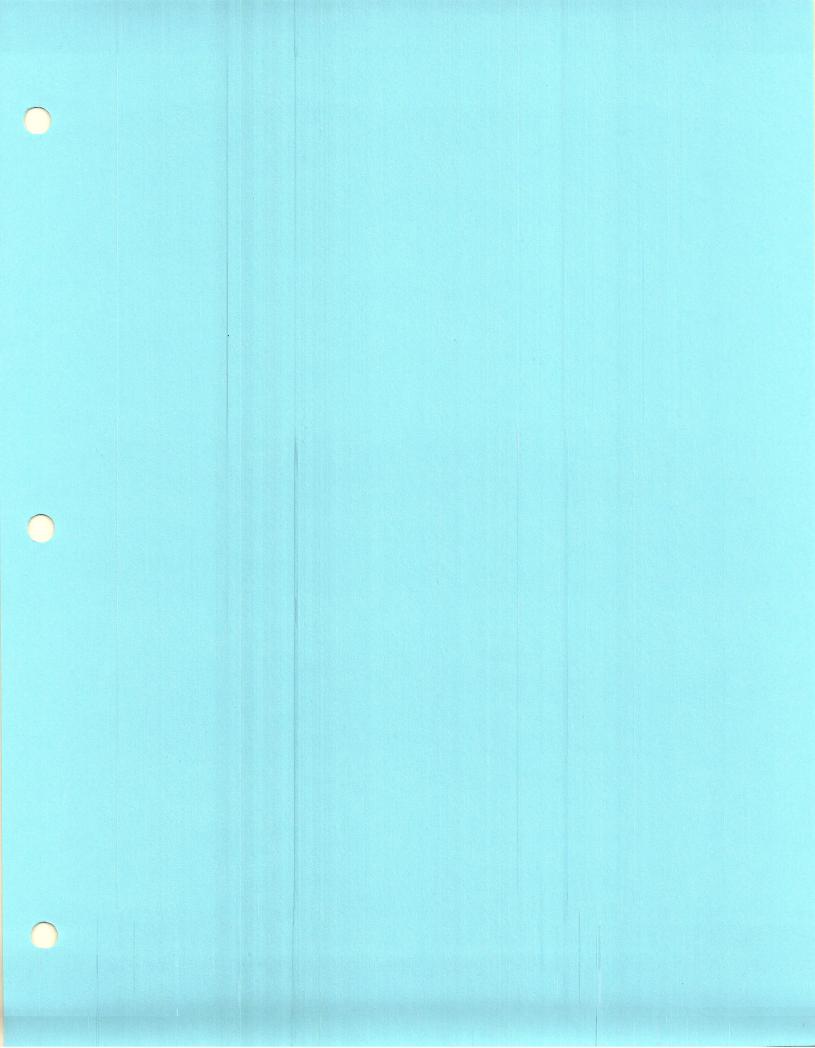
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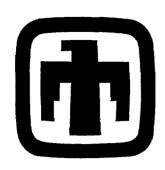
SWMU 136 Ecological			
Summary Statistics for	Chromium	Summary Statistics for	In(Chromium)
Number of Samples	9	Minimum	1.360977
Minimum	3.9	Maximum	2.753661
Maximum	15.7	Mean	2.00137
Mean	7.996667	Standard Deviation	0.40814
Median	6.5	Variance	0.166578
Standard Deviation	3.569993		
Variance	12.74485	Shapiro-Wilk Test Statisitic	0.925069
Coefficient of Variation	0.446435	Shapiro-Wilk 5% Critical Value	0.829
Skewness	1.398675	Data are Lognormal at 5% Significance Level	
95 % LICL (Assuming	n Normal Data)	Estimates Assuming Lognormal	Diotribustion
95 % UCL (Assuming Normal Data) Student's-t 10.20952		MLE Mean	8.041848
Ottudent 3-t	10.20932	MLE Standard Deviation	3.423748
OF 9/ LICL (Adjusted for Chaum)		MLE Coefficient of Variation	0.425741
95 % UCL (Adjusted for Skewness) Adjusted-CLT 10.54686		MLE Skewness	
Modified-t	10.34666		1.354392
Wodined-t	10.30199	MLE Median	7.399185
95 % Non-parametric UCL		MLE 80% Quantile	10.4463
CLT	9.954039	MLE 90% Quantile MLE 95% Quantile	12.50126
Jackknife	10.20952	MLE 99% Quantile	14.47988
Standard Bootstrap	9.90644	MLE 99% Quantile	19.11939
Bootstrap-t	11.94971	NAVILE Cation at a set Martin	7.000004
Chebyshev (Mean, Std)		MVU Estimate of Median	7.330964
Chebyshev (Mean, Std)	13.18375	MVU Estimate of Mean	7.963474
		MVU Estimate of Std. Dev.	3.285398
		MVU Estimate of SE of Mean	1.094262
		UCL Assuming Lognormal Distribution	
		95% H-UCL	10.99027
		95% Chebyshev (MVUE) UCL	12.73325
		99% Chebyshev (MVUE) UCL	18.85124
		Recommended UCL to use:	
		Student's-t or H-UCL	

SWMU 136 Ecological			
Summary Statistics for	Lead	Summary Statistics for	In(Lead)
Number of Samples	9	Minimum	1.458615
Minimum	4.3	Maximum	4.067316
Maximum	58.4	Mean	2.201793
Mean	13.40333	Standard Deviation	0.807195
Median	7.7	Variance	0.651563
Standard Deviation	17.18619		
Variance	295.3651	Shapiro-Wilk Test Statisitic	0.835385
Coefficient of Variation	1.282232	Shapiro-Wilk 5% Critical Value	0.829
Skewness	2.800981	Data are Lognormal at 5% Significance Le	
95 % UCL (Assuming	Normal Data)	Estimates Assuming Lognormal	Distribution
Student's-t	24.05618	MLE Mean	12.5231
		MLE Standard Deviation	12.00218
95 % UCL (Adjusted	for Skewness)	MLE Coefficient of Variation	0.958404
Adjusted-CLT	28.54141	MLE Skewness	3.755542
Modified-t	24.94763	MLE Median	9.04121
		MLE 80% Quantile	17.88339
95 % Non-parametric UCL		MLE 90% Quantile	25.50918
CLT	22.82626	MLE 95% Quantile	34.11129
Jackknife	24.05618	MLE 99% Quantile	59.10574
Standard Bootstrap	22.26489		
Bootstrap-t	63.76983	MVU Estimate of Median	8.718637
Chebyshev (Mean, Std)	38.37429	MVU Estimate of Mean	11.98302
		MVU Estimate of Std. Dev.	10.05561
		MVU Estimate of SE of Mean	3.317981
		UCL Assuming Lognormal Dis	etribution
		95% H-UCL	28.43478
		95% Chebyshev (MVUE) UCL	26.44576
		99% Chebyshev (MVUE) UCL	44.99652
		Recommended UCL to use:	44.55032
		H-UCL	
		I FOOL	

SWMU 136 Ecological			
Summary Statistics for	Mercury	Summary Statistics for	In(Mercury)
Number of Samples	9	Minimum	-4.76769
Minimum	0.0085	Maximum	0.122218
Maximum	1.13	Mean	-2.90012
Mean	0.175333	Standard Deviation	1.482144
Median	0.05	Variance	2.196752
Standard Deviation	0.36153		
Variance	0.130704	Shapiro-Wilk Test Statisitic	0.852805
Coefficient of Variation	2.061956	Shapiro-Wilk 5% Critical Value	0.829
Skewness	2.894159	Data are Lognormal at 5% Signi	ficance Level
95 % UCL (Assumin	g Normal Data)	Estimates Assuming Lognormal	Distribution
Student's-t	0.399427	MLE Mean	0.165011
		MLE Standard Deviation	0.466597
95 % UCL (Adjusted	for Skewness)	MLE Coefficient of Variation	2.827675
Adjusted-CLT	0.497778	MLE Skewness	31.09239
Modified-t	0.418804	MLE Median	0.055017
		MLE 80% Quantile	0.192491
95 % Non-parametric UCL		MLE 90% Quantile	0.369515
CLT	0.373554	MLE 95% Quantile	0.630027
Jackknife	0.399427	MLE 99% Quantile	1.728662
Standard Bootstrap	0.363465		
Bootstrap-t	2.833607	MVU Estimate of Median	0.048621
Chebyshev (Mean, Std)	0.700624	MVU Estimate of Mean	0.134961
		MVU Estimate of Std. Dev.	0.222681
		MVU Estimate of SE of Mean	0.068909
		UCL Assuming Lognormal Dis	stribution
		95% H-UCL	1.732001
		95% Chebyshev (MVUE) UCL	0.435327
		99% Chebyshev (MVUE) UCL	0.820593
		Recommended UCL to use:	
		95 % Chebyshev (MV	/UE) UCL

SWMU 136 Ecological		
Summary Statistics for	Silver	
Number of Samples	9	
Minimum	0.05	
Maximum	60.9	
Mean	7.363333	
Median	0.07	
Standard Deviation	20.11584	
Variance	404.6469	
Coefficient of Variation	2.731893	
Skewness	2.977957	
Shapiro-Wilk Test Statisitic	0.798688	
Shapiro-Wilk 5% Critical Value	0.829	
Data not Lognormal at 5% Signification	cance Leve	l
Data not Normal: Try Non-parame	etric UCL	
99 % UCL (Assuming No	rmal Data)	
Student's-t	26.7849	
99 % UCL (Adjusted for S	Skewness)	
Adjusted-CLT	36.07869	
Modified-t	27.89424	
99 % Non-parametric UC	L	
CLT	22.96214	
Jackknife	26.7849	
Standard Bootstrap	22.02239	
Bootstrap-t	994.8917	
Chebyshev (Mean, Std)	74.08002	





# Sandia National Laboratories/New Mexico Environmental Restoration Project

## REQUEST FOR SUPPLEMENTAL INFORMATION RESPONSE FOR DRAIN AND SEPTIC SYSTEMS SWMUs 48, 135, 136, 159, 165, 166, AND 167 AT TECHNICAL AREA II SOIL-VAPOR SAMPLING

June 2004



United States Department of Energy Sandia Site Office

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#### **ACRONYMS AND ABBREVIATIONS**

bas below ground surface

BTEX benzene, toluene, ethylbenzene, and xylene

DCB dichlorobenzene

DSS Drain and Septic Systems
ER Environmental Restoration

HE high explosive(s)

HWB Hazardous Waste Bureau MDL method detection limit

NFA no further action

NMED New Mexico Environment Department

NOD Notice of Deficiency PCE tetrachloroethene

RCRA Resource Conservation and Recovery Act
RSI Response for Supplemental Information
SNL/NM Sandia National Laboratories/New Mexico

SVOC semivolatile organic compound SWMU Solid Waste Management Unit

TA Technical Area

TAG Tijeras Arroyo Groundwater

TCA trichloroethane TCE trichloroethene

VOC volatile organic compound

#### 1.0 INTRODUCTION

#### 1.1 Investigation History

In August 1994, no further action (NFA) proposals were submitted for Solid Waste Management Units (SWMUs) 135 and 165 in Technical Area (TA)-II at Sandia National Laboratories/New Mexico (SNL/NM). In July 1995, NFA proposals were also submitted for TA-II SWMUs 48, 136, 159, 166, and 167. These seven SWMUs are shown on Figure 1.1-1.

In November 1995, the New Mexico Environment Department (NMED) Hazardous Waste Bureau (HWB) responded with comments on the NFA proposals submitted for SWMUs 48, 136, 159, 166, and 167 and recommended that a Resource Conservation and Recovery Act (RCRA) Facility Investigation Work Plan, which included these SWMUs, be developed for TA-II. At that time, the SNL/NM Environmental Restoration (ER) Project decided to undertake the investigation and cleanup of these sites and others in TA-II as Voluntary Corrective Actions, and formal work plans were not submitted.

On October 13, 1999, the NMED-HWB issued a Notice of Deficiency (NOD) for these seven SWMUs. Negotiations on November 17, 1999, further defined specific procedures for sampling these seven SWMUs and transferred a requirement for groundwater reporting for these SWMUs to the ongoing Tijeras Arroyo Groundwater (TAG) Investigation. The NOD subsequently was changed by NMED to a Request for Supplemental Information (RSI). The requirements negotiated to fulfill the RSI for these seven TA-II SWMUs were:

- Submit revised site maps showing septic and drain system component locations (as determined by backhoe excavation).
- Submit the results for passive soil-vapor surveys and active soil-vapor monitoring wells at TA-II.
- Collect soil samples at a depth equal to the base, and 5 feet below the base, of septic tanks, seepage pits, and drain lines. Sample locations in drainfields and system outfalls were approved by HWB personnel.
- Analyze soil samples for volatile organic compounds (VOCs), semivolatile organic compounds (SVOCs), high explosive (HE) compounds, polychlorinated biphenyls, RCRA metals, including hexavalent chromium, and total cyanide, radionuclides by gamma spectroscopy, and gross alpha/beta activity.
- Submit revised risk assessments for all seven SWMUs using all available soil data.

On January 26, 2000, the SNL/NM ER Project submitted a response to the NMED RSI, agreeing to excavations to locate system components below ground surface (bgs), confirm drainfield dimensions, pinpoint effluent release points, and investigate the SWMU 48 HE rinsewater drain line. SNL/NM also agreed to discuss additional sampling with the NMED-HWB when the maps were finalized and to submit the groundwater data requested in a subsequent TAG Investigation report.

For tracking purposes, these seven SWMUs are included with sites listed in the SNL/NM Drain and Septic Systems (DSS) program reporting schedule. In this RSI response, they will be referred to as the "Drain and Septic Systems SWMUs at TA-II."

#### 1.2 Additional Investigation Information

Although not specifically required as part of the RSI, this report presents additional information for several TA-II SWMUs as follows:

- In May 2003, soil-vapor monitoring wells were installed at SWMUs 159 and 165 as part of a separate site-wide DSS investigation. Additional details and sampling results for these wells are presented in the soil-vapor sampling chapter of this RSI response.
- Residual material in catch (settling) boxes for HE compound particulates located on HE rinse-water drain lines at SWMUs 48 and 136 was sampled as part of the site characterization process. The results are presented in the SWMU 48 and SWMU 136 chapters of this RSI response.

## 1.3 Report Organization

This RSI response presents the required information as follows:

- The soil-vapor survey information is presented as a whole and is not discussed on a site-by-site basis.
- Because NFA proposals were previously submitted for these SWMUs, only a brief
  description and history for each site is presented. Each SWMU is discussed as a
  separate report. The soil sampling analytical results and risk assessments are
  presented in separate annexes for each SWMU.

#### 2.0 SOIL-VAPOR SAMPLING AT TA-II

#### 2.1 Introduction

Soil-vapor data was collected using passive and active sampling methods at TA-II. The passive technique uses buried samplers to collect soil-vapor components onto suitable adsorbent material. After an appropriate period, the samplers are retrieved and analyzed for the components of interest. Active soil-vapor sampling analyzes soil-vapor volumes collected at discrete borehole depths while drilling or by specially designed vapor monitoring wells.

Passive soil-vapor surveys were conducted at TA-II in 1993 and 1994 to identify possible source areas. In November and December 1996, three boreholes were drilled in TA-II and sampled at 10-foot intervals for soil vapor. Two of these boreholes were later converted to permanent soil-vapor monitoring wells. In May 2003, soil-vapor monitoring wells were installed at SWMUs 159 and 165 as part of the SNL/NM site-wide DSS investigation. Additional details and the analytical results are presented in the following sections.

### 2.2 Passive Soil-Vapor Sampling at TA-II

## 2.2.1 Passive Soil-Vapor Sampling Methodology

PETREX™ passive soil-vapor sampling involves burying collectors containing activated carbon adsorption elements in shallow holes throughout the area to be investigated. After an appropriate period, usually approximately two weeks, the collectors are removed and submitted for analysis by thermal desorption-mass spectrometry. The methodology reports compound detections as relative intensities or response levels rather than the actual concentration of the compound in soil vapor. The data are best utilized as a semiquantitative measure, with an order of magnitude change in ion count values considered significant for interpreting potential source areas and migration/dispersion pathways versus background areas. Full details on the procedures, analytical methodology, and associated quality assurance/quality control measures are presented in the report prepared by Northeast Research Institute, Inc. (NERI 1994).

## 2.2.2 Passive Soil-Vapor Sampling Results

Two phases of passive soil-vapor sampling using PETREX™ collectors were conducted at TA-II (NERI 1994). The first phase, conducted between November and December 1993, installed 365 collectors throughout portions of TA-II. The second phase, conducted between January and February 1994, installed 99 additional collectors. Phase I was a broad reconnaissance survey to determine the types and locations of VOCs and SVOCs at the site. The Phase II survey further investigated potential areas of concern identified in Phase I.

As part of Phase I, eighteen collectors were installed west of TA-II in unimpacted areas assumed to represent background. It was later determined that these "background" areas may have been part of the old Oxnard Field runway where much activity occurred during the 1940s.

With the exception of some petroleum hydrocarbons, the areas provided suitable background data for VOCs and SVOCs.

The sample locations and identifications are shown on Figure 2.2.2-1. The analytical results for Phases I and II are presented in Tables A-1 and A-2 in Annex A. As shown in Table A-1, the majority of compounds detected in the soil-vapor samples were the chlorinated solvents trichloroethene (TCE) and tetrachloroethene (PCE) and the petroleum hydrocarbons benzene, toluene, ethylbenzene, and xylene(s) (BTEX). Table A-2 lists additional compounds detected during the surveys and shows that trichloroethane (TCA), dichlorobenzene (DCB), Freon-11 (trichlorofluoromethane), and Freon-113 (trichlorotrifluoroethane) were also detected in some samples.

Figures 2.2.2-2, 2.2.2-3, and 2.2.2-4 show the ion count contour plots for TCE, PCE, and BTEX respectively. Because TCA, DCB, Freon-11, and Freon-113 were detected only in a few samples, their distributions are not plotted. The highest TCE and BTEX ion counts were also identified near Buildings 913 and 914 at the southern end of TA-II. The survey concluded that the potential source area for these detections might exist southeast of TA-II (NERI 1994). No SVOCs were detected in any of the samples.

Figures 2.2.2-5 through 2.2.2-13 show the PETREX™ soil-vapor sample locations and, where appropriate, the soil-vapor monitoring wells for each of the TA-II SWMUs addressed in this RSI response. The analytical results for the passive soil-vapor samples at individual SWMUs are presented in Tables A-1 and A-2. The soil-vapor monitoring well analytical results are discussed Section 2.3.2.

### 2.3 Active Soil-Vapor Sampling at TA-II

#### 2.3.1 Active Soil-Vapor Sampling Methodology

Active soil-vapor sampling typically involves directly pumping soil-vapor from the subsurface for analysis. Vapor collection can be through simple open pipe systems analogous to groundwater monitoring wells screened in the interval of interest, or through sophisticated "down hole" systems with individual inlet port and collection tube sets placed at multiple depths. The extracted soil-vapor can be collected onto adsorbent media and analyzed immediately, or collected into special canisters for later laboratory analysis.

#### 2.3.2 Active Soil-Vapor Sampling Results

In November and December 1996, boreholes BH-020, BH-021, and BH-023 were drilled at TA-II (Figure 2.3.2-1) and sampled at 10-foot intervals during drilling for VOCs in soil-vapor. Permanent soil-vapor wells were constructed in boreholes BH-020 and BH-021 (TA2-VW-20 and TA2-VW-21). BH-023 was abandoned by backfilling with cuttings after drilling and sampling. Vapor well TA2-VW-020 was constructed so that vapor samples could be collected at 72 feet bgs. Vapor well TA2-VW-021 was constructed so that vapor samples could be collected at 52 and 92 feet bgs. The vapor wells were sampled for VOCs on an approximately quarterly basis between July 1997 and March 2002.

The July 1997 samples were collected onto adsorbent media both during and after purging of the collection system and were analyzed at the on-site Environmental Restoration Chemistry Laboratory. All subsequent samples were collected into special vacuum canisters and analyzed at various off-site laboratories. Sample results for wells TA2-VW-020 and TA2-VW-021 are presented in Tables A-3 and A-4, respectively. The results collected during the drilling of borehole BH-023 are presented in Table A-5. Method detection limits (MDLs) for the VOC analyses are presented in Table A-6.

The results for samples collected during the drilling for all three boreholes show the apparent widespread presence of VOCs in soil vapor at TA-II. Although the long-term monitoring data show a large amount of scatter, the results for vapor wells TA2-VW-20 and TA2-VW-21 indicate that VOC concentrations are somewhat steady with no apparent periodicity.

In May 2003, as part of the SNL/NM ER Project site-wide DSS investigation, soil-vapor monitoring wells were installed at SWMUs 159 and 165 (Figures 2.3.2-1, 2.2.2-10 and 2.2.2-11). Each vapor well was 150 feet deep and had vapor sampling ports at depths of 5, 20, 70, 100, 150 feet bgs. After installation, subsurface conditions were allowed to equilibrate over three months before the wells were sampled. The wells were sampled in September 2003 and the results are presented in Table A-7 for vapor well 159-VW-01 and Table A-8 for vapor well 165-VW-01. MDLs for the VOC analyses are presented in Table A-6.

In accordance with previous agreements with the NMED (SNL/NM October 1999), because the total VOC concentrations in the 150-foot-bgs sample from each well were less than 10 parts per million by volume, no additional soil-vapor sampling from these two wells, or soil-vapor or groundwater monitoring wells were required by the NMED at SWMUs 159 and 165.

#### 3.0 REFERENCES

EPA, see U.S. Environmental Protection Agency.

NERI, see Northeast Research Institute, Inc.

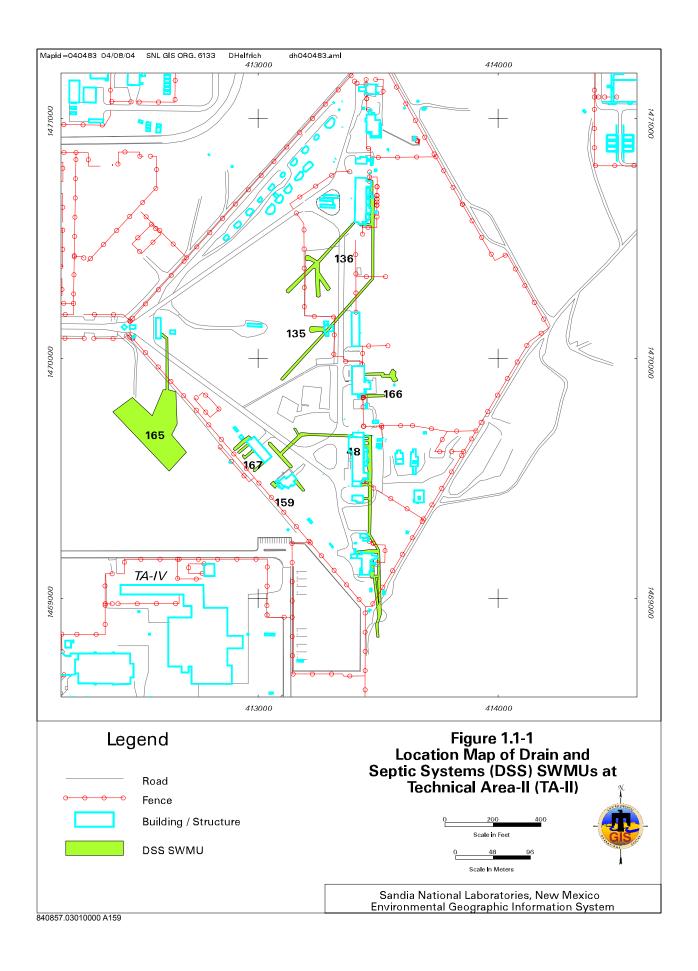
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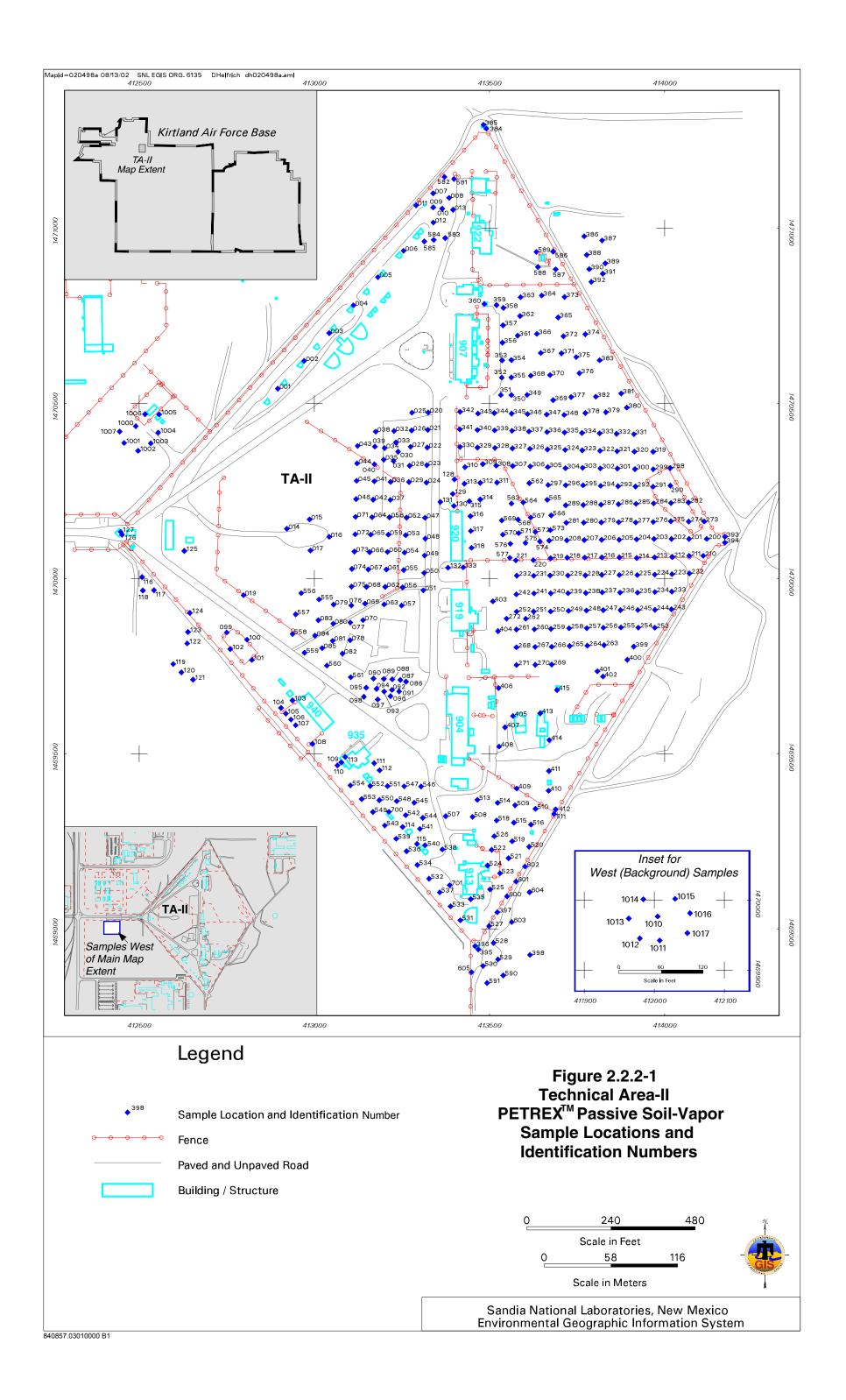
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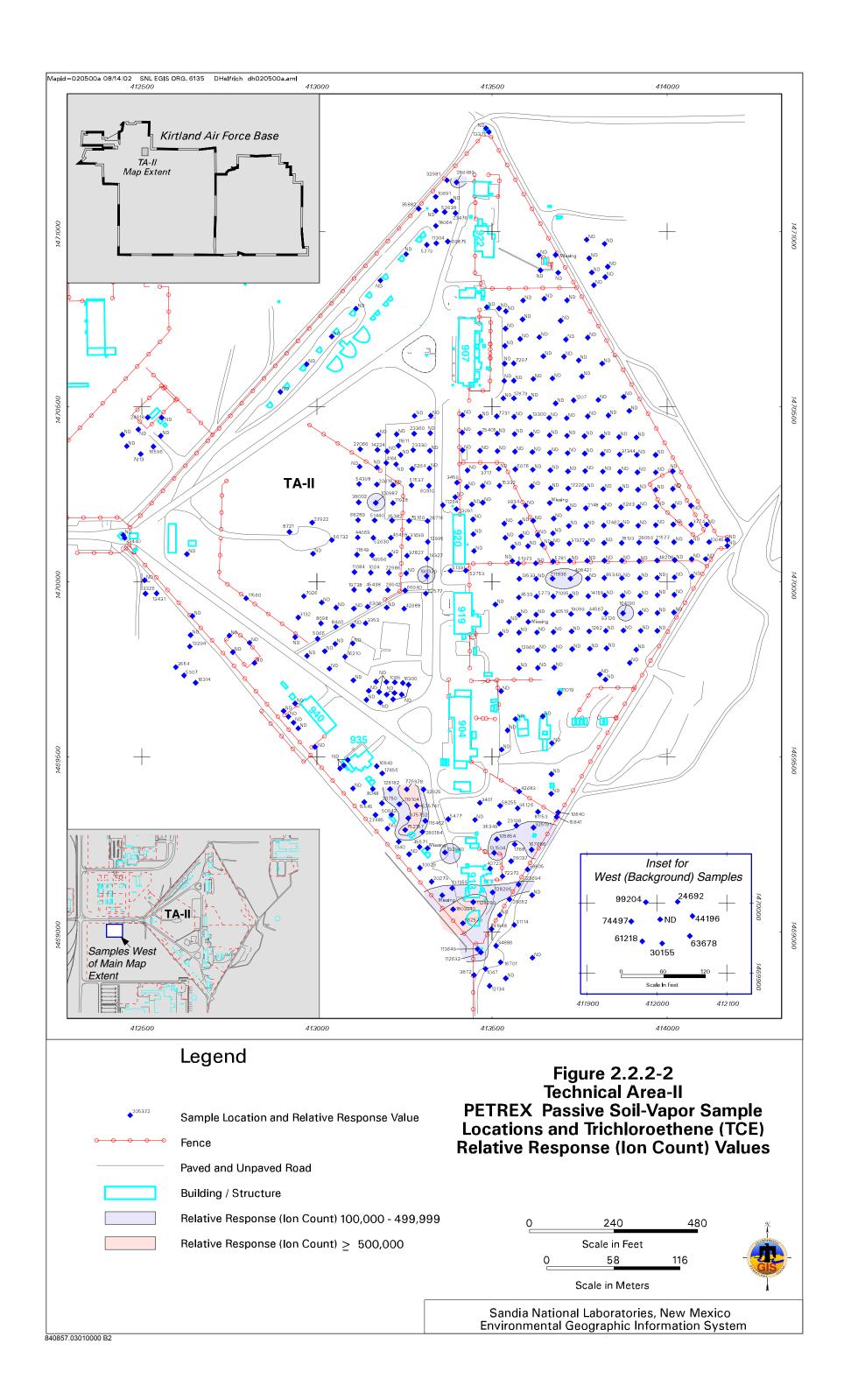
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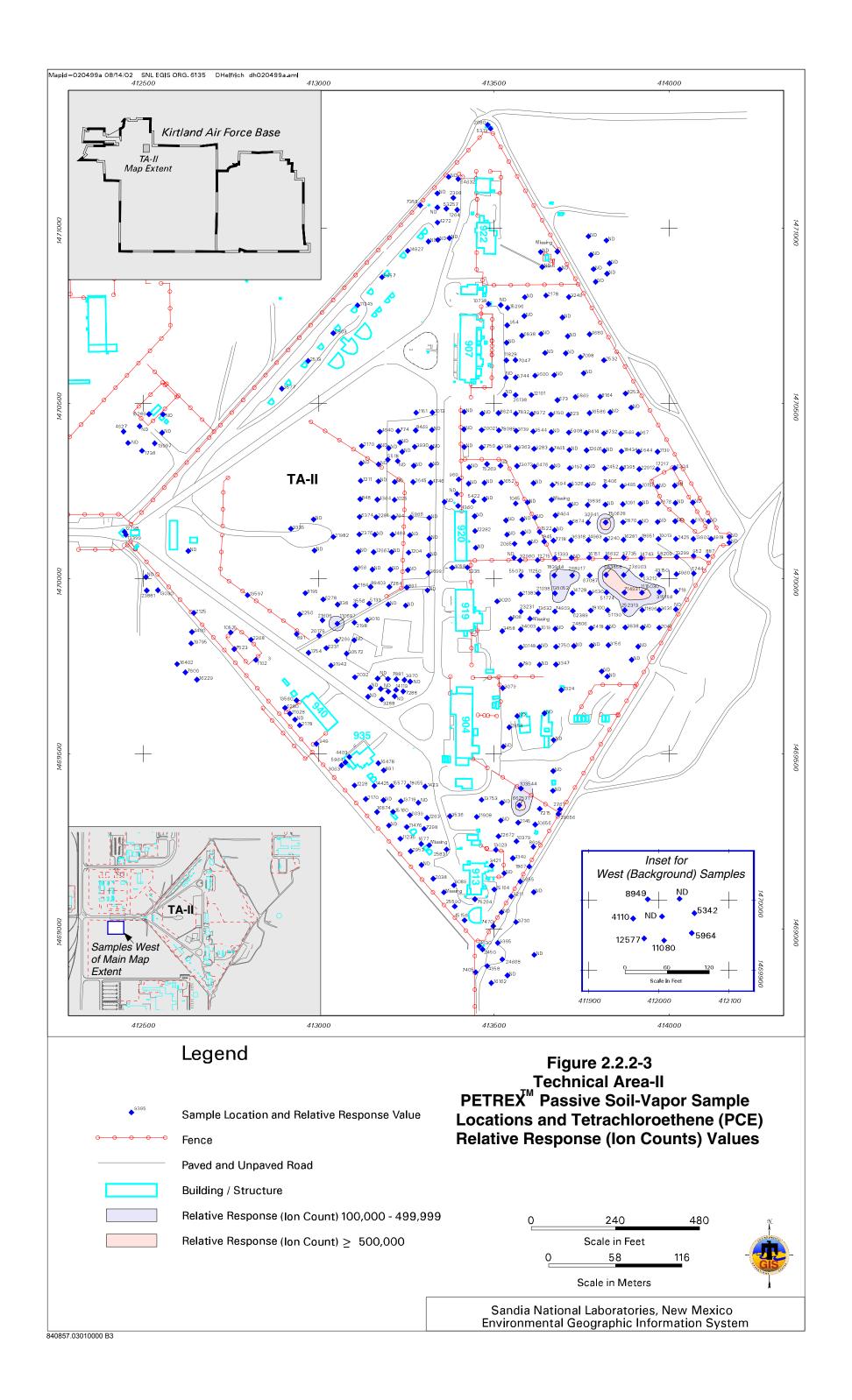
**FIGURES** 

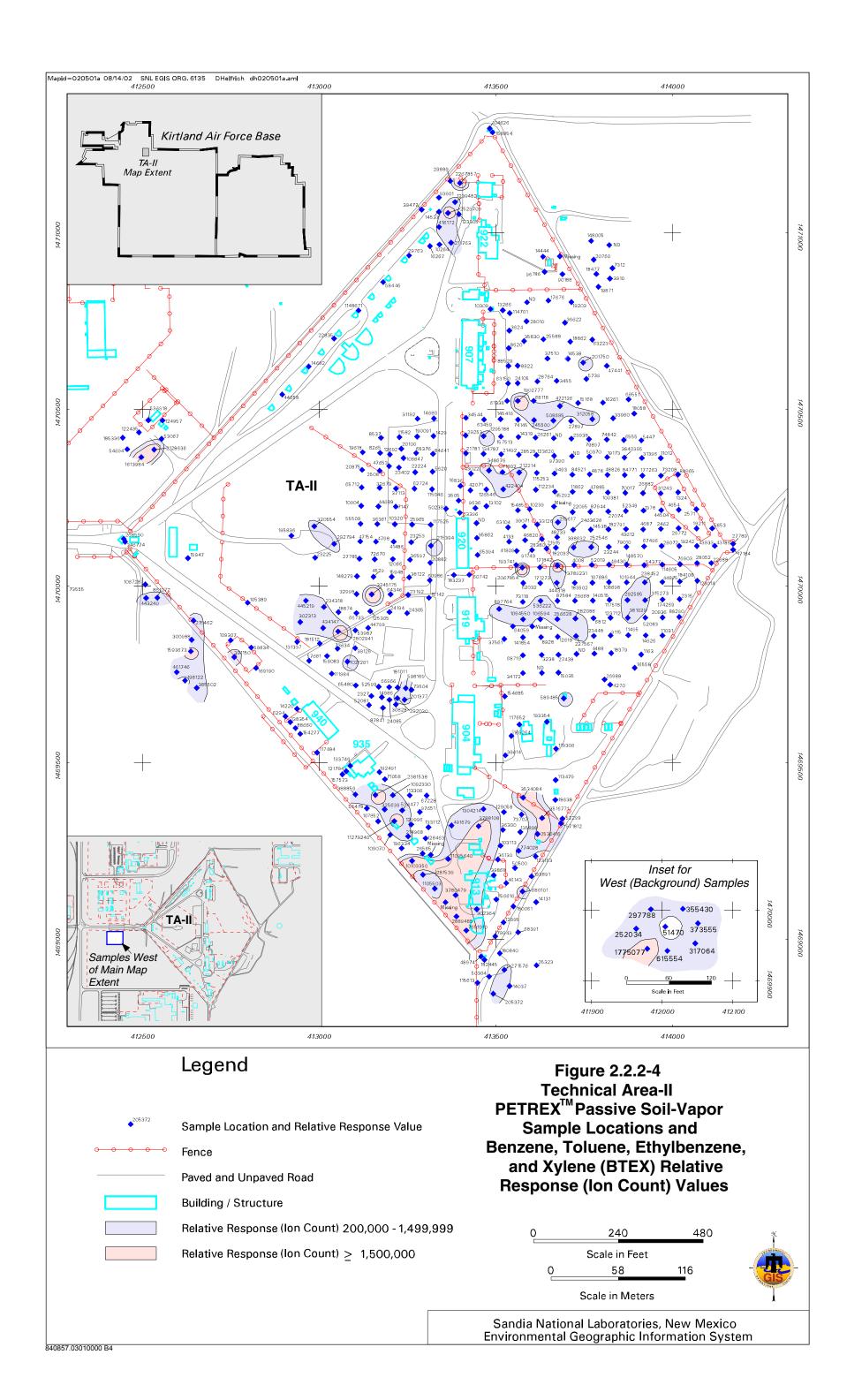
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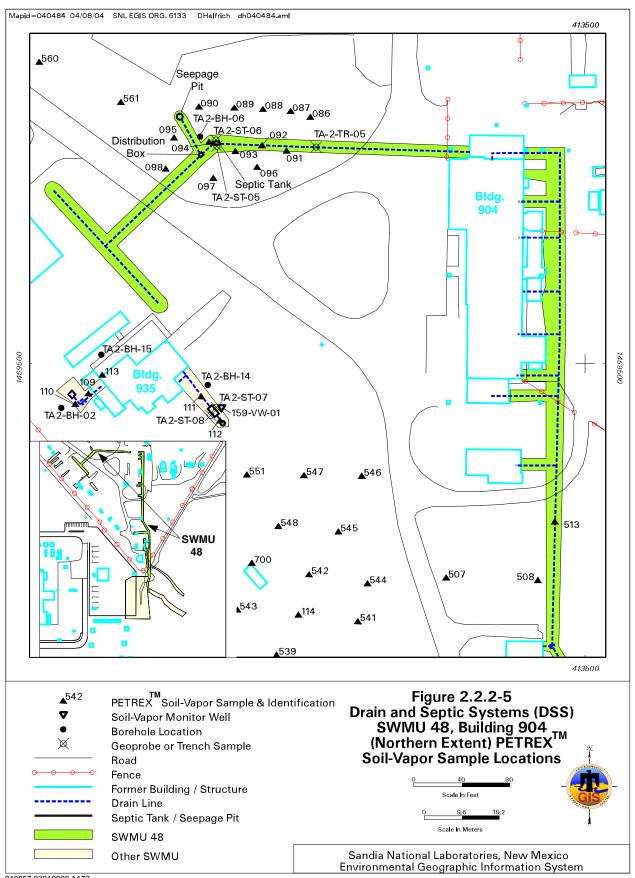


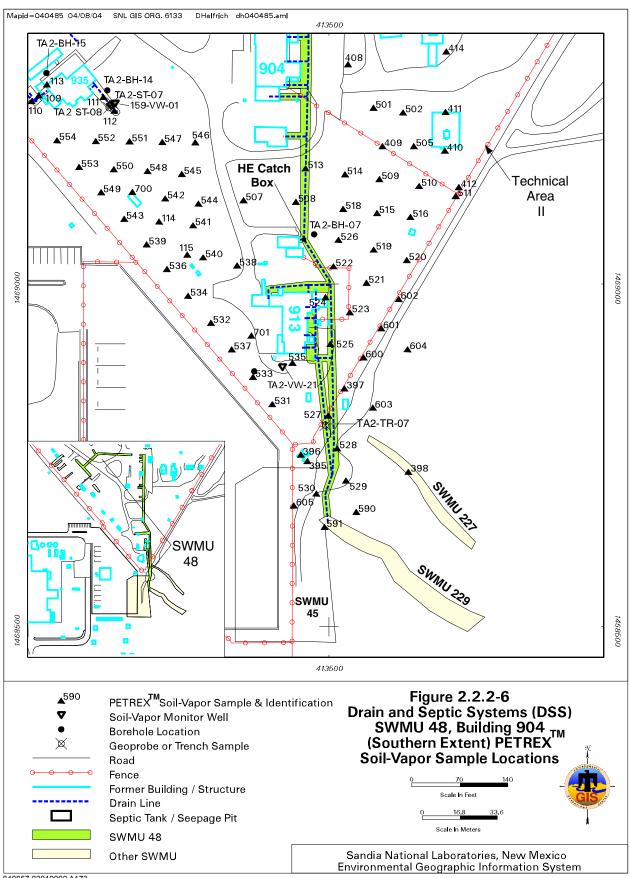


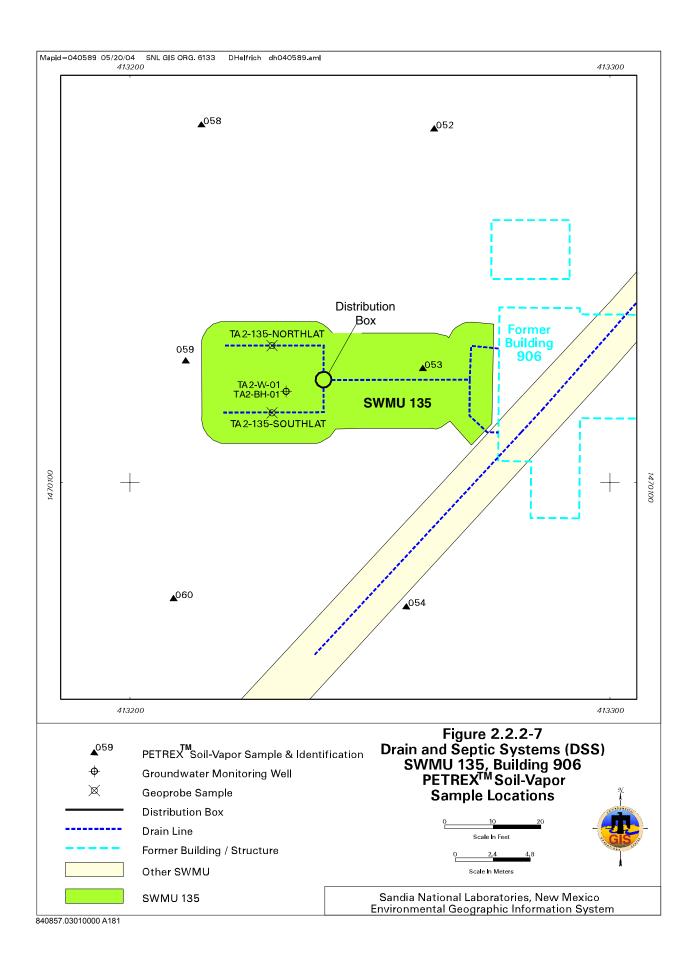


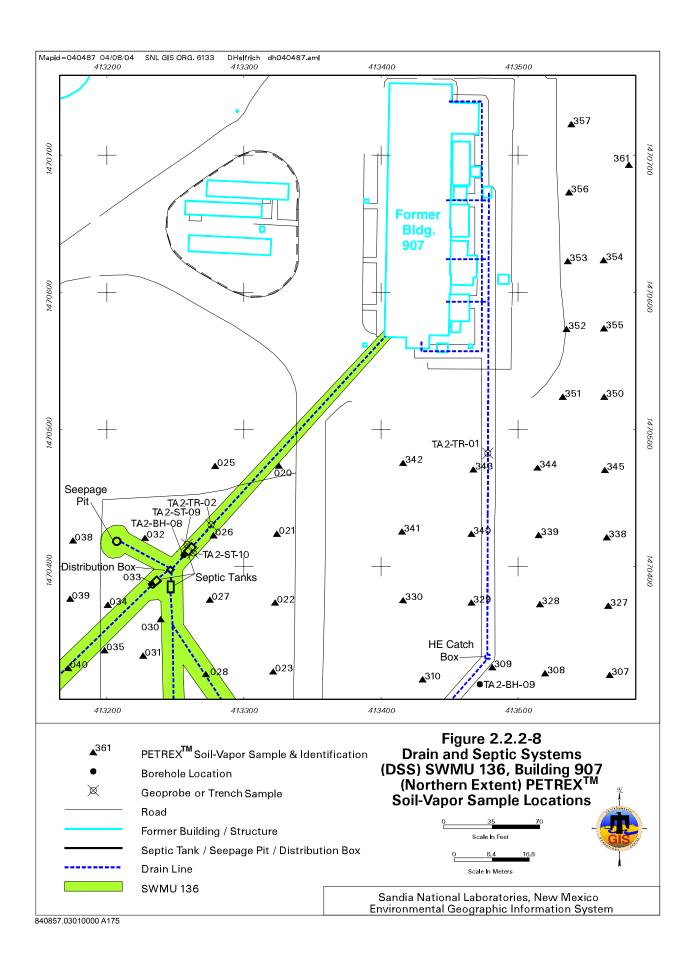


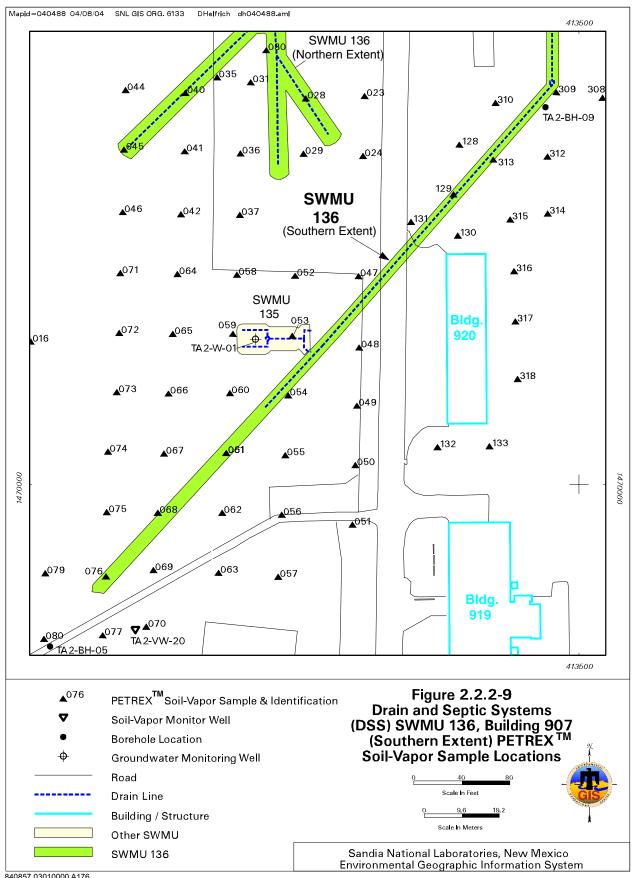


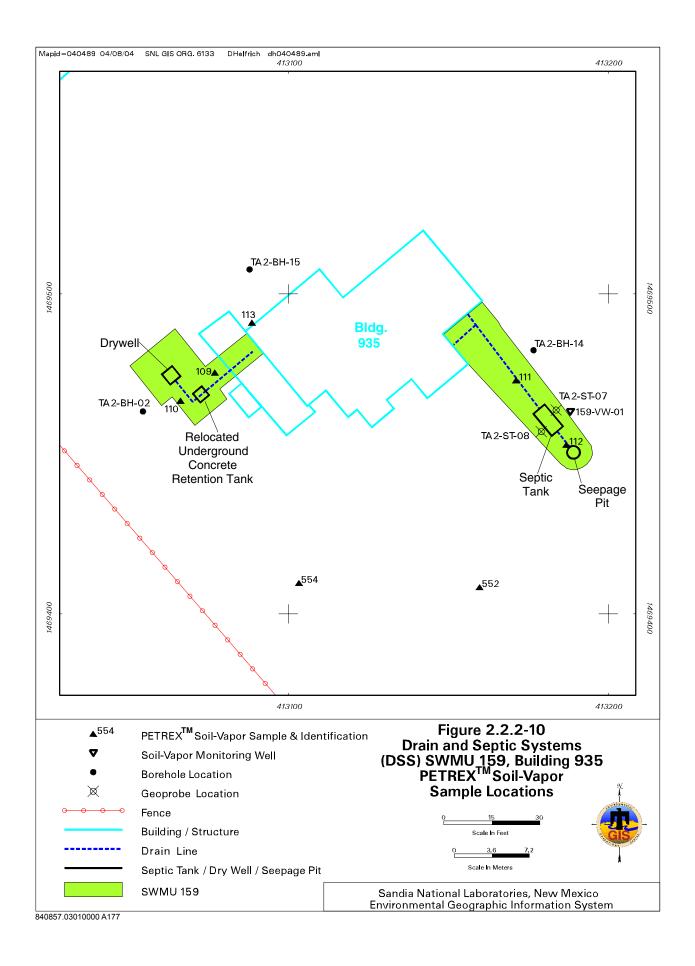


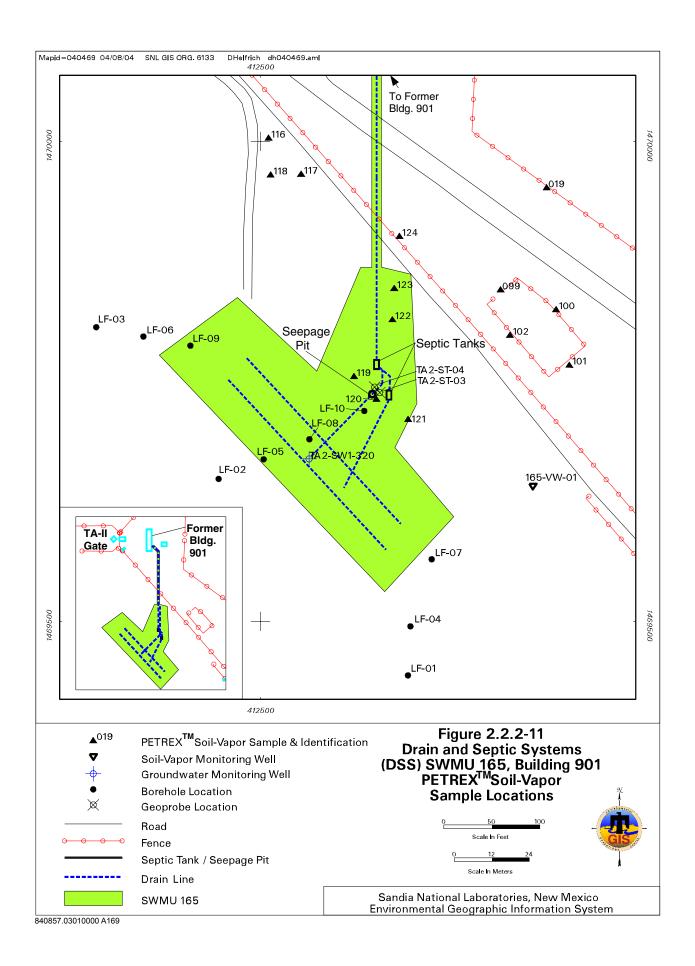


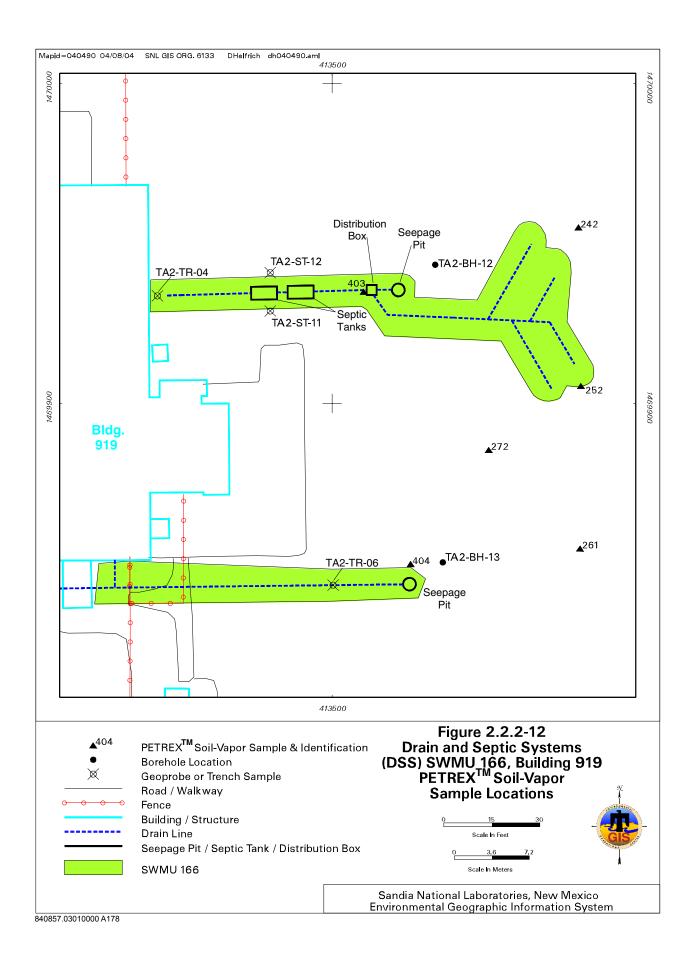


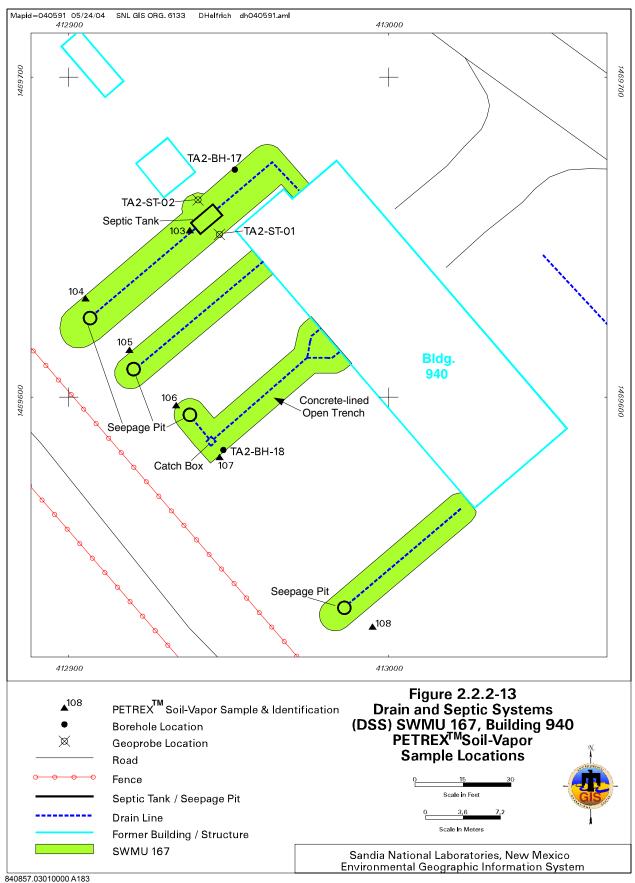


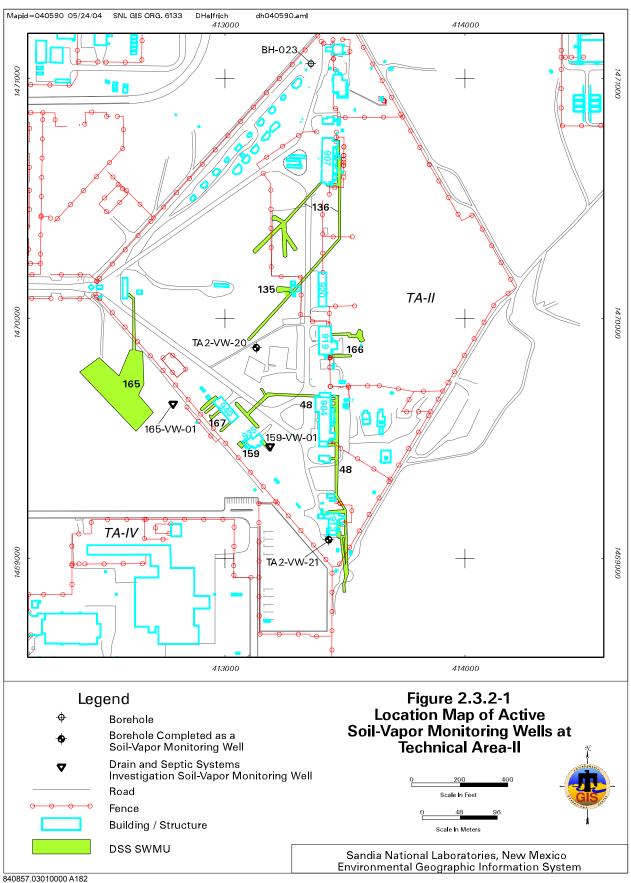


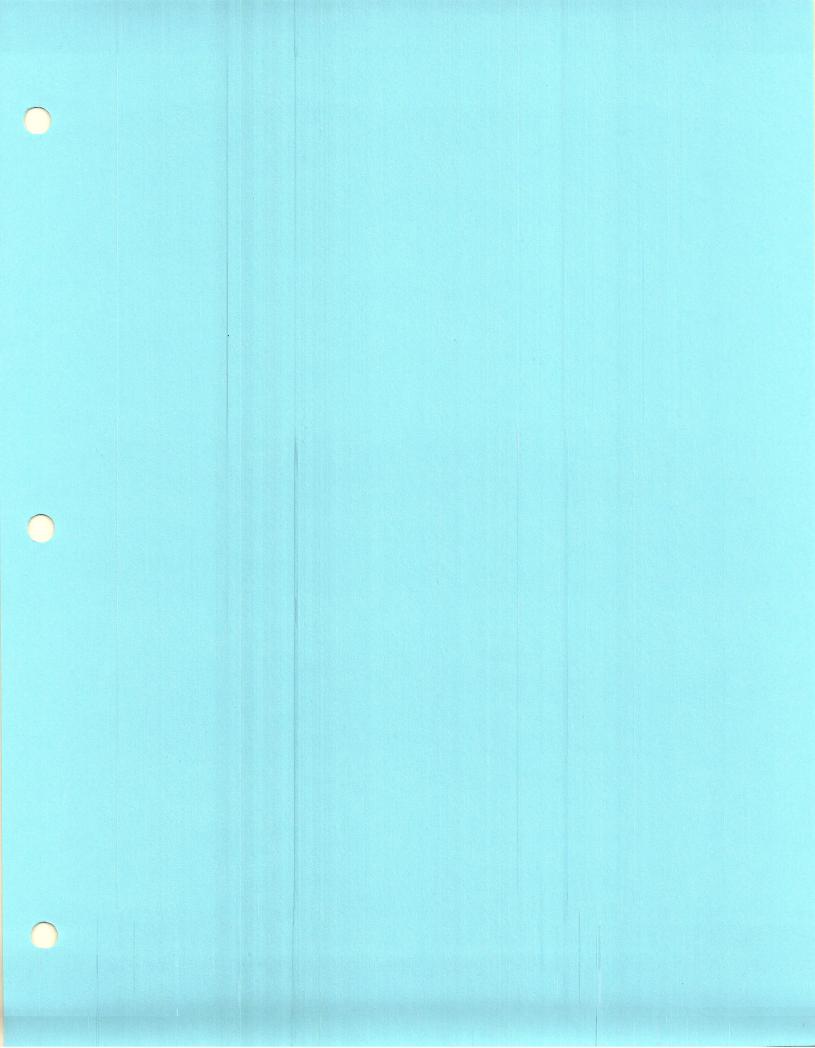












ANNEX A Soil-Vapor Analytical Data Summary Tables

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Table A-1
Summary of Technical Area II PETREX™ Passive Soil-Vapor Sampling
VOC Analytical Results
Phase I (November–December 1993) and Phase II (January–February 1994)

Sample	PETREX	™ Response Values	(ion counts)
Number	TCE	PCE	BTEX
		December 1993)	
001	ND	3577	44658
002	ND	2513	14682
003	ND	5903	22835
004	ND	11343	1148671
005	ND	5657	58446
006	ND	14927	29763
007	10691	ND	93901
008	ND	2398	1239482
009	ND	ND	14537
010	53938	53257	2529709
011	35882	7355	39477
012	18064	4272	416172
013	23476	1264	123305
014	8721	3335	195836
015	27922	ND	320554
016	56732	21982	292794
017	ND	ND	23225
018	ND	ND	41953
019	11560	13597	105389
020	ND	1013	14680
021	ND	ND	1429
022	ND	ND	84641
023	ND	ND	5620
024	60310	4746	115046
025	ND	1151	31192
026	23360	8469	190091
027	23330	3930	68376
028	6264	ND	22224
029	51537	2645	62724
030	ND	ND	106647
031	ND	ND	23402
032	ND	774	11582
033	11611	ND	20100
034	ND	ND	12692
035	4164	5518	47653
036	ND	ND	37113
037	11928	1025	7147
037-DU	17256	3829	11332
038	ND	4640	8537
038-DU	ND	3878	15434
039	14224	ND	8265
040	ND	ND	2508
041	20616	ND	32679

Phase I (November-December 1993) and Phase II (January-February 1994)

Sample	PETREX	™ Response Values	(ion counts)
Number	TCE	PCE	BTEX
042	100987	4364	44688
043	27066	2170	19618
044	ND	ND	20875
045	54398	1211	65712
046	38002	848	10004
047	28716	ND	117525
048	12995	ND	215394
049	19327	ND	20882
050	196370	3699	60986
051	62577	ND	15142
052	75160	5966	25965
053	31699	ND	23253
054	37827	1204	36597
055	ND	ND	38122
056	66940	891	23192
057	42688	ND	24305
058	35382	794	10320
059	46468	1484	41486
060	ND	ND	12066
061 <sup>a</sup>	22986	ND	15948
062	29942	7284	64346
063	ND	ND	24194
063-DU	ND	ND	31862
064	51440	2281	36361
065	52630	ND	4798
066	49956	17667	72670
067	1024	ND	4529
068	45438	88403	3245175
069	6306	5133	125305
070	3353	3010	44799
071	68289	6374	55509
072	44669	2376	47154
073	11849	ND	27785
074	11804	866	148279
074-DU	24355	ND	377027
075	19738	2184	32998
076	ND	3556	66733
077	ND	2198	35987
078	ND	ND	39125
078-DU	ND	ND	90355
079	ND	838	18874
080	8460	110697	2802941
082	16210	30572	1021261
083	8698	21606	434147

Table A-1 (Continued)

Summary of Technical Area II PETREX™ Passive Soil-Vapor Sampling

VOC Analytical Results

Phase I (November-December 1993) and Phase II (January-February 1994)

Sample	PETREX	™ Response Values	(ion counts)
Number	TCE	PCE	BTEX
084	5066	20179	181512
085	ND	6231	156083
086	ND	ND	73504
087	16300	3970	598169
088	1095	7881	181011
089	ND	ND	66966
090	ND	ND	52599
091	ND	7288	201377
092	ND	14119	292030
093	ND	ND	24085
093-DU	ND	ND	59981
094	ND	ND	14986
094-DU	ND	ND	3888
095	ND	ND	2327
095-DU	ND	ND	6303
096	ND	ND	30828
097	ND	3288	87841
098	ND	ND ND	52081
099	ND	10525	109307
100	ND	2268	59834
101	ND	11023	169190
102	ND	7523	391150
103	ND	13560	14220
103-DU	ND	30270	35014
104	ND	6280	6294
104-DU	ND	6622	9747
105	ND	11028	138354
105-DU	ND	2599	27159
106	ND	ND	88660
107	ND	2178	154277
108	ND	949	117484
109	ND	5964	121794
110	ND	3063	157573
111	16849	16478	192491
112	17865	891	71058
113	ND	4403	133746
114	752157	11476	214968
115	45571	1677	26565
116	ND	ND	106728
116-DU	ND	ND	168846
117	13431	13030	861072
118	3326	23881	443240
119	3554	16402	461746
120	5307	7600	198122

Phase I (November-December 1993) and Phase II (January-February 1994)

Sample	PETREX™ Response Values (ion counts)		
Number	TCE	PCE	BTEX
121	16314	16229	385502
122	13294	19795	1593673
123	ND	4400	300988
124	ND	2125	215462
125	ND	ND	15947
126	33440	13399	346774
127	ND	12798	268590
127-DU	ND	7324	179772
128	3459	969	16836
129	ND	ND	3505
130	13091	4360	63396
131	11264	ND	50239
132	51331	4058	183237
133	52753	5235	60742
180	ND	699	16063
200	10045	4918	43186
201	ND	13907	43933
201-DU	ND	3400	15394
202	ND	2425	19242
202-DU	ND	2600	15505
203	21577	10013	29072
204	28050	18951	67496
205	18193	16281	79010
206	ND	2240	23244
207	ND	14936	252546
208	31372	36318	398632
209-DU	ND	1020	48050
209	ND	7718	182033
210	ND	887	2688
211	ND	952	28052
212	ND	13299	76903
213	48200	58209	114605
214	ND	14743	54377
215	ND	37735	148570
215-DU	NDND	15626	57474
216	ND	16632	49430
217	ND	16151	52019
218	ND	ND	4008
219	5781	51333	3789231
220	ND	12715	171842
221	51973	32960	2067854
222	ND	6744	123014
223	ND	4960	184108
224	ND	43150	44875

Table A-1 (Continued)
Summary of Technical Area II PETREX™ Passive Soil-Vapor Sampling VOC Analytical Results

Phase I (November-December 1993) and Phase II (January-February 1994)

Sample	PETREX	™ Response Values	(ion counts)
Number	TCE	PCE	BTEX
225	ND	53212	238452
226	ND	276903	101944
227	85340	683858	108698
228	ND	67087	107686
229	408421	268917	183502
230	211836	183944	444318
231	ND	11250	171278
232	13533	55079	62032
233	ND	718	2316
234	ND	315158	174269
235	ND ND	1115030	315273
236	ND	714931	292585
237	ND ND	51774	117518
238	14158	9630	140515
239	ND	14728	26668
240	71095	135052	87584
241	5773	21038	
242	4533		536222
242	4555 ND	21383	73118
243		ND 0001	88290
	ND ND	3631	20936
245	ND	11896	52083
246	164190	252313	381029
247	93126	51130	123712
248	44567	18100	6812
249	19099	62389	282088
250	40519	74903	256628
251	ND	13632	106594
252	ND	23231	1094550
253	ND	2041	11031
254	ND	ND	14626
255	ND	3838	11495
256	ND	ND	9115
257	1282	6418	23448
258	ND	24806	227567
259	ND	ND	12618
260	ND	1019	8928
261	ND	34003	141854
261-DU	ND	10041	95682
263	ND	3156	18379
264	ND	ND	4488
265	ND	ND	ND
266	ND	2750	27438
267	ND	ND	3238
268	12966	10148	68719

Phase I (November-December 1993) and Phase II (January-February 1994)

Sample	PETREX	™ Response Values	(ion counts)
Number	TCE	PCE	BTEX
269	ND	3347	15035
270	ND	ND	ND
271	ND	780	34172
272	ND	808	64059
273	ND	ND	5853
274	3774	6100	19211
275	ND	ND ND	26772
276	ND	ND	2462
277	ND	ND	4687
278	ND	2870	43612
279	17462	750678	182791
280	ND ND	32941	14538
281 <sup>a</sup>	ND ND	6874	2463628
282	ND	ND	2571
283	ND	ND	4654
284	ND ND	7676	44504
285	ND	ND	1378
286	6263	1091	52349
287	ND	ND	27074
288	2148	13836	
			87944
289	ND ND	ND	22095
290	ND ND	ND ND	1324
290-DU	ND	ND ND	6664
291	ND ND	ND 10150	31243
292	ND ND	10156	26882
292-DU	ND	7440	18285
293	ND ND	9495	70017
294	ND ND	15406	100381
295	ND 47000	ND F000	47885
296	17226	5326	11862
296-DU	17964	10405	13166
297	ND ND	7594	45292
298	ND ND	6204	84965
299	ND ND	17217	73208
300	ND ND	22912	177263
301	ND ND	8395	84771
302	ND ND	2452	48826
303	ND	ND 0.10=	4878
304	ND	9197	84921
305	ND	ND	9493
305-DU	ND	ND	921
306	4973	6476	115253
307	6076	13073	212214
308	ND	ND ND	71802

Table A-1 (Continued)

Summary of Technical Area II PETREX™ Passive Soil-Vapor Sampling

VOC Analytical Results

Phase I (November-December 1993) and Phase II (January-February 1994)

Sample	PETREX	™ Response Values	(ion counts)
Number	TCE	PCE	BTÉX
309	3717	15269	348676
310	ND	ND	43022
311	15332	1652	422404
312	ND	ND	126546
313	ND	ND	42071
314	ND	ND	13102
315	ND	5422	9636
316	ND	ND ND	ND
317	ND	12282	95862
318	ND	ND ND	65394
319	ND	1139	11012
320	ND	5644	31396
320-DU	ND	926	16657
321 <sup>b</sup>	31144	18434	3840395
322	ND ND	ND	19173
323	ND	12605	50370
324	ND ND	ND	ND
324-DU	ND	ND ND	ND ND
325	ND	7465	97390
325-DU	ND		
325-00	ND	3465	63036
327	ND ND	9283	123620
	ND ND	9363	28528
328		3138	21492
329	ND	2750	134787
330	ND	ND ND	21781
331	ND ND	967	5447
332	ND	2569	6555
333	ND ND	8732	74842
333-DU	ND	22418	163380
334	ND	8414	79897
335	ND ND	5908	23938
336	ND ND	ND ND	ND ND
336-DU	ND	ND	ND
337	ND	3544	26261 -
338	ND	3139	14319
339	ND ND	78088	157513
340	25405	20021	1295188
341	ND	ND	29253
342	ND	ND	34544
343	ND	ND	63459
344	7731	8624	145416
345	ND	7832	74145
346	13300	8972	245500
347	ND	4190	508695

Phase I (November-December 1993) and Phase II (January-February 1994)

Sample	PETREX	™ Response Values	(ion counts)
Number	TCE	PCE	BTEX
347-DU	ND	6553	703828
348	ND	823	27897
349	ND	12101	68118
350	17873	26138	1902777
351	ND	ND	61338
352	ND	ND	63150
353	ND	11828	88529
354	7297	7047	8922
355	ND	5744	24105
356	ND	ND	8620
357	ND	954	3624
358	ND	15296	114701
359	ND	ND	13266
360	ND	40738	10309
361	ND	6838	35830
362	ND	ND	28010
363	ND	ND	ND
363-DU	ND	ND	ND
364	ND	2178	17676
365	ND	ND ND	36622
366	ND	ND	25588
367	ND	ND	37510
368	ND	9600	28764
369	ND	673	472126
369-DU	ND	650	404020
370	ND	ND	3455
371	ND	ND	14538
372	ND	ND	18862
373	ND	3249	19203
374	ND	4680	63223
375	ND	7098	201750
376	ND	ND	5739
376-DU	ND	ND	2554
377	1207	5869	15168
378	ND	16586	312058
379	ND	ND	33960
380	ND	ND	18058
381	2146	8253	69551
382	ND	8164	16261
383	ND	2532	47441
384	13375	5313	158854
385	ND	2080	104626
385-DU	ND	5144	116846
386	ND	6425	148005

# Table A-1 (Continued) Summary of Technical Area II PETREX™ Passive Soil-Vapor Sampling VOC Analytical Results Phase I (November–December 1993) and Phase II (January–February 1994)

Sample	DETREY	™ Response Values	(ion counte)
Number	TCE	PCE	
			BTEX ND
387	ND ND	ND ND	
387-DU	ND	ND	ND 00700
388	ND	ND	20760
389	ND	ND	7312
390	ND	ND	18477
391	ND	ND	3910
392	ND	ND	19871
393	3296	ND	27789
394	ND	ND	17184
395	112632	2450	182445
396	113649	4630	48974
397	ND	ND	12995
398	ND	ND	25323
399	ND	ND	1163
400	ND	ND	16558
401	ND	ND	26988
402	ND	ND	4270
403	ND	3020	897764
404	ND	3458	37561
405	ND	ND	117652
405-DU	ND	ND	59474
406	ND	3079	154685
407	ND	2858	168264
408	ND	ND ND	39614
408-DU	ND	ND	1618
409	42683	103544	3534084
410	ND	ND	18638
410-DU	ND	1943	26259
411	ND	ND	113479
412	10840	2761	52299
413	ND	ND ND	193354
414	ND	ND ND	119306
415	11019	1324	589485
1000			
<del> </del>	ND ND	ND ND	122435
1001	ND 7010	ND 1709	54694
1002	7913	1738	1613984
1003	16596	13997	2128636
1004	ND ND	ND	53067
1005	ND	ND 45000	124957
1006	24918	15289	536918
1007	ND	4627	185336
1010	ND	ND	51470
1011	30155	11080	615554
1012	61218	12577	1775077

# Table A-1 (Continued) Summary of Technical Area II PETREX™ Passive Soil-Vapor Sampling VOC Analytical Results Phase I (November–December 1993) and Phase II (January–February 1994)

Sample	PETREX	™ Response Values	(ion counts)
Number	TCE	PCE	BTEX
1013	74497	4110	252034
1014	99204	8949	297788
1015	24692	ND	355430
1016	44196	5342	373555
1017	63678	5964	317064
2000	ND	15002	143273
	mples (January-F		140270
42	7150	ND ND	73181
60	ND	ND	12066
66	140502	11934	18071
68	ND	ND	8666
80	ND	932	83164
81	ND	7299	73634
216	ND	ND ND	62478
227	84801	198874	24686
229°	178643	29509	40857
234	2585	11950	76896
262	4178	21257	192011
321	ND	ND ND	24278
339	22976	5489	200428
279	ND ND	ND	26977
507	ND	3536	491679
508	ND	ND	3788108
509	95126	662571	73707
510	81153	7215	3516771
511	15841	29656	971812
513	3401	13753	1304214
514	58255	ND	129058
515	23108	2145	135898
516	325191	10656	2536416
518	35349	ND	36300
519	1788	10379	774028
520	167686	8695	123153
521	26037	4349	50500
522	131504	13023	36130
523	72372	ND ND	46143
524	40723	3421	39868
525	128296	15164	156616
526	105854	12672	103113
527	41948	7470	79943
528	34886	9395	180840
529	16701	24668	271567
530	1047	4358	50304
531	662570	15156	7651910
	002070		7001010

Phase I (November-December 1993) and Phase II (January-February 1994)

Sample	PETREX	™ Response Values	(ion counts)
Number	TCE	PCE	BTEX
532	20273	2038	1105503
533	1803940	25590	2889485
534	10023	ND	281539
535	1128790	75264	302364
536	ND	2953	10103350
538	132940	25831	11095640
539	1340	11236	196334
541	280184	7298	128463
542	375732	6039	120996
543	ND	ND	109070
544	116462	1263	133112
545	575741	ND	97651
546	32925	1423	67228
547	779378	18055	113306
548	119104	13719	536477
549	27485	16874	107852
550	20750	ND	325639
551	128182	15577	1092330
552	8048	14425	2381538
553	15545	2170	56478
554	ND	1228	388859
555	ND	6278	234318
556	7026	8195	445219
557	3132	2250	302313
558	ND	891	131337
559	ND	1754	57661
560	ND	31942	111984
561	ND	2032	65480
562	ND	ND	112234
563	9934	10451	15485
564	ND	ND	10239
566	ND	8464	219617
567	ND	ND _	33126
568	ND	ND	39071
569	ND	ND	63104
570	ND	ND	4133
571	ND	ND	46820
572	ND	ND ND	21915
573	ND	ND	48233
574	ND	1845	28380
575	ND	ND ND	91749
576	ND	2065	41809
577	ND	ND	193741
581	286483	64632	2267857

Phase I (November-December 1993) and Phase II (January-February 1994)

Sample	PETREX	™ Response Values	(ion counts)
Number	TCE	PCE	BTEX
582	32981	ND	29889
583	69875	ND	211763
584	11304	ND	10264
585	5779	ND	16267
586	MISSING	MISSING	MISSING
587	ND	ND	90188
590	ND	ND	10437
591	12134	16162	205372
581	286483	64632	2267857
600	26652	3211	180061
601	123694	5955	680101
602	39805	1867	133891
603	31114	3730	68391
604	ND	ND	14131
605	3782	7405	115613
700	50042	15160	11279240
701	131355	9089	3780479
908	ND	ND	2494249
910	143838	9293	1755741
914	61556	ND	152262
916	378129	11156	2248256
922	88777	13378	25666
926	55521	1945	48394
929	18789	17065	291392
962	ND	ND	125956
968	ND	ND	24803
972	ND	ND	9635
	urance/Quality Co	ontrol Samples	
TB 2001	ND	ND	78838
TB 2002	ND	ND	3153
TB 2003	ND	ND	ND
TB 2004	ND	ND	ND

<sup>&</sup>lt;sup>a</sup>Value elevated due to interference with terpene compounds.

BTEX = Benzene, toluene, ethylbenzene, and xylene(s).

DU = Duplicate analysis. In laboratory reports, these samples are prefixed with a "3" before the sample number.

ND = Not detected above the PETREX™ background value.

PCE = Tetracholoroethene.

TB = Trip blank.

TCE = Trichloroethene.

VOC = Volatile organic compound.

<sup>&</sup>lt;sup>b</sup>Sampler integrity compromised; value may be elevated due to incidental cross-contamination.

<sup>&</sup>lt;sup>c</sup>Sampler exposed approximately 10 days longer than the remaining data set.

Table A-2
Summary of Technical Area II PETREX™ Passive Soil-Vapor Sampling
Additional VOCs Detected

Phase I (November-December 1993) and Phase II (January-February 1994)

Sample	PETI	REX™ Respons	e Values (ion co	unts)
Number	TCA	DCB	Freon-11	Freon-113
Phase I S	amples (Noven	nber-December	1993)	
5	ND	221746	ND	ND
10	ND	415426	ND	ND
68	ND	704865	ND	ND
80	ND	886514	ND	ND
83	ND	1742220	ND	ND
85	ND	271140	ND	ND
216	ND	ND	100532	ND
219	ND	2122370	ND	ND
225	ND	ND	227552	137744
226	ND	ND	558425	478299
227	ND	ND	204234	88984
228	ND	ND	408375	284606
231	ND	ND	554822	ND
235	ND	ND	379641	274423
236	ND	ND	500416	447926
238	ND	ND	600607	510369
239	ND	ND	222725	111590
240	ND	ND	146644	ND
246	ND	ND	174304	135227
247	109033	ND	ND	ND
251	ND	212469	ND	ND
346	ND	470719	ND	ND
406	ND	161433	ND	ND
409	ND	391198	ND	ND
Phase II	Samples (Janua	ary-February 19	94)	
531	ND	ND	247990	ND
801 <sup>a</sup>	ND	ND	183453	ND

<sup>a</sup>No known sample point with this designation, possibly referring to Sample 81.

DCB = Dichlorobenzene.

Freon-11 = Trichlorofluoromethane. Freon-113 = Trichlorotrifluoroethane.

ND = Not detected above the PETREX™ background value.

TCA = Trichloroethane.

VOC = Volatile organic compound.

Table A-3
Summary of Technical Area II Active Soil-Vapor Monitoring Well VW-20 Sampling
Soil-Vapor VOC Analytical Results
November 1996 to March 2002
(On- and Off-Site Laboratories)

Ouanterra 600796 Ouanterra 600796 Ouanterra 601237 Ouanterra 601237 Ouanterra 601237 Ouanterra 601638 Ouanterra 601638 Ouanterra 601623 Ouanterra 601623 Ouanterra 601623 Ouanterra 601623 Ouanterra 601623 Ouanterra 601623 Ouanterra 601623	ERCL 6173 TA2-VW-20-72-P ERCL 6173 TA2-VW-20-72-P ERCL 6942 TA2-VW-20-72-P ERCL 6942 TA2-VW-20-72-P ERCL 6942 TA2-VW-20-72-P ERCL 6942 TA2-VW-20-72-DU Gore TA2-VW-20-72 600308 Ouanierra TA2-VW-20-72 600796	Soll-vapor s ATL 9267 ATL 9267 ATL 9267 ATL 9268 ATL 9268 ATL 9268 ATL 9268 ATL 9268 ATL 9268 ATL 9268 ATL 9268 ATL 9268 ATL 9268	Sample Attributes  Sample Attributes  Laboratory and Record Aumber Sample Sampl
TA2-W-20-72-DU TA2-W-20-72-DU TA2-W-20-72-DU TA2-W-20-72-DU TA2-W-20-72-DU TA2-W-20-72-DU TA2-W-20-72-DU TA2-W-20-72-DU	TA2-VW-20-72-P TA2-VW-20-72-P TA2-VW-20-72-P TA2-VW-20-72-P TA2-VW-20-72-P TA2-VW-20-72 TA2-VW-20-72	TA2-BH-020-10-SV TA2-BH-020-20-SV TA2-BH-020-20-SV TA2-BH-020-40-SV TA2-BH-020-40-SV TA2-BH-020-60-SV TA2-BH-020-60-SV TA2-BH-020-60-SV TA2-BH-020-70-SV TA2-BH-020-90-SV TA2-BH-020-100-SV	ED.
20-72-DU 20-72-DU 20-72-DU 20-72-DU 20-72-DU 20-72-DU 20-72-DU	20-72-P 20-72-P 20-72-P 20-72-S 20-72-P 20-72-S 20-72-DU	mples collected during TA2-BH-020-10-SV TA2-BH-020-10-SV TA2-BH-020-30-SV TA2-BH-020-40-SV-DU TA2-BH-020-40-SV-DU TA2-BH-020-60-SV TA2-BH-020-60-SV TA2-BH-020-70-SV TA2-BH-020-90-SV TA2-BH-020-90-SV TA2-BH-020-100-SV-DU	Sample Attributes
	N)	DU DOL	ributes S.
72 72 72 72 72 72 72 72 72 72 72 72 72 7		hole drill 10 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	ample spth (tt)
12-02-98 12-02-98 03-22-99 03-22-99 06-24-99 06-24-99 09-07-99	07-17-97 07-17-97 09-04-97 09-04-97 09-04-98 05-14-98 05-14-98		ample Date
ND (2) (0.0 ND (2)	16 (1.0) ND	130 130 320 150 150 170 170 170 170 170 170 170 170	Acetone
(0.8) (0.8) (0.8)		21 ND 6.8 (7.0) 6.8 (ND 6.2) 7.7 ND 7.7 ND 7.7 ND 7.8 (6.2) 7.7 ND 7.9 (4.1) 31 ND 31 (3.7) 46 ND 31 (3.7) 47 ND (3.7) 48 ND (3.7) 41 ND (12) 32 ND (23) 86 ND (13) 94 ND 94 ND	Benzene
(0.27) (0.08) (0		21 ND 13 (7.0) 7.9 8 ND 7.9 8 (6.2) 36 (6.2) 38 (6.2) 38 (7.7) ND 9.4 11 ND 9.4 (3.7) 29 (3.7) 20 (3.7) 20 (3.5) 42 (3.7) 20 (3.7) 20 (3.5) 42 (3.7) 42	Bromodichloromethane
(0.57) (3.5) ND (1) ND (2) ND (1) ND (2) ND (1) ND (2) ND (1) ND (2) ND (1) ND (2) ND (1) 3 J (10) ND (1) ND (2) ND (1) ND (2)		13 7.9 7.9 9.4 9.4 9.4 9.4 9.6 9.6 9.6	1,3-Butadiene
	NR (1.2) NR (1.2) NR (1.2) NR (1.25) NR (1.25) NR (1.25) NR (1.25) NR (1.25) NR (1.25) NR (1.25) NR (1.25) NR (1.25) NR (1.25) NR (1.25)	28 ND 12 (7.0) 12 (4.4) 38 ND 14 (4.9) 14 (4.9) 14 (4.9) 24 ND 15 5. 50 6. 50 6. 62 1	2-Butanone Carbon disulfide
	D ND 2) (12) 0 ND 0 ND 0 ND 0 ND 0 ND 0 ND 0 ND 0 ND	9 8 9 5 8 8	Carbon tetrachloride
	(1.0 (1.0 (1.0 (1.0 (1.0 (1.0 (1.0 (1.0		Chloroform
(1.5) 5.8 (0.8) 5.4 (0.8) 6.8 (0.8) 7.7 (0.8) 5.8 (0.8) 5.9 (0.8) 5.9 (0.8) 5.9 (0.8) 6 (0.8) 6 (0.8)		9) (1.8) (1.9) (1.8) (1.9) (1.8) (1.9) (1.9) (1.9) (1.6) (1.	Chloroethane
	(1) (1,0) (1	0 ND 0 ND 0 ND 0 ND 0 ND 0 ND 0 ND 0 ND	Chloromethane
	NA NA NA NA NA	ND (13.7)	Cyclohexane
0 2 0 2 0 2 0 2 0 2 0 2 0 2 0 2 0 2 0 2	NA NA NA NA NA NA NA NA NA NA NA NA NA N	ND (1.8) ND (1.8) ND (1.8) ND (1.9) ND	1,2-Dibromoethane
(2) (0.8) (0.8) (0.8) (0.8) (0.8) (0.8) (0.8) (0.8) (0.8) (0.8) (0.8)	7 (1.0 NA NA NA NA NA NA NA NA NA NA NA NA NA	(1.8) ND (1.8) ND (1.8) ND (1.9) ND (1.9) ND (1.2)	1,2-Dichlorobenzene
0 1 0 1 0 1 0 1 0 1 0 1 0 1	NA NA NA NA NA NA NA NA NA NA NA NA NA N	(1.8) (1.8) (1.18) (1.19) (1.10) (1.1	1,3-Dichlorobenzene
(0.8) (0.8) (0.8) (0.8) (0.8) (0.8) (0.8) (0.8)		(1.8) (1.8) (1.8) (1.6) (1.2) (1.2) (1.2) (1.2) (1.2) (1.2) (1.2) (1.2) (1.2) (1.2) (1.2) (1.2) (1.2) (1.3) (1.3) (1.4) (1.4) (1.5) (1.5) (1.6)	1,4-Dichlorobenzene
20 1.5.1 1.5.1 1.5.2 21 1.5.2 22 1.6 23 1.6 24 1.6 25 1.6 26 1.6 27 1.6 28 1.6 29 1.6 20 1	0 0 Z Z Z Z Z Z Z	ND (1.8) (1.8) (1.8) (1.8) (1.6) (1.	Dichlorodifluoromethane
22 (0.5) 23 (0.5) 24 (0.5) 25 (0.5) 26 (0.5) 27 (0.5) 28 (0.5) 29 (0.5) 20 (0.5) 20 (0.5) 20 (0.5) 21 (0.5) 22 (0.5) 23 (0.5)		(1.8) NO (1.8) NO (1.8) NO (1.9) NO (1.	1,1-Dichloroethene
(0.54) (0.80) (0.80) (0.80) (0.80) (0.80) (0.80) (0.80) (0.80) (0.80) (0.80) (0.80) (0.80) (0.80) (0.80) (0.80) (0.80) (0.80)	<del></del>	(1.8) (1.8) (1.9) (1.2)	cis-1,2-Dichloroethene (F)
N N N N N N N N	NA NA NA NA	36 21 ND (6.2) 115 118 1430 2 430 177 55 55	Ethanol Ethyl benzene (FPA Method 8260 M3.
(0.5) (0.5) (0.5) (0.5) (0.5) (0.5) (0.5) (0.5) (0.5) (0.5) (0.5)	(1.2) (1.2) (1.2) (1.2) (1.2) (1.25) (1.25) (1.25) (1.25) (1.25) (1.25) (1.25) (1.25)	(1) (1) (2) (4) (4) (6) (7) (8)	Ethyl benzene
(0.5) (0.5) (0.5) (0.5) (0.5) (0.5) (0.5) (0.5) (0.5) (0.5) (0.5) (0.5) (0.5) (0.5)	0.01.00 NR NR NR NR NR NR NR NR NR NR NR NR NR	1.6 ND (7.0) 12 (4.9) 1.7 ND 1.8 5.5 (4.9) 1.1 9.3 111 9.3 5.5 1226 1226 1226 1226 1226 123 136 8.8 J (18) 336 8.8 J (18) 8.3 8.8 J	4-Ethyltoluene
N N N N N N N N	X X X X X X	35 38 38 26 26 27 27 28 29 100 20 3 190	4-Ethyltoluene 0-14/TO-14/TO-14A)
N N N N N N N N N N N N N N N N N N N	NA NA NA NA NA NA NA NA NA NA NA NA NA N		Hexane Units
ND (3) ND	NO 9 NR NR NR NR NR NR NR NR NR NR NR NR NR	111 ND 6.4 ND 19 ND 19 ND 6.6 (4.4) 19 ND 6.6 (A.9) 6.6 (A.9) 10 ND 10 (3.7) 54 4.50 20 ND 20 ND 21 (3.5) 44ND (12) 31ND (23) 86ND (18)	2-Hexanone as a sind of a
(0.8) (0.8) (0.8) (0.8) (0.8) (0.8) (0.8) (0.8)	23 18 18 23 23 5.3 5.3 ND ND	(1.8) ND (1.1) ND (1.1) ND (1.2) ND (1.2) ND (1.2) ND (1.2) ND (1.2) ND (1.2) ND (1.2) ND (1.2) ND (1.3) ND (1.4) ND (1.4) ND (1.5) ND (1.	Methylene chloride
ND (2) ND (2) ND (2) ND (2) ND (2) ND (2) ND (2) ND (2)	NA NA NA NA NA NA NA NA NA NA NA NA NA N	(7.0) (7.0) ND (7.0) (7.0) (7.0) (7.0) ND ND (4.4) ND (6.2) (6.2) ND ND (4.9) ND (4.9) ND (3.7) (3.7) (3.7) ND (3.5) (3.5) ND (12) ND (12) ND (12) ND (12) ND (12) ND (12) ND (12) ND (12) ND (12) ND (13) ND	4-Methyl-2-pentanone
NA NA NA NA NA	H H H N N N N	(7.0) (1.4)	2-Propanol
NA NA NA NA NA NA NA NA NA NA NA NA NA N	N N N N N N N N N N N N N N N N N N N	ND (7.0) ND (4.4) ND (6.2) ND (4.9) ND (4.1) ND (3.7) ND (3.7) ND (3.5) ND (23) ND (23)	Propylene
(0.5) (0.5) (0.5) (0.5) (0.5) (0.5) (0.5)	(1.25) (1.25) (1.25) (1.25) (1.25) (1.25) (1.25) (1.25) (1.26) (1.27) (1.28)	2.7 1.2 2.9 (1.2) (1.0) (1.0) (1.0) (1.0) (2.5 2.2 2.2 2.2 2.3 (0.88) (0.88) (0.88) (0.88) (0.88) (0.88) (0.88) (0.88) (0.88)	Styrene
0.51 J (2) (0.5) (0.57 J (2) (2) (2) 3.6 (0.5) (0.5)	(1.2) ND (1.2) ND (1.25) ND (1.25) ND (1.25) ND (1.25) ND (1.25)	(1.6) (1.1) (1.1) (1.1) (1.1) (1.6) (1.2) (1.2) (1.2) (1.2) (1.2) (1.2) (1.2) (1.2) (1.2) (1.2) (1.2) (1.2) (1.2) (1.3) (1.3) (1.4) (1.4) (1.4) (1.4) (1.4) (1.5) (1.4)	Tetrachloroethene
ND (0.5) ND (0.5) ND (0.5) ND (0.5) ND (0.5) ND (0.5) 2.5 (2) 2.5 (2) 3.8 J	(4.8) ND (1.2) ND (1.25) ND (1.25) ND (1.25) ND (1.25) ND (1.25)	300 300 260 290 1150 150 280 280 280 280 100 100 100 100 100 100 100 100 100	Toluene
ND (3.) ND (3.) ND (3.) ND (3.) ND (3.)	NA NA NA NA NA NA NA NA NA NA NA NA NA N	(1.9) (1.1) (1.1) (1.1) (1.1) (1.1) (1.1) (1.1) (1.1) (1.1) (1.2) (1.0) (1	
(0.5) (0.5) (0.5) (0.5) (0.5) (0.5) (0.5) (0.5) (0.5) (0.5) (0.5) (0.5) (0.5) (0.5)	(1.25) (1.25) (1.25) (1.25) (1.25) (1.25) (1.25) (1.25) (1.25)		1,1,1-Trichloroethane
21 21 21 21 25 25 27 27 27 28 28 28 28 28 28 28 28 28 28 28 28 28	(1.2) (1.2) (1.2) (1.2) (1.25) (1.25) (1.25) 8.0 8.0	N N N N N N N N N N N N N N N N N N	Trichloroethene
3.1 4.4 5.3 9.9 4	(1.0) NR NR NR NR NR NR NR NR NR NR NR NR NR		
120 110 110 120 130 95	(1.2) 1.8 J 1.8 J (1.48) ND (1.25) ND (1.25) ND (1.25) 38 38	18 24 17 17 60 60 80 120 120 220 220	1,1,2-Trichloro- 1,2,2-trifluoroethane
	0.688 J		1,2,4-Trimethylbenzene
(0.8) (0.8) (0.6)	0.42.1 0.42.1	0 0 0 0 0 0 0 0 0	1,3,5-Trimethylbenzene
NA NA NA NA NA NA NA NA NA NA NA NA NA N	(1.2) (1.2) (1.2) (1.2) (1.25) (1.25)		o-Xylene
NA NA NA NA NA NA NA NA NA NA NA NA NA N	(9.5) (2.5) (2.5) (2.5) (2.5) (2.5) (2.5) (2.5) (2.5) (2.5) (2.5) (2.5)	32 29 30 30 30 30 118 18 18 58 58 72 20 20 20 119 140	m-, p-Xylene
NA NA NA NA NA (0.8)	(1.0) (1.0) (1.0) (1.0) (1.0) (1.0)		Total Xylene

# Table A-3 (Concluded) Summary of Technical Area II Active Soil-Vapor Monitoring Well VW-20 Sampling Soil-Vapor VOC Analytical Results November 1996 to March 2002 (On- and Off-Site Laboratories)

ATL 92	000	O/STL	Q/STL 605162	604921	OST	604643	60443	Q/ST	0/STL	603898	603661	O/ST	0/STL	0/STL 603340	603340	603136	0/21	Q/STL	Quanterra 602989	602989	Chante	and Record	Laborat	
ATL 9268 TA2-BH-020-000-FB 11	-	_			4		1	4		یــــا	1	↲			<u> </u>	<del>,</del>	4	_		<del></del>	<del>al</del>		Ϋ́	
H-020-00	5	TA2-VW-20-72	A2-VW-20-72		CZ-02-MV-CA	(A2-VW-20-72		72-VW-20-72	.42-VM-20-72-DU	HC-VVV-20-12	20 70	A2-VW-20-72-DU	A2-VW-20-72	A2-VW-020-72-DU	HZ-VVV-020-12	200	110-c2-0c-Wv-ca	A2-VW-20-72	TA2-VW-20-72-DU		42.VW-20-72	FR Same		Sampl
)0-FB									Ď			ġ		2-DU	"		2	- 1	É			ĕ		Sample Attributes
	-	72	72	_	72	- 2		72	72	,	4	72	72	72	1 2	3	73	72	72	-	70	Sample Depth (ft)		- 18
11-				ļ_	4		_	_		-	-	4			ļ.,	↓_	1		_	Ι.	٠.			-
11-15-96		3-19-02	2-11-01	-	09-25-01	06-22-01 N		04-19-01	2-11-00	+	+-	09-13-00	09-13-00	06-20-00	0000	3	03-01-00	03-01-00		┷	-1	Sample		$\perp$
16ND (1.0)	-	10ND (0.8)	(13)	(10)	4.2 JND (0.8)	ND (2) ND (0.0)	(10)	9 JND (0.8)	(10)	(10)	70	6.3 JND (0.8)	7.4 JND (0.8) (10)	(10)		1	42ND (0.8)	100ND (0.8)	ND (2) ND (0.6)		2) ND		one	$\left  \cdot \right $
			(2.6) NI		7		1-			1	į	- 1			1	-	1		T	-	0.8		zene	-
2 Z		(0.8) - ND - ND	<u> </u>	-		(0.8)	-	ON ON	_	(0.8)				—			-	(S)	(0.8)		7		modichloromethane  Butadiene	-
1 N 0 N		ND (1)	<u>.</u>	$\overline{}$	ND (1) ND	(1)	2	ND (1) 2	אט (ז) אט (ב)		ND (1) ND (2)	ND (1) ND (2)	ND (1) ND (2)		3		ND (1) ND (2)	(1) NU (2)			D (1) ND (2)		utanone	-
(1.0) (1.0)		(10) (10)	(2.6) (2.6)		ND (2) ND (2)	F)		2.3 J ND (2)		1	(2) NO (2)	(2) ND (2)	(2) ND (2)		3 3	(2) ND (2)	(2) ND (2)	(2) ND (2)			(2) ND (2)	_	bon disulfide	-
0 D				T			T				2)			1	1	T				T	(S)		bon tetrachioride	1
0 0 (1.0)		0		1	4.1		3 7	3.7		Ā	<u>.</u>	2.7			T	<u>ω</u>	3.2		9 -	7	<u>-</u> 2	_	oratom	-
(1.0)		(0.8)	-	n (0.	6.2 ND	1	6.6 (0.0)	6.9 ND	T	71 (0.8)	N S	4.6 NO	(0.8)			5.2 (0.8)	6 N	(0.8)	(0.8)	Z (0.8	8.1 ND	_	oroethane	-
(1.0)			_	$\top$	(E) GIN	╄	NO (1)	OND (1)	↓		NO (3)	(E) GN	-		—	ND (1)	ND (1)	3)			ND (3)	-	oromethane	1
0 NO (1.0)		(4)	-	Z Z	(3) NA	$\top$	(1) NE	(T)	Т		E N	(T) NH	1	1	<u> </u>	NA NA	1) NH		+	=   -	_	1	clohexane	7
(1.0 (2.0 (3.0)		(0.5)	+			┸	_		-		4		<del> </del>	┷	ND (0.5)	ND (0.0)	S	(0.5)	(0.5)	NO (0.0)	2	1,2	-Dibromoethane	7
(1 N		(0.8)					- 1		L								-			_		-	-Dichlorobenzene	1
(1.0)	1							-							$\overline{}$	$\overline{}$						_	3-Dichlorobenzene	
(1.0)	┨	(0.8)	<del>-   -</del> -	_		—	-											+-			_	1	I-Dichlorobenzene	
(1.0)	1	1	(0.64)	$\sim$ L		23	2.3		1-	1100	1.9	_ :	1	= _	1 -	1.3	1.7		7	N		, Di	chlorodifluoromethane	
(1.0)	1	2) (0.5)	(0.6/ 0.6/	Z (		_			2) (0.5)		0.94	(0.5)	NO.5)	NO (0.5)	N (0.0	2	) Z	(0.5)	Z (0.5)	8	2 2	<b>3</b> 1.	1-Dichloroethene	
(1.0)	$\dashv$	(0.8)		_+		_																	s-1,2-Dichloroethene	
9.0			Ä	Y.		S	NR		S	Z S	NA		ZE I	NR.	NR R	NR	3		3	S		ı	hanol	
(1.0)		(0.5)	N (26	0.87	(0.5)	N (0.0	N	2	0.75	8	S N	(0.5)	N (0.5)	N (0.5)	N S	) N		ń	(0.5)	B	(2)	0.69 L	hyl benzene	
(1.0)	3		ND (2.6)	16	(0.6)	N S	S E	(0.6)	N G	8	) N	(0.6)	ND (0.6)	ND (0.5)	S	S N	(2)	(0.5)	ND (0.5)	8	(0.5)	<b>₹</b> 4	Ethyltoluene	
(1.0)	5	_	N N	N <sub>D</sub>		3	Z	1	¥	N.	¥	1	NH NH	Z.	¥	N.		ž	¥.	S.		_;	eptane	
(1.0)	3		¥	NR	:	N.	Z X		N <sub>D</sub>	NR	¥		Z Z	NR RN	S B	Z,		Z Z	ž	ž	_	_	lexane	
(1.0)	S		ND (1)	Š		ND (3)	ND (3)		ND (3)	ND (3)	ND (3)		Ν (ω)	ND (3)	ND (3)	ND (3)	į	N (3)	ND (3)	ND (3)		8 (3)	-Hexanone	
(1.0)	S	(0.8)	S	ND (3)	<u></u>	8		12	-	$\rightarrow$		٠.		S N				8 8	8 8	_	(2)	0.82 J	fletnylene chloride	
(1.0)	S	_	ND (2)	3 Z		ND (2)	(Z)		ND (2)	ND (2)	(2)		ND (2)	ND (2)	ND (2)	ND (2)		8	ND (2)	ND (2)		(2)	-Methyl-2-pentanone	
(1.0)	g		系	Z N		ž	Z Z		줆	Z Z	<u> </u>	;	줆	Ä	Ä	Z H	_	ž	품	Z Z		₹.	P-Propanol	
(1.0)	8		ž	Z	Ļ	Ä	ž	┸	₹	Ä	3	1.	¥	Z,	Ä	Z H	<u> </u>	系	N.	Z Z		4	Propylene	
(1.0)	8	(0.0)	S S	6 Z	(0.5)	8	(0.5)	(0.5)	Z Z	(0.5) OND		-					<del>-</del>	$\overline{}$			_	$\neg$	Styrene	
(1.0)	8	(0.0)	) N	_	•			5 5	0.64	(0, N 0, E	(2)	(0.0)	8	9 S	0.51 J (2)	(0.5)	(0.5)	8	0.7		1	ر 95.0	Tetrachloroethene	
	6.4	(0.0/	<u> </u>	_ <u>u</u>	(0.0)	200	(0.5) 	5	3.1	(0.5)	(0.5)	5 0	3	() N	(0.5)	(0.5)		427	20	0.69 (2)	2	ે જા	Toluene	
- h	8		O NO	2 2 2 3 4 5 7 7 7 7	5	¥D (3)	3	١	tD (3)	(3)	[ ]	3	4D (3)	(D)	(3)	(3)		D (3)	D (3)	(3)		D (3)	1,2,4-Trichlorobenzene	
(1.0)	8		6 S	(0.64)	5 3	9 N	(0.5)	5 3	) Z	(0.5) -	(0.5)	200	8	0 S	0.5 5	(0.5)	(0.5)	8	0 K	(0, N (5)	(0.0)	5 8	1,1,1-Trichloroethane	
(1.0)	Š		47	ن ف	3	34		20	21	20	نـــا	2	-54	_ <del>1</del> 3	19	-	-	19	16		-	25	Trichloroethene	
(1.0)	E		<u>ب</u>	_ <u>:</u>	-	<u> </u>		5.2	5.2	4.9		4	3.6	2.7	3.6		3	4.1	3.6		-	5.7	Trichlorofluoromethane	
(1.0)	2		67		8	<u>1</u>		120	130	120	Ţ-	120	96	73	90	$\overline{}$		120	98	2	3	130	1.1.2-Trichloro- 1.2.2-trifluoroethane	
(1.0)	2	4 1	(0.5)	_	_		$\overline{}$	$\neg$	0 8 5 0	·				(0.5)					0.75 J (2)			0.67 J	1,2,4-Trimethylbenzene	
(1.0)	É													(0.8)			+	S	(0.8)	├	+		1,3,5-Trimethylbenzene	
(1.0)	3 6	-ł i		Ε-	-+		┿	+	6 Z	<del> </del>	+-	-+		(0.6)	+	+	+	24	7.3	(0.5)	┽	0.99 <b>J</b>	o-Xylene	
(1.0)	2 6	5	(0.8)	(2.6)	2.3 J	(8.0 (8.0	(0.8)	용	N	(8.0)	(0.8)	Š	@ B	(0.8)	(0.8)	(0.8)	5	54	27	(0.8)	3	<u>ن</u> ن	m-, p-Xylene	
11:0	3 7	3	NH		NR.	3	1	Z D	ž		ā	N.	Z	ž	3	ő	3	S	Z		S	3	Total Xylene	

Note: Value FEPA Janue	A La B Q 8 Q 8 Q 8 Q 8 Q 8 Q 8 Q 8 Q 8 Q 8 Q
8 E 5 2	Laboratory and Record Number <sup>a</sup> Quanterra 602989 Quanterra 602989 Quanterra 602989 Quanterra 603340 QUSTL 603340 QUSTL 603661 QUSTL 603681 QUSTL 603898 QUSTL 603898 QUSTL 603898 QUSTL 603898 QUSTL 603898 QUSTL 603898 QUSTL 603898 QUSTL 603898 QUSTL 603898 QUSTL 603898 QUSTL 603898 QUSTL 603898 QUSTL 605812 QUSTL 605162 QUSTL 605407 QUSTL 605407 QUSTL 605407 QUSTL 605407 QUSTL 605407 QUSTL 605407 QUSTL 605407 QUSTL 605407 QUSTL 605407
in bold represent detected analytes. ber 1986. y 1897. y 1897. e Borehole Air Toxics Ltd. Laboratory Core Laboratories LuS. Environmental Protection Agency Environmental Restoration Environmental Restoration Chemistry Laboratory Field Blank Foot (feet) Identification The reported value is greater than or equal to the MDL but less than the practic - Method detection limit Miligrant(s) per cubic meter (air) Not Applicable - Not Applicable - Not Applicable - Not Applicable - Not detected above the MDL, shown in parentheses Not Applicable - Not exported Viganic vapor analyzer Viganic vapor analyzer Couanterra Laboratory	Sample Attributes  ER Sample ID Depth (ft)  TA2-VW-20-72 72  TA2-VW-20-72 72  TA2-VW-20-72-DU 72  TA2-VW-020-72-DU 72  TA2-VW-020-72-DU 72  TA2-VW-20-72-DU 72  TA2-VW-20-72-DU 72  TA2-VW-20-72-DU 72  TA2-VW-20-72 72  TA2-VW-20-72 72  TA2-VW-20-72 72  TA2-VW-20-72 72  TA2-VW-20-72 72  TA2-VW-20-72 72  TA2-VW-20-72 72  TA2-VW-20-72 72  TA2-VW-20-72 72  TA2-VW-20-72 72  TA2-VW-20-72 72  TA2-VW-20-72 72  TA2-VW-20-72 72  TA2-VW-20-72 72
to bld represent detected analytes. rer 1986. 1997. 1997. Servichole. Air Toxics Ltd. Laboratory. Core Laboratories. U.S. Environmental Protection Agency. Environmental Protection Agency. Foot (teet). Identification. Environmental Restoration Chemistry Laboratory. Field Blank. Foot (teet). Identification. The reported value is greater than or equal to the MDL but less than the practical quantitation limit, shown in parentheses. Not Applicable Not defected above the MDL, shown in parentheses. Not Applicable Not defected above the MDL, shown in parentheses. Not reported. Part(s) per billion on a volume per volume basis. Organic vapor analyzer. Sample collected while system tubing was being purged and before OVA readings stabilized. Quanterral Laboratory. Quanterral Severm Trent Laboratories. Sample collected after system tubing fully purged and OVA readings stabilized. Soil vapor. Technical Area. Volatile organic compound. Vapor well. Not applicable.	Sample Attributes  ER Sample ID  A2-VW-20-72  A2-VW-20-72  A2-VW-20-72  A2-VW-020-72  A2-VW-020-72  A2-VW-20-72
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Table A-4
Summary of Technical Area II Active Soil-Vapor Monitoring Well VW-21 Sampling Soil-Vapor VOC Analytical Results
November 1996 to March 2002
(On- and Off-Site Laboratories)

Soll-vapor conce Core 1/22 600308 1742 600308 1742 600308 1742 600308 1742 600796 1742 600796 1742 601237 1742 601237 Quanterra 1742	Soil-vapor of ERCL 6173 ERCL 6173 ERCL 6173 ERCL 6173 ERCL 6173 ERCL 6942 ERCL 6942 ERCL 6942 ERCL 6942	Soil-vapor Soil-vapor ATL 9264 ATL 9264 ATL 9264 ATL 9264 ATL 9264 ATL 9264 ATL 9264 ATL 9264 ATL 9265 ATL 9265 ATL 9265	Laboratory and Record
Soil-vapor concentration ppb(w/v) Core TA2-WW-21-50 600308 Core TA2-WW-21-92 600308 Core TA2-WW-21-92 600308 Core TA2-WW-21-92 600308 Core TA2-WW-21-92-DU 600308 Counterra TA2-WW-21-92 Counterra TA2-WW-21-92 Counterra TA2-WW-21-92 Counterra TA2-WW-21-92 Counterra TA2-WW-21-92 Counterra TA2-WW-21-92 Counterra TA2-WW-21-50 601237 Counterra TA2-WW-21-50 601638	Soil-vapor concentration mg/m ERCL 6173 TA2-VW-21-50-9 ERCL 6173 TA2-VW-21-50-9 ERCL 6173 TA2-VW-21-92-P ERCL 6173 TA2-VW-21-92-9 ERCL 6942 TA2-VW-21-50-P ERCL 6942 TA2-VW-21-50-S ERCL 6942 TA2-VW-21-92-P ERCL 6942 TA2-VW-21-92-P	Number at the Sample Depth (IV) Under Soli-vapor samples collected during drilling plot(VI) ATL 9264 TA2-BH-021-20-SV 10 11-11 ATL 9264 TA2-BH-021-20-SV-DU 20 11-11 ATL 9264 TA2-BH-021-30-SV 30 11-11 ATL 9264 TA2-BH-021-40-SV 40 11-11 ATL 9264 TA2-BH-021-40-SV 50 11-11 ATL 9264 TA2-BH-021-50-SV 50 11-11 ATL 9264 TA2-BH-021-70-SV 50 11-11 ATL 9264 TA2-BH-021-70-SV 70 11-11 ATL 9265 TA2-BH-021-90-SV 80 11-11 ATL 9265 TA2-BH-021-90-SV 90 11-11 ATL 9265 TA2-BH-021-90-SV 100 11-11 ATL 9265 TA2-BH-021-90-SV 100 11-11 ATL 9265 TA2-BH-021-90-SV 100 11-11 ATL 9265 TA2-BH-021-90-SV 100 11-11 ATL 9265 TA2-BH-021-100-SV 100 11-11 ATL 9265 TA2-B	Sample Attributes
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7.0 27 22 22 21 19 21 14 14 14 20 20 20 20	ND ND ND (1.2) (1.2) (1.2) (1.2) (1.2) (1.2) (1.2) (1.2) (1.2) (1.2) (1.25) (1.		Carbon tetrachloride
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NA NA (0.37) NA (0.55) NA (0.55)	N N N N N N N N N N N N N N N N N N N	7.4 NR 113 NR 113 NR 110 NR 111 NR	Cyclohexane 1,2-Dibromoethane
ND ND ND (1.0) (1.0) (1.0) (1.0) (1.0) (1.0) (1.0) ND ND (1.0) ND (1.0) (0.48) (0.48) ND (0.8) (0.5) ND ND (0.8) (0.5) ND ND ND (0.5) ND ND ND (0.5)	N	ND (1.0) (1.	1,2-Dichlorobenzene
ND ND ND (1.0) (1.	N N N N N N N N N N N N N N N N N N N	(0.79) 1.6 (0.79) 1.0 (1.0) (1.0) (1.0) (1.0) (1.0) (1.0) (1.0) (1.0) (1.0) (1.0) (1.0) (1.0)	1,4-Dichlorobenzene  Dichlorodifluoromethane
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(1.0) NA (1.	NA NA NA NA NA NA NA NA NA NA NA NA NA N	14 16 16 16 17 17 18 19 19 19 19 19 19	4-Ethyltoluene C
NA ND (3.9) NA ND (3.9) NA ND (3.9) NA ND (3.9) NA ND (3.9) NA ND (3.9)	N N N N N N N N N N N N N N N N N N N	7.3 ND 18 (3.2) 18 (4.1) 12 ND 39 (9.8) 39 (9.8) 375 ND (13 86 40 330 ND (40) ND (5.4) (6.4) (6.4) (6.4) (7.8) (7.8) (7.8) (7.8) (7.8) (8.4) (7.8) (7.8) (8.4) (8.4) (7.8) (8.4) (8.4) (7.8) (8.4) (8.4) (8.4) (8.4) (9.8)	Hexane
13 NA 10 NA ND NA (1,0) NA (0,44) (0,38) 13 NO (15) (0,38) 14 NO (15) (0,38) 14 NO (15) (0,38) 14 NO (15) (0,38) 14 NO (15) (0,38) 14 NO (15) (0,38) 17 NO (15) (0,38) (15) (0,38) (	(1.2) NA (1.2) NA (1.3) NA (4.8) NA (4.9) NA (4.9) NA (4.9) NA (4.9) NA (4.9) NA (4.9) NA	ND (0.79) (3.2) (1.0) (4.0) ND (1.0) (4.0) ND (1.0) (4.0) ND (1.0) (4.0) ND (1.0) (4.0) ND (1.0) (1.0) ND (1.0) (1.0) (1.0) ND (1.0) (1.0) (1.0) ND (1.0) (1.0) (1.0) (1.0)	Methylene chloride  4-Methyl-2-pentanone
NA NA NA NA NA NA NA NA NA NA NA NA NA N	Z Z Z Z Z Z Z Z Z Z Z Z Z Z Z Z Z Z Z	4.7 ND (3.2)  8.8 430  ND (4.0) (4.0) (9.8)	2-Propanol Propylene
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2.0 1.0 ND (1.0) ND (1.0) ND (1.0) ND (1.0) ND (1.0) ND (1.0) SJ 1.8J ND (1.5) ND (1.5) (2.3) AJ ND (2.3) (2.3) ND (3.3) (2.3) ND (3.3) (3.3) ND (3.3) (3.3) ND (3.3) (3.3) ND (3.3) (3.3) ND (3.3) ND (3.3) (3.3) ND (3.3) (3.3) ND (3.3) (3.3) ND (3.3) (3.3) ND (3.3) (2.3) ND (3.3) ND (3.3) (2.3) ND (3.3) ND (3.3) ND (3.3) (2.3) ND (3.3)	ND NA (1.2) NA (1.2) NA (1.2) NA (1.2) NA (1.2) NA (1.2) NA (1.2) NA (1.25)	9) 20 ND 119 28 ND 12.2 17 ND 2.2 17 ND 3.3 34 ND 3.3 (2.4) 5.0 94 (2.4) 9) 1500 ND 10) 250ND (10) 2.9 170 ND 1.9 (0.96) 4.3 560 ND 3.7 150 ND 1.9 (0.96) 4.1 560 ND 1.6 (0.96) 1.6 (0.96) 1.6 (0.96) 1.6 (0.96) 1.6 (0.96)	Toluene
(1.0) (1.0)	ND 2.9 J (1.2) (4.8) ND ND (1.2) (1.2) (1.2) ND ND (1.2) ND H2 J (1.25) (1.25) ND ND (1.25) (1.25) (1.25) ND ND ND (1.25) (1.25) (1.25) ND 45 J (1.25) (1.25) ND 8.9	(0.79) (0.79) (1.0	1,1,1-Trichloroethane Trichloroethene
230 ND ND 190 ND 190 ND 190 ND 190 ND 100 ND	9.J NR (1.2) 1.8) (2.1) 2.J NR (4.8) 2.J NR (0.2) 0 NR (1.2) 0 NR (1.25) 0 NR	56 2.3 ND 17 ND (0.79) 17 ND (1.0) 15 ND (1.0) 19 (1.0) 19 (1.0) 10 9.5 ND 10 (2.4) 1100 9.5 ND 120ND (10)ND (10) 640 3.5 ND 120ND (10)ND (10) 640 3.5 ND 120ND (10)ND (10) 640 3.5 ND 120ND (1.6) 120ND (1.6)	Trichlorofluoromethane  1,1,2-Trichloro- 1,2,2-trifluoroethane
(2) (0.5) (0.8)	8) NR NR NR NR NR NR NR NR NR NR NR NR NR	18 22 23 8 222 8 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7 7	1,2,4-Trimethylbenzene
9) NB NB NB NB NB NB NB NB NB NB NB NB NB	1 (1.2) (2.5) 1 (1.2) (2.5) 1 (1.2) (2.5) 1 (1.2) (2.5) 1 (1.2) (2.5) 24 J (24 J (2.5) 1 (1.25) (2.5) 1 (1.25) (2.5) 1 (1.25) (2.5) 1 (1.25) (2.5) 1 (1.25) (2.5) 1 (1.25) (2.5) 1 (1.25) (2.5) 1 (1.25) (2.5)		o-Xylene m-, p-Xylene
6) R R R R R R R R R R R R R R R R R R R		13 NR NR NR NR NR NR NR NR NR NR NR NR NR	Total Xylene

Table A-4 (Continued)
Summary of Technical Area II Active Soil-Vapor Monitoring Well VW-21 Sampling
Soil-Vapor VOC Analytical Results
November 1996 to March 2002
(On- and Off-Site Laboratories)

Refer to footnotes	0/STL 605162	O/STL	Q/STL	Q/STL	Q/STL 604643	Q/STL 604643	0/STL 604434	Q/STL 604434	Q/STL 603898	Q/STL 603898	Q/STL 603898	Q/STL 603661	Q/STL	Q/STL 603661	Q/STL 603340	Q/STL 603340	0/STL 603340	Q/STL 603136	Q/STL 603136	Q/STL 603136	Quanterra 602989	Quanterra 602989	Quanterra 602989	Quanterra 602830	Quanterra 602830	Quanterra 602830	Quanterra 601823	Quanterra 601823	Quanterra 601823	Quanterra 601638	Quanterra 601638	Laboratory and Record	
otes at end of table.	TA2-VW-21-92	TA2-VW-21-50	TA2-VW-21-92	TA2-VW-21-50	TA2-VW-21-92	TA2-VW-21-50	TA2-VW-21-92	TA2-VW-21-50	TA2-VW-21-92-DU	TA2-VW-21-92	TA2-VW-21-50	TA2-VW-21-92-DU	TA2-VW-21-92	TA2-VW-21-50	TA2-VW-021-92-DU	TA2-VW-021-92	TA2-VW-021-50	TA2-VW-21-92-DU	TA2-VW-21-92	TA2-VW-21-50	TA2-VW-21-92-DU	TA2-VW-21-92	TA2-VW-21-50	TA2-VW-21-92-DU	TA2-VW-21-92	TA2-VW-21-50	TA2-VW-21-92-DU	TA2-VW-21-92	TA2-VW-21-50	TA2-VW-21-DU	TA2-VW-21-92	ER Sample ID	Sample Attributes
	92	50	92	50	92	50	92	50	92	92	50	83	92	50	92	92		92	92	50	92	92	50	92	92	50	92	92	50			Sample	
	12-11-01	12-11-01	09-25-01	09-25-01	06-22-01 N	06-22-01			2-11-00 2		12-11-00	09-13-00	09-13-00	09-13-00	06-20-00	06-20-00	06-20-00	03-01-00 N	03-01-00 N		12-07-99		12-07-99	09-07-99 N	09-07-99 N	09-07-99	06-24-99	06-24-99	06-24-99		03-22-99	Sample Date	
- 1	(9.5) (3. N. (9.5)		13 J N			7.6 J N				5.9 J N (27) (2.	_	(7.4) (2. N)						ND (12) NI	ND (14) (5.		13 J N (40) (3.		(4.5) (1.		D (11) N (4.	(7.3) N	26 J N (77) (6.	25 J N (44) (3.	(22) (1 N	16 J N (26) (2.	16 (0. ⊻	Acetone	
- 1	8) ND (3.8)		4 N (4 N				1) ND (4.1)			D 2.8 ( 2) (5.4)		0 3.2 J 9) (7.4)										1.	i .	i I	_ 1	_	i. 1	_ !		0 3.4 J 1) (5.2)	ıμ	Benzena Bromodichloromethane	-
	(4.7)	ND (1)	(5.4)	ND (1)	(5.4)	ND (2.1)	(5.1)	ND (3)	ND (1)	ND (2.7)	ND (1)	(3.7)	(3.6)	ND (1.5)	ND (2.7)	ND (3.6)	(1.6)	ND (6)	ND (3)	ND (3)	ND (4)	(6.7)	(2.2)	(5.5)	ND (5.5)	(3.7)	ND (7.7)	ND (4.4)	(2.2)	(2.6)	ND (3)		
	(9.5)	ND (2) N	ND (11)	ND (2) N	IN(11) ON	(4.1)	무울	ND (2) N		ND (5.4)		(7.4)					<del>                                     </del>	Z	ND (14)ND (14)	(6) ND (2)	(8.1)	ND (13)N	(4.5)	ND (11) NE		(7.3)		(8.8) (8.8)		(5.2)	<del>                                     </del>	2-Butanone	_
	ND ND (9.5) (2.4)	ND (2) N	ND (11) N	ND (2) N						ND N (5.4) (1.		ND ND (7.4) (1.8)							(3.	D (6) ND (1.5)	(8.1) ND	-	(4.5)			(7.3) (1. N	<del></del>		<del> </del> -	(5.2) (1.3)		Carbon disulfide  Carbon tetrachloride	_
	4 0	5 0	70					5)									000	(3)							7 0	8 0	80	2) 0				Chloroform	
,	(3.8)	(0.8)	27 ND (4.4)	(0.8)	25 ND (4.3)			0.8)	L	(2.2)	(0.8)	(2.9)		(1.2)	(2.2)	(2.9)	1	<del> </del>		(2.4)	-	(5. <b>4</b> )	(1.8)	(4.4)	<b></b>	<del> </del>	·	1		+	(0.8)	Chloroethane	
	(4.7)	ND (1)	(5.4)	3	(5.4)	(2.1)	(5.1)		1	(2.7)		(3.7)		T			$\vdash$		<b>—</b>											(2.6)	1	Onlordienane	
	N 7	N A O z	NA (2	() Z	<del> </del>	<b>!</b> —		4	↓	<del> </del>	<b>!</b>	NA ND (1.8)	NA C Z	<u> </u>	-	—	-			(1.5)	L			ļ		ļ	ļ		↓	<del> </del>		Cyclohexane 1,2-Dibromoethane	_
	ND ND (2.4) (3.8				ļ	<b>↓</b> ~	1	↓	↓					1	<del> </del>				<del></del>	+	<del>-</del>		+	+	+		+	-		-	+	1,2-Dichlorobenzene	
	) (2.8)			<del></del>		+	+	+	+		-	+	<del> </del>	<del>i -</del>	+	+-	-	+	+	(1.5)	·		+	+	<del></del>	+		+	+		-	1,3-Dichlorobenzene	
	(3.8)	(0.8)	(4.4)	(0, N	(4.3)	(1.7)	(A) 2	1	1 .		1	(2.9)	(2.9)	(1.2)	(2.2)	(2.9)	(1.3)						+	<del> </del>	+	┼		+-	+		- تح	1,4-Dichlorobenzene	
	(2.4)	i	ì	١_			(2.6)			(1.3)	1	(1.8) (1.8)		L_	7	T	7		1	(6)		1	1.	1	1	_		1	1_	(1.3)	(2)	Dichlorodifluoromethane	_
	10 ND (3.8)			1	(11) (A.)		(10) (4		- E	~	<del>'-</del>	(7.4) (2.9)	_	-	-	-		-		_	+		7	7	7	7	(15) (6.2	7	7		*	cis-1,2-Dichloroethene	
	NA A	y Z					1		7						1	1	3 3	T		2 3	7	NA 3			T	"				2)		Ethanol	
	(2.4)	(0.5)	(2.7)	(0.5)	(2.7)	10 (1)	(2.6)	(0.5)	(0.5)	(1. g	(0.5)	(1 g	(1.8)	(0.74)	(1.3)	(1.8)	(0.8)	20 0	(3.5)	N (1.5)												Ethyl benzene	
	(3.3)	(0.7)	(3.3)	(0.6)	(3.2)	(1.2)	(3.1)	(0.6)	(0.61)	(1.6)	(0.6)	(2.2)	(2.2)	(0.88)	(1.3)	(1.8)	(0.8)	10 (3)	(3.5)	(1.5)	5 (	(3.4)	(1.1)	(2.7)	(2.7)	(1.8)	(3.8)	(2.2)	<b>3</b> []	Σ (1. g	(0.5)	4-Ethyltoluene	
	N >	<del> </del>	-	┼—	+-	+	╁	-	+-	2 2	┼	2 2	-	+	+-	+-	+-	+-	+-	+	+	+		2 2	+	+	2 2	┼─	+	NA S	+	Heptane  Hexane	
	NA ND ND (4.7)	١_,	L	L	i	(6.2)		NA NO (3)		NA (8.1)	<del></del>	NA ND (11)		1	(8.1)		(4.8)	- 1		NA ND (21)	1	- 1	(6.7)	NA SO (18)		1	NA NO (23)			N (7.8)		2-Hexanone	
	-	(0.8)		0 2	į	(1.7)	5	(0.8)		17	(0.8)	1	-	(1.2)			(1.3)	3		(2.4)	3		_			(2.9)	2		≘;	ξ		Methylene chloride	
	(9.5)	1 -		(2)		(4.1)	ND (10)	(01) (N	3	(5.4)		7.4	(7.2)	(2.9)	(5.4)	(7.2)	(3.2)	2	25 (19)	5ND (14)	(8.1)	N (	(1.8) (4.5)		ND (11)	(7.3)		(8.8)	(4.4)	(5.2)		3 4-Methyl-2-pentanone	
	Z	-	├	+-	+	╁╌	+	+-	+-		+	2 2	╁─╴	+-	+	+		+-	+	+	+-	-	+-	Z 2	-	+-	2 3	-	+	+	+	2-Propanol	
	NA ND ND (2.4	-	-	┼-	(2.7)							(1.8)	<u> </u>	Щ						(1.5)			_	Щ.			(3.8)					Styrene	
	9	†	3 (3)	+	+	1	(10) 2.5 J	_	25		+	7.4)		<del></del>		<del> </del>				-		$\top$		1-		T	29.1	1	_	T	f	Tetrachloroethene	
	-	(S) E	(2.7)	(S)	(2.2)	2 (	N (25)	3 (G) (	_	3 (3)	) (S)	3 3	(1.8)	S (2)	<u> </u>	Z (1)	S (0.3)	5 3	(3) (3) (3)	8 (3)	5	ND (3.4)	(1. (1.	2 S	3 3 3	(1.e)	(3.8)	(3.8)	(1.1) 2.5 J	N (1.3)	3	Toluene	
		(0.6)		<del></del>	$\neg$	(6.2)				ND (3)	-+	NO (1 1)			(8.1)		(4.8) (C	5			(9)		(6.7) (1							(7.8)		1,2,4-Trichlorobenzene	
		<del> -</del>	1	+-	(2.7)	1	1		T			(1.8)			Τ.		_			(1.5) ND 12		T								ND (1.3)		Trichloroethene	
	(9.5)		(2.7)	7	7		(2.6) 370 3.9 J	_	360 3.8	(5.4)		350 3.9	1	T			(3.2) 100 1.9 J	470 3	(3.5) 850 ND (3	200 ND (6)	3.6	500 4.4	500 3.7				(6.2) 460 3.7 J			(5.2) 410 4.6	$\top$	Trichlorofluoromethane	<del></del>
		1	1		3 (11)	39.	1.5 J	(2) 5 J (10)	1.5.1	(5.4)	4 (2)	7.4	(7.3)	(2.9)	(5.4)	(7.3) 2.9.J	(3.2) 2.9 J	122	3.1 J	6) (1.5) 3.8 J	) (8.1)	3) (13) J 7.7 J	J (4.5)									1,1,2-Trichloro- 1,2,2-trifluoroethane	
	(2.4)	(0.5)	27	(0.5)	(2.7)	3	(2.6) ND (1)	N (0.5)	Z (0.5)	χ Θω	(0.5)	3 6 8	(1.8)	(0.74)	نيا 6	(1.8)	N (0.8)	8	ND (3.5)	₹ (i.s)	8	N (3.4)	S (1	(2.7)	(2.7)	(1.8)	(3.8)	(2.2)	8 <u>1</u>	ĕ (± ω	N (0.5)	2 1,2,4-Trimethylbenzene	
		+	-						+-				+-		+-	+-	-+					<del>-</del>	+-	<del></del>		+-			+	+-		51,3,5-Trimethylbenzene	
		_										(2.2) (2.9) ND ND					i			(1.5) (2.4 ND ND					_							ဗြ ၀-Xylene ာ် m-, p-Xylene	
	B) N	1		+	_		_		3		┰	<del> </del>		$\neg$	$\top$			S G	N N		S S	$\top$		_		$\top$	_				1	∑ Total Xylene	

# Table A-4 (Concluded) Summary of Technical Area II Active Soil-Vapor Monitoring Well VW-21 Sampling Soil-Vapor VOC Analytical Results November 1996 to March 2002 (On- and Off-Site Laboratories)

		D)		П	
Q/STL	605407	and Record	Laborato		
A2-VW-21-92	FA2-VW-21-50	ER Sample ID		Samo	
23	č	pie ID		Sample Attributes	
	 	De o	)	utes	
92	2	<b>₹</b> °			
3-19-02	3-13-0	Date			
(6.9)		╁~~	tone	+	
) (2.8)	ω,		zene	1	
) (6.9)	+-	Bro	modichloromethane		
.9) (3.4)	(1.3	3 1,3	-Butadiene	-	
(6.9)	_	╁	Sutanone	1	
	===	1	rbon disulfide	-	
-	) (0.67)		rbon tetrachloride	7	
-	7)	-	loroform	+	
-	0 (1.1)	5	loroethane	4	
-		+	Noromethane	$\dashv$	
$\vdash$	NA NA	-	rclohexane	-	
	(0.67)	-	2-Dibromoethane	-	
-	N (1.1)	<del></del>	2-Dichlorobenzena	-	
			3-Dichlorobenzene	-	
-	NO CO	+	4-Dichlorobenzene	-	
8) (1.7)	-		ichlorodifluoromethane	_	
	- 6	25	1-Dichloroethene		
(6.9) (2.	ND ND	+	is-1,2-Dichloroethene	-	VOCs (EPA Method 8260-M3", TO-14/TO-14A <sup>b</sup> ) Units as indicated
8)	NA.		ithanol		PA Meth
1 (1.7)	+		Ethyl benzene		od 8260
() (2.4)			l-Ethyltoluene		M3, TC
4) !	N.		Heptane		-14/10-
-	NA NA		Hexane		14A <sup>b</sup> ) Ur
9			2-Hexanone		nits as in
		_	Methylene chloride		dicated
	14 ND		4-Methyl-2-pentanone		$\  \ $
-		-	2-Propanol		$\left  \cdot \right $
Ì	NA NA		Propylene		$\left\{ \left. \right  \right\}$
	IA ND (1.7)	_	Styrene		-
ŀ	_	-	Tetrachloroethene		$\left  \cdot \right $
		(2.7) (0.67	Toluene		-
	7)   ND		1,2,4-Trichlorobenzene		1
		1			-
			1,1,1-Trichloroethane		-
	(1.7				$\dashv$
	$\vdash$	7.	1,1,2-Trichloro-		-
	(6.9) (1.7	2:	1,2,2-trifluoroethane		$\dashv$
	7) (2.8)	13	1,3,5-Trimethylbenzene		$\dashv$
					-
	(2.8)	+	So-Xylene		-
	F	+	Total Xvlene		
	L	5	Total Xylene		

*EPA November 1986.	TIPA November 1986.
EPA January 1997.	ary 1997.
Analysis rec	Analysis request/chain-of-custody record.
AT.	= Air Toxics Ltd. Laboratory.
里	= Borehole.
Core	⇒ Core Laboratories.
5	= Duplicate sample.
EPA	= U.S. Environmental Protection Agency.
9	⇒ Environmental Restoration.
ERCL	= Environmental Restoration Chemistry Laboratory.
#	⇒ Foot (feet).
ō	= Identification.
	= The reported value is greater than or equal to the MUL out less than the province quantity of the most officers.
MD.	= Method detection limit.
mg/m³	= Milligram(s) per cubic meter (air).
N N	= Not analyzed.
ND()	= Not detected above the MDL, shown in parentneses.
S	= Not Reported.
ppb(v/v)	= Part(s) per billion on a volume per volume dasis.
OVA	= Organic vapor analyzer.
סד	= Sample collected while system tubing was being purged and before OVA redunits submission.
Quanterra	= Quanterra Laboratory.
Q/STL	■ Quanterra/Sevem Trant Laboratories.
S	= Sample collected after system tubing fully purged and OVA resumps statement.
STL	= Severn Trent Laboratories.
٧S	= Soil vapor.
ΑT	≂ Technical Area.
δ ດ	⊭ Votatile organic compound.
1747	

Table A-5
Summary of Technical Area II Active Soil Vapor Sampling During Borehole BH-023 Drilling
Soil-Vapor VOC Analytical Results
November 1996
(Off-Site Laboratory)

ATL 9267	ATL 9267	ATL 9267	ATL 9266	ATL 9266	ATL 9266	ATL 9266	ATL 9265	ATL 9265	ATL 9265	ATL 9265	ATL 9265	ATL 9265	Soil-vapor sa	Number	Laboratory		
TA2-BH-023-100-SV	TA2-BH-023-90-SV	TA2-BH-023-80-SV	TA2-BH-023-70-SV	TA2-BH-023-60-SV-DU	TA2-BH-023-60-SV	TA2-BH-023-50-SV-DU	TA2-BH-023-50-SV	TA2-BH-023-40-SV	TA2-BH-023-30-SV-DU	TA2-BH-023-30-SV	TA2-8H-023-20-SV	TA2-BH-023-10-SV	mples	ER Sample ID		Sample Attributes	
100	90	80	70	60	60	50	50	40	30	30	20	10	borehole drill	Depth (ft)	Sample	es	
11-14-96	11-13-96	11-13-96	11-13-96	11-13-96	11-13-96	11-13-96	11-13-96	11-13-96	11-13-96	11-13-96	11-13-96	11-13-96	ing	Date	Sample		1
59 ND	12	43	26 ND (0.86)	17 N	18 ND (0.84)	18	19	150	11	Zi	4	5	3	H	etone		
NO NO	7.2 ND (3.6)	3.2 NO	0 ND (3.4)		_	- 29	4.1 NO	(9.8)	$\overline{}$	1	(3.4)	(3.3)	20	┝	omodichloromethane	-	
3 N		├	(3.4)					+-	(3.4)	┼	(3.4)	;	1	1,:	3-Butadiene		
21	(3.6)	9	(3.4)	6.1	6.2	(3.3)	(3.5)		(3.4)	(3.4)			16	2.	Butanone		
3 Z	(3.6)	T	T	(4 ) (1)	(3.4)	0.4	_	6		*****		\$	11	C	arbon disulfide		
S N	4.2	<u>.</u>	(0.86)	- 1.5		(0.82)	0.88	-	(0.86)	(0.86)	+	1	S S	c	arbon tetrachloride	_	
() NO	1.5	, i			-	_				*	4	-	S	C	hloroform	_	
	<del></del>	+-	(0.86)				(0.88)			(0.86)	┰	(0.82)	ģ		hioroethane		
(0.88) (3.5)		(0.88)	5 86 (3.4)		(0.84)			(9.8)		86)	86) (3.4)	5 Z (3)	1.2 Z	ŀ	yclohexane	-	
	(0.9)	5) (0.8	4) (0.86)		(0.80	(0.8:	(0.88	8) (2.4 N)	(0.8c	7.4 (0.86	(0.86 ND	(0.82	NS NS	$\vdash$	,2-Dibromoethane	-	
			(0.86)	~		_						F	- 1	1	,2-Dichlorobenzene		
			(0.86)							-+				1	,3-Dichlorobenzene		
			Z (0.86)						-	_		_	_	1	,4-Dichlorobenzene		
(0.88)		(0.88)		2		1	1 9	(2.4)	$\neg \vdash$		(0.86) 1.1	$\neg$	0.85	E	Dichlorodifluoromethane		
(0.88)	2	(0.88)	(0.86)	Z (1.0)	(0.84)	(0.82)	(0.88)	N (2.4)	N (0.86)	N (0.86)	₹ (0.86)	N (0.82)	B	1	,1-Dichloroethene	_	
(0.88)		25	7.3	2.8	5.7	5.6	<u>ب</u>	(2.4) 4.5	N (0.86)	-	-	N (0.82)	g		sis-1,2-Dichloroethene	_ ,	ŕ
_	(3.6)	8	(3.4) (( 12						3.4)	8(3) 4)	S	7.9	7.8		Ethanol	[	VOCs (EPA Method IO-14") ppb(V/V)
	7.4	_	(0.86) (3						19 (3	1.9 2/3	7 (3	2.8	8.0 Z		Ethyl benzene	- 00	Method
3.5)	66	5 5	(3.4) (3.4)					-	ð <u>.</u>	<b>€</b>	0 4	To be	3 6		4-Ethyltoluene  Heptane		Q-14") E
_		4.2 (3	(3.4) 12 ND		(3.4) (3.4)				74	4	1 (3,	6.2 No	21		Hexane		(V/V)
(3.5)	-		(3.4) ND						20 ND	7.1 ND	읙	-	7.4 ND		2-Hexanone		
-			(0.86)		<del></del>		<del>-l-</del>					-			Methylene chloride		
F			(3.4)			+	-		+		+	-+	احت 8 د 9 د		4-Methyl-2-pentanone		
(3.5)	N (3.6)	N (3.5	ND (3.4)	N (4	NO (3.4)	ND (3.3)	ND (3.5)	N (9	NO.4	8 (4	S S	200	à Z		2-Propanol		
	(3.6)	8	34	(4.1) 18	(3.4) ND	ND (3.3)	NO (3.5)	N (9	S S (4	N S	S S	N	کا ف کا ف		Propylene		
	(0.90)	N (0.88	ND (0.86)	NO (1.0)	(0.84)	ND (0.82)	ND (0.88)	N ±	NO.	N S	$\overline{}$	$\overline{}$	<u>.</u>		Styrene		
(0.88)	8	6.9	2.5	3.0	5.2	5.2	3.4	4.9	2.8	22	2.7	S N			Tetrachloroethene		
	290	26	150	2	1.8	2.0	7.5	72	780	30	22	88	(C		Toluene	<b></b>	
(0.88)			NO S												1,2,4-Trichlorobenzene	,	
1.00)	8 8	38	NO SE	888	S	SS	8 8	6	5 6	g 6	3 6	3 6	<u>82</u>	5	1,1,1-Trichloroethane		
(0.	3.3	310 N	96 ND	120	150 N	160 N	100 S Z	130 2 Z	46 0 Z	43 0 Z	46 0 N	0 N	(0.8		Trichtoroethene  Trichtorofluoromethane		1
0.00/ 10.0	ND NO	9 0 8	g 0 8		⊋ Ö <b>2</b>	0	ğ Ö [8	ğ Ö				0.86) ON ON ON ON ON ON ON ON ON ON ON ON ON	32	1	1,1,2-Trichloro-		-
00/	9 0	59	17	9.8 ON O	25 NO	24 0 NO	-1 -0	2	7.0	5.7				2	1,2,2-trifluoroethane 1,2,4-Trimethylbenzen	е	_
1 (0.00)	1.5 NO 5	2.4 0 ND	1.8 ND (0.88)				0.82 0.80	1.0 ND 0.85	14 5.	9.1	9.7 3	1.9 0.90		75 2	1,3,5-Trimethylbenzen	е	4
7		= - <u>-                                 </u>	-+					- - -		3.5	5.5	90 3.6		27 8.2	o-Xylene		4
	7.8	Ċ	6	ND (0.86	ND (1.0)	ND (0.84)	_	<u>.</u>	<u> </u>	_ on	ັນ			2	m-, p-Xylene		_

Note: Values in bold represent detected analytes.

\*EPA January 1997.

\*Analysis request/chain-of-custody record.

ATL = Air Toxics Ltd. Laboratory.

BH = Borehole.

DU = Duplicate sample.

U.S. Environmental Protection Agency.

EPA = Environmental Restoration.

ft = Foxt (teat).

ID = Identification.

MDL = Method detection Limit.

ND () = Not detected above the MDL, shown in pare ppb(v/v/) = Part(s) per billion on a volume per volume bc SV = Soil vapor.

= Foot (feet).
= Identification.
= Method detection Limit.
= Not detected above the MDL, shown in parentheses.
= Part(s) per billion on a volume per volume basis.
= Soit vapor.
= Volatile organic compound.

Table A-6 Summary of Active Soil-Vapor VOC Analytical Method Detection Limits November 1996 to September 2003 (On- and Off-Site Laboratories)

	EPA Method	EPA Method	<b></b>
	8260-M3 <sup>a</sup>	TO-14 <sup>b</sup>	
	Detection Limit	Detection Limit	EPA Method
	On-Site	Off-Site	TO-14 <sup>b</sup>
	Laboratory	Laboratories	Detection Limit
	July and	November 1996–	Off-Site Laboratory
	September 1997	March 2002	September 2003
Analyte	(mg/m³)	[ppb(v/v)]	[ppb(v/v)]
Acetone	NA NA	2–560	2-4.1
Benzene	1.2-1.25	0.42-230	0.8-1.6
Benzyl chloride	NA	0.8-230	0.8-1.6
Bromodichloromethane	1.2–1.25	0.27-230	0.8-1.6
Bromoform	1.2-1.25	0.22-140	0.5-1
Bromomethane	NA	0.68-280	1-2
2-Butanone	NA	2-560	2-4.1
Carbon disulfide	1.2–1.25	0.73–560	2-4.1
Carbon tetrachloride	1.2–1.25	0.42-140	0.5–1
Chlorobenzene	1.2-1.25	0.5–140	0.5-1
Chloroethane	NA NA	0.8-230	0.8-1.6
Chloroform	1.2-1.25	0.39-230	0.8-1.6
Chloromethane	NA NA	1–280	1-2
Dibromochloromethane	1.2–1.25	0.23–140	0.5-1
1,2-Dibromoethane	NA NA	0.37-140	0.5-1
1,2-Dichlorobenzene	NA NA	0.69-230	0.8–1.6
1.3-Dichlorobenzene	NA NA	0.48-170	0.7-1.4
1,4-Dichlorobenzene	NA NA	0.68-230	0.8-1.6
Dichlorodifluoromethane	NA NA	0.45–140	0.5-1
1,1-Dichloroethane	1.2–1.25	0.5–140	0.5-1
1,2-Dichloroethane	1.2-1.25	0.76–230	0.8–1.6
1,1-Dichloroethene	1.2-1.25	0.5–140	0.5-1
cis-1,2-Dichloroethene	1.2-1.25	0.54-230	0.8-1.6
trans-1,2-Dichloroethene	1.2-1.25	0.5–140	0.5–1
1,2-Dichloropropane	1.2-1.25	0.8-230	0.8–1.6
cis-1,3-Dichloropropene	1.2–1.25	0.5–140	0.5-1
trans-1,3-Dichloropropene	1.2-1.25	0.8–230	0.8–1.6
Ethylbenzene	1.2–1.25	0.32-140	0.5-1
4-Ethyltoluene	NA NA	0.25-200	0.7–1.4
Hexachlorobutadiene	NA NA	0.57–280	1-2
2-Hexanone	NA NA	1-630	1-2
Methylene chloride	1.2–1.25	0.44-230	0.8-1.6
4-methyl-2-Pentanone	NA	0.38-560	2-4.1
Styrene	1.2–1.25	0.5–140	0.6-1.2
1,1,2,2-Tetrachloroethane	1.2–1.25	0.41-140	0.5-1
Tetrachloroethene	1.2-1.25	0.5–170	0.6-1.2
1,2-Dichloro-1,1,2,2-tetraffuoroethane	NA	0.36–230	0.8-1.6
Toluene	1.2–1.25	0.33-140	0.5-1
1,2,4-Trichlorobenzene	NA	0.6-630	1-2
1,1,1-Trichloroethane	1.2–1.25	0.49-140	0.5–1
1,1,2-Trichloroethane	1.2–1.25	0.5–170	0.5-1
Trichloroethene	1.2-1.25	0.28–140	0.5–1.2
1,1,2-Trichloro-1,2,2-trifluoroethane	0.25–1.25	0.5–140	0.5-1
1, 1,2-1 HOHIOTO-1,2,2-thilluoloethane	0.25-1.25	0.0-140	1 0.5-1

### Table A-6 (Concluded) Summary of Active Soil-Vapor VOC Analytical Method Detection Limits November 1996 to September 2003 (On- and Off-Site Laboratories)

	T		
	EPA Method	EPA Method	}
	8260-M3 <sup>a</sup>	TO-14 <sup>b</sup>	
	Detection Limit	Detection Limit	EPA Method
	On-Site	Off-Site	TO-14 <sup>b</sup>
}	Laboratory	Laboratories	Detection Limit
	July and	November 1996–	Off-Site Laboratory
	September 1997	March 2002	September 2003
Analyte	(mg/m <sup>3</sup> )	[ppb(v/v)]	[ppb(v/v)]
Trichlorofluoromethane	NA NA	0.27-140	0.5–1
1,2,4-Trimethylbenzene	NA	0.32-140	0.8-1.6
1,3,5-Trimethylbenzene	NA	0.29-230	0.8–1.6
Vinyl acetate	NA	1.2-560	2-4.1
Vinyl chloride	1.2-1.25	0.8-230	0.8-1.6
m-, p-Xylene	2.5	0.8-230	1-2
o-Xylene	1.2-1.25	0.5-170	0.6-1.2
Xylene (total)	NA	0.57-0.8	NA NA

<sup>&</sup>lt;sup>a</sup>EPA November 1986.

bEPA January 1997.

EPA = U.S. Environmental Protection Agency.

mg/m³ = Milligram(s) per cubic meter (air).

NA = Not analyzed.

ppb(v/v) = Part(s) per billion on a volume/volume basis.

VOC = Volatile organic compound.

Table A-7
Summary of Technical Area II Active Soil-Vapor Monitoring Well 159-VW-01 Sampling
Soil-Vapor VOC Analytical Results
September 2003
(Off-Site Laboratory)

Laboratory and Record Number STL 606760 11 STL 606760 11 STL 606760 11	- 1-1
59-VW 59-VW 59-VW	Se
Laboratory and Record Number STL 606760 159-VW-01-20-SV STL 606760 159-VW-01-70-SV STL 606760 159-VW-01-150-SV STL 606760 159-VW-01-150-SV STL 606760 159-VW-01-150-SV	Sample Attributes
	butes
Sample Specification (1) Sample Specification (1) Specification (1	
Sample Date 9-9-03 9-9-03 9-9-03 9-9-03	
ND (10) Acetone ND (10) Acetone (10) ND (10) (10) (10) (10) (10) (10) (10) (10)	_
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3 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2	
ND (10) ND (10	_
	_
5 7 1 1 1 1 1 2 2 2 2 2 2 2 2 2 2 2 2 2 2	-
(2.0) (1.6)	_
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	_
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9 0 2 0 9 0 9 0 9 0 9 0 9 0 9 0 9 0 9 0	-
(A) Z (R) Z	
O Dichlorodiffuoromethane	
(4.0) (4.0) (2.0) (2.0) (2.0) (3.0) (4.0)	
X X X X X Ethanol	
(4 N (2 N (2 O O O O O O O O O O O O O O O O O O	
(A N (2 N (2 N (2 N (2 N (4 - Etnyltroluene	 
N X X X X X Heptane	
N Z Z Z Z Z Hexane	VOS (Er o wented to the other)
(4 N (2 N (2 N (2 N (2 N (2 N (2 N (2 N	
4 N N N N N N N N N N N N N N N N N N N	
Z Z Z Z Z Z-Propanol	
₹ ₹ ₹ ₹ Propylene	
4 N (2 N (2 N (2 N (2 N (2 N (2 N (2 N (	
(A. 0.) (A. 0.	
1 1 1 1 1 1 1	
(3.0) (5.0) (1.2,4-Trichlorobenzens (10) (10)	<b></b>
0 0 0 1 1.1,1-Trichloroethane	
Trichloroethene	
24 1.1 Trichlerofluoromethan 2.1 1.9 2.1 1.1 Trichlerofluoromethan 1.9 2.1 1.1 Trichlerofluoromethan 1.9 1.9 1.0 1.0 1.0 1.0 1.0 1.0 1.0 1.0 1.0 1.0	е
1,1,2-Trichloro-	
440 440 825 57 29 1,2,2-trifluoroethane 440 8 255 57 29 1,2,2-trifluoroethane 8 25 57 27 20 1,2,4-Trimethylbenzer (4,0,0,0,0,0,0)	10
4 Z 2 2 0 0 0 0 1,3,5-Trimethylbenzer	
(A Z (2 Z (2 Z (2 Z (2 Z (2 Z (2 Z (2 Z	
(4) X (2) X (2) X (2) X (3) X (4) X (4) X (5) X (5) X (6) X (6) X (7) X	
되 및 및 및 및 Total Xylene	

Note: Values in bold represent detected analytes.

\*EPA January 1997.

\*Analysis request/chain-of-custody record.

DU = Duplicate sample.

EPA = Environmental Protection Agency.

ER = Environmental Protection Agency.

It = Foot (feet).

ID = Identification.

J ( ) = The reported value is greater than or equal to the MDL but loss than the practical quantitation limit, shown in parentheses.

ND ( ) = Not analyzed.

ND ( ) = Not detected above the MDL, shown in parentheses.

NR = Not analyzed.

NB = Part(s) par billion on a volume per volume basis.

STL = Severn Trent Laboratories.

Soli vapor.

VOC = Volatile organic compound.

Table A-8
Summary of Technical Area II Active Soil-Vapor Monitoring Well 165-VW-01 Sampling
Soil-Vapor VOC Analytical Results
September 2003
(Off-Site Laboratory)

STL 606761	STL 60676	STL 60676	STL 60676	STL 606761	and Record	Laboratory	
31 165-VW-01-150-SV	STL 606761 165-VW-01-100-SV	STL 606761 165-VW-01-70-SV	STL 606761 165-VW-01-20-SV	1 165-VW-01-5-SV	d ER Sample ID		Sam
1-150-SV	1-100-SV	1-70-SV	1-20-SV	1-5-SV			Sample Attributes
150	100	70	20	σı	Depth (ft)		
9-9-03	9-9-03	9-9-03	9-9-03 ND		┝╌	5	
	3.4 J ND (2.0)	15 ND (2.0)	ND (10) ND (2.0)	(10) (2.0)		zene	
) (2.0)	2.1	(2.0)	(2.0)	-	┼	modichloromethane	
-	(1.0) ND	ı			_	-Butadiene	-
ND (10) ND (10)	ND (10) ND (10)	ND (10) ND (10)	אט (וט) אט (יט	(10)		rbon disulfide	
6.9	6.1	(2 Z	Т	_	_	rbon tetrachloride	
120	140	2	2 -	į	Ch	leroform	_
(0.80) (4.0)	(0.80) (4.0)	+	-		-	loroethane	_
0) NA	-	$\vdash$	+	+	+	/clohexane	_
L	1					2-Dibromoethane	
	+			-	-	2-Dichlorobenzene	_
-	+-					3-Dichlorobenzene 4-Dichlorobenzene	_
- 1	- 1			- 1		ichlorodifluoromethane	_
.0) (2.0	0.88	0 (2.0)	7 (2.0) ND	(2.0)	짐	,1-Dichloroethene	
_						is-1,2-Dichloroethene	
-	Z A	ž	N.	Z	-	Ethanol	VOCs (EPA Methods TO-14/1 O-14A) ppp(VVV
					-+	Ethyl benzene	Methods
(2.0)	+	$\dashv$			_	I-Ethyltoluene	10-14/10
-	Ä	Ä	×	NA A	-	Heptane	-14A ) p
12	NA C	N Z	N N	NA NA	_	Hexane	
(0)	ð,					2-Hexanone Methylene chloride	
(2)	14 2 (S					4-Methyl-2-pentanone	-
	Ŋ.	Z.	Z A	Z		2-Propanol	$\neg$
-	N A	Z	2	N N		Propylene	
(2.5		_	-			Styrene	
						Tetrachloroethene	_
,	- 1	0 1.6 5	N	0.63	(2.0)	Toiuene	
10.0						1,2,4-Trichlorobenzene	•
	(2 N	(2 N (2.0)	(2.0)	(2.0)	(2.0)	1,1,1-Trichloroethane	
ļ	8.2	26	3.4			Trichloroethene	
	3.4	4.4	l	l l	Į.	Trichlorofluoromethan	9
	170	170	8.3		2	1,1,2-Trichloro- 1,2,2-trifluoroethane	
	-		├		┼	51,2,4-Trimethylbenze	
	ļ		-	+-	+	51,3,5-Trimethylbenze	ne
		┼	+	-		So-Xylene	
	20	-	╁┈	+-	+-	Sm-, p-Xylene	
	3	5		2	<u> </u>	≾ Total Xylene	

Note: Values in **bold** represent detected analytes.

\*EPA January 1997.

\*Analysis request/chain-of-custody record.

EPA = U.S. Environmental Protection Agency.

ER = Environmental Protection Agency.

It = Foot (teet).

ID = Identification.

ID = Identification.

I = Foot (teet).

ID = Identification.

I = Rot analyzed.

NO () = Method detection Limit.

NA = Not analyzed.

NO () = Not detected above the MDL, shown in parentheses.

NR = Not detected above the MDL, shown in parentheses.

STL = Severn Trent Laboratories.

SV = Soil vapor.

VOC = Volatile organic compound.

VW = Vapor well.