

ENVIRONMENTAL COST MANAGEMENT, INC. Managing Cost and Liability

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9:25 am, May 26, 2009

Alameda County
Environmental Health

May 19, 2009

Jerry Wickham, PG Alameda County Health Care Services Agency Environmental Health Services 1131 Harbor Bay Parkway, Suite 250 Alameda, CA 94502-6577

Re: Draft Corrective Action Plan

Carnation Dairy, 1310 14th Street, Oakland, CA

Fuel Leak Case No. RO0000018 and Geotracker Global ID T0600100262

Dear Mr. Wickham:

On behalf of Nestlé USA, Inc. (Nestlé), Environmental Cost Management, Inc. (ECM) has prepared this *Draft Corrective Action Plan* (CAP) for the site located at 1310 14th Street in Oakland, California.

This report is submitted as requested in the letter from the ACHS dated March 4, 2009. As requested in the letter from ACHS, this draft CAP Report addresses the provisions contained in section 2725 of the UST regulations (CCR, Title 23, Chapter 16) with regard to corrective action requirements. The draft CAP makes use of data and analyses previously presented in the November 2008 *Revised Site Conceptual Model Report* and subsequent February 2009 response to comments letter from ECM to ACHS.

Should you have any questions regarding this Draft Corrective Action Plan, please call me at (415) 282-1979.

Perjury Statement

I declare, under penalty of perjury, that the information and/or recommendations contained in the attached document or report is true and correct to the best of my knowledge.

Brent Searcy, P.E.

Senior Engineer

Environmental Cost Management, Inc.

Enclosure: Draft Corrective Action Report

Cc: Mike Desso, Nestlé USA (CD copy)

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Report to:

Nestlé USA, Inc. 800 North Brand Boulevard Glendale, California 91203

Draft Corrective Action Plan Former Nestlé USA, Inc. Facility 1310 14th Street, Oakland, CA

May 19, 2009

Prepared By:



ENVIRONMENTAL COST MANAGEMENT, INC.

Managing Cost and Liability

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Date: 05/19/09

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1 INTRODUCTION

On behalf of Nestlé USA, Inc. (Nestlé), Environnemental Cost Management, Inc. (ECM) prepared this Draft Corrective Action Plan (CAP) for the former Nestlé facility (Site) once located at 1310 14th Street, Oakland, California (Figure 1). This report responds to a request contained in a letter from the Alameda County Health Care Service's (ACHS) dated March 4, 2009. The March 4, 2009 letter acknowledged ECM's response to comments on the November 4, 2008 Revised Site Conceptual Model (SCM) and requested submission of a draft Corrective Action Plan to ACHS.

This draft CAP relies on the compilation and analysis of site investigation and remediation data as documented in the November 4, 2008 Revised SCM Report and the January 2001 Comprehensive Site Characterization Report¹.

2 SITE HISTORY

2.1 Operational History

This draft CAP addresses the northwest corner of 1310 14th Street, Oakland, California, indicated as "NW Parcel" in Figure 1 (Site), consistent with directives from ACHS. The ACHS issued closure letters for other portions of the property. The terms "facility" or "property" are used to discuss the full property, including all four parcels.

The former Nestlé facility, originally constructed by American Creamery in 1915, was used to manufacture ice cream and packaged milk. Carnation purchased the property in 1929 and made additions and improvements between 1946 and 1973 for dairy product processing and distribution. Nestlé USA, Inc. assumed operation of the property following its purchase of Carnation in 1985. Nestlé ceased operations at the property in 1991².

While it was operational, the facility was used for the distribution of ice cream and packaged fresh milk by trucks. The delivery trucks were fueled at dispensers near service bays located at the northwest corner and were repaired and maintained at the facility.

A chronological summary of historical operations at the facility and remedial actions and at the Site is provided below. In addition, Appendix A provides a series of historical aerial photos that illustrate changes in the development of the facility over time.

<u>1915 - 1979</u>

- Original facilities constructed by American Creamery in 1915.
- Facilities were further developed for ice cream manufacturing and distribution and milk packaging activities by Carnation between 1946 and 1973.
- Following development by Carnation, facilities included food processing equipment, large cooler/freezer rooms, and 5 underground storage tanks (USTs; 2 gasoline, 2 diesel, 1 waste oil) for delivery vehicles.

1980s

- Ice cream manufacturing and distribution activities declined in the late 1980s.
- In early 1988, ice cream and milk distribution ceased.

- In December 1988 and January 1989, five USTs were excavated and removed from the Site.
- Free product was observed in the tank excavation area during UST removal.
- 1,200 cubic yards of soil removed during UST excavation, treated on-site, and replaced in excavation.

1990s

- Additional remediation efforts were implemented in the early 1990s.
- A SVE system operated in the former UST area from January 1994 through 1995, removing approximately 34,000 pounds of hydrocarbons.
- A multi-phase extraction system operated from August 1997 through June 2000, removing 10,875 pounds of hydrocarbons.
- Active remediation terminated in November 1999, with the concurrence of ACHA and the RWQCB, in response to reductions in LPH measurements.
- Nestlé monitored a network of 11 monitoring wells for petroleum and HVOCs in groundwater semi-annually for two years, under the direction of the ACHA and the RWQCB.
- A Risk Based Corrective Action (RBCA) analysis addressed residual concentrations remaining on-site.

2000s

- Operation of the multi-phase extraction system terminated in June 2000.
- The ACHA accepted the RBCA, a Soil Management Plan (SMP), and a Deed Restriction by July 2000.
- The RBCA analysis concluded that no significant risk to human health exists as a result of residual chemicals in soil or groundwater for all applicable exposure pathways.
- The Risk Management Plan (RMP) and deed restriction were implemented to protect against any possible direct exposure routes to future construction workers.
- Nestlé sold the Site to Encinal 14th Street, LLC in July 2000, subject to the RBCA, the RMP and the legally binding deed restriction.
- Nestlé submitted a Comprehensive Site Characterization Report to ACHA in January 2001 in support of its request for Site closure.
- Nestlé submitted a Request for Case Closure to ACHA in February 2002.
- All unused wells were properly abandoned in December 2002, with approval from the ACHA.
- Nestlé requested Site closure with the submittal of the semiannual groundwater monitoring report dated February 23, 2005.
- Nestlé submitted Site closure request follow-up letters to the ACHA on June 12, 2006 and June 15, 2007.
- In response to ACHS' September 28, 2007 directive, Nestlé performed and submitted results of additional field sampling in March and July 2008.
- By letter dated August 8, 2008, ACHS established a separate case for the northwest portion of the property ("Parcel B"), thus separating regulatory oversight of the northwest portion, which is the subject of this report, from the other three subdivided parcels of the property.

2.2 On-Site Structures and Features

The underground storage tanks (USTs) listed below stored fuel for the operations on-site. :

- One used-oil tank (1,000-gallon capacity),
- Two gasoline tanks (10,000-gallon capacity each), and

Two diesel fuel tanks (12,000-gallon capacity each).

The fuel system included underground piping that connected the USTs to the dispensers outside of the service bays. Figure 2 shows the locations of the former USTs and piping.

Figure 2 illustrates the primary features located within the northwestern portion of the facility. The locations of the previous USTs and associated subsurface piping are indicated in Figure 2, as well as the footprint of the existing L-shaped (formerly truck maintenance) building, which extends along the northern and western edges of the property. This building is an open structure approximately 29,000 square feet in size and contains multiple roll-up vehicle access doors opening to the interior of the property.

ECM has searched for, but was unable to locate, final construction drawings for the L-shaped truck maintenance. Historic coring and drilling activities indicate that the concrete slab foundation varies in thickness from 4 to 8 inches. Building footings likely exist beneath the exterior and load bearing walls. The depth of these footings is unknown, although construction codes and practices at the approximate time of construction suggest a likely depth of 2 to 15 feet below grade, depending on the load distribution technique implemented. The possible presence of these load-bearing footings and/or consolidated soils beneath these footings may form a downgradient barrier to shallow groundwater flow in the subsurface along the northwestern edge of the Site boundary (see Section 6 for further discussion).

To better understand the distribution and movement of chemicals of concern (COCs), Subdynamic Corporation conducted a comprehensive survey of subsurface utility corridors in November 2007. The findings of this in-field survey are presented within the *Revised SCM Report*³. The results of this subsurface conduit survey confirmed the existence of on- and off-site utility trenches. All on-site utilities were either abandoned, or the utility corridors no longer convey, active utilities. Section 3 addresses the utility survey findings relative to the potential influence on subsurface distribution and transport of COCs.

The on-site utility survey also addresses concerns raised in the September 2007 ACHS directive regarding possible residual diary fat or detergent impacts in the subsurface. Information collected regarding the abandoned and non-operational sewer and/or storm drains on the site indicates there are no active sources of such substances within utility conduits beneath the Site.

3 GEOLOGY AND HYDROGEOLOGY

3.1 Regional Geology

The Site is located in an area of the San Francisco Bay region generally underlain by bay mud, the Merrit Sand, and Younger and Older Alluvium⁴. The San Francisco Bay is located approximately 2 miles to the west of the Site. The San Andreas Fault is located approximately 9 miles west, and the Calaveras/Hayward fault zone lies approximately 8 miles east. Shallow soils in this area of the eastern shore of the San Francisco Bay generally consist of clayey or silty sands. Due to its limited extent and thickness, the saturated portion of the Merritt Sand in this immediate region is not considered a drinking water resource⁵.

3.2 Site-Specific Geology and Hydrogeology

Soil borings extended and logged from 8 to 25 feet below ground surface (ft. bgs) provide geologic data for a thorough characterization of subsurface geology^{6,7,8}. Soils are predominantly well-

sorted sands (SP), with discontinuous areas of clayey or silty sands (SC, SM). Hydraulic conductivities of these soils have been estimated at approximately 30 ft/day⁹. Boring log information collected during historical soil boring investigations have been used in developing lithologic cross sections, as shown in Figures 3 through 5.

Groundwater has historically been encountered at depths ranging from approximately 5 to 12 ft. bgs. Most recently, static groundwater elevations encountered in borings extended during the May 2008 site investigation were in the 9 to 10 ft. bgs range. Groundwater generally flows to the northnorthwest (toward 16th Street) at an average gradient of 0.0027ft/ft¹⁰. Figure 6 shows the variation of groundwater flow directions from December 1995 through April 2000, and confirms the observation that the groundwater flow direction has been generally consistent and does not change significantly throughout the annual hydraulic cycle.

4 SITE CHARACTERIZATION AND REMEDIATION ACTIVITIES

4.1 Soil Gas Characterization

Soil gas samples were collected across the Site during two separate studies (in August 1999 and May 2008) to document the existence of any VOCs in shallow soil gas. Both soil gas investigations addressed soil vapor conditions following active remediation activities (see Section 5 for details of remedial activities). Tables 1a and 1b detail the results of these soil gas sampling investigations.

4.1.1 August 1999 Investigation

The August 1999 investigation supported the proposed shut down of the multi-phase extraction system and planned regulatory closure discussions. Table 1a documents soil gas concentrations from 15 soil borings sampled at 3 ft. bgs using direct-push equipment and summa canisters for vapor collection. Protocols for the collection of soil gas samples are provided in Appendix E of the January 2001 *Comprehensive Site Characterization*¹¹. Soil gas sampling locations represented areas thought to overlie the highest groundwater contaminant levels and the perimeter and downgradient edge of the soil and groundwater plume (see Appendix B, Figure 9). Active operation of the existing multi-phase extraction system ceased 24 hours prior to the soil gas sampling and remained off until the conclusion of soil gas sampling.

All soil gas samples were analyzed using USEPA Compendium Method TO-14/TO-14A for volatile organic compounds. Benzene and other fuel hydrocarbon compounds were measured in the soil vapor samples. Benzene concentrations ranged from 0.91 to 9,900 ppbv (Table 1a). Non-fuel hydrocarbon compounds detected in the soil vapor were acetone (10–260 ppbv), ethanol (23–1,400 ppbv), Freon-12 (0.93–630 ppbv), and tetrachloroethene (1.2–160 ppbv). 1,2-DCA was not detected in the soil vapor at or above the respective laboratory reporting limits.

4.1.2 May 2008 Investigation

The May 2008 investigation addressed the ACHS' directive dated September 28, 2007. Soil gas concentrations from this investigation are reported in Table 1b.

Soil gas sampling was performed as per the protocol recommended by the Los Angeles Regional Water Quality Control Board (LARWQCB)/California Department of Toxic Substances Control (DTSC) Advisory for Active Soil Gas Investigations¹². A California certified on-site mobile lab

(TEG, Inc) with full gas chromatography (GC) and mass spectrometry (MS) capabilities analyzed all soil gas samples immediately following sample collection ¹³.

Soil gas samples were collected from 12 sampling locations at a depth of 5 ft. bgs. Soil gas samples from each boring were analyzed according to the following methods:

- EPA 8015M for gasoline (TPH-g) and diesel (TPH-d) range organics, and
- EPA method 8260B for BTEX and VOCs.

Five of the 12 sampling locations exhibited detectable concentrations of hydrocarbons or VOC constituents. TPH-g concentrations ranged from below the laboratory reporting limit of 50 μ g/L to 2,600 μ g/L at boring SB-22. TPH-d was not detected in any soil gas samples. Benzene was detected at two of the 12 sampling locations, with the highest concentration at 40 μ g/L at boring SB-22. Ethylbenzene, toluene, and xylenes were detected at 3 of the 12 sampling locations. No detections of 1,2-DCA were reported in any of the soil gas samples. Detections of dichlorodifluoromethane (i.e., Freon-12) characterized soil gas samples from two soil borings (SB-22 and SB-26).

4.2 Soil Characterization

Three separate historical field investigations in 1991, 1999, and 2008 involved subsurface soil sampling and analysis. Results from these soil investigations are reported in Table 2. Soil data gathered after active remediation activities were integrated into the three dimensional model of hydrocarbon impacts presented in the Revised SCM Report (see Appendix B, Figures 16 through 27).

4.2.1 1991 Soil Sampling Investigation

An initial investigation conducted in 1991 yielded soil samples from 5, 10, 12.5, 15, and 20 ft. bgs (feet below ground surface) as documented in Table 2. The highest TPH-g and TPH-d concentrations were measured in the area immediately to the north and northwest of the former UST area, generally at 10 ft. bgs¹⁴.

The soil boring data indicated that TPH-g impacts were mainly limited to the 5 to 15 ft. bgs interval. The maximum TPH-g concentration at 5 ft. bgs was 2,500 mg/kg. At 10 ft. bgs, the maximum was 10,000 mg/kg. By 15 ft. bgs, the maximum concentration dropped to 1,900 mg/kg, and at 20 ft. bgs, the maximum TPH-g level decreased to 260 mg/kg.

The TPH-d distribution followed a similar pattern. The maximum TPH-d impact at 5 ft. bgs was 470 mg/kg. At 10 ft. bgs, the maximum increased to 940 mg/kg. By 20 ft. bgs, the maximum TPH-d level dropped to 23 mg/kg.

4.2.2 1999 Soil Sampling Investigation

An August 1999 investigation was performed to collect soil samples and characterize hydrocarbon impacts primarily at the site perimeter to the north and west of the former UST area. The August 1999 investigation collected soil samples from 13 soil borings (SB1 through SB15) (see Table 2 and Appendix B, Figure 9). The locations of the borings represented subsurface conditions in the area downgradient (NNW) of the UST source areas (see Figure 2) and assessed impacts beneath the footprint of the L-shaped building on the northwest edge of the property. Remediation equipment and aboveground piping restricted boring locations to the south side of the piping in the

former repair garage bay (SB6) and to the end bay at the east side of the building (SB14). Results of soil samples collected in August 1999 are shown in Table 2.

Low levels (at or below 2.7 micrograms per kilogram [ug/kg]) of 1,2-DCA, toluene, ethylbenzene and total xylenes characterized the 3.5 to 4.0 ft. bgs interval. Maximum TPH-d was 1,200 mg/kg in this interval.

Sporadic concentrations of hydrocarbons and halogenated VOCs (HVOCs) characterized the soil at the water table (6.5 to 7.0 ft. bgs). 1,2-DCA concentrations ranged from below laboratory reporting limits at multiple locations to 430 μ g/kg at SB6 within this vertical interval. Elevated concentrations of hydrocarbons were measured at sampling locations SB3, SB6, SB8, and SB12, and SB14, with TPH-g ranging from 2.25 to 10,100 mg/kg, TPH-d ranging from 60 to 2,900 mg/kg, and benzene ranging from 0.07 to 76 mg/kg for benzene.

4.2.3 2008 Soil Sampling Investigation

Following requests from the ACHS for additional delineation of potential residual hydrocarbons and polychlorinated biphenyls (PCBs) in the subsurface at the Site¹⁵, a May 2008 soil boring investigation was performed. This investigation collected soil samples from 15 borings (SB16 through SB27 and PCB4 through PCB6) (see Appendix B, Figure 10). Soil borings were located in areas of suspected residual hydrocarbon impacts and at the perimeter of impacted area to improve delineation of current residual COCs present beneath the Site.

For the May 2008 soil sampling investigation, soil borings were advanced using a 2-inch diameter direct-push Geoprobe® coring method and logged (ECC, 2008). At each boring, a soil sample was collected from immediately above the first-encountered saturated zone, typically between 6 and 10 ft. bgs, as documented in Table 2. Per prior agreement with the ACHS, soil boring SB-17 was to be extended up to 30 ft. bgs, collecting soil samples every 5 feet. The Geoprobe® was unable to drive sampling rods through saturated and consolidated sands encountered in this boring at approximately 20 ft. bgs. Samples were, therefore, collected and analyzed from 5, 10, 15, and 20 ft. bgs at soil boring SB-17.

Soil samples were analyzed for TPH-g, TPH-d, and TPH as motor oil (TPH-mo) via EPA Method 8015B modified and BTEX via EPA method 8260B. Soil samples were also analyzed for 1,2-DCA via EPA method 8260B and PCBs via EPA Method 8082 and some locations (see Table 2). Duplicate soil samples were collected to validate and verify soil sampling consistency and method.

TPH-g, TPH-d, and TPH-mo results were consistent with the impacts identified in previous soil and groundwater sampling efforts. Elevated levels of hydrocarbons were detected at borings located to the north and northwest of the former UST locations (see Figure 2). TPH-g ranged from ND up to 12,000 mg/kg. TPH-d ranged from ND up to 17,000 mg/kg. TPH-mo ranged from ND up to 13,000 mg/kg. All three maximum TPH fraction results came from SB-17 at 10 to 10.5 ft. bgs vertical interval, as did the highest benzene concentration of 140 mg/kg. TPH and benzene concentrations in soil samples collected above (8-8.5 ft. bgs) and below (15-15.5 ft. bgs) this interval were typically at least one order of magnitude lower than the preceding interval. All the most elevated results were reported from borings north and northwest of the former UST locations (see Table 2 and Appendix B, Figures 16 through 27). 1,2-DCA was not detected above detection limit at any of the soil boring sampling locations. Results from SB17 confirmed the absence of BTEX constituents below 10 ft. bgs and TPH-g, TPH-d, and TPH-mo below 15 ft. bgs (see Table 2 and Appendix B, Figures 16 through 27).

Per a request made by ACHS in the September 28, 2007 directive, soil samples were analyzed for PCBs at eight borings at depths ranging from 8.5 to 9.5 ft. bgs. No PCBs were detected above laboratory detection limits. These results are consistent with prior reports which stated that there were no sources of PCBs at the Site from dairy processing or distribution activities.

4.3 Groundwater Characterization

As many as 65 monitoring wells were sampled quarterly and semi-annually to characterize dissolved hydrocarbons and VOCs in groundwater between 1994 and 2004. The number of well monitored was reduced in 2004, consistent with ACHS approval in November 2002. Between December 2002 and late 2004, 11 monitoring wells were sampled. Grab groundwater samples were collected during the May 2008 soil boring investigation, as requested by the ACHS in the September 28, 2007 directive. Cumulative groundwater monitoring results (1993 through 2008) are shown in Table 3.

Historical groundwater results show that positive TPH-g and benzene detections are generally limited to the area immediately downgradient (NNW) of the former USTs. Dissolved hydrocarbons have historically existed immediately downgradient of the former UST location at wells PR50, PR53, and PR64. Historical and recent (May 2008) concentrations of benzene and TPH-g, as illustrated in the revised SCM Report, attenuate in the downgradient direction toward 16th Street. Monitoring wells farther downgradient from MW-26 (located at the southern edge of 16th Street) did not contain detectable TPH-g and benzene from 2002 through 2004, as shown in Table 3. Wells sampled to the west and east (cross-gradient) and south (upgradient) of the former UST area have not exhibited detectable TPH-g or BTEX, with the exception of 0.60 ug/L benzene and 0.90 ug/L ethylbenzene in upgradient well MW-33.

Historical groundwater sampling results and estimated TPH-g, TPH-d, and benzene isoconcentration contours are presented for sampling events in 2000, 2004, and 2008 in the *Revised SCM Report* ¹⁶. These plots indicate the stable condition of hydrocarbon constituents since the termination of active remedial activities (i.e., multi-phase extraction) in the second quarter of 2000.

HVOCs in groundwater have historically existed in the area immediately downgradient of the former UST locations and at lower levels further downgradient in the area of 16^{th} Street. The predominant HVOC is 1,2-DCA, which has a historical maximum concentration of 2,200 μ g/L found in a grab groundwater sample from boring SB18. Post-remediation monitoring revealed a maximum 1,2-DCA level of 83 μ g/L in downgradient wells within 16^{th} Street. Groundwater monitoring data do not indicate any predominant or persistent source of HVOCs.

Per regulatory requests made by ACHS in the September 28, 2007 directive, groundwater samples were analyzed for PCBs at eight borings during the May 2008 investigation. Laboratory reports show that no PCBs were detected ¹⁷. The absence of PCB detections in groundwater confirms that PCBs are not present at the Site.

4.4 Liquid Phase Hydrocarbon (LPH) Characterization

LPH has been present in the area of the USTs and maintenance bays since UST removal in 1988. More than 50 wells monitored the LPH since 1989. As indicated in Table 4, LPH levels in wells that historically reported the most significant LPH levels showed the most significant reductions in LPH thickness as a result of product removal by various technologies implemented at the site between 1989 and 2001. Following the cessation of regular LPH monitoring in August 2001, semi-

annual groundwater sampling was performed at 11 on- and off-site wells from November 2002 through November 2004. LPH was not observed in any of these 11 monitoring wells during these semi-annual sampling events.

Historical LPH gauging through August 2001 shows that LPH has not migrated following the termination of active remediation in the second quarter of 2000. This conclusion is based on the following facts:

- The number and location of wells containing LPH remained relatively constant prior to LPH recovery initiated in late 1997.
- The number of wells containing measurable LPH decreased after multi-phase extraction was initiated in late 1997.
- LPH has not been detected in any well outside the group of wells that have historically contained measurable LPH thickness, and
- LPH was never observed in any of the of the 11 post-remediation groundwater monitoring wells sampled from November 2002 through November 2004.

4.5 Remediation Activities

Four fuel USTs and associated dispensers and piping were excavated on December 19, 1988. One 1,000-gallon used-oil tank was removed on January 12, 1989. The former tank and fuel line excavation areas are shown in Figure 2. Each of the removal actions was documented in an 1989 report 18. Removal of the tanks and piping stopped the primary source of COC release to the subsurface.

Between January and March 1989, 1,200 cubic yards of soil were removed in the area of the former tanks and lines. This soil was treated onsite and placed back into the excavation. Following removal of the five USTs in December 1988 and January 1989, LPH skimming began in January 1989. LPH skimming removed approximately 1,800 gallons of LPH¹⁹.

A SVE system began treating residual hydrocarbons in 1994 in the vadose zone immediately to the north and northwest of the former UST area²⁰. This system operated until December 1995 and removed approximately 5,200 pounds of hydrocarbons.

A multiphase extraction system addressed both LPH and dissolved-phase hydrocarbons in the subsurface, starting in August 1997²¹. This system extracted LPH entrained above the groundwater table and floating directly on the groundwater table, as well as groundwater itself and subsurface vapors in areas with high dissolved hydrocarbon concentrations. This system operated until June 2000 and removed approximately 10,875 pounds of hydrocarbons and reduced residual LPH levels at the Site significantly, as documented in the *Revised SCM Report* (ECM, 2008)

LPH thickness in monitoring wells declined until the cessation of dual phase extraction activities. In November 1999, ACHS and RWQCB agreed that the Site conditions satisfied the criteria for consideration of closure as a low risk site with respect to petroleum hydrocarbons and residual HVOCs, and that the Site did not warrant further active remediation²². Multiphase extraction ceased in June 2000. Eleven groundwater monitoring wells were monitored on a semi-annual basis between 2002 and 2004, as directed by ACHS.

Since the cessation of active remediation and the completion of the required semi-annual confirmation groundwater monitoring, Nestlé made several requests that the site be considered for a no further action or case closure designation. Total costs due to investigation and remediation activities performed at the site since 1988 are approximated at almost 10 million dollars.

5 DEVELOPMENT OF CORRECTIVE ACTION PLAN ALTERNATIVES

In this section, ECM assesses several corrective action plan (CAP) alternatives, including descriptions of the targeted cleanup areas and scope for each alternative. ECM considered the following alternatives:

- Targeted Excavation of Impacted Soils
- 2. Excavation of all Soils Above TPH Tier I ESLs
- 3. Soil Vapor Extraction / Bioventing
- 4. In-situ Chemical Oxidation
- Institutional Controls

Each of the above alternatives is evaluated in terms of advantages, disadvantages, estimated costs, and the likelihood of success in reaching cleanup objectives for the site. Cleanup objectives are discussed below. Table 5 summarizes each alternative and the primary factors considered in this evaluation.

The May 2009 Screening Health Risk Evaluation (see Appendix A) was developed to assess and provide quantification of risks associated with residual COCs at the site and to assist in establishing cleanup objectives. Table 6, developed in conjunction with the risk evaluation process, indicates the exposure pathways which are considered complete in terms of potential exposure risks associated with residual COCs remaining at the site. The evaluation of the corrective action plan alternatives assesses each alternative and its likely success in providing cleanup of the site to levels such that these exposure risks are below acceptable (per USEPA guidance²³) levels.

Costs associated with each alternative have also been estimated through a thorough engineering cost assessment. An estimated cost for each alternative is calculated (see Table 5) considering all likely activities and system operation and maintenance if required associated with each alternative. Costs for the development of the May 2009 *Risk Evaluation* (see Appendix A) used in assessing these CAP alternatives are also included for each alternative evaluated. The itemized details of these cost estimates are available upon request.

Specific considerations and assumptions made in the development of the various CAP alternatives are presented in Sections 5.1 through 5.6 below. Section 6 discusses the advantages and disadvantages for each CAP alternative and recommends a corrective action plan for the Site. A summary of the primary features, advantages, and disadvantages of each CAP alternative is provided in Table 5.

Where necessary, and to establish the extent of areas to be targeted for remedial actions, May 2008 Tier 1 Environmental Screening Levels (ESLs)²⁴ drive the scope of the various alternatives. RWQCB guidance shows the following ESL criteria for shallow soil. ECM used these ESLs to target areas for the various remedial alternatives.

Environmental Screening Levels for Soil (RWQCB, 2008)

	Tier 1 ESLs (Table B)	Urban Area Eco- toxicity Criteria (Table B-2)	Odor Index (Table H-2)	Human Health Direct Exposure, Commercial/ Industrial Worker (Table K-2)	Direct Exposure, Construction / Trench Worker (Table K-3)	Vapor Intrusion into Buildings (Table B- 2)	Groundwate r Protection, Soil Leaching Concerns
Chemical	Soil ESL	Soil	Soil	Soil	Soil	Soil	Soil
	(mg/kg)	(mg/kg)	(mg/kg)	(mg/kg)	(mg/kg)	(mg/kg)	(mg/kg)
TPHd (nc)	180	-	355	450	4,200	*	180
TPHg (nc)	180	-	13,600	450	4,200	*	180
Benzene (c)	0.27	25.00	63.3	0.27	12.0	*	2.0
Toluene (nc)	9.3	-	3.5	210	650	*	9.3
Ethyl- benzene (c)	4.7	-	22.2	5.0	210	*	4.7
Xylenes (Total)(nc)	11.0	-	6.0	100	420	*	11.0
MTBE (c)	8.4	-	1,880	65	2,800	*	8.4

NOTE:

Table adapted from (RWQCB, 2008)

(c) carcinogenic effects.

(nc) noncarcinogen

Tables from ESLs where groundwater is not a current or potential source of drinking water and for commercial/industrial land use only.

- Not available.

Remediation alternatives focus on residual total petroleum hydrocarbon as gasoline (TPH-g) impacts at the site. Site characterization data (see Table 2) indicates that other COCs (TPH-d, benzene, etc.) impacts are coincident, or smaller in their extent, with TPH-g impacts to soil. Thus, TPH-g impacts were used when developing the extent of targeted areas for each CAP alternative. In addition, the May 2009 *Screening Health Risk Evaluation* (see Appendix A) is used to assess and provide quantification of risks associated with residual hydrocarbons.

This Screening Health Risk Evaluation provides a complete assessment of likely on- and offsite receptors. The risk evaluation notes the following complete exposure routes and receptors (see Table 6):

- vapor intrusion from soil gas to indoor air of the onsite commercial building;
- volatilization from soil gas to onsite outdoor; and

^{*} use soil gas.

 advective transport from onsite outdoor air to the outdoor or indoor air of offsite residential land uses

These potential receptors are considered when assessing the CAP alternatives in Sections 6.1.1 through 6.1.5 below. As noted in the Risk Evaluation (Appendix A) and the Revised SCM Report (ECM, 2008), there is no current use of groundwater from beneath the site as potable water supply. Well surveys conducted in 1997 and 2000 found no municipal or private groundwater supply wells located within a ¼-mile of the site 25. The current Deed Restriction (Appendix C) recorded for the site also prohibits the installation of any wells on site for the purpose of extracting groundwater for any potable or non-potable use.

Nestlé understands that the ACHS may request, as part the implementation of any of the selected CAP alternative, that the existing Deed Restriction be re-filed with the Alameda County Health Care Service office such that the document is recorded according to current protocols within the County.

In considering the following CAP alternatives, cleanup goals with respect to petroleum hydrocarbons were considered. As documented in the *Screening Health Risk Evaluation* (Appendix A), complete exposure routes for COCs at the site include exposure by onsite commercial workers to COCs in indoor and outdoor air. Cleanup goals for COC vapor concentrations which are protective of a cumulative carcinogenic risk of 10⁻⁵ or less (typical commercial/industrial point of departure; see Appendix A) are detailed in Table 16 of the attached *Screening Health Risk Evaluation* and summarized below.

Cleanup Goals for COC Vapor Concentrations (see Appendix A)

	Vapor Phase Cleanup Goals				
Chemical	(based on maximum concentrations in vapor at 5 ft. bgs protective of an indoor air cancer risk of 10 ⁻⁵ ; see Table 16, Appendix A)				
	(μg/m³)				
Benzene	1.0 x 10 ⁵				
Toluene	3.2 x 10 ⁷				
Ethyl-benzene	1.4 x 10 ⁶				
Xylenes (Total)	1.1 x 10 ⁷				
1,3-Butadiene	6.5 x 10 ³				
MTBE	9.1 x 10 ⁶				

Cleanup levels for other media (soil, groundwater) are dependent on complete exposure pathways and, with the exception to direct exposure to subsurface soil or groundwater by on-site intrusive construction workers, these pathways are documented as incomplete. The case of direct exposure to above cleanup-level concentrations in soil and groundwater are addressed through the existence of the Deed Restriction (Appendix C) and RMP (Appendix D) for the site.

5.1 Alternative 1: Targeted Excavation of Impacted Soils

Soil excavation is a rapid, conventional and reliable means of source reduction. Limited over-excavation in targeted zones of TPH-g contamination may pose a viable option for some specific areas. For this alternative, ECM selected soil excavation in areas where three-dimensional interpolation of historical soil sampling data indicates that TPH-g concentrations exceed 4,200 mg/kg. This contaminant level represents the ESL for direct exposure to a construction worker and is an approximate gauge of zones where an aggressive remedial action such as excavation may be most beneficial.

Soil excavation under this alternative involves removal of the surface cap which includes the existing building floor and foundation in the area where excavation is to occur, soil removal, potential dewatering, loading for transport to an offsite disposal facility, import and placement of clean soil, and compaction. When excavations encounter shallow water tables, dewatering can be necessary to facilitate further excavation and fill placement. Persistent constituents may require more extensive dewatering to flush the excavation, requiring an established pump-and-treat remedy. Excavation with dewatering requires favorable site conditions, most notably a shallow water table and a thin aquifer.

Groundwater has historically been encountered at depths ranging from approximately 5 to 12 ft. bgs. Static groundwater elevations encountered in borings extended during the May 2008 site investigation were in the 9 to 10 ft. bgs range. Groundwater generally flows to the north-northwest (toward 16th Street) at an average gradient of 0.0027ft/ft. The groundwater flow direction has been generally consistent and does not change significantly throughout the annual hydraulic cycle²⁶.

Excavation often extends one or two feet below the lowest observed water table elevation. Nearby stockpiles or roll-off containers can store the excavation spoils pending characterization and disposal arrangements. In the case of TPH constituents, local Class II landfills will normally accept impacted soil as non-hazardous, special waste. A mobile lab could help with spot excavations and stockpile characterization, if needed.

Soil sampling data is used to delineate the excavation area and to demonstrate successful removal. An understanding of likely source areas (near the former USTs and downgradient locations), repeated groundwater elevation measurements, and sample analyses provide additional basis upon which to determine potential excavation locations.

Estimated excavation volumes of soils above RWQCB-SF direct exposure limits (see Section 5 above) were derived using Rockworks Modeling Software and assume a universal depth of excavation of 15 ft. bgs in establishing the excavation boundaries. Soils were assumed to be predominately well-sorted sands for estimating purposes, with an estimated density of approximately 120 lbs/ft³. Rockworks modeling produced an estimated soil excavation volume of approximately 14,600 ft³ (541.7 cubic yards) to excavate soil with TPH-g levels above 4,200mg/kg (see Figure 7).

5.2 Alternative 2: Excavation of all Soils above TPH Tier I ESLs

As discussed in Section 6.3, soil excavation is a rapid, conventional and reliable means of source reduction. For this alternative, ECM selected soil excavation in areas where modeling indicates that TPH-g concentrations exceed 180 mg/kg (the RWQCB Tier 1 ESL, see Section 5 above).

Excavation to Tier I levels requires similar equipment and procedures as those discussed in Section 6.3, but results in a larger excavation volume. This alternative requires demolition of the existing L-shaped 29,000 square foot building (see Figure 2), engineered dismantling of all building related structures, excavation shoring to protect adjacent sidewalks and structures, soil removal, potential dewatering, loading soils for transport to an offsite disposal facility, import and placement of clean soil for fill and compaction. As this excavation alternative necessitates the demolition of the existing building, costs for reconstruction of a building of similar size and dimension are also included for this alternative.

Estimated excavation volumes were derived using Rockworks Modeling Software and assume a universal depth of excavation of 15 ft. bgs in establishing the excavation boundaries. Soils were assumed to be predominately well-sorted sands for estimating purposes, with an estimated density of approximately 120 lbs/ft³. Rockworks modeling produced an estimated total soil excavation volume of approximately 110,500 ft³ (4,090 cubic yards) to excavate soil with TPH-g levels above 180mg/kg. The small volume of soils located near SB12 is approximately 1,300ft³ (48 cubic yards) and is included in the total estimates soil excavation volume (Figure 8).

5.3 Alternative 3: Soil Vapor Extraction / Bioventing

The Soil Vapor Extraction (SVE) or Bioventing alternative target soils for remediation where TPH-g levels exceed the Tier I ESL. The following subsections discuss the specific options for SVE and Bioventing in more detail. Figure 9 shows the proposed SVE or Bioventing Layout.

5.3.1 Soil Vapor Extraction (SVE)

For SVE (including high-vacuum extraction (HVE)) a vacuum blower pumps air from the unsaturated part of the vadose zone, flushing out constituents in vapor form. SVE is the air-flow equivalent of groundwater pump and treat. Installation of an SVE system involves air permitting, extraction well installation, system construction, performance monitoring, and operations and maintenance.

The SVE process capitalizes on the volatility of constituents to partition from the aqueous and soil-adsorbed phases into the vapor phase. Thus, SVE may treat contamination in both soil and groundwater. These mechanisms work best on constituents with low water solubility, a high Henry's Law constant, and a high vapor pressure. Gasoline may be considered to have a relatively high vapor pressure with its flash point being below typical room temperature. Pavement or other semi-permeable ground cover is often necessary to reduce short circuiting through the vadose zone to the atmosphere when applying SVE.

Extracted vapor typically requires treatment prior to discharge to the atmosphere to meet any Bay Area Air Quality Management District's air permitting requirements. The methods of treatment depend upon the concentrations of the various vapor constituents and include GAC adsorption, internal combustion engines, or thermal processes (catalytic and thermal oxidation).

5.3.2 Bioventing

Bioventing treats the vadose zone and may treat the uppermost 2 or 3 feet of the aquifer using an air injection system. Bioventing replenishes the fresh air supply in the subsurface formation around injection wells, enhancing remediation efforts in two ways:

- Exposing fuel impacts to fresh air, thereby promoting aerobic biodegradation and hence, in situ destruction of fuel hydrocarbons; and
- Smearing any remaining free product along the well borehole as the water level drops due
 to increased air pressure inside the injection well. The smear zone's higher exposure to air
 increases oxygen demand, thus accelerating the remediation of the smear zone and the
 free product within it.

5.4 Alternative 4: In-situ Chemical Oxidation

In-situ chemical oxidation (ISCO) treatment injects oxidizers into the subsurface to oxidize organic constituents in place. Oxidizers can include hydrogen peroxide, ozone, Fenton's Reagent (iron-catalyzed hydrogen peroxide), potassium permanganate, sodium persulfate, and other chemicals. The technology is implemented by drilling wells so that the oxidizer can be injected into the contaminated zone. ISCO treatment may reduce organic pollutant concentrations, biological oxygen demand (BOD), chemical oxygen demand (COD), and/or improve odor and color.

Successful application of ISCO is sensitive to site conditions such as natural organic matter content and hydrogeology. ISCO using chemicals like potassium permanganate and sodium persulfate require a slurry or solution of the reagent in water and injection of the mix into the subsurface in batches.

5.5 Alternative 5: Institutional Controls

This CAP alternative involves documentation of the risks associated with residual hydrocarbons remaining at the site, and the implementation of institutional controls required to address any exposure risks documented in this evaluation (Appendix A). Documentation of existing residual hydrocarbon impacts is complete and documented in the Revised Site Conceptual Model (SCM) and the *February 18, 2009 Response Letter to SCM Comments*, which provide the basis for the assessment of exposure risks in the *Screening Health Risk Evaluation* (see Appendix A).

Institutional controls are currently in place to address direct exposure risks pursuant to the Covenant and Environmental Restriction (i.e., a Deed Restriction) recorded in June 2000. The deed restriction controls land use and on-site activities and is binding on the current and future property owner. The deed restriction requires a surface cap to be maintained and notice to be provided to the regulatory agencies prior to disturbing subsurface soils or seeking changes to current land use restrictions at the Site.

Nestlé understands that the ACHS may request, as part of this proposed CAP alternative, that the existing Deed Restriction be re-filed with the Alameda County Health Care Service office such that the document is recorded according to current protocols within the County.

6 RECOMMENDED REMEDIAL ACTION ALTERNATIVE

This section provides a detailed evaluation of each CAP alternative and, based on evaluation of their technical feasibility, likelihood of success, and estimated cost, recommends the most feasible alternative for implementation.

6.1 Evaluation and Comparison of Viable Alternatives

Each of the alternatives described above have been considered in terms of advantages, drawbacks, likelihood of success, technical challenges, and estimated cost of implementation. Table 5 presents a summary of these considerations and provides an overview of the critical considerations assessed for each alternative.

6.1.1 Alternative 1: Targeted Excavation of Impacted Soils

CAP Alternative 1, soil excavation of targeted areas (areas of soil with THP-g or TPH-d above RWQCB-SF direct exposure limits) provides the following advantages:

- Provides direct removal of COC mass from the Site,
- The excavation of soils is likely to reduce risks of direct exposure associated with residual COCs, and
- No on-going discharge of treated groundwater or vapor is necessary

Excavation of soils cannot guarantee complete removal of targeted residual hydrocarbons. The typical limitations of the environmental sampling performed to date are such that defining excavation limits must rely on the interpretation and interpolation of a reasonable population of soil sampling data. Thus, post-excavation levels of residual hydrocarbons in soil and groundwater can be anticipated to be significantly reduced through targeted excavation, but complete removal of all hydrocarbons above direct exposure limits is not certain.

Excavation of soils under this alternative will also require disruption and replacement of the concrete and asphalt surface cap currently present at the site. The presence this surface cap is currently required by the existing Deed Restriction at the site, and its removal and appropriate replacement would need approval from the ACHS and/or RWQCB, as required by the terms of the Deed Restriction.

Costs of excavation of areas above direct exposure limits are estimated from \$478,600 to \$574,400, and could exceed this range depending on the groundwater levels and dewatering volumes required to reach the target depth of 15 ft. bgs across the soil excavation area.

6.1.2 Alternative 2: Excavation of all Soils above TPH Tier I ESLs

CAP Alternative 2, soil excavation of soils above Tier I ESLs (areas above RWQCB-SF commercial/industrial Tier I ESLs) provides similar advantages and drawbacks to Alternative 1 However, as illustrated in Figure 8 and detailed in Section 5.2, the total estimated soil volume required to remove the estimated area of soil impacts above Tier I ESLs (180 mg/L TPHg) is 110,500 ft³ (4,090 yds³). This is over 7 times the volume of soil estimated for excavation under Alternative 1.

In addition to the advantages and drawbacks of an excavation approach detailed for Alternative 1 above, the excavation of soils within areas estimated to contain TPH-g levels above Tier I ESLs would likely require the demolition of the existing L-shaped building located on the northwestern portion of the Site. Additional engineering controls and excavation shoring would likely be required to ensure stability of sidewalks and off-site structures along the northern edge of the property.

These engineering and structural considerations would result in significant additional costs for Alternative 2. As with Alternative 1, the complete removal of all soils above the targeted (Tier 1 ESLs) limits is not certain following excavation activities due to the necessary estimation and interpolation of impacted areas using all available soil sampling data from the Site (see Table 2 and Figure 8).

Costs of excavation of areas above direct exposure limits are estimated from \$2,856,600 to \$3,427,900, and could exceed this range depending on the groundwater levels and dewatering volumes required to reach the target depth of 15 ft. bgs across the soil excavation area. The significant increase in cost in comparison to Alternative 1 is primarily related to the need to demolish and replace the existing building and disposal and backfilling costs associated with the additional soil volumes under this alternative.

6.1.3 Alternative 3: Soil Vapor Extraction / Bioventing

Alternative 3, as detailed in Section 5.3 above, involves extraction and treatment of soil vapor from the subsurface areas of highest residual hydrocarbon impacts to soil as outlined in Figure 9. Bioventing of soils (see Section 5.3 above) is a means of promoting vadose zone air flow and enhancing in-situ biodegradation of hydrocarbons.

Soil vapor and/or bioventing at the Site offer the following advantages:

- SVE and/or bioventing provides for direct removal of vapor phase COCs from the subsurface (primarily vadose zone).
- SVE and/or bioventing can target areas of highest hydrocarbon concentrations in soil.
- SVE and/or bioventing Involves minimal disruption of existing concrete/asphalt cap and building.

Groundwater has been encountered at depths ranging from approximately 5 to 12 feet below ground surface (bgs) at the Site. Where shallow aquifers exist, SVE can lift the water table high enough to saturate the part of the vadose zone around the well where most of the airflow occurs, effectively cutting off airflow. SVE is generally less effective for sites where shallow groundwater is present. In May 2008, the recent monitoring event indicated that the vadose zone ranged from 9 to 10 feet in thickness, which is relatively shallow and may present challenges for removal of hydrocarbons entrained in soil below the groundwater surface.

A tight formation may restrict flow and require a higher well density. A thin vadose zone may allow short circuiting to atmosphere; thus, testing and sealing the surface with an impermeable material and seal coat may be necessary with SVE or bioventing.

Previous dual phase (high vacuum) extraction efforts implemented at the Site from August 1997 through June 2000 provide insight into the likely effectiveness of SVE for removal of hydrocarbon mass via extracted vapors. This dual phase system extracted hydrocarbons in groundwater and

vapor phases, over a three year period. Operation of the system was terminated, with regulatory approval, when ongoing groundwater and LPH monitoring (see Table 4) indicted that hydrocarbon levels in the areas of highest impact reached asymptotic levels and were no longer declining.

Given this experience, any newly installed extraction system likely result in a rapid return to this asymptotic condition. Outside of an initial and short-lived elevated rate of hydrocarbon mass removal during extraction system start-up, it is unlikely that operation of an SVE system would produce any significant and consistent reductions in hydrocarbon mass beneath the Site.

Estimated costs for the installation and operation (for one year) of a combined SVE and bioventing system are \$593,700 to \$712,400, depending on the number of extraction points and the mass of hydrocarbons requiring treatment in extracted vapors. Cost will increase if the operation of the SVE is continued beyond one year.

6.1.4 Alternative 4: In-situ Chemical Oxidation

Alternative 4, as detailed in Section 5.4 above, involves subsurface injection of chemical oxidants to effect the degradation of hydrocarbons to non-toxic (carbon dioxide and water) endpoints in groundwater and saturated soil beneath the Site. Typical oxidizers include hydrogen peroxide, ozone, Fenton's Reagent (iron-catalyzed hydrogen peroxide), potassium permanganate, sodium persulfate, and other chemicals. Figure 9 illustrates a proposed oxidizer injection network and layout which targets areas of highest hydrocarbon impacts groundwater and soil beneath the Site.

Use of in-situ chemical oxidation offers the following advantages:

- It provides for conversion of residual hydrocarbon mass to non-toxic (carbon dioxide, water) by-products.
- It may accelerate reduction of residual hydrocarbon mass beneath Site.
- It creates minimal disruption of existing concrete/asphalt cap and existing building.

Successful implementation of in-situ chemical oxidation depends on adequate distribution of injected oxidizer throughout the hydrocarbon-impacted area. It may be difficult to achieve good mixing between the groundwater and injected oxidant solutions due to the presence of discontinuous silt and silty-sand zones beneath the Site. The injected solution tends to displace the impacted groundwater and then react with natural organic matter before it reacts with impacted groundwater. Oxidant solution injection also has the potential to displace the plume and increase chemical migration. Injection of a chemical oxidation solution is regulated and permits are required by the regulatory agencies which may delay the implementation of this alternative²⁷.

ISCO may have disadvantages such as increasing the corrosion potential of pipeline infrastructure at the Site. The effectiveness of ISCO may be limited in areas of separate phase product or other more-impacted areas. ISCO may require special planning and operational constraints, especially when using ozone as an oxidizer. These constraints include permitting and inspections, additional monitoring for ozone or degradation by-products, and preparation of a hazardous materials business plan.

The effectiveness of ISCO remediation may be limited by the ability to directly apply oxidizer. Aqueous solutions or slurries of oxidizing chemicals are injected into wells and the effectiveness is limited by soil conditions and well spacing. Ozone-oxygen gas blends can be sparged into wells

(ozone sparging). Ozone sparging may reach more target zones than other oxidizing schemes because ozone is very mobile. However, ozone sparging is subject to the same distribution limitations as the traditional air sparging oxidizers and delivers weaker reagent doses than permanganate or persulfate slurries. Liquid or slurried oxidizers can deliver more concentrated reagent doses and react with source areas more effectively. In either case, naturally occurring carbon (total organic carbon) from various sources will be oxidized, reducing the effect of the oxidizer and requiring a high volume of oxidizer.

Estimated costs for the installation and operation (for one year) of a ozone-based oxidation system are \$917,600 to \$1,101,100, depending on the number of required injection points and the number of injection cycles required to effect degradation of hydrocarbons across the areas of highest impact.

6.1.5 Alternative 5: Institutional Controls

Alternative 5, as discussed in Section 5.5 above, involves documenting risks associated with residual hydrocarbons, and implementing institutional controls to address any exposure risks documented in the Risk Assessment.

The National Contingency Plan (NCP) is cited by USEPA²⁸ as the basis for defining acceptable incremental risk levels. According to the NCP, lifetime incremental cancer risk levels posed by a site should be within the risk range of one in a million (1×10^{-6}) to 100 in a million (1×10^{-4}) . Thus, USEPA and Cal/EPA agencies typically consider the 1×10^{-6} risk level to be an insignificant risk, and consider a calculated excess cancer risk between 1×10^{-6} and 1×10^{-4} to be within the acceptable risk range. For commercial exposure scenarios, a typical point of departure is a risk level of 1×10^{-5} ; i.e., if risks are at or below 1×10^{-5} , the agency of record will generally accept no further action.

The risk evaluation conducted for the Site (Appendix A) concludes that indoor air exposure risks associated with the concentrations of COCs in soil and groundwater are within the risk management range for carcinogens as determined by the United States Environmental Protection Agency (USEPA) and below the typical point of departure (1x10⁻⁵) for commercial exposure scenarios. The risk evaluation documents the potential for exposure risks above Tier I screening levels in the case of on-site construction workers that may come into direct contact with impacted soils or groundwater during any future construction activities at the Site which involve trenching or excavation. All other potential exposure pathways are incomplete and/or below Tier I screening levels.

Institutional controls currently in place address the actions to be taken to prevent direct exposure risks. These controls can be found in the Covenant and Environmental Restriction (i.e., a Deed Restriction, Appendix C) recorded in June 2000 and the 2001 Risk Management Plan (Appendix D). These documents control land use and limit on-site activity to protect against risks posed by the presence of residual hydrocarbons beneath the Site. They are legally binding on all future owners of the property. The deed restriction requires a surface cap at the Site and notice to regulatory agencies if disturbance to subsurface soils or changes in land use are proposed for the Site. Section 3.1 of the Deed Restriction (see Appendix C) details the implementation of Risk Management Plan for any future development and new construction.

Nestlé understands that the ACHS may request, as part of this proposed CAP alternative, that the existing Deed Restriction be re-filed with the Alameda County Health Care Service office such that the document is recorded according to current protocols within the County.

Estimated costs for Alternative 5, including the cost of the risk evaluation (Appendix A) and any modifications deemed necessary to the existing Deed Restriction and Risk Management Plan for the Site, are estimated at \$127,200 to \$182,600.

6.2 Draft Corrective Action Plan Recommendation

After evaluation and comparison of the Corrective Action Plan options (see Table 5), Alternative 5 (Institutional Controls) is recommended as the Corrective Action Plan for implementation at the Site.

Data from previously implemented remediation efforts indicate that additional extraction or in-situ technologies are not likely to produce significant additional mass removal. This conclusion is supported by the asymptotic rates of mass removal observed prior to the termination of previous active remediation through dual phase extraction ^{29,30}. As illustrated in Appendix B (see Figure 24), wells that historically reported the most significant LPH levels showed significant reductions in LPH thickness as a result of the dual phase extraction system operated at the site. Following the removal of over 10,800 pounds of hydrocarbons via dual phase extraction from August 1997 through June 2000, asymptotic levels of groundwater concentrations and mass removal rates were achieved, resulting in the recommendation and approval by ACHS to terminate active remediation at the site in June 2000.

Any newly installed extraction system would likely result in a return to this asymptotic condition within a very short period of time. Outside of a short-lived initial rate of hydrocarbon mass removal during extraction system start-up, it is unlikely that further groundwater or soil vapor extraction would produce any significant and consistent reductions in hydrocarbon mass beneath the Site.

Local subsurface lithology, site hydrogeology, and compliance groundwater monitoring data indicate that hydrocarbons remaining at the site are stable and not migrating³¹. Two years of semi-annual compliance monitoring were then requested by the ACHS, and subsequently performed between January 2002 and December 2004. This compliance monitoring has confirmed the stability of dissolved hydrocarbons and the lack of off-site migration of dissolved hydrocarbons (see revised SCM Report³²).

The assessment of potential excavation activities (see Sections 6.1.1 and 6.1.2 above) have resulted in the conclusion that direct removal of soils from large areas of the site is logistically and economically infeasible. These alternatives likely pose considerable engineering risk and financial cost due to the presence of the existing building and likely shoring requirements to protect adjacent and off-site sidewalk and street infrastructure. Costs estimated for the demolition of the existing 29,000 square foot building, engineered dismantling of all building related structures, excavation shoring to protect adjacent sidewalks and structures, and building reconstruction costs are estimated to exceed \$2,800,000 (for Alternative 2).

The risk evaluation (Appendix A) analyzes the risks for existing exposure pathways, including indoor air and future onsite workers. Based on the thorough documentation of residual hydrocarbons and the assessment of potential exposure risks associated with the impacts

remaining at the site (Appendix A), the implementation of institutional controls via the existing Deed Restriction (Appendix C) and Risk Management Plan (Appendix D) is the most viable corrective action alternative evaluated for the site.

7 SCHEDULE OF UPCOMING ACTIONS AND PATH TO CLOSURE

Following acceptance by ACHS of the recommended Draft Corrective Action Plan for the site, and/or the documentation of any additions or changes to the draft recommendation made within this report, Nestlé will submit a Final Corrective Action Plan (Final CAP) to ACHS. This report will be submitted within 90 days of any comments on this Draft Corrective Action Plan.

Any requested changes, or re-filing, of the Deed Restriction for the Site will also be addressed following agreement and acceptance of the recommended corrective action plan for the site. Any required provisions for public review and comment on the Final CAP will be scheduled and accommodated in accordance with ACHS or requirements for such a review process.

8 END NOTES

¹ ETIC (ETIC Engineering, Inc.). 2001. *Comprehensive Site Characterization Report, Former Nestlé USA, Inc. Facility, 1310 14th Street, Oakland, California*. ETIC, Pleasant Hill, California. January.

² Harding Lawson Associates (HLA). 1991. *Site Characterization Report, Carnation Facility, Oakland, California*. HLA, Novato, California. September

³ Environmental Cost Management (ECM, Inc.). 2008. *Revised Site Conceptual Model Report, Former Nestlé USA, Inc. Facility, 1310 14th Street, Oakland, California.* ECM, Costa Mesa, California. November.

⁴ Harding Lawson Associates (HLA). 1991. *Site Characterization Report, Carnation Facility, Oakland, California*. HLA, Novato, California. September

⁵ ACFCD (Alameda County Flood Control and Water Conservation District). 1988. Geohydrology and Groundwater –Quality Overview. East Bay Plain Area, Alameda County, California. 205J Report. 83 pp. (as cited in Dames & Moore 1988, Site Contamination Assessments, Carnation Dairy Facility, 1310 14th Street, Oakland, CA; and Carnation Distribution Center, 891 Laurelwood Road, Santa Clara, CA, 2 August).

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⁷ ETIC (ETIC Engineering, Inc.). 2001. *Comprehensive Site Characterization Report, Former Nestlé USA, Inc. Facility, 1310 14th Street, Oakland, California*. ETIC, Pleasant Hill, California. January.

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¹¹ ETIC (ETIC Engineering, Inc.). 2001. *Comprehensive Site Characterization Report, Former Nestlé USA, Inc. Facility, 1310 14th Street, Oakland, California.* ETIC, Pleasant Hill, California. January.

¹² Los Angeles Regional Water Quality Control Board (LARWQCB)/California Department of Toxic Substances Control (DTSC). 2003. *Advisory for Active Soil Gas Investigations*, LARWQCB/ DTSC. Los Angeles, California. January

¹³ Environmental Cost Management (ECM, Inc.). 2008. *Supplemental Soil, Soil Gas, and Groundwater Investigation Report, Former Nestlé USA, Inc. Facility, 1310 14th Street, Oakland, California*. ECM, Costa Mesa, California. July.

¹⁴ Harding Lawson Associates (HLA). 1991. *Site Characterization Report, Carnation Facility, Oakland, California*. HLA, Novato, California. September

¹⁵ Alameda County Health Care Services Agency. 2007. *September 28th letter directive from Jerry Wickham, P.G. to Mr. Mike Desso (Nestlé) and Mr. Mark Hall (Encinal)*, Fuel Leak Case No. ROO000018 and Geotracker Global ID T0600100262, Carnation Dairy, 1310 14th Street, Oakland, CA 94607, Alameda, California.

¹⁶ Environmental Cost Management (ECM, Inc.). 2008. *Revised Site Conceptual Model Report, Former Nestlé USA, Inc. Facility, 1310 14th Street, Oakland, California*. ECM, Costa Mesa, California. November.

¹⁷ Environmental Cost Management (ECM, Inc.). 2008. Supplemental Soil, Soil Gas, and Groundwater Investigation Report, Former Nestlé USA, Inc. Facility, 1310 14th Street, Oakland, California. ECM, Costa Mesa, California. July.

¹⁸ AGE (Anania Geologic Engineering). 1989. Remedial Action Plan for the Carnation Oakland Dairy Facility Located at 1310 14th Street, Oakland, California, Alameda County. AGE, Rancho Cordova, California. April.

¹⁹ ETIC (ETIC Engineering, Inc.). 2001. *Comprehensive Site Characterization Report, Former Nestlé USA, Inc. Facility, 1310 14th Street, Oakland, California.* ETIC, Pleasant Hill, California. January

²⁰ ETIC (ETIC Engineering, Inc.). 2001. *Comprehensive Site Characterization Report, Former Nestlé USA, Inc. Facility, 1310 14th Street, Oakland, California.* ETIC, Pleasant Hill, California. January

²¹ ETIC (ETIC Engineering, Inc.). 2001. *Comprehensive Site Characterization Report, Former Nestlé USA, Inc. Facility, 1310 14th Street, Oakland, California.* ETIC, Pleasant Hill, California. January

²² ETIC (ETIC Engineering, Inc.). 2001. *Comprehensive Site Characterization Report, Former Nestlé USA, Inc. Facility, 1310 14th Street, Oakland, California.* ETIC, Pleasant Hill, California. January

²³ . Revised May 2008. *Screening for Environmental Concerns at Sites with Contaminated Soil and Groundwater*, Interim Final - November 2007, California RWQCB, San Francisco Bay Region.

²⁴ RWQCB. Revised May 2008. *Screening for Environmental Concerns at Sites with Contaminated Soil and Groundwater*, Interim Final - November 2007, California RWQCB, San Francisco Bay Region.

²⁵ ETIC (ETIC Engineering, Inc.). 2001. *Comprehensive Site Characterization Report, Former Nestlé USA, Inc. Facility, 1310 14th Street, Oakland, California.* ETIC, Pleasant Hill, California. January

²⁶ Environmental Cost Management (ECM, Inc.). 2008. *Revised Site Conceptual Model Report, Former Nestlé USA, Inc. Facility, 1310 14th Street, Oakland, California.* ECM, Costa Mesa, California. November

²⁷ Batelle, *Final Air Sparging Guidance Document*, Contract N47408-95-D-0730, Prepared for Naval Facilities Engineering Service Center, 1100 23rd Avenue, Port Hueneme, California, August 31, 2001.

²⁸ United States Environmental Protection Agency (USEPA). 1989. *Risk Assessment Guidance for Superfund Volume I: Human Health Evaluation Manual (Part A)*. Interim Final. Office of Emergency and Remedial Response. December.

²⁹ ETIC (ETIC Engineering, Inc.). 2001. *Comprehensive Site Characterization Report, Former Nestlé USA, Inc. Facility, 1310 14th Street, Oakland, California.* ETIC, Pleasant Hill, California. January

³⁰ Environmental Cost Management (ECM, Inc.). 2008. *Revised Site Conceptual Model Report, Former Nestlé USA, Inc. Facility, 1310 14th Street, Oakland, California*. ECM, Costa Mesa, California. November.

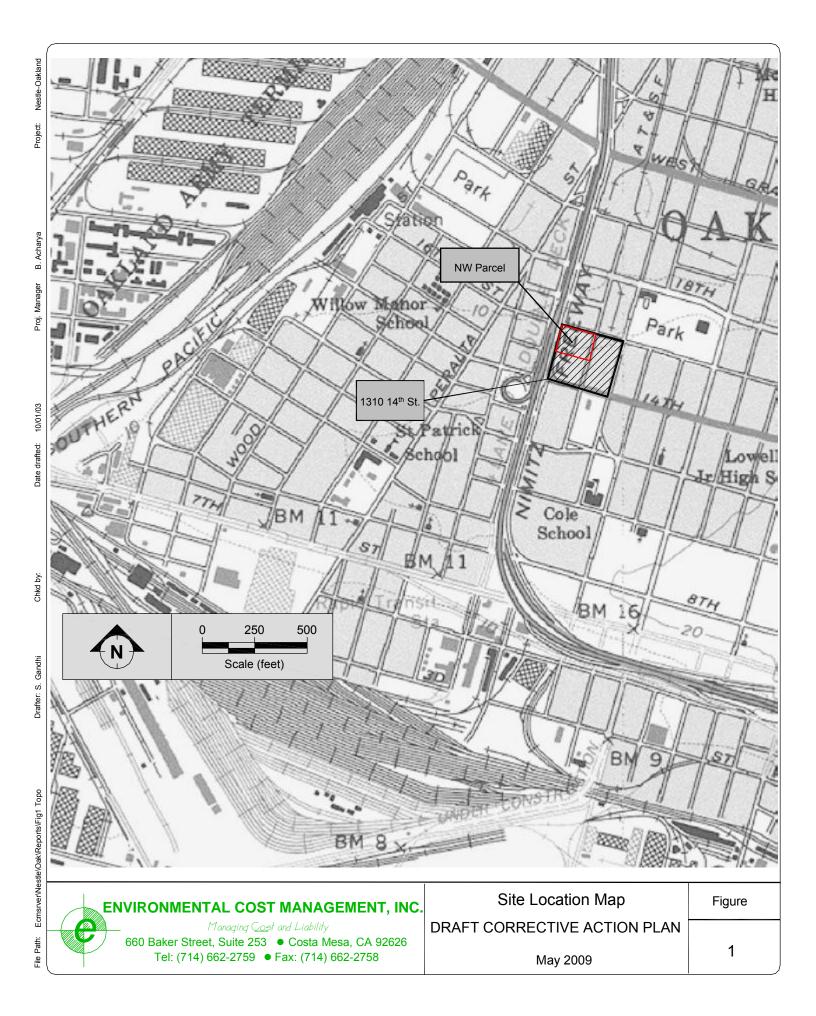
³¹ Environmental Cost Management (ECM, Inc.). 2008. *Revised Site Conceptual Model Report, Former Nestlé USA, Inc. Facility, 1310 14th Street, Oakland, California*. ECM, Costa Mesa, California. November

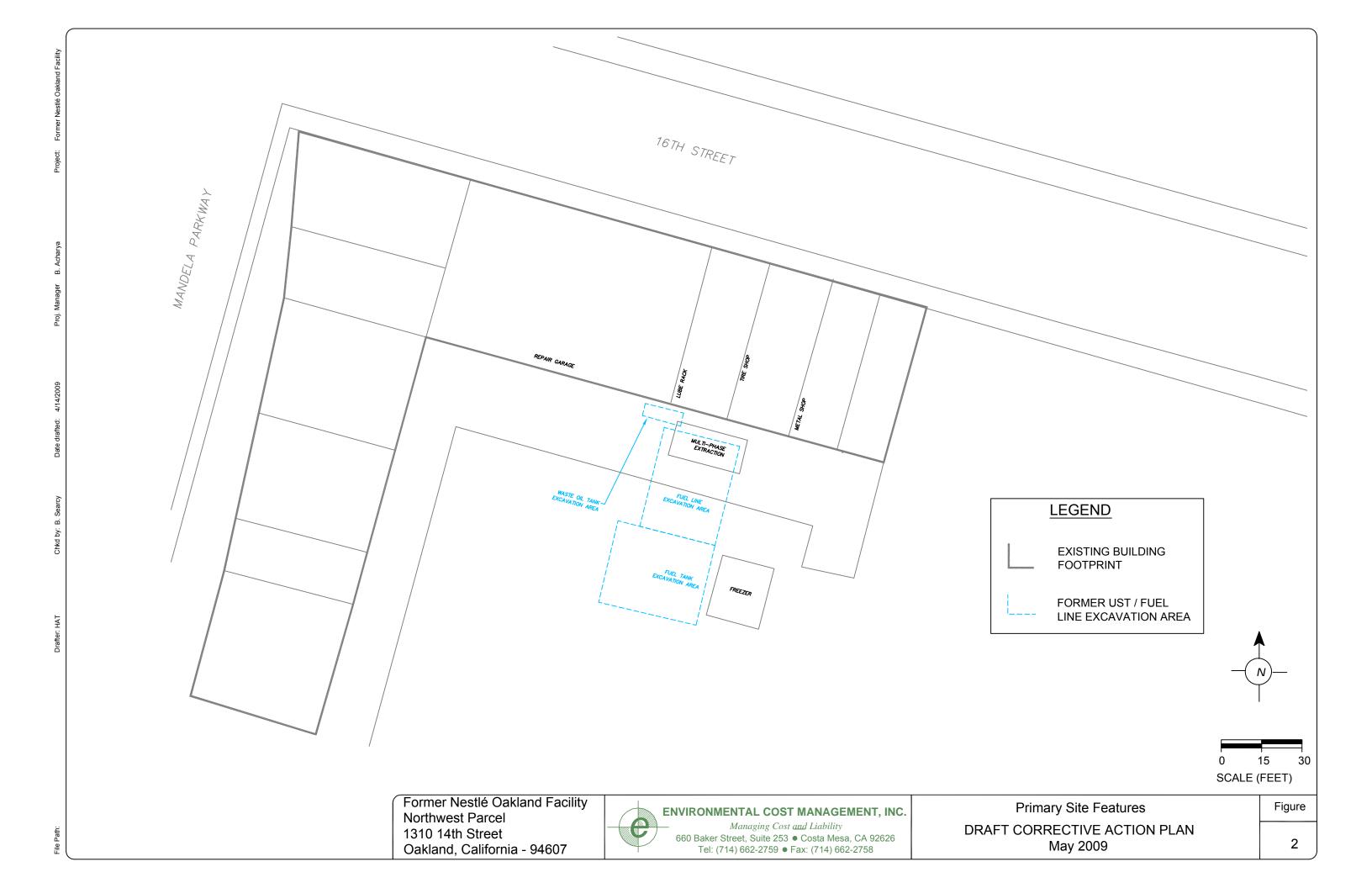
³² Environmental Cost Management (ECM, Inc.). 2008. *Revised Site Conceptual Model Report, Former Nestlé USA, Inc. Facility, 1310 14th Street, Oakland, California.* ECM, Costa Mesa, California. November.

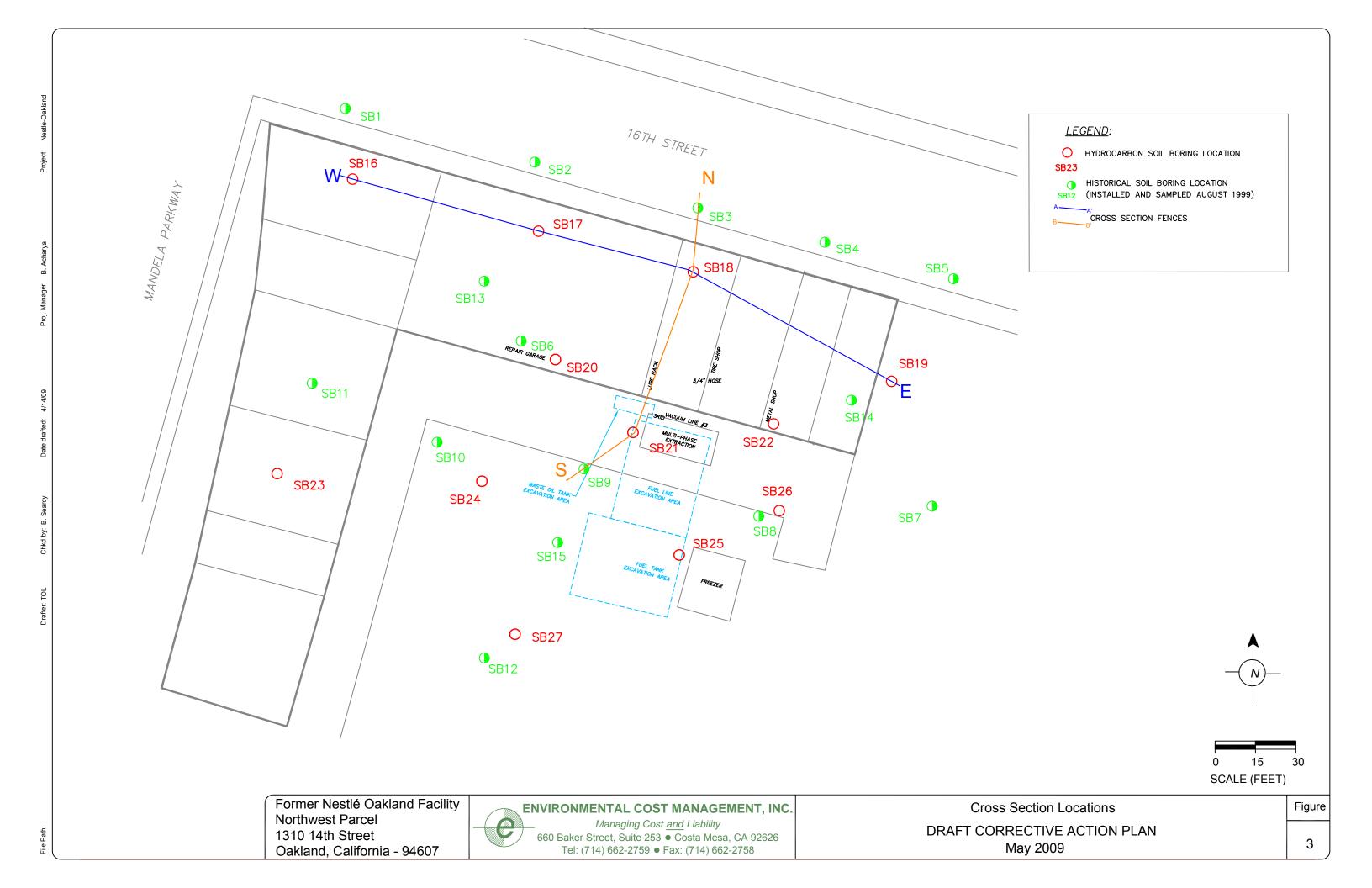
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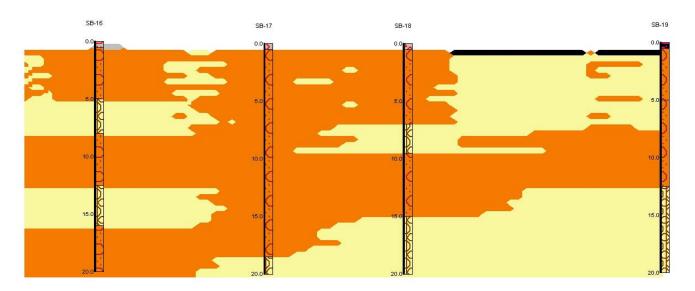
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Former Nestlé USA, Inc. Facility Northwest Parcel 1310 14th Street Oakland, California 94607



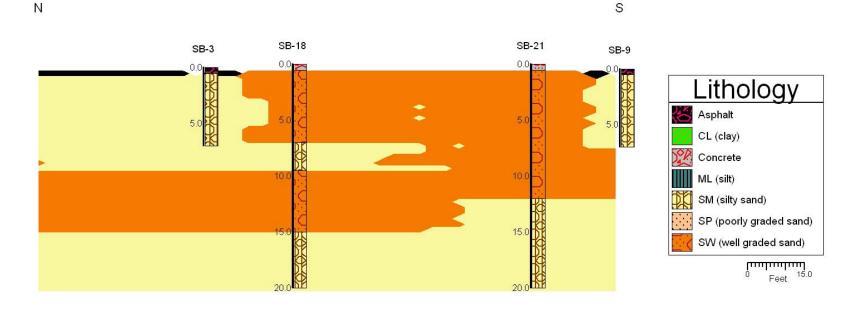
ENVIRONMENTAL COST MANAGEMENT, INC.

Managing Cost <u>and</u> Liability
660 Baker Street, Suite 253 ● Costa Mesa, CA 92626
Tel: (714) 662-2759 ● Fax: (714) 662-2758

E-W Cross Section

DRAFT CORRECTIVE ACTION PLAN May 2009 Figure

4



Former Nestlé USA, Inc. Facility Northwest Parcel 1310 14th Street Oakland, California 94607

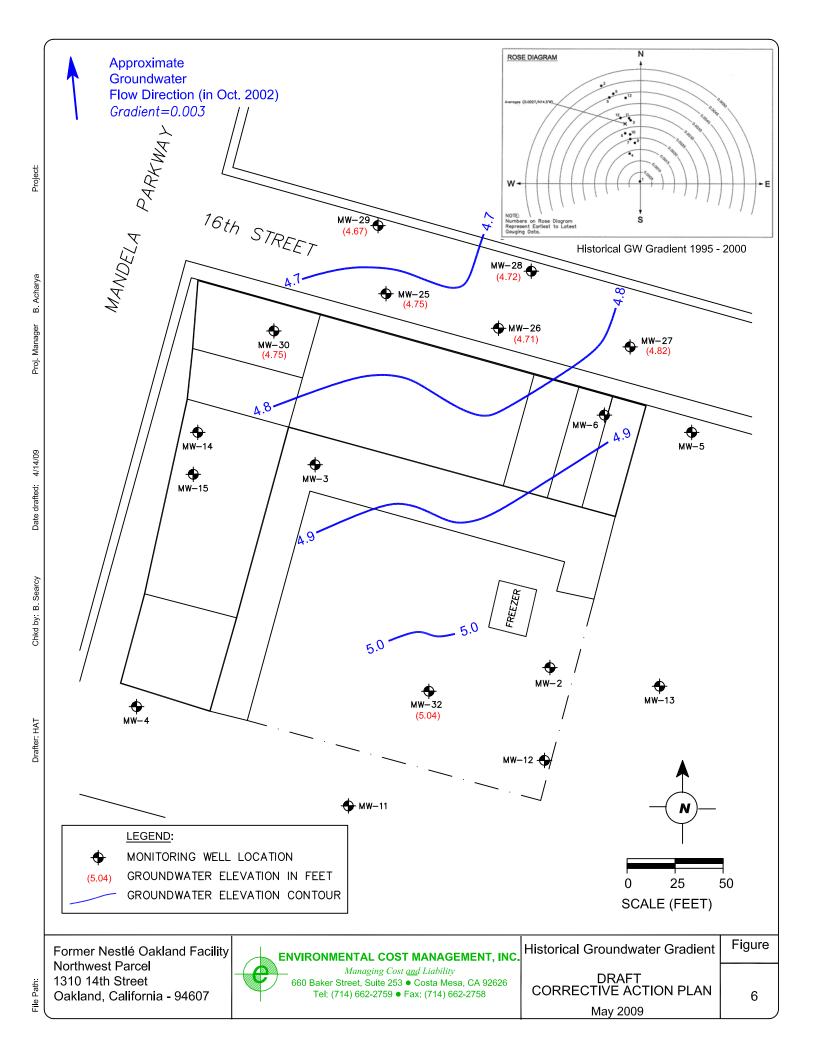


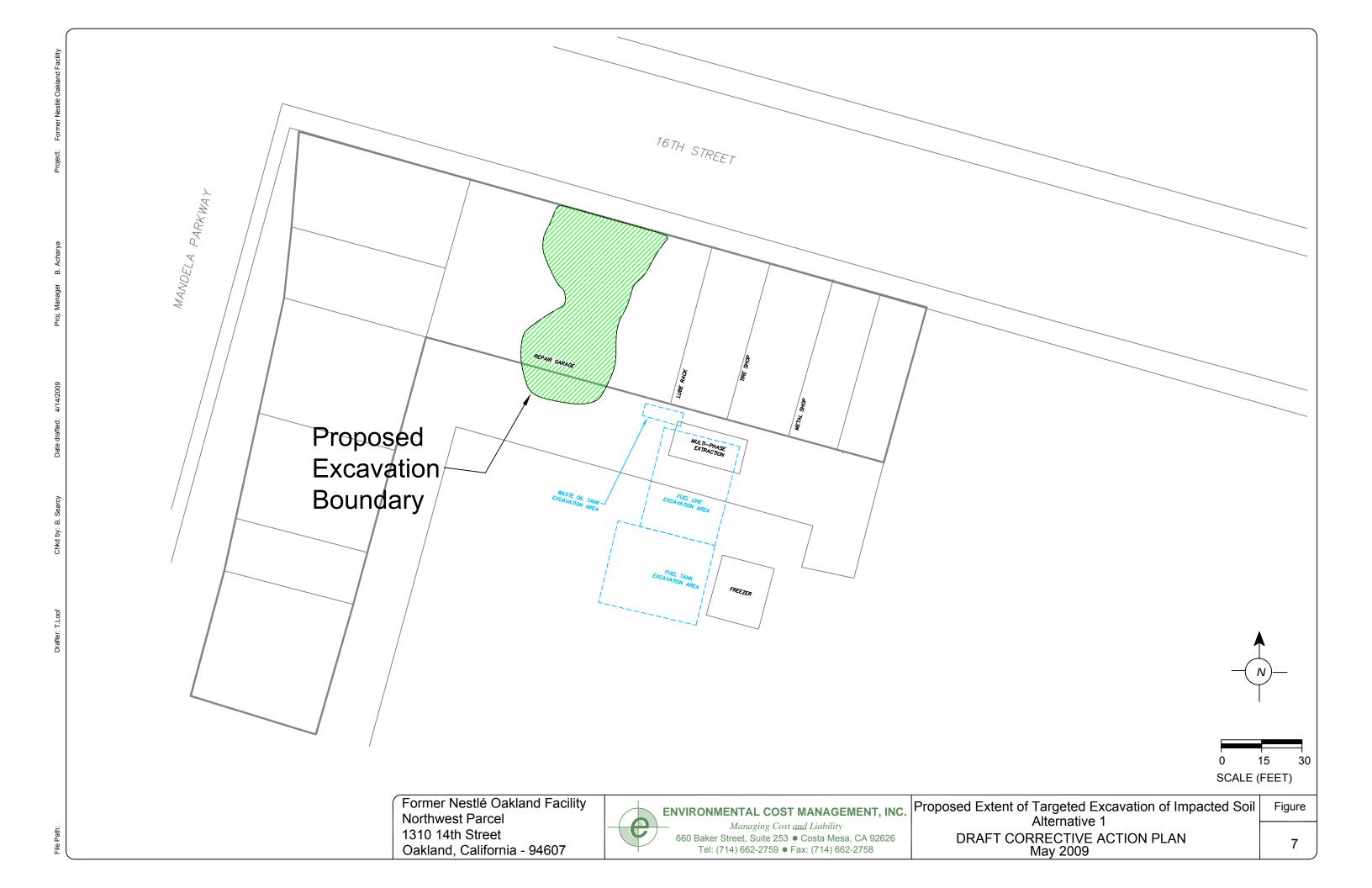
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Tel: (714) 662-2759 ● Fax: (714) 662-2758

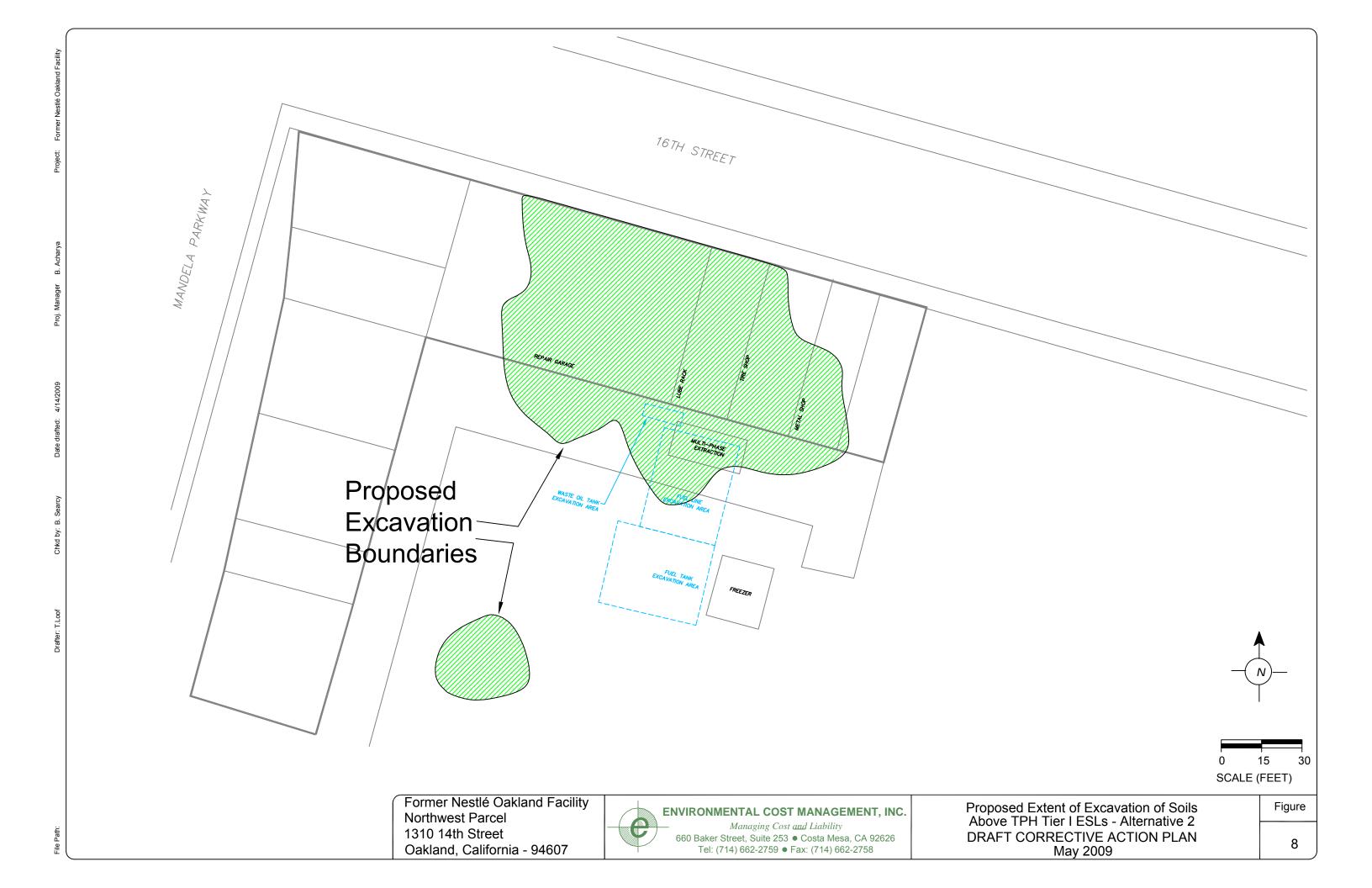
N-S Cross Section

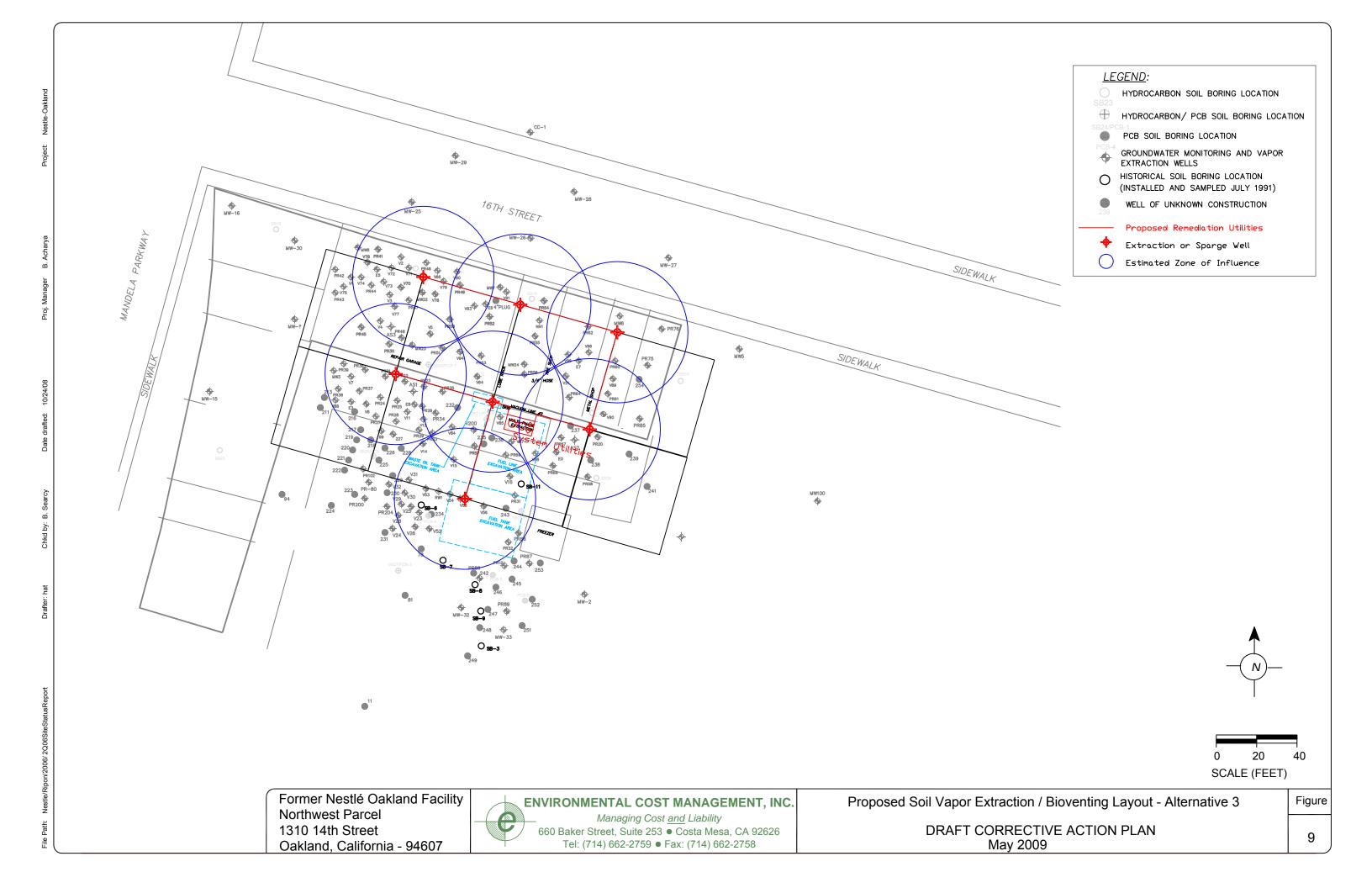
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Table 1a: Soil Vapor Sampling Results, August 1999

Corrective Action Plan Report, May 2009 Former Nestle USA, Inc. Facility 1310 14th Street, Oakland, CA

Concentration (ppbv)

													ition (ppbv	,										
															1,2-Di-	1,3-Di-	1,4-Di-	1,1-Di-	1,2-Di-	1,1-Di-	cis-1,2-			
Sample			Ethyl-	Total				1,3-Bu-	2-Bu-	Carbon	Chloro-	Chloro-	Chloro-	Cyclo-	chloro-	chloro-	chloro-	chloro-	chloro-	chloro-	Dichloro-	1,4-Di-		4-Ethyl-
ID	Benzene	Toluene	benzene	Xylenes	TPH-g	TPH-d	Acetone	tadiene	tanone	Disulfide	benzene	form	methane	hexane	benzene	benzene	benzene	ethane	ethane	ethene	ethene	oxane	Ethanol	toluene
SB-1	4.3	3.1	<0.65	2.74	800	NA	77 a	2.8	13	6.2	<0.65	<0.65	<0.65	<2.6	<0.65	<0.65	0.77	<0.65	<0.65	<0.65	<0.65	<2.6	63	<2.6
SB-2	7.5	12	3.6	17.6	1,100	NA	260 a	<2.7	24	9.0	<0.67	3.9	<0.67	12	<0.67	<0.67	1.8	<0.67	<0.67	<0.67	<0.67	<2.7	110	<2.7
SB-3 SB-3 (dup)	9,900 9,500	230 240	68 <140	67 <140	36,000 40,000	NA NA	<190 <580	<190 <580	<190 <580	<190 <580	<48 <140	<48 <140	<48 <140	<190 <580	<48 <140	<190 <580	<190 <580	<190 <580						
SB-4	1,200	76	8.1	18.7	4,600	NA	200 a	19	<14	<14	<3.5	<3.5	<3.5	32	<3.5	<3.5	<3.5	<3.5	<3.5	<3.5	<3.5	<14	1,400	<14
SB-5	7.6	5.6	0.80	1.9	1,900	NA	45 a	61	12	18	<0.71	<0.71	0.77	8.2	<0.71	<0.71	<0.71	<0.71	<0.71	<0.71	<0.71	3.3	55	<2.8
SB-6	3.0	4.2	<0.68	2.52	560	NA	11 a	<2.7	4.0	<2.7	<0.68	<0.68	<0.68	<2.7	<0.68	<0.68	<0.68	<0.68	<0.68	<0.68	<0.68	<2.7	35	<2.7
SB-7	5.9	6.2	0.87	4.3	780	NA	43 a	3.4	7.9	3.3	<0.73	<0.73	<0.73	5.1	<0.73	<0.73	2.0	<0.73	<0.73	<0.73	<0.73	8.2	94	<2.9
SB-8	10	12	3.8	15.7	1,300	NA	42 a	<11	<11	<11	<2.8	<2.8	<2.8	<11	<2.8	<2.8	<2.8	<2.8	<2.8	<2.8	<2.8	<11	62	<11
SB-9	12	18	1.7	9.9	690	NA	19 a	<2.7	6.0	<2.7	<0.68	1.1	<0.68	4.9	<0.68	<0.68	<0.68	<0.68	<0.68	<0.68	<0.68	<2.7	47	<2.7
SB-10	3.5	2.8	<0.80	1.7	610	NA	39 a	<3.2	9.7	<3.2	<0.80	1.6	<0.80	<3.2	<0.80	<0.80	<0.80	<0.80	<0.80	<0.80	<0.80	<3.2	40	<3.2
SB-11	2.7	1.9	<0.82	0.91	520	NA	38 a	<3.3	9.9	<3.3	<0.82	<0.82	3.7	<3.3	<0.82	<0.82	<0.82	<0.82	<0.82	<0.82	<0.82	22	23	<3.3
SB-12	250	<70	<70	610	750,000	NA	<280	<280	<280	<280	<70	<70	<70	<280	480	<70	76	<70	<70	<70	<70	<280	<280	760
SB-13	0.91	8.5	<0.67	1.3	550	NA	49 a	<2.7	5.5	6.4	<0.67	<0.67	<0.67	<2.7	<0.67	<0.67	<0.67	<0.67	<0.67	<0.67	<0.67	4.3	410 b	<2.7
SB-14	2.7	5.3	0.87	4.7	620	NA	10 a	<2.8	3.5	<2.8	<0.70	<0.70	<0.70	<2.8	<0.70	<0.70	1.6	<0.70	<0.70	<0.70	<0.70	<2.8	67	<2.8
SB-15	42	12	1.6	6.7	2,100	NA	51 a	13	13	<5.8	<1.4	<1.4	<1.4	<5.8	<1.4	<1.4	<1.4	<1.4	<1.4	<1.4	<1.4	<5.8	190	<5.8

Notes:

All soils vapor samples were collected at 3 feet bgs ppbv Parts per billion volumetric.

a Compound present in laboratory blank greater than reporting limit (background subtraction not performed).

Exceeds instrument calibration range.

NA Not analyzed.

TPH-g Total Petroleum Hydrocarbons as gasoline.
TPH-d Total Petroleum Hydrocarbons as diesel.

Table 1a CAP_soil_vapor_99 Page 1 of 2

Table 1a: Soil Vapor Sampling Results, August 1999

Corrective Action Plan Report, May 2009 Former Nestle USA, Inc. Facility 1310 14th Street, Oakland, CA

Concentration (ppbv)

								onicenti a	tion (ppbv)						
						4-Methyl-		Methyl			Tetra-	Tetra-	1,1,1-Tri-	Tri-	1,2,4-Tri-	1,3,5-Tri-
Sample	Freon	Freon	Freon	Нер-	Hex-	2-penta-	Methylene	t-butyl	2-Pro-	Sty-	chloro-	hydro-	chloro-	chloro-	methyl-	methyl-
ID	11	12	113	tane	ane	none	Chloride	ether	panol	rene	ethene	furan	ethane	ethene	benzene	benzene
SB-1	0.74	0.93	27	<2.6	4.4	3.8	3.7	<2.6	5.6	<0.65	1.2	<2.6	<0.65	<0.65	1.1	<0.65
SB-2	1.2	200	<0.67	3.3	5.3	8.1	2.2	<2.7	<2.7	3.0	<0.67	<2.7	<0.67	<0.67	2.0	0.77
SB-3 SB-3 (dup)	<48 <140	180 160	<48 <140	<190 <580	590 580	<190 <580	<48 <140	<190 <580	<190 <580	<48 <140	<48 <140	<190 <580	<48 <140	<48 <140	<48 <140	<48 <140
SB-4	<3.5	100	<3.5	<14	19	15	340	<14	22	<3.5	160	<14	21	<3.5	<3.5	<3.5
SB-5	4.4	1.2	3.4	<2.8	<2.8	<2.8	<0.71	<2.8	<2.8	<0.71	<0.71	<2.8	<0.71	<0.71	<0.71	<0.71
SB-6	<0.68	<0.68	<0.68	<2.7	<2.7	<2.7	<0.68	<2.7	<2.7	<0.68	<0.68	<2.7	<0.68	<0.68	1.1	<0.68
SB-7	0.74	1.1	<0.73	<2.9	6.8	4.4	<0.73	<2.9	3.8	1.0	2.0	<2.9	<0.73	<0.73	1.8	<0.73
SB-8	6.5	630	<2.8	<11	<11	<11	<2.8	<11	<11	<2.8	<2.8	<11	<2.8	<2.8	5.3	<2.8
SB-9	1.5	20	<0.68	<2.7	4.3	<2.7	<0.68	<2.7	<2.7	<0.68	<0.68	<2.7	<0.68	<0.68	2.3	0.77
SB-10	<0.80	1.4	<0.80	<3.2	3.9	<3.2	<0.80	<3.2	<3.2	<0.80	<0.80	<3.2	<0.80	<0.80	1.2	<0.80
SB-11	4.6	<0.82	<0.82	<3.3	<3.3	<3.3	1.2	<3.3	<3.3	<0.82	<0.82	<3.3	<0.82	<0.82	0.85	<0.82
SB-12	<70	<70	<70	<280	18,000	<280	<70	<280	<280	<70	<70	<280	<70	<70	580	740
SB-13	<0.67	<0.67	<0.67	3.4	<2.7	<2.7	5.6	<2.7	26	<0.67	<0.67	58	<0.67	<0.67	1.1	<0.67
SB-14	<0.70	<0.70	<0.70	<2.8	<2.8	2.8	1.3	2.9	<2.8	0.82	<0.70	<2.8	<0.70	<0.70	2.0	0.81
SB-15	<1.4	46	<1.4	<5.8	50	<5.8	4.8	<5.8	<5.8	<1.4	2.1	<5.8	<1.4	<1.4	1.8	<1.4

Notes:

All soils vapor samplesAll soils vapor samples were collected at 3 feet bgs

ppbv Parts per billion volumetric.

a Compound present in laboratory blank greater than reporting limit (background subtraction not performed).

b Exceeds instrument calibration range.

NA Not analyzed.

TPH-g Total Petroleum Hydrocarbons as gasoline.
TPH-d Total Petroleum Hydrocarbons as diesel.

Table Ia CAP_soil_vapor_99 Page 2 of 2

Table 1b: Soil Vapor Sampling Results, May 2008

Corrective Action Plan Report, May 2009 Former Nestlé USA, Inc. Facility 1310 14th Street, Oakland, CA

Boring Location	Sample Depth (feet	Date of Sample				Analytical res	ults of Vap	or, μg/l		
	bgs)	Collection	TPH g	TPH d	Benzene	Ethylbenzene	Toluene	Xylenes, Tot	1,2-DCA	Others
SB-16	5	19-May-08	<10	<50	<0.10	<0.10	<0.20	< 0.30	<0.10	
SB-17	5	19-May-08	<10	<50	<0.10	<0.10	<0.20	< 0.30	<0.10	
SB-18	5	19-May-08	630	<50	2.2	<0.10	0.44	< 0.30	<0.10	
SB-19	5	19-May-08	<10	<50	<0.10	<0.10	<0.20	< 0.30	<0.10	
SB-20/ PCB-7	5	19-May-08	19	<50	<0.10	<0.10	<0.20	< 0.30	<0.10	
SB-21/ PCB-8	5	19-May-08	25	<50	<0.10	<0.10	<0.20	< 0.30	<0.10	
SB-22	5	19-May-08	2,600	<50	40	7.7	32	19.1	<0.10	Dichlorodifluoromethane: 0.39
SB-23	5	19-May-08	<10	<50	<0.10	<0.10	<0.20	< 0.30	<0.10	
SB-24/ PCB-1	5	19-May-08	<10	<50	<0.10	<0.10	0.22	< 0.30	<0.10	
SB-25/ PCB-2	5	19-May-08	<10	<50	<0.10	<0.10	<0.20	< 0.30	<0.10	
SB-26	5	19-May-08	<10	<50	<0.10	<0.10	<0.20	< 0.30	<0.10	Dichlorodifluoromethane: 10
SB-27/ PCB-3	5	19-May-08	<10	<50	<0.10	<0.10	<0.20	< 0.30	<0.10	
SB-22 dup	5	19-May-08	2,600	<50	40	7.5	32	18.0	<0.10	Dichlorodifluoromethane: 0.38
Probe Blank	NA	19-May-08	<10	<50	<0.10	<0.10	<0.20	< 0.30	<0.10	

Notes:

EPA Method 8260B for VOC Analyses of soil vapor EPA Mentod 8015m for TPH-g and TPH-d analyses of soil vapor

Table 2: Historical Soil Sample Results (1999 - 2008)

Boring Location	Sample Depth (feet	Date of Sample					Analytica	ıl results (mg	/kg)		
	bgs)	Collection	TPH g	TPH d	TPH mo	Benzene	Toluene	Ethylbenzene	Xylenes, Tot	1,2-DCA	Others
SB-1	3.5-4.0	08/12/99	<0.13	1,200	NA	<0.0013	<0.0013	<0.0013	<0.0013	<0.0011	
SB-1	6.5-7.0	08/12/99	<0.10	<5.9	NA	<0.001	<0.001	<0.001	<0.001	<0.0008	
SB-2	3.5-4.0	08/12/99	<0.09	<5.6	NA	<0.0009	<0.0009	<0.0009	<0.0009	<0.001	
SB-2	6.5-7.0	08/12/99	<0.10	<5.9	NA	<0.001	<0.001	< 0.001	<0.001	0.001	
SB-3	3.5-4.0	08/12/99	<0.10	<5.6	NA	<0.001	<0.001	<0.001	<0.001	0.0007	
SB-3	6.5-7.0	08/12/99	6,160	<5.7	NA	11	190	100	460	0.0018	MTBE: 0.073
SB-4	3.5-4.0	08/12/99	<0.10	<5.5	NA	<0.001	<0.001	<0.001	<0.001	<0.0007	
SB-4	6.5-7.0	08/12/99	1	94	NA	0.082	0.0085	0.0073	0.013	0.001	
SB-5	3.5-4.0	08/12/99	<0.09	<5.5	NA	<0.0009	<0.0009	<0.0009	<0.0009	0.0006	
SB-5	6.5-7.0	08/12/99	<0.08	<5.9	NA	<0.0008	<0.0008	<0.0008	<0.0008	0.0009	
SB-6	3.5-4.0	08/13/99	<0.10	<5.5	NA	<0.001	<0.001	<0.001	<0.001	<0.0008	
SB-6	6.5-7.0	08/13/99	10,100	1,100	NA	76	490	170	990	0.43	
SB-7	3.5-4.0	08/12/99	<0.10	<5.4	NA	<0.001	<0.001	<0.001	<0.001	<0.0008	
SB-7	6.5-7.0	08/12/99	<0.11	<5.8	NA	<0.0011	<0.0011	<0.0011	<0.0011	<0.0009	
SB-8	3.5-4.0	08/12/99	<0.10	<5.6	NA	<0.001	<0.001	<0.001	<0.001	<0.0007	
SB-8	6.5-7.0	08/12/99	13	<5.8	NA	0.43	0.36	0.12	0.83	0.0012	MTBE: 0.022
SB-9	3.5-4.0	08/13/99	<0.09	<5.6	NA	<0.0009	<0.0009	< 0.0009	<0.0009	<0.001	
SB-9	6.5-7.0	08/13/99	<0.61	<5.8	NA	0.024	<0.0061	< 0.0061	<0.0061	<0.0011	
SB-10	3.5-4.0	08/13/99	<0.09	<5.6	NA	<0.0009	<0.0009	<0.0009	<0.0009	<0.0008	
SB-10	6.5-7.0	08/13/99	<0.13	<6.4	NA	< 0.0013	< 0.0013	< 0.0013	< 0.0013	<0.001	
SB-11	3.5-4.0	08/13/99	<0.20	<5.5	NA	<0.002	<0.002	<0.002	<0.002	<0.0011	
SB-11	6.5-7.0	08/13/99	<0.11	<5.7	NA	<0.0011	<0.0011	< 0.0011	<0.0011	<0.001	
SB-12	3.5-4.0	08/12/99	<0.10	<5.5	NA	<0.001	<0.001	<0.001	<0.001	<0.0006	
SB-12	4.5-5.0	08/12/99	496	2,900	NA	0.07	0.032	4	6.7	<0.0009	Chlorobenzene: 0.0017 1,2-DCB: 3.1 1,3-DCB: 0.038 1,4-DCB: 0.33 MTBE:
SB-12	6.5-7.0	08/12/99	2	60		<0.001	<0.001	0.023	0.0098	<0.0011	MTBE: 0.001
SB-13	3.5-4.0	08/13/99	1	390	NA	< 0.0012	0.002	0.0027	0.0027	0.0025	
SB-13	6.5-7.0	08/13/99	12	65	NA	0.25	0.048	0.15	0.49	0.0014	
SB-14	3.5-4.0	08/12/99	<0.08	<5.5	NA	<0.0008	<0.0008	<0.0008	<0.0008	<0.0008	MTBE: 0.084
SB-14	6.5-7.0	08/12/99	29	450	NA	0.56	0.29	0.33	1.7	0.0097	
SB-15	3.5-4.0	08/12/99	<0.51	140	NA	<0.0054	<0.0054	<0.0054	<0.0054	<0.0091	
SB-15	6.5-7.0	08/12/99	<0.57	81	NA	<0.0061	0.012	<0.0061	0.0085	<0.0098	
SB-16	6-6.5	05/19/08	<0.22	30	<50	<0.0043	< 0.0043	<0.0043	<0.0087	<0.0043	
SB-17	8-8.5	05/22/08	2,500	3,600	2,900	30	130	27	120	ND	
SB-17	10-10.5	05/22/08	12,000	17,000	13,000	140	580	120	620	<8.3	
SB-17	15-15.5	05/22/08	64	1,400	1,300	<0.89	<0.89	<0.89	<1.8	<0.89	
SB-17	20-20.5	05/22/08	<0.21	<0.99	<49	<0.0042	< 0.0042	<0.0042	<0.0084	<0.0042	
SB-18	8-8.5	05/21/08	1,900	67	<49	41	110	28	130	<19	
SB-19	8-8.5	05/21/08	<0.25	<0.99	<49	<0.0050	<0.0050	<0.0050	<0.010	<0.0050	
SB-20/ PCB-7	8-8.5	05/22/08	5,600	390	51	86	280	54	280	<8.3	
SB-21/ PCB-8	8-8.5	05/21/08	3,800	2,500	<49	40	210	69	360	<19	

Table 2: Historical Soil Sample Results (1999 - 2008)

Corrective Action Plan Report, May 2009 Former Nestle USA, Inc. Facility 1310 14th Street, Oakland, CA

Boring Location	Sample Depth (feet	Date of Sample					Analytica	ıl results (mg/	/kg)		
	bgs)	Collection	TPH g	TPH d	TPH mo	Benzene	Toluene	Ethylbenzene	Xylenes, Tot	1,2-DCA	Others
SB-22	8-8.5	05/21/08	3,200	1,100	<500	<47	140	<47	190	<47	
SB-23	11.5-12	05/22/08	<0.21	1.2	<49	<0.0041	<0.0041	<0.0041	<0.0082	<0.0041	
SB-24/ PCB-1	9-9.5	05/20/08	<0.19	1.6	<50	< 0.0039	< 0.0039	< 0.0039	<0.0078	<0.0039	
SB-25/ PCB-2	8-8.5	05/20/08	<0.19	1.1	<50	< 0.0037	< 0.0037	< 0.0037	< 0.0075	<0.0037	
SB-26	8.5-9	05/21/08	<0.23	10	<50	<0.0047	< 0.0047	< 0.0047	< 0.0093	<0.0047	
SB-27/ PCB-3	8.5-9	05/20/08	<0.27	< 0.99	<49	< 0.0054	< 0.0054	< 0.0054	<0.011	<0.0054	
SB-20/ PCB-7 Dup	8-8.5	05/22/08	4,900	610	<250	99	300	64	340	<21	
SB-25/ PCB-2 Dup	8-8.5	05/20/08	NA	<1.0	<50	NA	NA	NA	NA	NA	

Notes:

NA = Not Analyzed

EPA Method 8260 for BTEX and 1,2-DCA analyses of soil EPA Mentod 8015m for TPH-g, TPH-d, and TPM-mo analyses of soil

Well Number	Date Sampled	Benzene µg/L	Toluene μg/L	Ethyl- Benzene µg/L	Xylenes μg/L	TPH-G μg/L	TPH-D μg/L	1,1- DCA µg/L	1,2- DCA µg/L	1,1,1- ΤCA μg/L	TCE µg/L	MTBE µg/L	Notes
MW-2	03/23/93	ND	ND	ND	ND	ND	ND						
	07/27/93	ND	ND	ND	ND	ND	ND						
	11/05/93												
	02/25/94 06/03/94	<1 <0.5	<1 <0.5	<1 <0.5	<1 <0.5	<100 <50	<1,000 <20,000						
	08/31/94	<0.3	<0.3	<0.3	<0.6	<500	<500						
	12/22/94	<0.5	<0.5	<0.5	<0.5	<50	<50						Non-diesel peak reported.
	03/13/95	0.8	<0.5	<0.5	<0.5	<50	<400						
	06/09/95	<0.5	<0.5	<0.5	<0.5	<100	<50						
	09/21/95	0.7	<0.5	<0.5	<0.5	<50	<50						
	12/12/95 03/12/96	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<1.0 <0.5	<100 <100	<50 <50						
	06/21/96		~0.5 	~0.5 	~0.5 								
	08/29/96	<0.5	<0.5	<0.5	<0.5	<50	<150						
	01/16/97	<0.5	<0.5	<0.5	<0.5	<50	<150	0.7	<0.5	<0.5	<0.5		
	07/07/97	<0.5	<0.5	<0.5	<0.5	<50	<150					<0.5	
	01/27/98	<0.5	<0.5	<0.5	<0.5	100	<150					<0.5	
	07/22/98 07/22/99	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<50 <50	<200	 <0.5	 <0.5	<0.5	<0.5	<0.5 <0.5	
MW 0													
MW-3	03/23/93 07/27/93	35 97	2.9 1	2 4	3.2 1.1	300 220	ND ND						
	11/05/93	4.9	ND	ND	1.2	170	ND						
	02/25/94	42	<1	<1	<1	100	<1,000						
	06/03/94	120	8.2	8.4	4.5	320	<20,000						
	08/31/94	83	1.1	5.3	2.9	<500	<500						
	12/22/94	1,460	18	100	50	3,800	270						
	03/13/95	3,600	260	270	280	14,000	1,700						
	06/09/95 09/21/95	4,700 9,800	58 58	140 600	71 95	3,700 14,000	120 300						
	12/12/95	330	2.1	47	5.3	700	<50						
	03/12/96	350	4.6	23	8.7	600	<50						
	06/21/96	940	76	98	57	1,900	<50						
	08/29/96	420	29	44	28	900	<150						
	01/16/97	1,600	270	120	194	3,600	700	<0.5	9.2	<0.5	<0.5		
	04/15/97 07/07/97	1,300 100	300 84	180 100	160 67	4,300	800	<0.5	16	<0.5	1.1	6.9 3.8	
	10/27/97	1,030	60	54	40	1,900 2,200	350	<0.5	2.4	<0.5	<0.5	3.0	
	01/27/98	1,070	98	73	69	3,200						3.9	
	04/22/98	610	56	49	54	1,800		<0.5	3.0	<0.5	<0.5	1.1	
	07/22/98	1,800	230	160	180	3,600	370					5.0	
	10/21/98	78	1.0	3.8	0.6	110	<250	<0.5	0.6	<0.5	<0.5	<0.5	
	07/23/99 10/28/99	1,500 1,100	140 43	76.0 58	260 102	4,000 3,000	790 600	<0.5 <0.5	1.0 0.9	<0.5	<0.5 <0.5	5.60	
	02/10/00	690	22	36	49	1,400	520	<0.5	<0.5	<0.5	<0.5	2.20	
	04/27/00	1,100	140	73	163	2,400	250	<0.5	0.6	<0.5	<0.5	<0.5	
	08/03/00	520	7.7	21	27	1,100	750	<0.5	0.6	<0.5	<0.5	<0.5	
	10/23/00	2,000	16	22	46	3,800	760	<0.5	0.7	<0.5	<0.5	<0.5	
	01/31/01	360	8.6	14	28	860	300	<0.5	0.6	<0.5	<0.5	<0.5	
	04/26/01	808 788	60.6	46.8	115 80.7	1,530	280	<0.5	0.8	<0.5	<0.5	<0.5	
	07/30/01 10/29/01	788 852	23.3 14.3	44.6 24.5	80.7 38.6	1,400 1,730	350 500	<0.5 <0.5	0.6 0.5	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	
	01/29/02	1,250	85.3	64.7	95.7	4,240	490	<0.5	1.4	<0.5	<0.5	<0.5	
	04/29/02	1,120	51.5	84.4	117	5,710	700	<0.5	1.1	<0.5	<0.5	<0.5	
MW-5	02/05/99	<0.5	<0.5	<0.5	<0.5	<50	<150	<0.5	<0.5	<0.5	<0.5	<0.5	
MW-6	03/23/93	ND	ND	ND	ND	ND	ND						
	07/27/93	ND	ND	ND	ND	ND	ND						
	11/05/93	ND	ND	ND	ND	ND	ND						
	02/25/94 06/03/94	<1 2.7	<1 <0.5	<1 <0.5	3.5 <0.5	<100 69	<1,000 <20,000						
	08/31/94	<0.3	8.7	1.6	3.5	<500	<500						
	12/22/94	<0.5	<0.5	<0.5	<0.5	<50	<50						Non-diesel peak reported.
	03/13/95	1.2	<0.5	<0.5	<0.5	<50	<400						
	06/09/95	0.6	<0.5	<0.5	<0.5	<100	<50						
	09/21/95	<0.5	<0.5	<0.5	<0.5	<50	<50						
	12/12/95 03/12/96	<0.5	<0.5	<0.5 <0.5	<1.0 <0.5	<100 <100	<50						
	03/12/96	<0.5 	<0.5 	<0.5	<0.5 	<100 	<50 						
	08/29/96	<0.5	<0.5	<0.5	<0.5	<50	<150						
	01/16/97	5.5	16	2.9	16	140	220	<0.5	6.3	<0.5	<0.5		
	07/07/97	<0.5	<0.5	<0.5	<0.5	<50	<150					<0.5	
	07/22/98	<0.5	<0.5	<0.5	<0.5	<50	<250					<0.5	
	10/24/00 01/31/01	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<50 <50	<250 <250	<0.5 <0.5	7.7 6.9	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	
	01/31/01	~U.5	~U.5	\U.5	~U.5	~ 3∪	~200	\U.5	ს.9	~ U.5	~U.5	\U.5	

			l	Ethyl-			1	1,1-	1,2-	1,1,1-			T
Well Number	Date Sampled	Benzene µg/L	Toluene µg/L	Benzene µg/L	Xylenes µg/L	TPH-G µg/L	TPH-D μg/L	DCA µg/L	DCA µg/L	TCA µg/L	TCE µg/L	MTBE µg/L	Notes
MW-6	04/27/01	<0.5	<0.5	<0.5	<0.5	<200	<250	<0.5	6.6	<0.5	<0.5	<0.5	
(cont.)	07/30/01	<0.5	<0.5	<0.5	<0.5	<200	<250	<0.5	9.2	<0.5	<0.5	<0.5	
	10/30/01 01/29/02	<0.5 0.54	<0.5 <0.5	<0.5 <0.5	<1.0 <1.0	<200 <200	<500 <250	<0.5 <0.5	10 10	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	
	04/30/02	<0.5	<0.5	<0.5	<1.0	<200	<250	<0.5	14	<0.5	<0.5	<0.5	
B.854.4.4													
MW-11	02/05/99	<0.5	<0.5	<0.5	<0.5	<50	<150					<0.5	
MW-12	02/05/99	<0.5	<0.5	<0.5	<0.5	<50	<150				-	<0.5	
MW-13	02/05/99	<0.5	<0.5	<0.5	<0.5	<50	<150				-	<0.5	
MW-15	02/05/99	<0.5	<0.5	<0.5	<0.5	<50	430	<0.5	<0.5	<0.5	<0.5	<0.5	
	07/22/99	<0.5	<0.5	<0.5	<0.5	<50	<200	<0.5	<0.5	<0.5	<0.5	<0.5	
MW-25	03/23/93	ND	ND	ND	ND	ND	ND						
	07/27/93	ND	ND	ND	ND	ND	ND						
	11/05/93	4.2	4.4	2.5	20	170	ND						
	02/25/94	2.1	<1	<1	<1	<100	<1,000						
	06/03/94	2.4	14	<0.5	3.4	97	<20,000						
	08/31/94 12/22/94	0.5 0.5	<0.3 <0.5	<0.3 <0.5	<0.6 <0.5	<500 <50	<500 <50						Non-diesel peak reported.
	03/13/95	0.58	<0.5	<0.5	<0.5	150	950						INOTI-uleser peak reported.
	06/09/95	0.8	<0.5	<0.5	<0.5	<100	60						
	09/21/95	<0.5	<0.5	<0.5	<0.5	50	<50						
	12/12/95	<0.5	<0.5	<0.5	<1.0	<100	<50						
	03/12/96	<0.5	<0.5	<0.5	<0.5	120	<50						
	06/21/96												
	08/29/96 01/16/97	<0.5 0.6	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	90 80	<150 <150	 25	 41	<0.5	 <0.5		
	07/07/97	<0.5	<0.5	<0.5	<0.5	140	<150					11	
	01/27/98	<0.5	<0.5	<0.5	<0.5	<100						10	
	07/22/98	<0.5	<0.5	<0.5	<0.5	<50	<250					24	
	02/05/99	<0.5	<0.5	<0.5	<0.5	<50	340	28	59	<0.5	<0.5	28	1,1-DCE detected, 0.9 µg/L.
	04/07/99	<0.5	<0.5	<0.5	<0.5	<50	<250	27	72	<0.5	<0.5	27	1,1-DCE detected, 1.6 μg/L.
	07/23/99	1.80	<0.5	<0.5	<0.5	<50	<200	30	58	<0.5	<0.5	23.0	
	10/27/99	<0.5	1.4	<0.5	1.0	<100	<200	35	47 41	 -0 F	<0.5		4.4 Dishlaresthans datasted at 2.4 cm/l
	02/08/00 04/26/00	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	100 <100	<250 <250	39 51	38	<0.5 <0.5	<0.5 <0.5	29.0 18	1,1-Dichloroethene detected at 3.1 µg/L. 1,1-Dichloroethene detected at 4.2 µg/L.
	08/03/00	<0.5	<0.5	<0.5	<0.5	<50	<250	40	57	<0.5	<0.5	27	1,1-Dichloroethene detected at 4.2 µg/L.
	10/23/00	<0.5	<0.5	<0.5	<0.5	<50	<250	54	68	<0.5	<0.5	38	1,1-Dichloroethene detected at 3.5 µg/L.
	01/31/01	<0.5	<0.5	<0.5	<0.5	90	<250	52	46	<0.5	<0.5	22	1,1-Dichloroethene detected at 6.5 µg/L.
	04/26/01	<0.5	0.62	<0.5	<0.5	<200	<250	49	37	<0.5	<0.5	15.8	1,1-Dichloroethene detected at 6.0 µg/L.
	07/30/01	<0.5	<0.5	<0.5	<0.5	<200	<250	33	36	<0.5	<0.5	10.9	Chloromethane detected at 0.8 µg/L; 1,1-Dichloroethene detected at 4.6 µg/L.
	10/29/01	<0.5	<0.5	<0.5	<1.0	<200	<500	22	38	<0.5	<0.5	10.5	Chloromethane detected at 4.6 µg/L. 1,1-Dichloroethene detected at 0.5 µg/L; 1,1-Dichloroethene detected at 1.8 µg/L.
	01/28/02 04/29/02	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<1.0 <1.0	<200 <200	<250 <250	25 14	56 44	<0.5 <0.5	<0.5 <0.5	8.90 6.92	1,1-Dichloroethene detected at 2.8 µg/L. 1,1-Dichloroethene detected at 1.7 µg/L; 1,1,2,2-Tetrachloroethane detected at 0.5 µg/L.
	10/22/02	7.64	248	133	843	4,790	1,240	9.6	34	<0.5	<0.5	1,410	1,1-Dichloroethene detected at 0.9 μg/L.
	11/15/02	<0.5	<0.5	<0.5	<1.0	<200	<250	11	35	<0.5	<0.5	7.3	Chloroethane detected at 22 µg/L.
	05/06/03 10/14/03	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<1.0 <1.0	<200 <200	<250 <250	8.5 7.6	34	<0.5 <0.5	<0.5 <0.5	5.7	1,1-Dichloroethene detected at 0.8 μg/L.
	04/27/04	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<1.0 <1.0	<200 <200	<250 <250	7.6 5.1	27 18	<0.5 <0.5	<0.5 <0.5	6.3 5.2	
	11/17/04	<0.50	<0.50	<0.50	<0.50	<50	190	6.7	25	<0.50	<0.50	6.1	1,1-Dichloroethene detected at 0.51 µg/L.
MW-26	03/23/93	180	190	55	330	7,000	1,300	ND	ND	ND	ND		
	07/27/93	470	96	30	80	1,800	ND	ND	140	ND	ND		
	11/05/93	4,700	1,300	9	1,400	19,000	ND	ND	120	ND	ND		
	02/25/94	4,800	570	200	860	14,000	<1,000	<1	28	<1	<1		
	06/03/94	4,100	300	120	230	12,000	<20,000	1.7	140	<0.5	<0.5		Bromodichloromethane detected, 0.84 μg/L.
	08/31/94 12/22/94	4,100	360 170	170 85	450 200	93,000	1,400	<4.0	<4.0	<4.0	<4.0		9 other velatiles detected by 9260
	03/13/95	1,030 320	170 19	85 23	290 66	5,000 3,000	560 810	<2.0 53	<2.0 5.8	<2.0 <0.5	<2.0 <0.5		8 other volatiles detected by 8260.
	06/09/95	14,000	64	31	230	10,800	310	240	3.1	1	<0.5		
	09/21/95	1,900	160	160	330	8,000	200	1.3	120	<0.5	<0.5		
	12/12/95	13,000	38	36	120	25,000	0.6	1.4	180	<0.5	<0.5		No diesel pattern detected; result due to high gasoline concentration.
	03/12/96	9,000	33	30	65	4,400	<50	<0.5	180	<0.5	<0.5		gasonia concentration.
	06/21/96	14,000	27	16	66	5,400	<50	3.2	170	<0.5	<0.5		
	08/29/96	8,500	26	28	74	19,000	<150	<0.5	160	<0.5	<0.5		
	01/16/97	6,500	21	31	47	4,600		4.3	>50	<0.5	<0.5	26	
	04/15/97	16,000	33	40	160	26,000	2,200	3.5	97	<0.5	2.4	40	cis-1,2-DCE detected, 0.7 μg/L.
	07/07/97	22,000	44	170	200	28,000	1,100	<5.0	<5.0	<5.0	<5.0	95	
	10/27/97 01/27/98	16,000 23,600	26 <5.0	100 <5.0	37 <5.0	30,000 26,000	420	3.6 8.3	92 100	<0.5 <0.5	<0.5 <0.5	38 100	
	04/22/98	5,000	4.3	9.2	16	14,000		13	130	<0.5	<0.5	27	

Well Number	Date Sampled	Benzene µg/L	Toluene μg/L	Ethyl- Benzene µg/L	Xylenes μg/L	TPH-G μg/L	TPH-D μg/L	1,1- DCA µg/L	1,2- DCA µg/L	1,1,1- ΤCA μg/L	TCE µg/L	MTBE μg/L	Notes
MW-26	07/22/98	3,800	5.7	6.9	11	5,200	750	10	110		<1.0	33	
(cont.)	10/21/98	420	<0.5	2.1	2.7	820	<250	24	82	<0.5	<0.5	31	
	02/05/99	20	<0.5	0.60	0.80	230	230	10	51	<0.5	<0.5	29	
	04/07/99	<0.5	<0.5	<0.5	<0.5	80	<250	15	54	<0.5	<0.5	25	
	07/23/99	7.10	<0.5	<0.5	0.80	180	<200	12	32	<0.5	<0.5	12.0	
	10/27/99	14	1.4	2.9	7.8	400	<200	13	30		<0.5		
	02/08/00	<0.5	<0.5	<0.5	<0.5	80	<250	13	32	<0.5	<0.5	28.0	
	04/26/00	0.7	<0.5	0.6	<0.5	200	340	7.5	39	<0.5	<0.5	22	
	08/03/00	6.8	<0.5	0.6	1.4	<50	<250	7.4	19	<0.5	<0.5	19	
	10/23/00	10	0.8	1.7	1.7	80	<250	5.1	37	<0.5	<0.5	26	
	01/31/01	26	0.70	2.4	2.2	390	320	5.7	51	<0.5	<0.5	33	
	04/26/01 07/30/01	10.6 107	<0.5 <0.5	0.70 1.42	1.04 1.06	400 1,920	350 380	16 22	39 44	<0.5 <0.5	<0.5 <0.5	28.5 31.4	
	10/29/01	31.6	<0.5	< 0.5	<1.00	2,020	500	26	25	<0.5	<0.5	27	
	01/28/02	30.0	<0.5	0.70	<1.0	450	380	43	<0.5	<0.5	<0.5	14.5	1,1-Dichloroethene detected at 1.8 µg/L.
	04/29/02	394	<0.5	<0.5	<1.0	1,870	550	50	23	<0.5	<0.5	8.62	1,1-Dichloroethene detected at 1.5 µg/L.
	10/22/02	1,440	25.7	6.60	20.4	4,440	890	53	26	<0.5	<0.5	168	1,1-Dichloroethene detected at 2.5 μg/L.
	11/15/02	1,630	0.56	3.22	3.86	5,590	780	18	33	<0.5	<0.5	49.2	1,1-dichloroethene detected at 3.7 μg/L.
	05/06/03	1,250	<0.5	2.42	<1.0	3,730	380	46	24	<0.5	<0.5	13.1	1,1-Dichloroethene detected at 1.0 µg/L.
	10/14/03	51	<0.5	1.38	<1.0	3,100	<250	83	28	<0.5	<0.5	23.8	1,1-Dichloroethene detected at 3.3 µg/L.
	04/27/04	467	<0.5	1.24	<1.0	1,380	<250	82	33	<0.5	<0.5	<0.5	1,1-Dichloroethene detected at 5.2 µg/L.
	11/17/04	120	<1.0	2.50	1.3	740	820	31	44	<0.50	<0.50	120	1,1-Dichloroethene detected at 1.1 µg/L.
MW 27													
MW-27	06/21/96 08/29/96	<0.5	<0.5	<0.5	<0.5	<50	<50	<0.5	6.8	<0.5	<0.5		
	08/29/96	 12	 5.0	 <0.5	2.6	 70	 <150	<0.5	5.7	 <0.5	 <0.5		
	07/22/98	<0.5	<0.5	<0.5	<0.5	<50	<250	<1.0			<1.0	<0.5	
	02/05/99	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<50 <50	<250 <150	<0.5	1.4 0.7	 <0.5	<0.5	<0.5	
	07/23/99	<0.5	<0.5	<0.5	<0.5	<50	<200	<0.5	0.7	<0.5	<0.5	<0.5	
	10/27/99	<0.5	<0.5	<0.5	<0.5	<100	<200	<0.5	<0.5		<0.5		
	02/08/00	<0.5	<0.5	<0.5	<0.5	<50	<250	<0.5	<0.5	<0.5	<0.5	<0.5	
	04/27/00	<0.5	<0.5	<0.5	<0.5	<100	250	<0.5	<0.5	<0.5	<0.5	<0.5	
	08/16/00	<0.5	<0.5	<0.5	<0.5	<50		<0.5	<0.5	<0.5	<0.5	<0.5	
	10/23/00	<0.5	<0.5	<0.5	<0.5	<50	<250	<0.5	<0.5	<0.5	<0.5	<0.5	
	01/31/01	<0.5	<0.5	<0.5	<0.5	<50	<250	<0.5	<0.5	<0.5	<0.5	<0.5	
	04/26/01	<0.5	<0.5	<0.5	<0.5	<200	<250	<0.5	<0.5	<0.5	<0.5	<0.5	
	07/30/01	<0.5	<0.5	<0.5	<0.5	<200	<250	<0.5	<0.5	<0.5	<0.5	<0.5	
	10/29/01	<0.5	<0.5	<0.5	<1.0	<200	<500	<0.5	<0.5	<0.5	<0.5	<0.5	
	01/28/02	<0.5	<0.5	<0.5	<1.0	<200	<250	<0.5	0.5	<0.5	<0.5	<0.5	
	04/29/02	<0.5	<0.5	<0.5	<1.0	<200	<250	<0.5	<0.5	<0.5	<0.5	<0.5	
	10/22/02	8.56	56.2	9.37	59.3	650	600	<0.5	<0.5	<0.5	<0.5	331	
	11/15/02	<0.5	<0.5	<0.5	<1.0	<200	<250	<0.5	<0.5	<0.5	<0.5	<0.5	
	05/06/03	<0.5	<0.5	<0.5	<1.0	<200	<250	<0.5	<0.5	<0.5	<0.5	<0.5	
	10/14/03	<0.5	<0.5	<0.5	<1.0	<200	<250	<0.5	<0.5	<0.5	<0.5	<0.5	
	04/27/04	<0.5	<0.5	<0.5	<1.0	<200	<250	<0.5	<0.5	<0.5	<0.5	<0.5	
	11/17/04	<0.50	<0.50	<0.50	<0.50	<50	64	<0.50	<0.50	<0.50	<0.50	<5.0	
MW-28	03/23/93	ND	ND	ND	ND	110	ND						
	07/27/93	ND	ND	ND	ND	ND	ND						
	11/05/93	ND	ND	ND	2.1	ND	ND						
	02/25/94	<1	<1	<1	<1	<100	<1						
	06/03/94	3.1	<0.5	<0.5	<0.5	<50	<20,000						
	08/31/94	1.4	<0.3	<0.3	<0.6	<500	<500						
	12/22/94	<0.5	<0.5	<0.5	<0.5	<50	<50						Non-diesel peak reported.
	03/13/95	0.91	<0.5	<0.5	<0.5	<50	<400						
	06/09/95	<0.5	<0.5	<0.5	<0.5	<100	<50						
	09/21/95	<0.5	<0.5	<0.5	<0.5	<50	<50						
	12/12/95	<0.5	<0.5	<0.5	<1.0	<100	<50						
	03/12/96	<0.5	<0.5	<0.5	<0.5	<100	<50						
	06/21/96	<0.5	<0.5	<0.5	<0.5	<100	<50						
	08/29/96	<0.5	<0.5	<0.5	<0.5	<50	<150						
	01/16/97	18	20	2.2	13	220	<150	5.1	85	<0.5	<0.5	8.2	
	04/15/97	<0.5	<0.5	<0.5	<0.5	120	<150	1.1	150	<0.5	<0.5	7.1	
	07/07/97	<0.5	<0.5	<0.5	<0.5	110	<150	<5.0	170	<5.0	<5.0	7.2	
	10/27/97	3.6	<0.5	<0.5	<0.5	300		6.2	120	<0.5	<0.5	36	
	01/27/98	7.6	<0.5	<0.5	<0.5	500	<150					56	
	04/22/98	<0.5	<0.5	<0.5	<0.5	<50		1.0	89	<0.5	<0.5	8.6	
	07/22/98	<0.5	<0.5	<0.5	<0.5	<50		<1.0	85		<1.0	18	
	10/21/98	<0.5	<0.5	<0.5	<0.5	<50	<250	0.5	80	<0.5	<0.5	12	
	02/05/99	<0.5	<0.5	<0.5	<0.5	<50	<150	32	29	<0.5	<0.5	5.0	1,1-DCE detected, 0.9 μg/L.
	04/07/99	<0.5	<0.5	<0.5	<0.5	<50	<250	<0.5	62	<0.5	<0.5	4.5	
	07/23/99	<0.5	<0.5	<0.5	<0.5	<50	<200	<0.5	50	<0.5	<0.5	1.80	
	10/27/99						<200						
	11/02/99	0.7	<0.5	<0.5	<0.5	<100		<0.5	32		<0.5		
	02/08/00	<0.5	<0.5	<0.5	<0.5	<50	<250	<0.5	39	<0.5	<0.5	4.30	
	04/26/00	<0.5	<0.5	<0.5	<0.5	<100 <50	<250	<0.5	50	<0.5	<0.5	1.5	
	08/03/00	<0.5	<0.5	<0.5	<0.5	<50	<250	<0.5	47	<0.5	<0.5	3.7	

Well Number	Date Sampled	Benzene µg/L	Toluene μg/L	Ethyl- Benzene µg/L	Xylenes μg/L	TPH-G μg/L	TPH-D μg/L	1,1- DCA µg/L	1,2- DCA µg/L	1,1,1- TCA μg/L	TCE µg/L	MTBE μg/L	Notes
MW-28	10/23/00	<0.5	<0.5	<0.5	<0.5	<50	<250	<0.5	57	<0.5	<0.5	4.7	
(cont.)	01/31/01	<0.5	<0.5	<0.5	<0.5	<50	<250	<0.5	46	<0.5	<0.5	4.4	
	04/26/01	<0.5	<0.5	<0.5	<0.5	<200	<250	<0.5	26	<0.5	<0.5	1.98	
	07/30/01	0.5	<0.5	0.64	2.58	<200	<250	<0.5	38	<0.5	<0.5	3.0	Chloromethane detected at 3.3 μg/L.
	10/29/01	<0.5	<0.5	<0.5	<1.0	<200	<500	<0.5	29	<0.5	<0.5	3.74	
	01/28/02	6.20	<0.5	<0.5	<1.0	<200	<250	2.8	50	<0.5	<0.5	6.00	
	04/29/02	1.64	<0.5	<0.5	<1.0	<200	<250	3.7	44	<0.5	<0.5	4.81	
	10/22/02	25.0	<0.5	<0.5	<1.0	750	<250	2.0	59	<0.5	<0.5	<0.5	
	11/15/02	13.4	<0.5	1.29	<1.0	610	<250	1.3	54	<0.5	<0.5	<0.5	Chloromethane detected at 1.0 μg/L.
	05/06/03	3.1	<0.5	<0.5	<1.0	390	<250	0.8	70	<0.5	<0.5	9.29	Chloroethane detected at 0.8 µg/L.
	10/14/03	< 0.5	<0.5	<0.5	<1.0	<200	<250	<0.5	38	<0.5	<0.5	6.44	
	04/27/04	<0.5 <0.50	<0.5	< 0.5	<1.0	<200	<250	<0.5	<0.5	<0.5	<0.5	9.29 <5.0	
	11/17/04		<0.50	<0.50	<0.50	<50	<50	<0.50	4.7	<0.50	<0.50	\5.0	
MW-29	03/23/93	ND	ND	ND	ND	ND	ND						
	07/27/93	ND	ND	ND	ND	ND	ND						
	11/05/93	ND	ND	2.1	11	ND	ND						
	02/25/94	<1	<1	<1	<1	<100	<1,000						
	06/03/94	<0.5	<0.5	<0.5	<0.5	<50	<20,000						
	08/31/94	<0.3	<0.3	<0.3	<0.6	<500	<500						No. Post of sold of
	12/22/94	< 0.5	<0.5	<0.5	<0.5	<50 <50	<50						Non-diesel peak reported.
	03/13/95	0.59	<0.5	<0.5	<0.5	<50 <100	<400 <50						
	06/09/95	<0.5	<0.5	<0.5	<0.5	<100 <50	<50						
	09/21/95	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<0.5 <1.0	<50 <100	<50 <50						
	12/12/95 03/12/96	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<1.0 <1.0	<100 <100	<50 <50						
	06/21/96			~0.5 	~1.0								
	08/29/96	<0.5	<0.5	<0.5	<0.5	<50	<150						
	01/16/97	6.6	8.9	0.6	9.3	120	<150	47	24	<0.5	<0.5	1.8	
	07/07/97	<0.5	<0.5	<0.5	<0.5	<50	<150	52	21	<5.0	<5.0	1.2	
	01/27/98	<0.5	<0.5	<0.5	<0.5	100	<150					8.0	
	07/22/98	<0.5	<0.5	<0.5	<0.5	<50	<250	12	29		<1.0	7.8	
	02/05/99	<0.5	<0.5	<0.5	<0.5	<50	<150	<0.5	68	<0.5	<0.5	8.5	
	04/07/99	<0.5	<0.5	<0.5	<0.5	<50	<250	30	38	<0.5	<0.5	4.9	1,1-DCE detected, 1.4 µg/L.
	07/23/99	<0.5	<0.5	<0.5	<0.5	<50	<200	44	33	<0.5	1.9	4.70	1,1-Dichloroethene detected at 2.3 μg/L; cis-1,2-Dichloroethene detected at 2.3 μg/L.
	10/27/99	<0.5	<0.5	<0.5	<0.5	<100	<200	36	23		<0.5		
	02/08/00	<0.5	<0.5	<0.5	<0.5	<50	<250	87	25	<0.5	<0.5	18.0	1,1-Dichloroethene detected at 9.6 µg/L.
	04/26/00	<0.5	<0.5	<0.5	<0.5	<100	<250	61	38	<0.5	<0.5	12	1,1-Dichloroethene detected at 5.2 µg/L.
	08/16/00	<0.5	<0.5	<0.5	<0.5	<50		49	21	<0.5	<0.5	17	1,1-Dichloroethene detected at 6.0 µg/L.
	10/23/00	<0.5	<0.5	<0.5	<0.5	<50	<250	94	40	<0.5	<0.5	34	1,1-Dichloroethene detected at 14 μg/L.
	01/31/01	<0.5	<0.5	<0.5	<0.5	60	<250	100	35	<0.5	<0.5	26	1,1-Dichloroethene detected at 13 µg/L.
	04/26/01	<0.5	<0.5	<0.5	<0.5	<200	270	87	38	<0.5	<0.5	39.1	1,1-Dichloroethene detected at 12 µg/L.
	07/30/01	1.25	1.28	1.1	5.99	220	<250	120	42	<0.5	<0.5	42.3	1,1-Dichloroethene detected at 13 µg/L.
	10/29/01	<0.5	<0.5	<0.5	<1.0	<200	<500	120	34	<0.5	<0.5	28.0	1,1-Dichloroethene detected at 14 µg/L.
	01/28/02	< 0.5	<0.5	<0.5	<1.0	<200	<250	120	44	<0.5	<0.5	28.9	1,1-Dichloroethene detected at 26 µg/L.
	04/29/02	4.95	<0.5	<0.5	<1.0	<200	<250	130	29	<0.5	<0.5	20.9	1,1-Dichloroethene detected at 23 µg/L.
	10/22/02	<0.5	<0.5	<0.5	<1.0	<200	<250	140	26	<0.5	<0.5	18.1	1,1-Dichloroethene detected at 19 µg/L.
	11/15/02 05/06/03	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<1.0 <1.0	<200 <200	<250 <250	120 140	26 31	<0.5 <0.5	<0.5 <0.5	13.9 13.1	1,1-dichloroethene detected at 15 µg/L. 1,1-Dichloroethene detected at 24 µg/L.
	10/14/03	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<1.0 <1.0	<200 <200	<250 <250	110	22	<0.5	<0.5	11.9	Chloromethane detected at 24 µg/L.
	04/27/04	<0.5	<0.5	<0.5	<1.0	<200	<250	160	28	<0.5	<0.5	15.3	1,1-Dichloroethene detected at 31 µg/L.
	11/17/04	<1.0	<1.0	<1.0	<1.0	120	<50	33	6.5	<0.50	<0.50	120	1,1-Dichloroethene detected at 5.5 µg/L.
MANA 20													., detected at 0.0 pg/L.
MW-30	03/23/93	ND	ND	ND	ND	ND	ND						
	07/27/93	ND	ND	ND	ND	ND	ND						
	11/05/93	ND 13	ND	ND	2.8 <1	ND <100	ND <1,000						
	02/25/94 06/03/94	1.3 1.1	<1 <0.5	<1 <0.5	<1 <0.5	<100 <50	<1,000						
	08/31/94	0.8	<0.5	<0.5	<0.5 <0.6	<500 <500	<20,000 <500						
	12/22/94	0.6	<0.5	<0.5	<0.5	<500 <50	<500 <50						Non-diesel peak reported.
	03/13/95	0.0	<0.5	<0.5	<0.5	<50 <50	<400						Tron Globel peak reported.
	06/09/95	<0.5	<0.5	<0.5	<0.5	<100	<50						
	09/21/95	<0.5	<0.5	<0.5	<0.5	<50	<50						
	12/12/95	<0.5	<0.5	<0.5	<1.0	<100	<50						
	03/12/96	<0.5	<0.5	<0.5	<0.5	<100	<50						
	06/21/96												
	08/29/96	<0.5	<0.5	<0.5	<0.5	<50	<150						
	01/16/97	<0.5	<0.5	<0.5	0.6	80	<150	<0.5	<0.5	<0.5	0.9		
	07/07/97	<0.5	<0.5	<0.5	<0.5	<50	<150					<0.5	
	01/27/98	5.4	<0.5	<0.5	<0.5	100						<0.5	
	07/22/98	<0.5	<0.5	<0.5	<0.5	<50						<0.5	
	04/07/99	<0.5	<0.5	<0.5	<0.5	<50	<250					<0.5	
	07/22/99	<0.5	<0.5	<0.5	<0.5	<50		<0.5	<0.5	<0.5	<0.5	<0.5	
	01122133					i e		1	1			i	1
	10/28/99	<0.5	<0.5	<0.5	<0.5	<100	<200	<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 <0.5	<100 <50 <100	<200 <250 250	<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	

Well Number	Date Sampled	Benzene µg/L	Toluene μg/L	Ethyl- Benzene µg/L	Xylenes µg/L	TPH-G μg/L	TPH-D μg/L	1,1- DCA µg/L	1,2- DCA µg/L	1,1,1- ΤCA μg/L	TCE µg/L	MTBE µg/L	Notes
MW-30	08/04/00	<0.5	<0.5	<0.5	<0.5	<50	<250	<0.5	<0.5	<0.5	<0.5	<0.5	
(cont.)	10/24/00	5.4	<0.5	<0.5	<0.5	<50	<250	<0.5	<0.5	<0.5	<0.5	<0.5	
	01/31/01	<0.5	<0.5	<0.5	<0.5	<50	<250	<0.5	<0.5	<0.5	<0.5	<0.5	
	04/27/01 07/30/01	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<200 <200	<250 <250	<0.5	<0.5 <0.5	<0.5	<0.5 <0.5	<0.5 <0.5	
	10/29/01	<0.5	<0.5	<0.5	<0.5 <1.0	<200	<500	<0.5 <0.5	<0.5	<0.5 <0.5	<0.5	<0.5	Chloroethane detected at 1.3 µg/L.
	01/29/01	<0.5	<0.5	<0.5	<1.0	<200	<250	<0.5	<0.5	<0.5	<0.5	<0.5	Chloroetrane detected at 1.5 µg/L.
	04/30/02	<0.5	<0.5	<0.5	<1.0	<200	<250	<0.5	<0.5	<0.5	<0.5	<0.5	
	10/22/02	<0.5	<0.5	<0.5	<1.0	<200	<250	<0.5	<0.5	<0.5	<0.5	<0.5	
	05/06/03	<0.5	<0.5	<0.5	<1.0	<200	<250	<0.5	<0.5	<0.5	<0.5	<0.5	
	10/14/03	<0.5	<0.5	<0.5	<1.0	<200	<250	<0.5	<0.5	<0.5	<0.5	<0.5	
	04/27/04	<0.5	<0.5	<0.5	<1.0	<200	<250	<0.5	<0.5	<0.5	<0.5	<0.5	
	11/17/04	<0.50	<0.50	<0.50	<0.50	<50	140	<0.50	<0.50	<0.50	<0.50	<5.0	
MW-32	03/23/93	391	6.2	3.1	9	440	ND	ND	60	ND	ND		
	07/27/93	ND	ND	ND	ND	ND	ND	ND	14	ND	ND		
	11/05/93	20	ND	1.8	2.1	170	ND	ND	7.9	ND	ND		
	02/25/94	5.6	<1	<1	<1	<100	<1,000	<1	<1	<1	<1		
	06/03/94	120	1.3	<0.5	1.4	350	<20,000	<0.5	11	<0.5	<0.5		
	08/31/94	39	0.5	2.2	1.2	<500	<500	<4.0	10	<4.0	<4.0		
	12/22/94	4.8	<0.5	<0.5	<0.5	<50	<50	<2.0	4.6	<2.0	<2.0		Non-diesel peak reported.
	03/13/95	220	3.6	6.5	5.8	1,100	<400	<0.5	16	<0.5	<0.5		
	06/09/95	1,500	7.9	43	14	2,200	180	0.7	<0.5	0.5	<0.5		
	09/21/95 12/12/95	1,200	2.4	72	4.5	2,300	60	<0.5	6.7	<0.5	1.4		
	03/12/96	230 40	<0.5 <0.5	8.9 1.7	<1.0 <0.5	500 110	<50 <50	<0.5 <0.5	28 6.8	<0.5 <0.5	<0.5 <0.5		
	06/21/96										~0.5		
	08/29/96	150	<0.5	49	<0.5	700	<150	<0.5	27	<0.5	<0.5		
	01/16/97	14	<0.5	1.9	<0.5	150	<150	<0.5	10	<0.5	0.7		cis-1,2-DCE detected, 0.8 μg/L.
	07/07/97	370	11	110	21	1,600	190					11	· · · · · · · · · · · · · · · · · · ·
	01/27/98	13	<0.5	1.0	<0.5	300		<0.5	7.5	<0.5	<0.5	2.5	
	07/22/98	700	55	88	66	2,300						14	
	07/22/99	59.0	0.80	1.80	<0.5	900	220	<0.5	5.9	<0.5	<0.5	8.70	
	10/28/99	95	2.5	2.1	1.6	500	<200	<0.5	12		<0.5		
	02/10/00	7.0	<0.5	<0.5	<0.5	120	<250	<0.5	4.3	<0.5	<0.5	1.10	
	04/27/00	240	7.0	12	18.8	800	250	<0.5	9.8	<0.5	<0.5	<0.5	
	08/03/00	620	3.0	14	4.1	1,300	<250	<0.5	3.0	<0.5	<0.5	<0.5	
	10/23/00	430	4.30	5.50	8.80	1,200	260	<0.5	7.8	<0.5	<0.5	<0.5	
	01/31/01	42 268	1.5	0.90	2.8 22.0	280 780	<250 <250	<0.5 <0.5	5.7 6.3	<0.5 <0.5	<0.5 <0.5	3.6	
	04/26/01 07/30/01	29.4	13.0 <0.5	22.1 0.52	0.51	320	<250	<0.5	6.6	<0.5	<0.5	<0.5 <0.5	
	10/29/01	16.1	2.01	1.14	3.96	<200	<500	<0.5	5.4	<0.5	<0.5	<0.5	
	01/29/02	12.0	<0.5	0.70	<1.0	<200	<250	<0.5	4.9	<0.5	2.0	<0.5	cis 1,2-Dichloroethene detected at 1.3 μg/L.
	04/29/02	188	5.52	9.70	13.0	680	<250	<0.5	6.0	<0.5	<0.5	<0.5	5.5 1,2 2.6.116.166.16.16 dottotted at 1.6 pg/2.
	10/22/02	4.84	<0.5	<0.5	<1.0	<200	<250	<0.5	4.8	<0.5	<0.5	<0.5	
	05/06/03	20.72	0.76	0.86	2.08	<200	<250	<0.5	5.8	<0.5	<0.5	<0.5	
	10/14/03	6.02	<0.5	<0.5	<1.0	<200	<250	<0.5	3.2	<0.5	<0.5	<0.5	
	04/27/04	23.60	1.68	0.67	3.91	<200	<250	<0.5	3.0	<0.5	<0.5	<0.5	
	11/17/04	2.0	<0.50	<0.50	<0.50	<50	<50	<0.50	2.1	<0.50	<0.50	<5.0	
MW-33	04/07/99	0.60	<0.5	0.90	<0.5	<50	<250					<0.5	
	07/22/99	8.90	<0.5	1.00	<0.5	<50	<200	0.6	0.7	<0.5	<0.5	<0.5	
	10/28/99	40	0.9	21	3.8	200	<200	0.8	1.3		<0.5		
	02/10/00	20	0.7	12	10.0	380	<250	0.9	0.6	<0.5	<0.5	1.30	
	04/27/00	6.9	<0.5	6.4	<0.5	<100	250	4.3	0.9	<0.5	<0.5	<0.5	
	08/03/00	31	0.5	20	1.0	150	550	<0.5	0.6	<0.5	<0.5	<0.5	
	10/23/00	89	1.5	36	3.9	350	<250	<0.5	2.1	<0.5	<0.5	<0.5	
	01/31/01	6.8	<0.5	2.0	<0.5	<50	<250	1.9	0.6	<0.5	<0.5	0.7	
	04/26/01	6.61	0.56	1.63	0.61	<200	<250	2.6	<0.5	<0.5	<0.5	<0.5	
	07/30/01	4.43	2.61	1.34	6.6	<200 <200	<250 <500	2.2	0.5	<0.5	<0.5	<0.5	Dichlorodifluoromethane detected at 0.6 μg/L.
	10/29/01 01/28/02	14.2	<0.5	0.63	<1.0	<200 <200	<500 <250	1.3	0.7	<0.5 <0.5	<0.5	<0.5 <0.5	Dichlorodiflouromethane detected at 1.9 µg/L.;
	01/20/02	<0.5	<0.5	<0.5	<1.0	<200	<250	1.1	0.5	~0.5	3.8	~U.5	cis 1,2-Dichloroethene detected at 8.9 µg/L.;
	04/29/02	14.6	<0.5	1.41	<1.0	<200	<250	0.8	0.9	<0.5	<0.5	<0.5	Dichlorodiflouromethane detected at 1.9 μg/L.
MW-100	07/06/01	<0.5	<0.5	<0.5	<0.5	<200	<250	<0.5	<0.5	<0.5	<0.5	<0.5	
	07/30/01	<0.5	<0.5	<0.5	<0.5	<200	<250	<0.5	<0.5	<0.5	<0.5	<0.5	Chloromethane detected at 1.8 μg/L.
	10/30/01	<0.5	<0.5	<0.5	<1.0	<200	<500	<0.5	<0.5	<0.5	<0.5	<0.5	
	01/28/02	<0.5	<0.5	<0.5	<1.0	<200	<250	<0.5	<0.5	<0.5	<0.5	< 0.5	
	04/29/02	<0.5	<0.5	<0.5	<1.0	<200 <200	<250	<0.5	<0.5	<0.5	<0.5	<0.5	
	10/22/02	<0.5	<0.5	<0.5	<1.0	<200	<250 <250	<0.5	<0.5	<0.5	<0.5	<0.5	
	05/06/03 10/14/03	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<1.0 <1.0	<200 <200	<250 <250	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	
	04/27/04	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<1.0 <1.0	<200 <200	<250 <250	<0.5	<0.5	<0.5	<0.5	<0.5	
	11/17/04	<0.50	<0.50	<0.50	<0.50	<50	<50	<0.50	<0.50	<0.50	<0.50	<5.0	
		0.00	.0.50	0.00	0.00	-50	-00	3.50	0.00	0.00	3.50	-0.0	

Well Number	Date Sampled	Benzene µg/L	Toluene µg/L	Ethyl- Benzene µg/L	Xylenes µg/L	TPH-G μg/L	TPH-D μg/L	1,1- DCA µg/L	1,2- DCA µg/L	1,1,1- ΤCA μg/L	TCE µg/L	MTBE µg/L	Notes
MW-?	02/05/99	<0.5	<0.5	<0.5	<0.5	<50	430		-			<0.5	
PR-26	07/26/99 10/26/99	20,000 28,000	15,000 25,000	1,100 2,300	7,250 8,400	82,500 110,000	11,000 60,000	 <0.5	 24		 <0.5	33.0	
PR-45	07/26/99 10/28/99	13,200 12,000	8,200 8,200	2,600 1,700	15,600 8,500	82,500 45,000	39,000 25,000	 <0.5	 <0.5		 <0.5	35.0	
	02/09/00	24,000	25,000	10,000	53,000	360,000	82,000	<0.5	4.0	<0.5	<0.5	1,000	
	04/27/00	17,000	9,500	16,000	92,000	1,300,000	20,300	<5.0	<5.0	<5.0	<5.0	<5.0	
	08/04/00	20,000	8,800	2,600	16,000	73,000	54,500	<0.5	1.0	<0.5	<0.5	<0.5	
	10/23/00	26,000	12,000	4,000	20,000	96,000	36,000	<0.5	1.2	<0.5	<0.5	<5.0	Chloroethane detected at 6.0 µg/L.
	04/27/01 07/30/01	16,200 14,500	8,600 8,900	3,220 4,400	19,000 24,700	178,000 132,000	22,700 29,700	<0.5 <0.5	14 11	<0.5 <0.5	<0.5 <0.5	<25 <50	Chloroethane detected at 4.6 µg/L. Chloromethane detected at 0.6 µg/L;
	07/30/01	14,500	0,900	4,400	24,700	132,000	29,700	~0.5	11	~0.5	~ 0.5	\50	Chloroethane detected at 11 µg/L; Methylene chloride detected at 0.5 µg/L.
	10/29/01	12,600	6,650	2,260	12,400	86,100	50,000	<0.5	7.8	<0.5	<0.5	<25	Chloroethane detected at 6.0 µg/L.
	01/29/02	8,930	4,860	2,640	12,700	114,000	19,400	<0.5	30	<0.5	<0.5	<0.5	Chloroethane detected at 7.5 µg/L.
	05/16/02	14,300	2,630	1,580	7,780	125,000	15,600	<0.5	1.0	<0.5	<0.5	<0.5	Chloroethane detected at 7.3 µg/L.
PR-52	07/26/99	12,000	1,720	750	12,400	172,000	40,000	<0.5	1.8	<0.5	<0.5	217	Methylene chloride detected at 7.9 μg/L.
	10/28/99	19,000	530	1,800	5,800	40,000	450,000	<0.5	<0.5		<0.5		,
	02/09/00	22,000	1,600	4,100	15,800	200,000	140,000	<0.5	1.3	<0.5	<0.5	430	
	04/28/00	20,000	2,200	4,700	18,600	270,000	88,000	<1.0	<1.0	<1.0	<1.0	<5.0	
	08/04/00 10/24/00	26,000	1,600	2,900	15,000 180,000	150,000 650,000	110,000 280,000	<0.5 <5.0	2.3 <5.0	<0.5 <5.0	<0.5 <5.0	<0.5 <5.0	
	01/31/01	52,000 81,000	13,000 840	41,000 57,000	210,000	5,300,000	276,000	<0.5	1.0	<0.5	<0.5	<5.0 500	Chloroethane detected at 2.4 µg/L;
													Methylene chloride detected at 0.6 μg/L.
	04/27/01 07/30/01	25,000	16,300	14,700	55,000 51,700	886,000	134,000	<0.5 <0.5	<0.5 1.3	<0.5 <0.5	<0.5	1,040	Chloroethane detected at 1.5 μg/L. Chloromethane detected at 13 μg/L;
	07/30/01	31,100	2,480	13,500	51,700	340,000	185,000	<0.5	1.3	V 0.5	<0.5	2,510	Chloroethane detected at 46 µg/L;
	10/29/01	22,700	1,630	3,070	11,500	126,000	140,000	<0.5	0.9	<0.5	<0.5	<50	Methylene chloride detected at 0.6 μg/L. Chloromethane detected at 0.6 μg/L;
	10/29/01	22,700	1,030	3,070	11,500	120,000	140,000	~ 0.5	0.9	\0.5	~ 0.5	\ 50	Chloroethane detected at 4.0 µg/L;
													Methylene chloride detected at 0.7 μg/L.
	01/29/02 05/16/02	21,500 31,600	1,840 53,600	4,540 43,800	16,800 216,000	517,000 2,020,000	272,000 75,000	<0.5 <5.0	<0.5 <5.0	<0.5 <5.0	<0.5 <5.0	44.1 63.5	Chloroethane detected at 1.5 µg/L. Chloroethane detected at 8.3 µg/L.
PR-53	07/26/99	31,000	12,000	1,900	8,800	110,000	98,000	<0.5	43	<0.5	<0.5	43.0	Methylene chloride detected at 6.2 µg/L.
	10/27/99	17,000	3,900	890	3,320	54,000	16,000	<0.5	18		<0.5		Wellylone chloride detected at 0.2 pg/2.
	02/09/00	21,000	5,000	1,200	5,300	65,000	9,400	0.6	20	<0.5	<0.5	67.0	Methylene chloride detected at 0.8 μg/L.
	04/28/00	34,000	30,000	9,300	51,000	730,000	104,000	<1.0	<1.0	<1.0	<1.0	340	
	08/04/00	35,000	17,000	3,800	24,000	180,000	69,500	<0.5	1.7	<0.5	<0.5	110	
	10/24/00 01/31/01	99,000 66,000	110,000 15,000	80,000 28,000	640,000 140,000	580,000 2,400,000	380,000 960,000	<5.0 <0.5	5.0 1.5	<5.0 <0.5	<5.0 <0.5	380 660	Chloroethane detected at 1.7 μg/L;
	04/27/01	55,500	10,000	23,700	137,000	4,240,000	806,000	<0.5	<0.5	<0.5	<0.5	<5,000	Methylene chloride detected at 1.7 μg/L, Chloroethane detected at 1.7 μg/L,
	10/29/01	46,500	9,520	12,900	74,000	1,630,000	130,000	<0.5	0.8	<0.5	<0.5	<500	Methylene chloride detected at 1.1 µg/L. Chloroethane detected at 3.0 µg/L;
													Methylene chloride detected at 0.9 μg/L.
	01/29/02	33,000	7,340	10,300	41,800	495,000 3.280.000	462,000	<0.5 <5.0	1.8 <5.0	<0.5 <5.0	<0.5 <5.0	122 242	Chloroethane detected at 3.2 μg/L.
	05/16/02	35,800	10,500	18,700	130,000	-,,	113,000						
PR-54	07/26/99	32,000	22,000	1,500	21,800	170,000	28,000 350,000	<0.5	3.0	<0.5	<0.5 <0.5	56.0	Methylene chloride detected at 2.5 μg/L.
	10/26/99 02/09/00	27,000 27,000	10,000 23,000	3,700 9,900	19,500 50,000	190,000 960,000	110,000	<0.5 <0.5	<0.5 3.9	<0.5	<0.5	1,000	
	04/28/00	24,000	14,000	1,200	9,000	76,000	80,000	<1.0	1.6	<1.0	<1.0	300	
	08/04/00	27,000	7,600	1,400	11,000	120,000	54,500	<0.5	2.0	<0.5	<0.5	200	
	10/24/00	23,000	4,400	2,000	13,000	140,000	96,000	<0.5	2.3	<0.5	<0.5	<100	Chloroethane detected at 5.3 µg/L; Methylene chloride detected at 2.3 µg/L.
	01/31/01	30,000	8,300	3,300	21,000	220,000	236,000	<0.5	2.6	<0.5	<0.5	480	Chloroethane detected at 2.8 µg/L; Methylene chloride detected at 1.7 µg/L.
	04/27/01	26,100	8,650	2,120	15,900	51,300	108,000	<0.5	<0.5	<0.5	<0.5	<500	Chloroethane detected at 1.7 µg/L.
	07/30/01	31,700	18,000	9,880	58,400	320,000	71,200	<0.5	3.9	<0.5	<0.5	2,750	Chloromethane detected at 2.2 µg/L;
													Chloroethane detected at 22 µg/L; Methylene chloride detected at 2.6 µg/L.
	10/30/01	25,400	11,300	3,500	18,800	222,000	530,000	<0.5	1.2	<0.5	<0.5	276	Chloroethane detected at 7.4 μg/L;
	01/29/02	13,300	9,850	4,240	33,100	108,000	48,000	<0.5	7.5	<0.5	<0.5	51.3	Methylene chloride detected at 2.0 μg/L. Chloroethane detected at 6.2 μg/L.
	05/16/02	27,900	34,500	5,630	36,400	324,000	172,000	<5.0	43	<5.0	<5.0	251	Chloroethane detected at 9.8 μg/L.
PR-64	07/26/99	22,000	18,000	1,700	10,300	110,000		<0.5	130	<0.5	<0.5	35.0	Methylene chloride detected at 1.4 μg/L.
	10/27/99 02/09/00	11,000 22,000	7,400 20,000	1,200 6,000	3,900 17,000	66,000 120,000	50,000 40,000	<0.5 <0.5	110 >50	<0.5	<0.5 <0.5	 110	
	04/28/00	19,000	16,000	1,800	13,900	130,000	78,000	<1.0	67	<1.0	<1.0	300	
	05/16/02	18,300	40,100	10,400	104,000	30,600,000	419,000	<5.0	<5.0	<5.0	<5.0	<500	
PR-65	07/26/99	12,000	1,400	1,300	13,000	68,000	16,500	<0.5	2.6	<0.5	<0.5	20.0	
	10/26/99	14,000	2,300	1,800	11,000	65,000	50,000	<0.5	<0.5		<0.5		1

| Well Number | Date
Sampled | Benzene
µg/L | Toluene
μg/L | Ethyl-
Benzene
µg/L

 | Xylenes
μg/L | TPH-G
μg/L | TPH-D
μg/L | 1,1-
DCA
µg/L | 1,2-
DCA
µg/L
 | 1,1,1-
ΤCA
μg/L | TCE
μg/L | MTBE
µg/L | Notes |
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| PR-68 | 07/26/99
10/26/99 | 1,900
2,800 | 24.0
36 | 27.0
86

 | 62.0
62 | 4,900
8,000 | 11,000
2,800 | <0.5
<0.5 | 1.2
<0.5
 | <0.5
 | <0.5
<0.5 | 4.40 | |
| PR-76 | 04/07/99 | <0.5 | <0.5 | <0.5

 | <0.5 | <50 | <250 | | -
 | | | <0.5 | |
| | 10/22/02 | <0.5 | <0.5 | <0.5

 | <1.0 | <200 | <250 | <0.5 | <0.5
 | <0.5 | <0.5 | <0.5 | |
| | 05/06/03
10/14/03 | <0.5
<0.5 | <0.5
<0.5 | <0.5
<0.5

 | <1.0
<1.0 | <200
<200 | <250
<250 | <0.5
<0.5 | <0.5
<0.5
 | <0.5
<0.5 | <0.5
<0.5 | <0.5
<0.5 | |
| | 04/27/04 | <0.5 | <0.5 | <0.5

 | <1.0 | <200 | <250 | <0.5 | <0.5
 | <0.5 | <0.5 | <0.5 | |
| | 11/17/04 | <0.50 | <0.50 | <0.50

 | <0.50 | <50 | 85 | <0.50 | <0.50
 | <0.50 | <0.50 | <5.0 | |
| V-24 | 04/07/99 | <0.5 | <0.5 | <0.5

 | <0.5 | 120 | <250 | |
 | | | 0.5 | |
| V-31 | 07/26/99 | 7,000 | 600 | 550

 | 1,370 | 17,500 | 5,350 | |
 | | | 19.0 | |
| V-51 | 10/26/99 | 7,000 | 120 | 850

 | 950 | 18,000 | 3,000 | <0.5 | <0.5
 | | <0.5 | | |
| V-46 | 02/05/99 | <0.5 | <0.5 | <0.5

 | <0.5 | <50 | 270 | <0.5 | <0.5
 | <0.5 | <0.5 | <0.5 | |
| V-55 | 07/22/99 | 8,000 | 480 | 740

 | 2,880 | 30,000 | 2,100 | <0.5 | <0.5
 | <0.5 | <0.5 | 13.0 | |
| | 10/28/99 | 11,000 | 59 | 1,200

 | 317 | 28,000 | 38,000 | <0.5 | <0.5
 | | <0.5 | | |
| | 02/09/00 | 2,200 | 59
540 | 760

 | 350 | 7,900 | 10,000 | <0.5 | <0.5
 | <0.5 | <0.5 | 9.70 | |
| | 04/28/00
08/03/00 | 2,900
9,400 | 510
380 | 440
720

 | 2,340
2,200 | 14,000
28,000 | 26,500
70,000 | <5.0
<0.5 | <5.0
<0.5
 | <5.0
<0.5 | <5.0
<0.5 | <5.0
<0.5 | |
| | 10/23/00 | 11,000 | 140 | 900

 | 1,300 | 30,000 | 51,000 | <0.5 | <0.5
 | <0.5 | <0.5 | <12 | |
| | 01/31/01 | 4,600 | 57 | 550

 | 1,200 | 34,000 | 88,500 | <0.5 | <0.5
 | <0.5 | <0.5 | 44 | |
| | 04/26/01 | 6,400 | 61.5 | 250

 | 336 | 34,200 | 227,000 | <0.5 | <0.5
 | <0.5 | <0.5 | <25 | |
| | 10/30/01 | 5,360 | 70.0 | 1,090

 | 1,450 | 32,700 | 78,000 | <0.5 | <0.5
 | <0.5 | <0.5 | <25 | |
| | 01/29/02 | 1,660 | 140 | 492

 | 818 | 12,000 | 4,100 | <0.5 | <0.5
 | <0.5 | <0.5 | <0.5 | |
| | 04/29/02 | 5,170 | 95.1 | 572

 | 523 | 30,600 | 35,100 | <0.5 | <0.5
 | <0.5 | <0.5 | 1.06 | |
| V-72 | 07/26/99 | 13,500 | 6.80 | 1.10

 | 3.90 | 3,900 | 12,900 | <0.5 | 11
 | <0.5 | <0.5 | <0.5 | |
| | 10/28/99
02/09/00 | 2,900
670 | 58
8.2 | 21
<0.5

 | 47.7
17.8 | 6,000
890 | 48,000
6.100 | <0.5
<0.5 | 3.4
3.0
 |
<0.5 | <0.5
<0.5 |
<0.5 | |
| | 04/28/00 | 130 | 8.2
<0.5 | <0.5
<0.5

 | 17.8
<0.5 | 200 | 6,100
5,950 | <0.5 | 0.7
 | <0.5 | <0.5 | <0.5 | |
| | 08/04/00 | 460 | 0.8 | <0.5

 | 0.6 | 440 | 4,120 | <0.5 | 2.8
 | <0.5 | <0.5 | <0.5 | |
| | 10/24/00 | 2,700 | 3.2 | 0.5

 | 2.3 | 3,500 | 17,000 | <0.5 | 4.0
 | <0.5 | <0.5 | <0.5 | |
| | 04/27/01 | 1,240 | 2.05 | <0.5

 | 2.78 | 1,310 | 6,290 | <0.5 | 5.1
 | <0.5 | <0.5 | <0.5 | Dichlorodifluoromethane detected at 0.8 μg/L. |
| | 07/30/01 | 1,790 | 69.8 | 1.22

 | 2.50 | 1,490 | 4,290 | <0.5 | 6.2
 | <0.5 | <0.5 | <0.5 | Chloromethane detected at 1.5 µg/L. |
| | 10/29/01 | 1,330 | 4.38 | 0.55

 | 3.32 | 1,960 | | <0.5 | 5.6
 | <0.5 | <0.5 | <0.5 | Chloromethane detected at 1.1 µg/L. |
| | 01/29/02 | 655 | 6.40 | <0.5

 | 8.00 | 1,840 | 2,250 | <0.5 | 3.9
 | <0.5 | < 0.5 | < 0.5 | Chloromethane detected at 1.8 µg/L. |
| | 05/16/02 | 43.8 | 1.09 | <0.5

 | 4.36 | 230 | 5,120 | <0.5 | <0.5
 | <0.5 | <0.5 | <0.5 | Chloromethane detected at 1.8 μg/L. |
| V-84 | 07/26/99
10/26/99 | 2,400
1,100 | 440
130 | 80.0
46

 | 340
108 | 8,700
4,000 | 2,350
700 | <0.5
<0.5 | 2.4
<0.5
 | <0.5 | <0.5
<0.5 | 6.40 | |
| | 02/09/00 | 300 | 30 | 8.9

 | 53 | 2,300 | 1,100 | <0.5 | 1.2
 | <0.5 | <0.5 | <0.5 | |
| | 04/28/00 | 30 | 1.9 | <0.5

 | <0.5 | 100 | 550 | <5.0 | <5.0
 | <5.0 | <5.0 | <0.5 | |
| | 08/04/00 | | |

 | | | 1,380 | | 1.0
 | <0.5 | <0.5 | <0.5 | |
| | 06/04/00 | 900 | 110 | 34

 | 120 | 2,700 | 1,300 | <0.5 | 1.0
 | | | 40 F | |
| | 10/24/00 | 900
2,000 | 110
480 | 34
24

 | 120
110 | 2,700
48,000 | 1,900 | <0.5 | 1.0
 | <0.5 | <0.5 | <0.5 | |
| | 10/24/00
01/31/01 | 2,000
68 | 480
1.3 | 24
5.3

 | 110
8.2 | 48,000
970 | 1,900
1,820 | <0.5
<0.5 | 1.0
<0.5
 | <0.5 | <0.5 | <0.5 | |
| | 10/24/00
01/31/01
04/26/01 | 2,000
68
925 | 480
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8.2
59.7 | 48,000
970
2,360 | 1,900
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1,180 | <0.5
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<0.5 | |
| | 10/24/00
01/31/01
04/26/01
07/30/01 | 2,000
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01/31/01
04/26/01
07/30/01
10/30/01
01/29/02
04/29/02
07/23/99
10/28/99
02/08/00
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Corrective Action Plan Report, May 2009 Former Nestle USA, Inc. Facility 1310 14th Street, Oakland, CA

Well Number	Date	Benzene	Toluene	Ethyl- Benzene	Xylenes	TPH-G	TPH-D	1,1- DCA	1,2- DCA	1,1,1- TCA	TCE	MTBE	Notes
well number	Sampled	μg/L	μg/L	μg/L	μg/L	μg/L	μg/L	μg/L	μg/L	μg/L	μg/L	μg/L	Notes
30 (CC-2)	07/30/01	<0.5	1.43	<0.5	1.63	<200	<250	<0.5	1.6	<0.5	<0.5	<0.5	Dichlorodifluoromethane detected at 2.8 µg/L.
(cont.)	10/29/01	<0.5	<0.5	<1.0	<0.5	<200	<500	<0.5	<0.5	<0.5	<0.5	<0.5	
	01/28/02	<0.5	<0.5	<0.5	<1.0	<200	<250	<0.5	1.9	<0.5	<0.5	<0.5	Dichlorodifluoromethane detected at 3.8 µg/L.
	04/29/02	<0.5	<0.5	<0.5	<0.5	<200	<250	<0.5	2.5	<0.5	<0.5	0.86	Dichlorodifluoromethane detected at 3.6 µg/L.
	10/10/02	<0.5	<0.5	<0.5	<1.0	<200	<250	<0.5	<0.5	<0.5	<0.5	<0.5	Chloroform detected at 0.6 µg/L.
	11/15/02	<0.5	<0.5	<0.5	<1.0	<200	<250	<0.5	<0.5	<0.5	<0.5	<0.5	Chloroform detected at 0.5 µg/L.
	05/06/03	<0.5	<0.5	<0.5	<1.0	<200	<250	<0.5	<0.5	<0.5	<0.5	<0.5	
81	02/05/99	<0.5	<0.5	<0.5	<0.5	<50	<150	<0.5	<0.5	<0.5	<0.5	<0.5	
	07/22/99	0.70	<0.5	<0.5	<0.5	<50	<200	<0.5	<0.5	<0.5	<0.5	<0.5	
94	02/05/99	<0.5	<0.5	<0.5	<0.5	<50	170					<0.5	
	07/22/99	<0.5	<0.5	<0.5	<0.5	<50	<200	<0.5	<0.5	<0.5	<0.5	<0.5	
210	02/05/99	<0.5	<0.5	<0.5	<0.5	<50	960					<0.5	
223	10/26/99	<0.5	<0.5	<0.5	<0.5	<100	<200	<0.5	<0.5		<0.5		
	02/10/00	<0.5	<0.5	<0.5	<0.5	<50	640	<0.5	<0.5	<0.5	<0.5	<0.5	
	04/27/00	<0.5	<0.5	<0.5	<0.5	<100	250	<0.5	<0.5	<0.5	<0.5	<0.5	
	08/03/00	<0.5	<0.5	<0.5	<0.5	<50	680	<0.5	<0.5	<0.5	<0.5	<0.5	
	10/23/00	1.30	<0.5	<0.5	<0.5	<50	<250	<0.5	<0.5	<0.5	<0.5	<0.5	Chlorobenzene detected at 0.9 µg/L.
	01/31/01	<0.5	<0.5	<0.5	<0.5	<50	<250	<0.5	<0.5	<0.5	<0.5	<0.5	
	04/26/01	<0.5	<0.5	<0.5	<0.5	<200	390	<0.5	<0.5	<0.5	<0.5	<0.5	1,2-Dichlorobenzene detected at 0.5 μg/L.
	07/30/01	<0.5	<0.5	<0.5	<0.5	<200	<250	<0.5	<0.5	<0.5	<0.5	<0.5	Dichlorodifluoromethane detected at 0.5 µg/L.
	10/30/01	<0.5	<0.5	<0.5	<1.0	<200	<500	<0.5	<0.5	<0.5	<0.5	<0.5	Chloromethane detected at 0.8 µg/L.
	01/29/02	<0.5	<0.5	<0.5	<1.0	<200	<250	<0.5	<0.5	<0.5	<0.5	<0.5	
	04/29/02	<0.5	<0.5	<0.5	<1.0	<200	<250	<0.5	<0.5	<0.5	<0.5	<0.5	
224	07/26/99	<0.5	<0.5	<0.5	<0.5	<50	640	<0.5	<0.5	<0.5	<0.5	<0.5	
239	07/26/99	55,000	85.0	1,500	190	30,000		<0.5	<0.5	<0.5	<0.5	5.30	
	10/26/99	23,000	53	1,500	103.2	28,000	10,000	<0.5	<0.5		<0.5		
	02/10/00	40,000	48	1,900	52	44,000	21,000	<0.5	1.0	<0.5	<0.5	14.0	
	04/28/00	25,000	540	2,000	710	36,000	12,500	<5.0	<5.0	<5.0	<5.0	<5.0	
	08/04/00	25,000	220	1,900	920	45,000	32,500	<0.5	0.6	<0.5	<0.5	<0.5	
	10/24/00	24,000	100	1,500	390	50,000	50,000	<0.5	<0.5	<0.5	<0.5	<5.0	
	01/31/01	23,000	84	1,900	200	52,000	112,000	<0.5	0.9	<0.5	<0.5	<0.5	
	04/26/01	23,900	113	1,990	590	298,000	143,000	<0.5	<0.5	<0.5	<0.5	<25	
	07/30/01	30,200	384	2,000	966	66,500	19,100	<0.5	<0.5	<0.5	<0.5	<0.5	
	10/30/01	41,200	273	1,470	215	54,300	120,000	<0.5	<0.5	<0.5	<0.5	<50	
	01/28/02 04/29/02	24,500 25,900	228 280	1,670 1,380	352 491	112,000 71,600	6,900 9,400	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	Chloroethane detected at 0.6 µg/L.
241	04/29/02	<0.5	<0.5	<0.5	<0.5	<50	<250					<0.5	
271	04/07/99	\0. 5	~ 0.5	~ 0.5	~ 0.5	\ 00	~250					~0.3	
249	07/22/99	<0.5	<0.5	<0.5	<0.5	<50	<200	<0.5	<0.5	<0.5	<0.5	<0.5	
SB-16	05/20/08	<0.50	<0.50	<0.50	530	<50	530	NA	<0.50	NA	NA	NA	
SB-17	05/22/08	12,000	3,200	17,000	560,000	120,000	560,000	NA	<0.50	NA	NA	NA	
SB-18	05/22/08	50,000	2,300	46,000	23,000	190,000	23,000	NA	2,200	NA	NA	NA	
SB-19	05/22/08	<12	220	<12	1,600	8,200	1,600	NA	<12	NA	NA	NA	
SB-20/ PCB-7	05/22/08	41,000	3,000	30,000	47,000	170,000	47,000	NA	930	NA	NA	NA	
SB-21/ PCB-8	05/23/08	12,000	2,600	20,000	3,500	110,000	3,500	NA	<250	NA	NA	NA	
SB-22	05/22/08	27,000	13,000	39,000	73,000	870,000	73,000	NA	<2,500	NA	NA	NA	
SB-24/ PCB-1	05/21/08	1.1	<0.50	<0.50	360	<50	360	NA	<0.50	NA	NA	NA	
SB-25/ PCB-2	05/21/08	<0.50	<0.50	<0.50	140	<50	140	NA	<0.50	NA	NA	NA	
SB-26	05/22/08	<0.50	<0.50	<0.50	270	<50	270	NA	<0.50	NA	NA	NA	
SB-27/ PCB-3	05/20/08	< 0.50	< 0.50	< 0.50	NA	NA	NA	NA	<0.50	NA	NA	NA	

Notes:

ND Not detected.

NA Not analyzed or not sampled. Micrograms per liter.

μg/L TPH-G

Total Petroleum Hydrocarbons as gasoline.
Total Petroleum Hydrocarbons as diesel.
1,1-Dichloroethane. TPH-D 1,1-DCA 1,2-DCA

1,2-Dichloroethane. 1,1-DCE 1,1-Dichloroethene. 1,1,1-TCA 1,1,1-Trichloroethane. cis1,2-DCE cis 1,2-Dichloroethylene. TCE Trichloroethene. MTBE Methyl tertiary butyl ether.

10/22/02 Data was confirmed anomalous by resampling on 11/15/02.

Table 4: Historical LPH Monitoring (Product Thickness in Feet)

Corrective Action Plan Report, May 2009 Former Nestle USA, Inc. Facility 1310 14th Street, Oakland, CA

Well	11/4/1993	2/24/1993	3/18/1994	6/2/1994	8/31/1994	12/22/1994	3/13/1995	6/9/1995	7/27/1995	9/22/1995	12/6-28/95	2/27/1996	2/29/1996	6/20/1996	8/30/1996	9/18/1996	10/4/1996	10/11/1996	10/18/1996	10/22/1996	11/22/1996	12/6/1996	12/17/1996	12/21/1996	1/3/1997	1/14/1997	2/10/1997	2/17/1997	2/28/1997	3/7/1997	3/14/1997
MW-3 MW-7 MW-8	 0.79 0.47	 1.14 0.44	2.82 0.30	 0.26 0.31	 0.01 0.31	0.04 0.26	 <0.01 0.08	 <0.01 0.09	 0.23	 0.21 0.24	 0.24	 <0.01 <0.01		0.02 0.03	 0.20 0.04	0.04 0.03			 	 	 				 					 	
MW-22 MW-23	1.83 1.21	1.54 0.07	>3.0 1.40	1.14 1.79	0.19 0.68	0.03 0.41	<0.01 <0.01	<0.01 0.31	<0.01 0.44	0.32 0.71	0.30 0.30	<0.01 0.19	0.15	0.01 1.00	0.04 0.24	0.22 0.63															
MW-24 E-0	1.77	12.10	>3.0	0.97	0.39	<0.01	<0.01	<0.01	2.72	1.41	<0.01 <0.01	<0.01 3.92	0.07	2.46 0.18	1.45 <0.01	1.15 <0.01	 <0.01	 <0.01	 <0.01	 <0.01	0.38	1.55	1.45	0.3	0.39		 <0.01	 <0.01	<0.01	 <0.01	 <0.01
E-3 E-5											1.50	0.27	0.03	0.10	0.01	0.04															
E-6 E-8									0.10		0.42	0.19	0.02	<0.01	<0.01	<0.01															
PR-12 PR-20	 0.91	 1.15	3.41	 1.45	0.88	1.04	0.14	0.16	2.54	1.12	 <0.01	3.5	2.65	3.50	0.69	0.47	0.36	0.2													
PR-21 PR-22	0.63 0.98	1.43	2.76 >3.0	1.39 0.90	0.42 0.47	2.01 0.04	4.11 0.60	2.42 0.71	1.93 0.68	0.70 0.71	0.60 0.23	2.99 1.57	0.77 0.94	1.50 1.20	0.86 0.47	0.54 0.42															
PR-23 PR-24	0.67	0.36	1.06	0.38 <0.01	0.17 <0.01	0.06 <0.01	0.34 <0.01	0.06 <0.01	0.08 0.01	0.12 <0.01	0.11 <0.01	<0.01		<0.01	0.09	<0.01															
PR-25 PR-26	0.6	0.54	2.05	0.39	0.17	<0.01	 <0.01	<0.01		0.13	0.12	0.27	 <0.01	0.01	0.07	0.03															
PR-27 PR-28		<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	0.01	<0.01	<0.01																				
PR-29 PR-30				2.81	1.21	 1.97	 <0.01	 <0.01		 Dry	 Dry	 Dry		 Dry	 Dry	 Dry															
PR-32 PR-33																															
PR-34 PR-35	0.66	1.17	2.81	1.07	0.37	2.45	4.06	3.54	2.30	1.03	0.58	5.10	1.22	1.95	1.14	0.48	0.33	0.23	0.01	<0.01	<0.01	0.26	0.59	0.25	<0.01	<0.01	0.75	0.67	0.98	1.15	1.23
PR-36 PR-37	0.41	1.13 1.29	1.43 2.35	1.13 0.96	0.37 0.14	0.19 0.22	0.15 0.83	0.23 0.82	0.22 0.58	Dry 0.58	Dry 0.18	0.20 1.14	0.05 0.32	0.01 0.20	Dry 0.19	Dry 0.11															
PR-38 PR-41	0.59	0.53	0.42	0.13	0.43	0.03	 <0.01	 <0.01		 Dry	 Dry	 Dry		 Dry	 Dry	 Dry															
PR-42 PR-43																															
PR-44 PR-45	0.24 0.17	0.22 5.27	0.19 0.10	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01		Dry <0.01	- <0.01	<0.01 <0.01		Dry <0.01	Dry <0.01	Dry <0.01															
PR-46 PR-47	0.75	0.41	 sheen	 <0.01	<0.01	0.01	 <0.01	<0.01		0.08	0.08	 <0.01		 <0.01	0.08	0.02															
PR-48 PR-49	1.12	0.20 3.24	>3.0 <0.01	0.83 <0.01	0.07 <0.01	1.43 < 0.01	0.64 <0.01	0.65 <0.01	0.94	0.50 <0.01	0.54 <0.01	0.11 <0.01	0.06	2.06 Dry	1.36 Dry	0.38 <0.01															
PR-50 PR-51	1.08	1.58 6.57	0.89 >3.0	<0.01 <0.01	<0.01 0.72	<0.01 2.02	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 Dry		<0.01 Dry	<0.01 Dry	<0.01 <0.01															
PR-52 PR-53	1.01 1.15	5.09 3.01	1.16 >3.0	0.45 0.61	0.05 0.49	0.03 1.52	<0.01 <0.01	<0.01 1.55	<0.01 1.47	<0.01 1.08	0.17	<0.01	0.27	<0.01	<0.01	<0.01 0.38															
PR-54 PR-55	0.97 1.48	0.99 0.07	1.20 1.31	<0.01 0.87	0.08 <0.01	0.01	<0.01 <0.01	<0.01 Dry	 Dry	<0.01 Dry	<0.01	<0.01 Dry		<0.01 Dry	<0.01 Dry	<0.01 Dry															
PR-56 PR-57	0.90	1.30 6.40		0.89 <0.01	0.15 <0.01	1.48 < 0.01	<0.01 <0.01	<0.01	0.01	<0.01	-	<0.01		<0.01	<0.01	Dry <0.01															
PR-58 PR-60	0.96	0.85 <0.01		1.48 < 0.01	0.89 <0.01	2.15 <0.01	1.41 < 0.01	1.34 < 0.01	2.40 0.01	1.18 < 0.01	0.57 <0.01	2.67	1.25	2.79	1.47	1.01		0.52	0.23	0.11	<0.01	<0.01	<0.01	<0.01	0.2	1.04	2.3	2.4	2.21	2.45	
PR-61 PR-62	0.25 0.04	0.39	0.35 0.07	1.03 0.09	<0.01 <0.01	0.01 <0.01	<0.01 <0.01	<0.01 <0.01	1.30	<0.01 <0.01	<0.01 <0.01	1.48 < 0.01	0.45	1.96	0.93	0.38				<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	0.45	0.34	0.6	0.55	0.77
PR-64 PR-65	1.49 0.04	0.11 0.02	>3.0 0.09	0.08	1.06 < 0.01	2.15 <0.01	1.03 < 0.01	1.17 < 0.01	2.12 <0.01	1.15 < 0.01	0.58 <0.01	3.08 <0.01	0.4	3.15	1.01		0.82	0.69	0.68												
PR-67 PR-68	1.05	0.65	0.81						0.05		<0.01	<0.01		0.03	0.10	0.07															
PR-70 V-2			1.59											*																	
V-3 V-4																															
V-5 V-7																															
V-8 V-11									0.01		<0.01																				
V-12 V-21																															
V-54 V-55											0.04			<0.01	<0.01	*															
V-56 V-64																															
V-66 V-70																															
V-71 V-72																															
V-73 V-74																															
V-77 V-78A									0.78 0.01	Dry 	 <0.01	 <0.01		 <0.01	<0.01	<0.01															
V-78B V-79									0.01		<0.01	<0.01		<0.01	<0.01	<0.01															
V-80 V-84																															
V-85 V-89																															
V-90 V-91		1.41		0.94	0.16	1.68	0.02	0.02	Dry 	Dry 	<0.01	Dry 		Dry 	Dry 	Dry 															
V-93 V-94									0.01																						
232 235																															
239 243																															
244 247																															
252 253																															
	- ***	all not n	٠.																												

⁻⁻ Well not monitored.

* Well inaccessible.

Table 4 CAP_LPH monitoring

 $a = Monitored \ on \ 2/5/99; \ b = mud \ in \ well \ at \ 3.80 \ feet; \ c = well \ was \ dry, \ but \ probe \ showed \ oil \ on \ it; \ d = black, \ oily \ substance \ on \ probe; \ e = interface \ probe \ not \ responding \ accurately \ to \ LPH; \ f = Amount \ verified \ on \ 8/10/01 \ for \ wells \ MW-8, \ E-5, \ PR-32, \ PR-53, \ PR-55, \ PR-67, \ and \ 244.$

Table 4: Historical LPH Monitoring (Product Thickness in Feet)

Corrective Action Plan Report, May 2009 Former Nestle USA, Inc. Facility 1310 14th Street, Oakland, CA

Well	3/28/1997	4/11/1997	4/17/1997	4/25/1997	5/2/1997	5/9/1997	5/16/1997	6/6/1997	10/25/1999	11/8/1999	12/1/1999	12/20/1999	1/17/2000	2/7/2000	2/28/2000	3/20/2000	4/10/2000	5/1/2000	5/22/2000	6/12/2000	10/25/2000	11/16/2000	12/11/2000	1/31/2001	2/28/2001	3/28/2001	4/30/2001	5/18/2001	6/29/2001	7/31/01 f	8/29/2001
MW-3 MW-7 MW-8									 <0.01 <0.01	 0.01 <0.01	 <0.01 <0.01	 <0.01 <0.01	 <0.01 <0.01	 <0.01 <0.01	<0.01 <0.01	 <0.01 <0.01	<0.01 <0.01	 <0.01 <0.01	 <0.01 <0.01	 <0.01 <0.01	 0.02 <0.01	 <0.01 <0.01	 <0.01 <0.01	<0.01 <0.01	 <0.01 <0.01	<0.01 <0.01	 <0.01 <0.01	 <0.01 <0.01	<0.01 <0.01	 0.02 <0.01	<0.01 <0.01
MW-22 MW-23									0.01	<0.01	0.04	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01
MW-24 E-0	 <0.01	 <0.01	 <0.01	 <0.01	<0.01	<0.01	 <0.01	 <0.01	0.13 <0.01	0.13 <0.01	0.13 <0.01	<0.01 <0.01	0.04 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	0.01 <0.01	<0.01 <0.01	<0.01	0.40 <0.01	0.41 <0.01	0.41 <0.01	0.41 <0.01	0.31 <0.01	0.34 <0.01	0.38 <0.01	0.45 <0.01	0.60 <0.01	0.62 <0.01	0.74 <0.01
E-3 E-5 E-6									<0.01 <0.01	0.11 <0.01	<0.01 <0.01	<0.01 <0.01	0.01 <0.01	0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	0.10 <0.01	0.10 0.01	0.02 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	0.04 <0.01	e <0.01	0.01 <0.01	0.01 <0.01
E-8 PR-12									<0.01	<0.01	<0.01	<0.01	<0.01	<0.01 <0.01 <0.01	<0.01	<0.01	<0.01	<0.01	<0.01 <0.01 <0.01	<0.01	<0.01 <0.01	<0.01 <0.01 <0.01	<0.01 <0.01 <0.01	<0.01 <0.01 <0.01	<0.01	<0.01 <0.01 <0.01	<0.01	<0.01 <0.01 <0.01	<0.01	<0.01	<0.01
PR-20 PR-21									<0.01 Dry	 Dry	<0.01 Dry	<0.01 Dry	<0.01 Dry	<0.01 <0.01	<0.01 Dry	<0.01 Dry	<0.01 Dry	<0.01 Dry	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 Dry	<0.01 Dry	<0.01 Dry						
PR-22 PR-23									<0.01	Ory <0.01	<0.01	<0.01	Sheen <0.01	<0.01	<0.01 <0.01	<0.01	<0.01	<0.01	<0.01	<0.01 <0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01 <0.01
PR-24 PR-25 PR-26									<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01
PR-27 PR-28									<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01
PR-29 PR-30									<0.01	<0.01	<0.01	<0.01	<0.01	<0.01		<0.01	<0.01	<0.01	<0.01	<0.01	<0.01 Dry	<0.01 Dry	<0.01 Dry	<0.01 Dry	<0.01 Dry	<0.01 Dry	<0.01 Dry	<0.01 Dry	<0.01 Dry	<0.01 Dry	<0.01 Dry
PR-32 PR-33									<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01
PR-34 PR-35 PR-36	0.65	1.31	0.8	1.06	0.7 	0.66	0.64	0.75	<0.01 <0.01 Dry	<0.01 <0.01 Dry	<0.01 <0.01 Dry	<0.01 <0.01 Dry	<0.01 <0.01 Dry	<0.01 <0.01 <0.01	<0.01 <0.01 Dry	<0.01 <0.01 Dry	<0.01 <0.01 Dry	<0.01 <0.01 <0.01	<0.01 <0.01 <0.01	<0.01 <0.01 <0.01	<0.01 <0.01 <0.01	<0.01 <0.01 <0.01	<0.01 <0.01 <0.01	Ory <0.01 Dry	<0.01 <0.01 <0.01						
PR-37 PR-38									<0.01	Dry 	<0.01	<0.01	<0.01	<0.01		<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01
PR-41 PR-42																															
PR-43 PR-44 PR-45																					Dry <0.01	Dry <0.01	Dry <0.01	Dry <0.01	<0.01 Dry	Dry <0.01	Dry <0.01	<0.01 <0.01	<0.01 <0.01	Dry <0.01	Dry <0.01
PR-46 PR-47									0.04	 Sheen	Sheen	<0.01	Sheen	<0.01	<0.01	<0.01	Sheen	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	 <0.01	<0.01	<0.01	<0.01	<0.01	 Dry	<0.01
PR-48 PR-49									0.02	0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	0.11	0.06	0.12	0.07	<0.01	<0.01	<0.01	0.03	<0.01	0.06	<0.01
PR-50 PR-51									<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01 Dry	<0.01 Dry	<0.01 Dry	<0.01 Dry	<0.01 Dry	<0.01 Dry	<0.01 Dry	<0.01 Dry	<0.01 Dry	Dry Dry	<0.01
PR-52 PR-53 PR-54									<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01 Sheen	<0.01	<0.01	<0.01 <0.01 <0.01	<0.01 0.01 <0.01	<0.01 0.03 <0.01	<0.01 0.06 <0.01	<0.01 0.06 <0.01	<0.01 <0.01 <0.01	<0.01 0.02 <0.01	<0.01 <0.01 <0.01	<0.01 <0.01 <0.01	<0.01 0.04 <0.01	<0.01 0.06 <0.01	<0.01 0.04 <0.01
PR-55 PR-56									0.02 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	0.03 Sheen	<0.01 Sheen	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	0.02 <0.01	<0.01	<0.01	0.10	0.11	0.07	<0.01	<0.01	0.03	0.02	<0.01 <0.01	0.06	0.06
PR-57 PR-58	2.45	2.14	1.8	2.06	 1.79	1.64	 1.49	1.44	0.06	0.01	 <0.01	 <0.01	0.06	<0.01	<0.01	<0.01	0.01	<0.01	<0.01	<0.01	<0.01 0.03	<0.01 0.03	<0.01 0.07	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 0.02	<0.01 0.06
PR-60 PR-61 PR-62	0.02	0.17	0.33	0.42	0.27	0.4	0.34	0.32	0.02	0.01	0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	0.15	<0.01	<0.01 0.26 <0.01	<0.01 0.25 <0.01	<0.01 0.24 <0.01	<0.01 0.22 <0.01	<0.01 0.19 <0.01	<0.01 0.18 <0.01	<0.01 0.22 <0.01	<0.01 0.22 <0.01	<0.01 0.24 <0.01	<0.01 0.27 <0.01	<0.01 <0.01 <0.01
PR-64 PR-65									0.08	0.02	0.01	0.04	<0.01	<0.01	<0.01	<0.01	0.02	<0.01	0.17	<0.01	0.51	0.52	0.49	0.48	0.50	0.64	0.30	0.45	0.63	0.64	0.60
PR-67 PR-68									<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	 <0.01	<0.01 <0.01	<0.01 <0.01	0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01
PR-70 V-2 V-3																															
V-3 V-4 V-5																			 			 	 								
V-7 V-8																					 <0.01	 <0.01	 <0.01	<0.01	 <0.01	<0.01	<0.01	 <0.01	 <0.01	 <0.01	 <0.01
V-11 V-12																							 	 	 	 	 	 			
V-21 V-54 V-55									Sheen	Sheen	 <0.01	 Sheen	Sheen		 <0.01	Sheen	 <0.01	 <0.01	Sheen	 <0.01	<0.01 <0.01	<0.01	Dry <0.01	Dry <0.01	Dry <0.01	Dry <0.01	Dry <0.01	Dry <0.01	<0.01 <0.01	<0.01 <0.01	Ory <0.01
V-56 V-64									0.02	0.03	Sheen 	Sheen 	Sheen 	Sheen 	Sheen 	Sheen 	<0.01	Sheen 	Sheen 	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01
V-66 V-70																					<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01
V-71 V-72 V-73																					<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01
V-74 V-77									 Dry												0.01	 <0.01	 <0.01	 <0.01							
V-78A V-78B										<0.01	<0.01		<0.01			<0.01					<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01
V-79 V-80 V-84																							<0.01								
V-84 V-85 V-89																															
V-90 V-91																															
V-93 V-94									Dry 	Dry 	Dry 												Dry								
232 235 239																															
243 244										<0.01 <0.01	<0.01 Sheen	<0.01 <0.01	<0.01 <0.01	<0.01 0.08	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	 <0.01
247 252										<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	
253		 ell not n								< 0.01	< 0.01	< 0.01	< 0.01	< 0.01	< 0.01	< 0.01	< 0.01	< 0.01	< 0.01	< 0.01	< 0.01	< 0.01	< 0.01	Dry	< 0.01	< 0.01	< 0.01	< 0.01	< 0.01	< 0.01	< 0.01

⁻⁻ Well not monitored.

Table 4 CAP_LPH monitoring Page 2 of 2

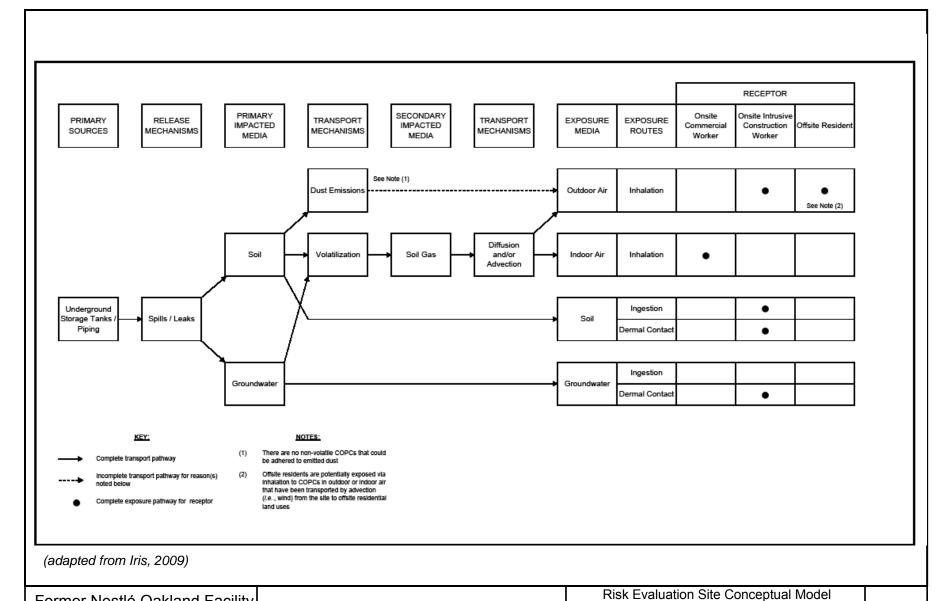
^{*} Well inaccessible. $a = Monitored \ on \ 2/5/99; \ b = mud \ in \ well \ at \ 3.80 \ feet; \ c = well \ was \ dry, \ but \ probe \ showed \ oil \ on \ it; \ d = black, \ oily \ substance \ on \ probe; \ e = interface \ probe \ not \ responding \ accurately \ to \ LPH; \ f = Amount \ verified \ on \ 8/10/01 \ for \ wells \ MW-8, \ E-5, \ PR-32, \ PR-53, \ PR-55, \ PR-67, \ and \ 244.$

TABLE 5 Comparative Matrix of Corrective Action Plan Approaches Draft Corrective Action Plan 1310 14th Street Oakland, CA

Corrective Action Plan Alternative	Alternative Details	Advantages	Drawbacks	Cost
1. Targeted Excavation of Impacted Soils Targeting areas for excavation identified in SCM above RWQCB-SF direct exposure limits for TPHg (4,200 mg/kg) and TPHd (4,200 mg/kg)	Excavation of soils to be performed from areas above RWQCB-SF direct exposure limits for TPHg and TPHd Extent of excavation to be defined by COC soil concentration data and interpolation as presented in November 2008 SCM Report (see Figure 7)	 Provides direct removal of COC mass from site Excavation of soils likely to provide reduction of direct exposure risks associated with residual COCs No on-going discharge of treated groundwater or vapor 	 All exposure risks associated with remaining hydrocarbons not likely to be eliminated Concrete/asphalt cap will need to be breeched in order to perform excavation Engineering protections and shoring likely necessary to protect against any compromises to integrity of existing building and surrounding sidewalks/streets 	• \$478,600 to \$574,400
2. Excavation of all Soils Above TPH Tier I ESLs Targeting areas for excavation identified in SCM above Commercial/Industrial Tier I ESLs for TPHg (180 mg/kg) and TPHd (180 mg/kg)	 Excavation of soils to be performed from areas above RWQCB-SF Tier I commercial/industrial screening levels for TPHg and TPHd Extent of excavation to be defined by COC soil concentration data and interpolation as presented in November 2008 SCM Report (see Figure 8) 	 Provides direct removal of COC mass from site Excavation of soils likely to provide reduction of direct exposure risks associated with residual COCs May result in elimination of direct exposure risks at site As result of possible elimination of direct exposure risks, Deed Restriction and RMP requirements could be reduced or eliminated Soil excavation activities not likely to remove all hydrocarbon mass dissolved in groundwater No on-going discharge of treated groundwater or vapor 	 All exposure risks associated with remaining hydrocarbons not certain to be eliminated Concrete/asphalt cap will need to be breeched in order to perform excavation Complete demolition of existing building necessary to access full extent of soils above Tier I levels Engineering protections and excavation shoring likely necessary to protect against any compromises to integrity of surrounding sidewalks/streets Soil excavation activities not likely to remove all hydrocarbon mass dissolved in groundwater 	• \$2,856,600 to \$3,427,900
3. Soil Vapor Extraction / Bioventing Extraction of vapors from areas identified in SCM above Tier I ESLs for TPHg (180 mg/kg) and TPHd (180 mg/kg)	 SVE or bioventing system to be installed with extraction points focused at areas of highest residual hydrocarbon impacts SVE or bioventing system designed to remove mass via vapor phase, and potentially promote dissolution of hydrocarbon mass from groundwater Bioventing may be implemented to promote additional circulation of air in subsurface and enhance biodegradation of hydrocarbons 	 Provides for removal of COC from subsurface (primarily vadose zone) Ability to target areas of highest hydrocarbon concentrations in soil Minimal disruption of existing concrete/asphalt cap and building 	 Shallow groundwater reduces effectiveness of SVE and bioventing Does not directly address submerged residual hydrocarbon All exposure risks associated with remaining hydrocarbons not likely to be eliminated Future construction at site will require consideration of any soil vapor extraction and treatment equipment present at site 	• \$593,700 to \$712,400

TABLE 5 Comparative Matrix of Corrective Action Plan Approaches Draft Corrective Action Plan 1310 14th Street Oakland, CA

4. In-situ Chemical Oxidation In-situ treatment through injection of oxidizer (hydrogen peroxide, Fenton's Reagent, potassium permanganate, or sodium persulfate, ozone) at areas identified in SCM above Tier I ESLs for TPHg and TPHd	 In-situ injection of oxidizer to be performed at injection points installed at areas of highest residual hydrocarbon impact (see Figure 9) Injection of oxidizer designed to promote degradation/destruction of hydrocarbons in groundwater and (saturated) soils 	 Provides for reduction of residual hydrocarbon mass to non-toxic (carbon dioxide, water) by-products Possible accelerated reduction of residual hydrocarbon mass beneath site Minimal disruption of existing concrete/asphalt cap and building 	 Subsurface distribution of chemical oxidant may be impeded by discontinuous silty-sand zones Previous high vacuum extraction efforts have revealed challenges in effecting removal of hydrocarbons within soil matrix; similar difficulty likely in generating oxidizer contact with hydrocarbons bound within submerged soils Future construction at site will require consideration of any injection equipment present at site On-site worker and building integrity may be an issue due to vigorous oxidizer reaction under building foundation 	• \$917,600 to \$1,101,100
5. Institutional Controls Revised Risk Assessment, in conjunction with Deed Restriction & Risk Management Plan for site outlining required institutional controls	 Risk Assessment (Appendix A) documents risks associated with residual hydrocarbons remaining at site Risk Assessment concludes that the risks associated with the concentrations of COC in soil and groundwater to indoor and outdoor air are within the USEPA's risk management range for carcinogens. Direct exposure risks (on-site construction workers) to be addressed through Risk Management Plan and existing Deed Restriction Institutional controls outlined in Risk Management Plan and Deed Restriction (site to remain commercial/industrial, surface cap to remain in place, etc.) remain applicable to property as part of site closure request 	 Consistent with RWQCB/ACEH requirements for risk-based case closure No disruption of concrete/asphalt cap, as required by existing Deed Restriction for site Approach provides protection for future occupants and construction workers to any residual impacts at site via following the approved Deed Restriction and RMP 	Requires ongoing notification and approval by oversight agencies of future site development activities, per requirements of RMP and Deed Restriction	• \$127,200 to \$182,600



Former Nestlé Oakland Facility Northwest Parcel 1310 14th Street Oakland, California - 94607



660 Baker Street, Suite 253 ● Costa Mesa, CA 92626 Tel: (714) 662-2759 • Fax: (714) 662-2758

(exposure pathway analysis) DRAFT CORRECTIVE ACTION PLAN May 2009

Table

6

APPENDICES

Appendix A: Screening Health Risk Evaluation Appendix B: Site Conceptual Model Figures Appendix C: Covenant and Environmental Restriction (Deed Restriction) Appendix D: Risk Management Plan Appendix A: Screening Health Risk Evaluation

IRIS ENVIRONMENTAL

Via Email and FTP

May 18, 2009

Jerry Wickham Senior Hazardous Materials Specialist Alameda County Environmental Health 1131 Harbor Bay Parkway, Suite 250 Alameda, California 94502-6577

Re: Screening Health Risk Evaluation Former Nestlé USA, Inc. Facility 1310 14th Street, Oakland, California

Dear Mr. Wickham:

On behalf of Nestlé USA, Inc., Iris Environmental is pleased to submit this Screening Health Risk Evaluation for the former Nestlé USA, Inc. facility located at 1310 14th Street in Oakland, California (the former Carnation Dairy).

We declare, under penalty of perjury, that the information and recommendations contained in the attached report are true and correct to the best of our knowledge.

Please don't hesitate to call us at (510) 834-4747 if you have any questions regarding this report.

Sincerely,

IRIS ENVIRONMENTAL

Robert Balas

Principal, Air Sciences

Gregory S. Noblet, P.E.

Senior Manager

Attachments: Screening Health Risk Evaluation, Former Nestlé USA, Inc. Facility, 1310 14th Street, Oakland, California

SCREENING HEALTH RISK EVALUATION

Former Nestlé USA, Inc. Facility

1310 14th Street, Oakland, California

May 18, 2009

Prepared for:

Nestlé USA, Inc. 800 North Brand Boulevard Glendale, California 91203

Prepared by:

IRIS ENVIRONMENTAL 1438 Webster Street, Suite 302 Oakland, California 94612 (510) 834-4747

EXECUTIVE SUMMARY

This report describes the methodology and results of a screening level human health risk evaluation for the northwestern portion of the commercial property located at 1310 14th Street in Oakland, California. The potential health impacts to onsite and offsite populations, associated with exposures to site-related chemicals, have been quantified. Potentially exposed populations which have been considered in this evaluation are onsite indoor commercial/industrial workers who are assumed to work full-time in the onsite commercial building for 25 years, onsite outdoor intrusive construction workers who are assumed to work onsite for 4 weeks, and offsite residents who are assumed to live near the site for 30 years. Estimated potential cancer risks for residential and commercial receptor populations are compared to the typical points of departure, with respect to risk management, of one in a million (1×10^{-6}) and 10 in a million (1×10^{-5}) , respectively. Estimated potential noncancer hazard indices for all receptor populations are compared to the threshold noncancer hazard index of 1.

The main conclusions of the screening health risk evaluation are as follows.

- The estimated potential cancer risk for onsite indoor commercial/industrial workers is 8.0×10^{-6} . The estimated potential noncancer hazard index for onsite indoor commercial/industrial workers is 0.051. Both the estimated cancer risk and the noncancer hazard index are below levels of concern.
- The estimated potential cancer risk for onsite outdoor intrusive construction workers is 9.8×10⁻⁵. The estimated potential noncancer hazard index for onsite outdoor intrusive construction workers is 21. Both the estimated cancer risk and the noncancer hazard index are above levels of concern. However, this cancer risk and noncancer hazard are attributable entirely to assumed dermal contact with COPCs in groundwater at the bottom of a construction trench, and do not account for personal protective equipment that intrusive construction workers would be required to use. Actual exposures after implementation of a site-specific Environmental Health and Safety Plan are highly likely to much lower than estimated here, and the actual cancer risk and hazard are likely to be below levels of concern.
- The estimated potential cancer risk for offsite residents is 4.1×10^{-7} . The estimated noncancer hazard index for offsite residents is 0.0040. Both the estimated cancer risk and the noncancer hazard index are below levels of concern.

The human health risk evaluation presented in this report is a screening-level evaluation that is based on a combination of conservative assumptions – regarding exposure point concentrations (including vapor intrusion modeling assumptions), exposure assumptions, toxicological data, and summation of health effects across chemicals and exposure routes – and therefore it is likely that actual health risks to exposed populations would be lower, or significantly lower, than those estimated in this analysis.

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1.0 INTRODUCTION

This report describes the methodology and results of a screening level human health risk evaluation for the northwestern portion of the commercial property located at 1310 14th Street in Oakland, California (the site). The site is the former location of the Carnation Dairy. Historical operations at the site are known to have resulted in release of petroleum hydrocarbons to soil and groundwater. Remedial activities have resulted in the removal of hydrocarbon product and a reduction in hydrocarbon concentrations in site soil, soil gas, and groundwater, however residual concentrations of hydrocarbons remain in these site media, as documented by a recent May 2008 site investigation. The purpose of this screening level human health risk evaluation is to conservatively estimate the potential health impacts to onsite and offsite populations, associated with exposures to site-related chemicals.

2.0 SITE DESCRIPTION

2.1 Site History

The 1310 14th Street site (see Figure 1) was formerly occupied by the Carnation Dairy. The primary activities conducted at the site were the manufacturing and distribution of ice cream and packaged milk. Delivery trucks were fueled and maintained onsite; the fuel storage and dispensing system was located in the northwest portion of the site, and consisted of underground storage tanks and associated underground piping. These activities were conducted at the site until 1988. The underground storage tanks and associated piping are now known to have leaked petroleum products into site soils, resulting in petroleum contamination of subsurface soils, a layer of petroleum product floating on the groundwater table, and dissolved petroleum hydrocarbons in site groundwater (ETIC, 2001a). These impacts have been partially addressed by various remedial activities, as described below.

2.2 Remedial Activities

Five underground storage tanks and associated underground piping were removed from the site between December 1988 and January 1989, including: two 12,000-gallon diesel tanks, two 10,000-gallon gasoline tanks, and one 1,000-gallon used oil tank. At that time, 1,200 cubic yards of petroleum hydrocarbon-impacted soil were excavated, treated onsite, and replaced into the excavation. Noted at the time of this removal action was the presence of petroleum hydrocarbons in the soil and floating on the groundwater table outside of the excavation area (ETIC, 2001a).

Various site investigations and remedial activities have been conducted at the site since the initial underground storage tank excavations. Remedial activities have included the following (COFS, 2000; ETIC, 2001a).

• Approximately 1.5 million gallons of groundwater were extracted from the subsurface following removal of the underground storage tanks.

- Product skimming was conducted between January and March 1989.
 Approximately 1,800 gallons of liquid phase hydrocarbons were removed from the subsurface.
- A soil vapor extraction system was operated from January 1994 to February 1995. An estimated 5,200 gallons of hydrocarbon equivalent were removed from the subsurface.
- A multi-phase extraction system was operated from August 1997 through June 2000. A total of 10,875 pounds of hydrocarbons were removed during this period.

Current site conditions have been characterized through soil, soil gas, and groundwater sampling conducted in May of 2008, as described below.

2.3 Site Investigations

Impacts to site soil, groundwater, and soil gas, associated with leaks of petroleum hydrocarbons from underground storage tanks and piping, have been documented in several site investigations performed since 1991. Soil gas investigations were performed in 1999 and in May of 2008. Soil investigations were performed at the time of underground storage tank excavation in 1991, in 1999, and most recently in May of 2008. Groundwater monitoring was performed on a regular basis from 1993 to 2004 and in May of 2008.

As noted in the Supplemental Soil, Soil Gas, and Groundwater Investigation Report (ECM, 2008a), components of the May 2008 site investigation consisted of

- soil sampling for total petroleum hydrocarbons (TPH) and volatile organic compounds (VOCs) at five locations (SB-16 through SB-20), at various depths, to provide current characterization of residual hydrocarbon impacts in the area downgradient from the former underground storage tanks;
- soil sampling for TPH and VOCs at seven locations (SB-21 through SB-27), at various depths, to provide delineation of hydrocarbon impacts in areas of the site which had not been thoroughly characterized;
- soil sampling for polychlorinated biphenyls (PCBs) at seven locations (PCB-1 through PCB-7), at various depths, to document the presence or absence of PCBs at the site;
- soil gas sampling for TPH and VOCs at 12 locations (SB-16 through SB-27), including seven locations in the area downgradient from the former underground storage tanks (SB-20 through SB-27), at a depth of 5 feet, to provide a complete set of soil gas data for use in evaluating vapor intrusion; and
- grab groundwater sampling for TPH and VOCs at 11 locations (SB-16 through SB-27 exclusive of SB-23).

The results of all previous site investigations, as summarized in Table 1a (1999 soil gas data), Table 1b (2008 soil gas data), Table 2 (soil TPH and VOC data), Table 3 (soil PCB

data), and Table 4 (groundwater data) of the *Revised Site Conceptual Model Report* (ECM, 2008b), are provided in Appendix A.

In addition to the investigations performed to characterize impacts to site soil, soil gas, and groundwater, noted above, a site investigation was conducted in January 2009 to characterize site-specific soil properties, specifically to support transport modeling (see Sections 4.5.1 and 4.5.2). The number and location of soil properties samples were determined by the project geologist, in order to ensure that the soil at the site is well characterized. The soil properties data collected during this investigation are documented in Figure 2.

3.0 CHEMICALS OF POTENTIAL CONCERN

Chemicals of potential concern (COPCs) for the screening-level human health risk evaluation are defined for each of three impacted site media: soil gas, soil, and groundwater. In general, any chemical that has been detected above the laboratory reporting limit in any sample from any site investigation is designated as a COPC in that medium. The COPCs in soil gas, soil, and groundwater are identified below.

3.1 Soil Gas

As discussed above, site soil gas has been characterized by investigations conducted in 1999 and 2008. The soil gas data from the 2008 investigation are considered to be generally more appropriate for evaluation of future vapor transport into the onsite commercial building, for one primary reason: the data collected in 2008 are likely more representative of current and future conditions than the data collected in 1999, particularly since remedial activities were conducted after 1999 to remove product from the subsurface. As shown in the historical soil gas data summaries presented in Appendix A, VOCs were generally detected at higher concentrations in 1999 (ECM Table 1a) than in 2008 (ECM Table 1b). Also of note, the 2008 soil gas data were collected at a depth of 5 feet below ground surface (bgs), whereas the 1999 soil gas data were collected at 3 feet bgs.

Current DTSC soil gas sampling guidance (Cal/EPA, 2005b) states that, "soil gas samples should not be collected depths shallower than 5 feet in order to minimize barometric pumping effects." That the 1999 soil gas samples were collected at depths shallower than 5 feet provides a secondary rationale for favoring the 2008 data. However, the 1999 dataset includes detections of 1,3-butadiene, a potentially significant carcinogen which was not sampled for in 2008. Therefore, to be comprehensive, both the 1999 and the 2008 soil gas data are used to characterize COPCs in soil gas for this screening level health risk evaluation. The COPCs in soil gas are defined as those 38 VOCs which were detected above their respective laboratory reporting limit in at least one soil gas sample from either the 1999 or 2008 soil gas investigation:

- acetone;
- benzene;
- 1,3-butadiene;

- 2-butanone (methyl ethyl ketone);
- · carbon disulfide;
- · chlorobenzene;
- · chloroform;
- chloromethane (methyl chloride);
- · cyclohexane;
- 1,2-dichlorobenzene;
- 1,3-dichlorobenzene;
- 1,4-dichlorobenzene;
- dichlorodifluoromethane (Freon 12);
- 1,1-dichloroethane;
- 1,2-dichloroethane;
- 1,1-dichloroethene (1,1-DCE);
- *cis*-1,2-dichloroethene (*cis*-1,2-DCE);
- 1,4-dioxane;
- ethanol;
- ethylbenzene;
- 4-ethyltoluene;
- heptane;
- hexane;
- methyl tertiary butyl ether (MTBE);
- methylene chloride;
- 4-methyl-2-pentanone (methyl isobutyl ketone);
- 2-propanol;
- styrene;
- tetrachloroethene (PCE);
- tetrahydrofuran;
- toluene;
- 1,1,1-trichloroethane;
- trichloroethene (TCE);
- trichlorofluoromethane (Freon 11);
- 1,1,2-trichloro-1,2,2-trifluoroethane (Freon 113);
- 1,2,4-trimethylbenzene;

- 1,3,5-trimethylbenzene; and
- xylenes.

The derivation of representative concentrations of these COPCs in soil gas is described in Section 4.5 below.

3.2 Soil

As discussed above, site soils have been characterized by several investigations conducted since 1991. Results of these soil investigations are included in Appendix A. The COPCs in soil for this screening risk evaluation are defined as those ten VOCs which have been detected above their respective laboratory reporting limit in at least one site soil sample from any investigation:

- benzene;
- chlorobenzene;
- 1,2-dichlorobenzene;
- 1,3-dichlorobenzene;
- 1,4-dichlorobenzene;
- 1,2-dichloroethane;
- ethylbenzene;
- methyl tertiary butyl ether (MTBE);
- toluene; and,
- xylenes.

The derivation of representative concentrations of these COPCs in soil is described in Section 4.5 below.

3.3 Groundwater

Groundwater monitoring has been performed at the site since 1993. Results of these groundwater investigations are included in Appendix A. The COPCs in groundwater for this screening risk evaluation are defined as those 20 VOCs which have been detected above their respective laboratory reporting limit in at least one site groundwater sample from any investigation or sampling event:

- benzene;
- bromodichloromethane;
- chlorobenzene;
- chloroethane (ethyl chloride);
- chloroform;
- chloromethane (methyl chloride);

- 1,2-dichlorobenzene;
- dichlorodifluoromethane (Freon 12);
- 1,1-dichloroethane;
- 1,2-dichloroethane;
- 1,1-dichloroethene (1,1-DCE);
- *cis*-1,2-dichloroethene (*cis*-1,2-DCE);
- ethylbenzene;
- methyl tertiary butyl ether (MTBE);
- methylene chloride;
- 1,1,2,2-tetrachloroethane;
- toluene;
- 1,1,1-trichloroethane;
- trichloroethene (TCE); and,
- xylenes.

The derivation of representative exposure point concentrations of these COPCs in groundwater is described in Section 4.5 below.

4.0 EXPOSURE ASSESSMENT

The purpose of exposure assessment is to estimate the type and magnitude of exposures to the chemicals of potential concern that are present at, or migrating from, the site. The results of the exposure assessment are combined with chemical-specific toxicity information (see Section 5) to characterize potential risks (see Section 6).

4.1 Physical Setting

The 1310 14th Street property is located in West Oakland, approximately 2 miles from San Francisco Bay. The deed-restricted portion of the 1310 14th Street property (the site) is approximately 1.3 acres in size. The site contains one existing unoccupied L-shaped building of approximately 29,000 square feet which was formerly used for maintenance of delivery trucks. The site is relatively flat, and is covered with concrete and asphalt pavement in addition to the large building. There are no onsite surface water bodies. Groundwater has been historically detected at the site at depths ranging from 5 to 12 feet. The site is surrounded by a mix of light industrial, commercial, and residential land uses.

4.2 Sources of Contamination

As noted above, the underground storage tanks and associated piping which comprised the fuel storage and distribution system are known to have leaked gasoline, diesel fuel, and waste oil into the subsurface, resulting in hydrocarbon impacts to soil and groundwater and a layer of hydrocarbon product floating on the groundwater table. The various remedial activities performed at the site have apparently been successful in extracting the free-phase hydrocarbons, and in reducing hydrocarbon concentrations in soil and groundwater.

4.3 Potential Transport Mechanisms

As noted above, the primary impacted media at the site are soil and groundwater. Direct exposures to impacted site soil and groundwater are evaluated for intrusive onsite construction workers, as described below in Section 4.4. Described in this section are the transport pathways by which COPCs in site soil and groundwater may migrate to other media, where exposures may also occur.

Volatilization of chemicals from soil and groundwater has resulted in impacts to soil gas in the vadose (unsaturated) soil zone above the groundwater table. Given the presence of volatile COPCs in site soil, groundwater, and soil gas, there are several transport mechanisms whereby COPCs could potentially migrate to another medium. These potential transport mechanisms are evaluated below. Transport pathways which are determined to be complete and are included in the screening health risk evaluation are shown in the conceptual site model depicted in Figure 3.

4.3.1 Particulate Emission into Outdoor Air

This transport pathway comprises the release of particulate matter (*i.e.*, dust) from the ground surface to ambient air, either by wind erosion or by mechanical disturbance. The dust-inhalation pathway is evaluated when there are non-volatile chemicals present in site soils; the non-volatile chemicals may be adhered to the dust which is emitted from soil into ambient air. Based on the site characterization, however, it appears that COPCs in site media are primarily volatile, and thus would generally be emitted into ambient air in the vapor phase, not the particulate phase. Further, the site is essentially capped with pavement and the existing commercial building, and the site deed restriction (COFS, 2000) and Risk Management Plan (ETIC, 2001b) require that future site development maintains a surface cap of site soil, exclusive of minor landscaped areas, by buildings or pavement. Because site soils are capped, the potential for dust emissions is very low. Therefore, based on the volatile nature of identified site-related COPCs and the presence of the cap, the inhalation of respirable particulate matter (*i.e.*, dust) is not included in this screening level health risk evaluation.

4.3.2 Vapor Intrusion from Soil Gas into Indoor Air

The conceptual site model (see Figure 3) assumes that volatile COPCs which are present in the subsurface may migrate upwards via diffusion through the vadose (unsaturated) soil zone, and be transported by advection through cracks, conduits, or seams in the building foundation and into the indoor air space of the onsite building (a transport phenomenon known as "vapor intrusion"), where building occupants may be exposed to the COPCs via the inhalation route.

4.3.3 Volatilization from Soil Gas to Outdoor Air

The conceptual site model (see Figure 3) assumes that volatile COPCs which are present in the subsurface may migrate upwards via diffusion through the vadose soil zone and be emitted from the ground surface into ambient air, where outdoor workers may be exposed to the COPCs via the inhalation route.

4.3.4 Advective Transport to Offsite Locations

The conceptual site model (see Figure 3) assumes that, once volatile COPCs have been emitted from the ground surface into onsite outdoor air (see Section 4.3.2 above), they may be transported downwind to offsite locations. As described below, the offsite populations which are evaluated in this screening risk evaluation are child and adult residents. Thus, the conceptual site model assumes the advective transport of volatile COPCs from the site to downwind offsite residential land uses.

4.3.5 Groundwater and Soil Gas Transport to Offsite Locations

The SCM Report (ECM, 2008b) concludes that the plume of dissolved hydrocarbons in site groundwater is not migrating offsite. The SCM Report notes that chemical concentrations measured in groundwater wells located downgradient of the primary area of impact have stabilized at low or non-detect levels. The SCM Report further notes that a review of subsurface utilities indicates that subsurface utilities do not act as conduits for migration of chemicals in the subsurface. Based on these findings, it appears that significant offsite transport of COPCs is not occurring, via either groundwater transport or diffusive transport of soil gas. Accordingly, these subsurface transport pathways are considered to be incomplete based on the information provided in the SCM Report.

4.3.6 Surface Water Transport

The potential for COPCs to migrate offsite via storm water runoff has been considered. The 1310 14th Street site is essentially capped with pavement and the existing commercial building. The Covenant and Environmental Restriction on Property ("deed restriction") between Nestlé USA, Inc. and City of Oakland Fire Services (COFS, 2000) and the site Risk Management Plan (ETIC, 2001b) require that future site development maintains a surface cap of site soil, exclusive of minor landscaped areas, by buildings or pavement. Because site soils are capped, the potential for storm water runoff to become impacted by site-related COPCs is likely to be low. Accordingly, the offsite transport of COPCs via storm water runoff is considered to be incomplete.

4.3.7 Onsite Extraction of Groundwater

While not a considered a "transport" pathway in the formal sense, the extraction and onsite use of impacted site groundwater as a potable water supply could lead to exposures for onsite indoor commercial/industrial workers to COPCs present in the groundwater via ingestion and dermal contact. However, because use of site groundwater (via an extraction well) is prohibited by the deed restriction (COFS, 2000), this transport

pathway is considered to be incomplete, and no ingestion or dermal contact exposures to groundwater (*i.e.*, tap water) are possible.

4.3.8 Summary of Complete Transport Pathways and Exposure Media

In summary, the primary impacted media at the site are soil, groundwater, and soil gas. The site deed restriction (COFS, 2000) effectively breaks certain transport pathways, including transport from site soil to other media via surface water runoff or dust emissions, and extraction of groundwater for use as potable water supply. Other potential transport pathways are considered to be incomplete for the reasons described above. The transport pathways which are considered to be potentially complete are:

- vapor intrusion from soil gas to indoor air of the onsite commercial building;
- volatilization from soil gas to onsite outdoor; and
- advective transport from onsite outdoor air to the outdoor or indoor air of offsite residential land uses.

Thus, the exposure media which are included in the quantitative screening risk evaluation are:

- onsite soil;
- onsite groundwater;
- onsite soil gas;
- indoor air of the existing onsite commercial building;
- onsite outdoor air; and
- outdoor or indoor air at offsite residential land uses.

Potentially exposed human populations and routes of exposure (ingestion, etc.) are discussed below in Section 4.4.

4.4 Potentially Exposed Populations and Exposure Routes

This section describes the potentially exposed populations and associated exposure routes that are included in the screening health risk evaluation of the 1310 14th Street site. These populations and routes are summarized in the conceptual site model depicted in Figure 3.

4.4.1 Onsite Commercial/Industrial Worker

The intended future land use for the 1310 14th Street site is commercial/industrial, consistent with the site deed restriction (COFS; 2000). Thus, the primary population of potential concern is future commercial/industrial workers, who are assumed to work full-time in the onsite building. Because the site is effectively capped by the existing building and by pavement as required by the deed restriction, no direct contact between the commercial/industrial worker and site soil or groundwater is possible, and therefore these exposure routes are considered to be incomplete. Because use of site groundwater (via an

extraction well) is prohibited by the deed restriction, no ingestion or dermal contact exposures to groundwater are possible, and therefore these exposure routes are also considered to be incomplete. Therefore, as indicated in the conceptual site model depicted in Figure 3, the only complete exposure pathway/route for the onsite indoor commercial/industrial is inhalation of volatile COPCs that are present in indoor air as a result of transport from soil gas to indoor air (*i.e.*, vapor intrusion).

4.4.2 Onsite Intrusive Construction Worker

Also included in this screening level health risk evaluation are onsite, outdoor intrusive construction workers who are assumed to engage in excavation of site soils (e.g., trenching for utility installation) over a relatively short period of time. These intrusive construction workers are assumed to contact impacted soils via dermal contact and incidental ingestion, and are assumed to contact impacted groundwater via dermal contact with groundwater that could be present at the floor of a utility trench. Intrusive construction workers are also assumed to be exposed via inhalation to volatile COPCs present in onsite outdoor air as a result of volatilization from soil gas to outdoor air. For the purpose of quantifying exposures, it is assumed that intrusive construction workers are onsite and in a utility trench for 8 hours per day, for a total of 20 working days (i.e., four weeks). It should be noted that intrusive construction work would necessarily puncture the site cap and likely lead to dust emissions; because the identified site-related COPCs are volatile in nature, however, exposure of intrusive construction workers to fugitive dust is not evaluated (USEPA, 2002).

4.4.3 Offsite Resident

The current land uses in the vicinity of the 1310 14th Street site include commercial, industrial, and residential uses. In this screening analysis, no distinction is made with respect to the actual locations of these various land uses, relative to the site. As explained above (see Section 4.3), the only potentially complete offsite transport pathway is advective transport (*i.e.*, by wind) of volatile COPCs, and the only complete exposure route for offsite populations is inhalation of volatile COPCs in air. Due to the greater exposure frequency and exposure duration associated with residential exposures compared with commercial and industrial exposures, the offsite receptor that is included in the analysis is the offsite resident. As discussed below, it is conservatively assumed that the concentrations of volatile COPCs that an offsite residential building occupant may be exposed to are equal to the concentrations of those volatile COPCs in onsite outdoor air.

4.5 Exposure Point Concentrations

An exposure point is defined as a location of potential contact between an organism (e.g., human receptor) and a physical or chemical agent. The exposure point concentration is defined as the average concentration of the physical or chemical agent in the exposure medium over the period of exposure. The exposure point concentration does not represent the maximum concentration that could be contacted at any one time, but rather represents a reasonable estimate of the concentration likely to be contacted over time

(USEPA, 1989). In this screening level health risk evaluation, however, exposure point concentrations are conservatively based on historical maximum detected concentrations in site media. Exposure point concentrations in each of the relevant exposure media – indoor air of the onsite commercial building, onsite outdoor air, indoor or outdoor air inhaled by an offsite resident, onsite soil, and onsite groundwater – are discussed below.

4.5.1 Indoor Air of Onsite Commercial Building

The conceptual site model (see Figure 3) assumes that volatile COPCs present in the subsurface may migrate upwards through the vadose soil zone and into the indoor air space of the overlying onsite commercial building, where workers may be exposed to the volatile COPCS via inhalation; this transport phenomenon is referred to as vapor intrusion. The transport of COPCs from soil gas to the indoor air of the onsite building is modeled using the United States Environmental Protection Agency (USEPA)-recommended Johnson & Ettinger Model for soil gas (SG-SCREEN Version 2.0), as modified by the California Environmental Protection Agency (Cal/EPA) Department of Toxic Substances Control (DTSC) Human and Ecological Risk Division (HERD) (Johnson and Ettinger, 1991; USEPA, 2004a; Cal/EPA, 2005a), and as modified by Iris Environmental to allow for the input of multiple chemicals and site-specific building parameters. As recommended by DTSC (Cal/EPA, 2005c), soil gas data, rather than soil or groundwater data, are used to evaluate the vapor intrusion pathway, because soil gas data represent a direct measurement of the volatile chemicals that may potentially migrate into indoor air.

The Johnson and Ettinger model is a conservative, screening-level model that incorporates both convective and diffusive mechanisms for estimating the transport of vapor-phase chemicals from soil gas to an indoor air space located directly above the source of contamination. The Johnson and Ettinger model is described in detail in the *User's Guide for Evaluating Subsurface Vapor Intrusion into Buildings* (USEPA, 2004a). Inputs to the Johnson and Ettinger model include depth and concentration of contamination, physicochemical properties of the chemicals being transported, lithology and building parameters, and soil properties. Model input data are documented in Tables 1 through 3 and are discussed below.

Source Characterization

The soil gas data collected during both the 1999 and 2008 site soil gas investigations are used to characterize the contaminant source in soil gas. The maximum detected concentration of each COPC in soil gas, from the combined 1999 and 2008 datasets, is conservatively assumed to be representative of the contaminant source strength. This assumption is consistent with current DTSC vapor intrusion guidance (Cal/EPA, 2005b), which recommends the use of maximum detected soil gas concentrations in vapor intrusion screening risk evaluations.

As noted above, the 1999 soil gas samples were collected at 3 feet bgs, while the 2008 soil gas samples were collected at 5 feet bgs. For the purpose of modeling vapor intrusion transport, all COPCs are conservatively assumed to be present at the shallower

3 feet bgs sampling depth. The concentration in soil gas and depth to contamination of each COPC, used as Johnson and Ettinger model inputs, are documented in Table 1.

Physicochemical Properties

Physicochemical properties that are used by the Johnson and Ettinger model to simulate the transport of volatile chemicals through the subsurface include: diffusivity in water, diffusivity in air, Henry's Law constant, molecular weight, and other properties. These data are used by the model to calculate the effective diffusivity of the volatile chemical through the vadose zone, which varies slightly from chemical-to-chemical. These input data are documented in Table 2.

Lithology and Building Parameters

The screening-level Johnson and Ettinger model for soil gas used in this analysis (SG-SCREEN Version 2.0) is a one-soil-layer model; assumed soil properties are documented below. The existing on-site commercial building is assumed to be of slab-on-grade construction. Building-parameter inputs to the screening-level Johnson and Ettinger model are the area of the building footprint, the depth below grade of the bottom of the foundation slab, the building air exchange rate, and the flow rate of soil gas into the building. The values assigned to these building parameters include site-specific values of building area and height, and Cal/EPA default values of depth below grade of the bottom of the foundation slab, building air exchange rate, and flow rate of soil gas into the building. The modeled site lithology and building geometry are depicted in Figures 4 and 5. These input data are documented in Table 3.

Soil Properties

Soil-property inputs to the screening-level Johnson and Ettinger model are total porosity, water-filled porosity, bulk density, and temperature. Soil properties were measured in eight soil samples collected from four soil borings (Appendix A, Figure 3) on January 22, 2009; as noted above, that investigation was performed specifically to obtain soil properties data to support transport modeling, and the number and location of soil samples were determined by the project geologist to adequately characterize site soil conditions. Results of that investigation are presented in Figure 2. The site average bulk density, total porosity, and water-filled porosity from these eight soil samples are assigned to the soil property inputs in the model. The assumed soil temperature of the soil layer is the USEPA-recommended default value for the San Francisco Bay area (USEPA, 2004a). These input data are documented in Table 3.

Modeling Results

The modeled concentrations of volatile COPCs in the indoor air space of the onsite commercial building, predicted by the Johnson and Ettinger model, are presented in Table 4. These modeling results represent the exposure point concentrations of volatile COPCs in indoor air that onsite commercial/industrial workers are assumed to be exposed to via the inhalation route.

4.5.2 Onsite Outdoor Air

The conceptual site model (see Figure 3) assumes that volatile COPCs present in soil gas may migrate upwards through the vadose soil zone and into ambient (outdoor) air, where onsite intrusive construction workers may be exposed to the volatile COPCS via inhalation. This transport process is similar to vapor intrusion, except that the volatile COPCs are emitted from the ground surface into outdoor air rather than into the indoor air space of an overlying building. The transport of volatile COPCs from soil gas to outdoor air is modeled using the USEPA methodology presented in *Supplemental Guidance for Developing Soil Screening Levels for Superfund Sites* (USEPA, 2002). In this methodology, the transport of volatile COPCs from soil gas to outdoor air is comprised of two components: the diffusive transport of volatile chemicals from soil gas to the ground surface; and the dispersion of volatile chemicals from the ground surface into the ambient air (USEPA, 2002; ASTM, 1995):

$$C_{OA} = \left[\frac{D_{eff} \times C_{SG}}{d} \times CF1 \times CF2\right] \times \left[\frac{1}{(Q/C)_{vol}}\right]$$
 (Eq. 1)

where:

 C_{OA} = concentration of volatile COPC in outdoor air (mg/m³);

 D_{eff} = effective diffusivity of COPC through vadose soil zone (cm²/s) (see

Equation 2, below);

 C_{SG} = concentration of COPC in soil gas (mg/m³);

d = depth of COPC contamination in soil gas (m);

CF1 = units conversion factor (m^2/cm^2) ;

CF2 = units conversion factor (g/kg);

 $(Q/C)_{vol}$ = dispersion factor $(g/m^2/s \text{ per kg/m}^3)$.

The first bracketed term of Equation 1 represents the steady-state flux of the volatile COPC from the ground surface into ambient air. This flux is a function of the concentration of the COPC in soil gas (C_{SG}), the effective diffusivity of the COPC through the vadose soil zone (D_{eff}), and the length (*i.e.*, height of soil column) over which the COPC must diffuse to reach the ground surface (d). The concentration of the COPC in soil gas and the depth to contamination are as-assumed in the evaluation of vapor intrusion transport, discussed above in Section 2.4.1 and documented in Table 1. The effective diffusivity of the COPC through the vadose soil zone (D_{eff}) is calculated as follows (USEPA, 2002; ASTM, 1995):

$$D_{eff} = \frac{(D_a \times q_a^{10/3})}{\eta^2} + \frac{(D_w \times q_w^{10/3})}{\eta^2 \times H'}$$
 (Eq. 2)

where:

 D_a = diffusivity of COPC in air (cm²/s);

q_a = air-filled soil porosity (cm³/cm³);
 η = total soil porosity (cm³/cm³);

 $D_{\rm w}$ = diffusivity of COPC in water (cm²/s);

 q_w = water-filled soil porosity (cm³/cm³); and

H' = Henry's law constant (cm^3/cm^3) .

Values of the physicochemical properties appearing in Equation 2 – diffusivity in air (D_a) , diffusivity in water (D_w) , and Henry's law constant – are taken from the DTSC/HERD Johnson and Ettinger model (Cal/EPA, 2005a); these physicochemical data are documented in Table 2. Values of the soil properties appearing in Equation 2 – air filled soil porosity, water-filled soil porosity, and total soil porosity – are the average site-specific values measured during the January 2009 site investigation, and are the same values used as inputs to the Johnson and Ettinger modeling analysis of vapor intrusion transport to indoor air (see Section 4.5.1). As noted above, that investigation was performed specifically to obtain soil properties data to support transport modeling, and the number and location of soil samples were determined by the project geologist to adequately characterize site soil conditions. These soil properties data are documented in Table 3.

The second bracketed term of Equation 1 represents the dispersion of the volatile COPC, from the point of release at the ground surface into the larger body of onsite ambient air. The dispersion of volatile chemicals from the ground surface to ambient air is estimated as recommended in the *Supplemental Guidance for Developing Soil Screening Levels for Superfund Sites* (USEPA, 2002). The USEPA soil-screening guidance document defines an annual-average dispersion factor, (Q/C)_{vol}, which represents the reciprocal of the ratio of the geometric mean air concentration at the center of a square source area to the emission flux from the square source area; *i.e.*, the (Q/C)_{vol} dispersion factor is specifically designed to estimate the annual-average concentration of a volatile COPC in ambient air at the center of an area source from which it is emitted. The dispersion factor is a function of the source size and of empirical coefficients which are based on air dispersion modeling for specific climate zones (USEPA, 2002):

$$(Q/C)_{vol} = A \times exp \left[\frac{(\ln A_{site} - B)^2}{C} \right]$$
 (Eq. 3)

where:

A = empirical dispersion coefficient (unitless);

 A_{site} = area of source (acres);

B = empirical dispersion coefficient (unitless); and

C = empirical dispersion coefficient (unitless).

The site area from which volatilization of COPCs is modeled to occur (A_{site}) is conservatively assumed to be the size of the deed-restricted portion of the property.

Values assigned to the empirical dispersion coefficients (A, B, and C) are USEPA-recommended conservative default values specific for the San Francisco Bay region (USEPA, 2002). The values assigned to the variables in Equation 3 and the resulting dispersion factor are documented in Table 5.

In summary, the effective diffusivity (D_{eff}) of each volatile COPC is calculated by Equation 2, based on soil properties (porosity, etc.) and the physicochemical properties of the COPC (diffusivity in air, etc.). A site-specific dispersion factor [(Q/C)_{vol}] is calculated by Equation 3, based on the site area (A_{site}) and based on default regional dispersion coefficients (A, B, and C). These two derived parameters, and the depth and concentration of the COPC in soil gas(d and C_{SG}), are plugged into Equation 1 to calculate the concentration of the COPC in ambient air at the center of the site, resulting from volatilization from soil gas to ambient air. These exposure point concentrations of COPCs in onsite outdoor air, used to quantify potential health impacts associated with inhalation by intrusive construction workers, are documented in Table 6.

4.5.3 Indoor and Outdoor Air of Offsite Residences

The conceptual site model (see Figure 3) assumes that volatile COPCs present in the subsurface may migrate upwards through the vadose soil zone and into ambient (outdoor) air at the site (as described in the previous section). The volatile COPCs are then assumed be transported downwind to offsite residential land uses, where residents may be exposed to the VOCs via the inhalation route. In this screening level health risk evaluation, it is conservatively assumed that the concentrations of volatile COPCs that offsite residents are exposed to in outdoor or indoor air are equal to those in onsite ambient air. In other words, the exposure point concentrations of COPCs that offsite residents are potentially exposed to are assumed equal to the exposure point concentrations of COPCs in onsite outdoor air, as documented in Table 6. This is a highly conservative assumption, as the concentrations of COPCs in air would be diluted during downwind transport from the site to offsite residential land uses.

4.5.4 Onsite Soil

The conceptual site model (see Figure 3) assumes that onsite outdoor intrusive construction workers are exposed to COPCs in site soils via ingestion and dermal contact. (Direct soil exposures for the indoor commercial/industrial worker are assumed to be incomplete, due to the cap provided by the building and pavement.) For this screening level risk evaluation, it is assumed that intrusive construction workers are exposed to the maximum detected concentrations of COPCs in site soils, from any site soil investigation, at any sampling location, at any depth below ground surface. The exposure point concentrations of COPCs in site soils, used to quantify potential health impacts associated with ingestion and direct contact exposures, are documented in Table 7.

4.5.5 Onsite Groundwater

The conceptual site model (see Figure 3) assumes that onsite outdoor intrusive construction workers are exposed to COPCs in site groundwater via dermal contact with

the groundwater, such as at the bottom of a utility trench. For this screening level risk evaluation, it is assumed that the intrusive construction workers are exposed to the maximum detected concentrations of COPCs in site groundwater, from any site groundwater investigation or sampling event, at any sampling location. The exposure point concentrations of COPCs in site groundwater, used to quantify potential health impacts associated with direct contact exposures, are documented in Table 8.

4.6 Quantification of Potential Chemical Intakes

Exposure is defined as contact between an organism (e.g., human body) and a chemical or physical agent. In accordance with USEPA risk assessment methodology (USEPA, 1989), exposures are averaged over time and to the body weight of the receptor, and are referred to as intakes. In this formulation, chemical intake is a function of: the concentration of the chemical in the exposure medium (e.g., soil), the contact rate between the receptor and the exposure medium, the frequency and duration of the exposure, the body weight of the receptor, and the time period over which the exposure is normalized. Intakes are averaged over long periods of time are referred to as chronic daily intakes.

The generic equation for estimating the chronic daily intake of a chemical in air, soil, or water is as follows (USEPA, 1989):

$$CDI = \frac{C \times CR \times EF \times ED}{BW} \times \frac{1}{AT}$$
 (Eq. 4)

where:

CDI = chronic daily intake (mg/kg/d);

exposure point concentration of chemical in soil (mg/kg), groundwater (mg/L), or air (mg/m³);

CR = contact rate with soil (kg/d), groundwater (L/d), or air (m^3/d) ;

EF = exposure frequency (d/yr);

ED = exposure duration (yr);

BW = body weight (kg); and

AT = averaging time (d).

Chronic daily intakes are specific to the exposure pathway, the receptor, and the type of health effect being evaluated (*i.e.*, cancer or noncancer). The pathway-specific equations used to quantify chronic daily intakes are documented in Table 9. The values assigned to the intake parameters are documented in Table 10, with the exception of dermal permeability coefficients which are documented in Table 2, and exposure point concentrations which are documented in Table 4 (onsite indoor air), Table 6 (onsite outdoor air and offsite indoor or outdoor air), Table 7 (onsite soil), and Table 8 (onsite groundwater). As documented in Table 10, the values assigned to the intake parameters are standard, conservative default values recommended by USEPA and/or Cal/EPA.

In evaluating the potential health impacts to residential populations, both child and adult receptors are considered. For noncancer health hazard, the child receptor experiences the largest intakes (due to lower body weight), and thus the noncancer hazard assessment for the offsite resident is based on the child receptor. Per DTSC guidance (Cal/EPA, 1994), cancer risks for residential populations are calculated using an age-adjusted approach, to account for the higher per-body weight exposures that occur during the childhood years; accordingly, the cancer risk assessment for the offsite resident is based on an "age adjusted" receptor who is assumed to be a child for the first 6 years of exposure and an adult for the remaining 24 years of an assumed 30-year residential exposure duration.

Chronic daily intakes calculated for all exposure pathways and exposed populations are presented in Table 11. In conjunction with toxicity data (see Section 5.0), these chronic daily intakes are used to characterize potential cancer risks and noncancer hazards, as discussed in Section 6.0.

5.0 TOXICITY ASSESSMENT

Toxicity assessment characterizes the relationship between the magnitude of exposure to a chemical and the potential for adverse effects. More specifically, toxicity assessment identifies or derives toxicity values that can be used to estimate the likelihood of adverse effects occurring in humans at different exposure levels. Consistent with regulatory risk assessment policy, adverse health effects resulting from chemical exposures are evaluated in two categories: carcinogenic effects and noncarcinogenic effects. The hierarchy of sources for the toxicity criteria used for this health risk evaluation generally corresponds to DTSC guidelines (Cal/EPA, 1994); all carcinogenic and noncarcinogenic toxicity values used to evaluate the potential health effects associated with exposure to COPCs in groundwater are presented and documented in Table 13.

5.1 Toxicity Assessment for Carcinogenic Effects

Current health risk assessment practice for carcinogens is based on the assumption that, for most substances, there is no threshold dose below which carcinogenic effects do not occur. This no-threshold assumption for carcinogenic effects is based on an assumption that the carcinogenic processes are the same at high and low doses. This approach has generally been adopted by regulatory agencies as a conservative practice to protect public health. The no-threshold assumption is used in this risk assessment for evaluating carcinogenic effects. Although the magnitude of the risk declines with decreasing exposure, the risk is believed to be zero only at zero exposure.

The response potency of a potential carcinogen is quantified by the cancer slope factor (SF). The slope factor represents the excess lifetime cancer risk due to a continuous, constant exposure to a specified level (*i.e.*, unit dose) of a carcinogen. Slope factors are generally reported as excess incremental cancer risk per milligram of chemical per kilogram body weight per day (per mg/kg/day). Separate slope factors are generally reported for inhalation and oral exposures; these slope factors are referred to as the inhalation slope factor (SF_i) and the oral slope factor (SF_o), respectively. Both dermal and oral exposures are generally evaluated using the oral slope factor.

The inhalation and oral slope factors (SF_i and SF_o values) used in this risk evaluation are taken from the following hierarchy of sources:

- 1) Cal/EPA Office of Environmental Health Hazard Assessment (OEHHA) on-line *Toxicity Criteria Database* (Cal/EPA, 2008); and
- 2) USEPA on-line Integrated Risk Information System (IRIS) (USEPA, 2008).

The slope factors, and their sources, used to evaluate the potential carcinogenic toxicity of COPCs are documented in Table 12.

5.2 Toxicity Assessment for Noncarcinogenic Effects

The toxicity assessment for noncarcinogenic effects is based on the assumption that there exists a threshold level of exposure below which no adverse health effects occur. This threshold level varies from individual to individual. In developing a toxicity parameter for noncarcinogenic effects, the approach is to identify a threshold value that is protective of sensitive individuals in the population. For most chemicals, this level can only be estimated, and the developed toxicity value incorporates uncertainty factors indicating the degree of extrapolation used to derive the estimated value. The developed toxicity level is generally considered to have uncertainty spanning an order of magnitude or more, and should not be viewed as a strict scientific demarcation between what level is toxic and nontoxic (USEPA, 1989).

The toxicity parameter that is typically used to evaluate noncarcinogenic effects is the reference dose (RfD). The reference dose represents an intake level, expressed in milligrams of chemical per kilogram of body weight per day (mg/kg/day), that would not be expected to cause adverse noncancer health effects in potentially exposed populations, including sensitive subpopulations (USEPA, 1989). Thus, the reference dose is often referred to as the "acceptable dose." The chronic reference dose specifically represents the daily exposure level that is unlikely to produce adverse noncancer health effects in potentially exposed populations, including sensitive subpopulations, over a lifetime. Analogous to slope factors, chronic reference doses are generally reported for inhalation and oral exposures; these chronic reference doses are referred to as the inhalation chronic reference dose (RfD_i) and the oral chronic reference dose (RfD_o), respectively. Both dermal and oral exposures are generally evaluated using the oral chronic reference dose.

The inhalation and oral chronic reference doses (RfD_i and RfD_o values) used in this risk assessment are taken directly from the following hierarchy of sources:

- 1) Cal/EPA Office of Environmental Health Hazard Assessment (OEHHA) on-line *Toxicity Criteria Database* (Cal/EPA, 2008);
- 2) USEPA on-line Integrated Risk Information System (IRIS) (USEPA, 2008); and
- 3) USEPA Regional Screening Levels (RSLs) for Chemical Contaminants at Superfund Sites (USEPA, 2008).

The chronic reference doses, and their sources, used to evaluate the potential noncarcinogenic toxicity of COPCs are documented in Table 13.

6.0 RISK CHARACTERIZATION

6.1 Cancer Risk

For carcinogens, risk is estimated as the incremental probability of an individual developing cancer over a lifetime as a result of daily exposure to the potential carcinogen. The cancer slope factor (SF) (see Section 5.1) converts the estimated daily intake averaged over a lifetime of exposure directly to incremental risk of an individual developing cancer. Because relatively low intakes are likely to result from exposure to chemicals at contaminated sites (compared to those experienced by laboratory test animals), it is assumed the dose-response relationship is linear. Under this assumption, the slope factor is constant and risk is directly related to intake (USEPA, 1989):

$$RISK = SF \times CDI$$
 (Eq. 5)

where:

RISK = cancer risk, *i.e.*, the probability of an individual developing cancer over a lifetime as a result of daily exposure to a particular carcinogen (unitless);

SF = cancer slope factor, *i.e.*, the lifetime incremental cancer risk per unit dose of the carcinogen (per mg/kg/d); and

CDI = chronic daily intake of the carcinogen (mg/kg/d).

Cancer risks are estimated by Equation 5 for each relevant exposure pathway and receptor and each carcinogenic chemical. As a matter of policy, USEPA (1989b) considers the cancer risk for contact with multiple carcinogens to be additive, regardless of the carcinogens' mechanisms of toxicity or sites (organs of the body) of action. Therefore, within each exposure pathway (*e.g.*, inhalation of indoor air), the chemical-specific cancer risks are summed to produce an estimate of the cumulative (multichemical) risk associated with the exposure pathway. In addition, cancer risk for a given receptor across multiple exposure routes is also considered to be additive (USEPA, 1989b). Therefore, the pathway-specific risks are summed to produce an estimate of the cumulative (multi-chemical and multi-pathway) risk to each specific receptor. This cumulative risk estimate represents the incremental probability of an individual within that receptor population (*e.g.*, onsite indoor commercial/industrial workers) developing cancer over a lifetime as a result of exposure to site-related carcinogenic COPCs. Estimated cancer risks are presented in Table 14.

The estimated incremental cancer risks presented in Table 14 are compared to the "acceptable" cancer risk level, as defined and endorsed by relevant state and federal agencies. The National Contingency Plan (NCP) is cited by USEPA (1989) as the basis for defining acceptable incremental risk levels. According to the NCP, lifetime incremental cancer risk levels posed by a site should be within the risk range of one in a million (1×10^{-6}) to 100 in a million (1×10^{-4}). Thus, USEPA and Cal/EPA agencies typically consider the 1×10^{-6} risk level to be an insignificant risk, and consider a calculated excess cancer risk between 1×10^{-6} and 1×10^{-4} to be within the acceptable risk

range. For commercial exposure scenarios, a typical point of departure is a risk level of 1×10^{-5} ; *i.e.*, if risks are at or below 1×10^{-5} , the agency of record will generally accept no further action.

6.2 Noncancer Hazard

The reference-dose approach (see Section 5.2) is based on the theory that there exists a threshold level of exposure below which it is unlikely for even sensitive subpopulations to experience adverse noncancer health effects. If the actual exposure level (*i.e.*, the chronic daily intake) exceeds this threshold value (*i.e.*, the chronic reference dose), there may be concern for potential noncancer health effects. Generally, the larger ratio of chronic daily intake to chronic reference dose, the greater the level of concern. This ratio is not to be interpreted as a probability of developing noncancer health effects, however, and the level of concern does not increase linearly with this ratio USEPA (1989).

The ratio of the chronic daily intake of a chemical to the chronic reference dose for that chemical is referred to as the noncancer hazard quotient:

$$HQ = \frac{CDI}{RfD}$$
 (Eq. 6)

where:

HQ = noncancer hazard quotient, *i.e.*, the potential (not probability) for an individual to develop adverse noncancer health effects over a lifetime as a result of daily exposure to a particular chemical (unitless);

CDI = chronic daily intake of the chemical (mg/kg/d); and

RfD = chronic reference dose, *i.e.*, the threshold level of exposure that would not be expected to cause adverse noncancer health effects in potentially exposed populations, including sensitive subpopulations (unitless).

Noncancer hazard quotients are estimated by Equation 6 for each relevant exposure pathway and receptor and each chemical. Within each exposure pathway (*e.g.*, inhalation of indoor air), the chemical-specific noncancer hazard quotients are summed to produce an estimate of the cumulative (multi-chemical) "hazard index" associated with the exposure pathway. It should be noted here that the summation of hazard quotients across chemicals, independent of the target organ which is affected by each chemical, is conservative, as chemicals that impact different target organs (*e.g.*, liver, kidney) are not truly additive in their potential to cause the adverse impact. The pathway-specific hazard indices are then summed to produce an estimate of the cumulative (multi-chemical and multi-pathway) noncancer hazard index for the specific receptor. This cumulative hazard index estimate represents the incremental potential (not probability) of an individual within that receptor population (*e.g.*, onsite indoor commercial/industrial workers) developing adverse noncancer health effects as a result of exposure to site-related COPCs. Estimated noncancer hazards are presented in Table 15.

The estimated incremental noncancer hazards presented in Table 15 are compared to the threshold level of 1. Chemical exposures that yield hazard indices of less than 1 are not expected to result in adverse noncancer health effects (USEPA, 1989).

7.0 DISCUSSION OF RESULTS

7.1 Cancer Risk

A summary of estimated cancer risks, for all potentially exposed populations and associated routes of exposure, is presented in Table 14. These results are discussed in this section by receptor population.

As indicated in Table 14, the total (summed across all COPCs) estimated cancer risk for onsite indoor commercial/industrial workers, associated with inhalation of volatile COPCs that are present in indoor air as a result of vapor intrusion (the only complete exposure pathway for this receptor), is 8.0×10^{-6} . This estimated risk is below the 1×10^{-5} risk level that is the typical point of departure for commercial exposure scenarios. Nearly all of this risk may be attributed to inhalation of benzene (85 percent of total risk) and 1,3-butadiene (10 percent).

As indicated in Table 14, the total (summed across all COPCs and four exposure pathways) estimated cancer risk for onsite outdoor intrusive construction workers is 9.8×10^{-5} , which is approximately 10 times greater than the 1×10^{-5} risk level that is the typical point of departure for commercial/industrial exposure scenarios. This risk level is attributable to assumed dermal contact with COPCs in groundwater at the bottom of a construction trench – the estimated risks associated with inhalation of volatile COPCs in outdoor air, ingestion of COPCs in soil, and dermal contact with COPCs in soil are all well below levels of concern. The groundwater-dermal contact risk may be attributed to dermal contact with benzene (77 percent of total risk) and ethylbenzene (23 percent) in groundwater.

It should be noted that dermal exposures to soil and groundwater for the intrusive construction worker conservatively assume that the hands and forearms are exposed, *i.e.*, that the worker is wearing a short-sleeved shirt and no gloves. The site Risk Management Plan (ETIC, 2001b), however, requires that constructor contractors develop a site-specific Environmental Health and Safety Plan that specifies appropriate safety equipment to minimize contact between the construction worker and site soil and groundwater. Therefore, actual exposures after implementation of a site-specific Environmental Health and Safety Plan are highly likely to much lower than estimated here, and actual cancer risks are likely to be below levels of concern.

As indicated in Table 14, the total (summed across all COPCs) estimated cancer risk for offsite residents, associated with inhalation of volatile COPCs in indoor or outdoor air that have migrated downwind from the site (the only complete exposure pathway for this receptor), is 4.1×10^{-7} . This estimated risk is below the 1×10^{-6} risk level that is typically considered to be the point of departure for residential exposure scenarios.

7.2 Noncancer Hazard

A summary of estimated noncancer hazard indices, for all potentially exposed populations and associated routes of exposure, are presented in Table 15. These results are discussed in this section by receptor population.

As indicated in Table 15, the total (summed across all COPCs) estimated noncancer hazard index for onsite indoor commercial/industrial workers, associated with inhalation of volatile COPCs that are present in indoor air as a result of vapor intrusion (the only complete exposure pathway for this receptor), is 0.051. This estimated hazard is well below the threshold hazard index of 1, and thus may be considered negligible.

As indicated in Table 15, the total (summed across all COPCs and four exposure pathways) estimated noncancer hazard index for onsite outdoor intrusive construction workers is 21, which is 21 times greater than the threshold hazard index of 1. This hazard level is attributable to assumed dermal contact with COPCs in groundwater at the bottom of a construction trench – the estimated hazards associated with inhalation of volatile COPCs in outdoor air, ingestion of COPCs in soil, and dermal contact with COPCs in soil are all well below levels of concern. The groundwater-dermal contact hazard may be attributed primarily to dermal contact with benzene (63 percent of total hazard) and xylenes (23 percent) in groundwater.

As noted above, dermal exposures to soil and groundwater for the intrusive construction worker conservatively assume that the hands and forearms are exposed, *i.e.*, that the worker is wearing a short-sleeved shirt and no gloves. The site Risk Management Plan (ETIC, 2001b), however, requires that constructor contractors develop a site-specific Environmental Health and Safety Plan that specifies appropriate safety equipment to minimize contact between the construction worker and site soil and groundwater. Therefore, actual exposures after implementation of a site-specific Environmental Health and Safety Plan are highly likely to much lower than estimated here, and actual noncancer hazards are likely to be below levels of concern.

As indicated in Table 15, the total (summed across all COPCs) estimated noncancer hazard index for offsite residents, associated with inhalation of volatile COPCs in indoor or outdoor air that have migrated downwind from the site (the only complete exposure pathway for this receptor), is 0.0040. This estimated hazard is well below the threshold hazard index of 1, and may be considered negligible.

8.0 DEVELOPMENT OF RISK-BASED SOIL GAS CLEANUP GOALS

Risk-based soil gas cleanup goals are calculated in anticipation of potential site cleanup. Following the risk evaluation methodologies of Sections 5.0 and 6.0, risk-based indoor air concentrations protective of commercial workers are calculated for both carcinogenic and noncarcinogenic endpoints. These indoor air concentrations are presented in Table 16.

Transport of soil gas from 5.0 feet below ground surface into indoor air of the onsite commercial building is modeled with the Johnson and Ettinger Model as described in

Section 4.5.1. Site-specific soil and building parameters are used. The output parameter of the Johnson and Ettinger model is the attenuation factor (α). By definition, the attenuation factor is the ratio of the COPC concentration in indoor air (resulting from vapor intrusion) to the COPC concentration in soil gas beneath the building:

$$\alpha = \frac{C_{IA}}{C_{SG}}$$
 (Eq. 7)

where:

 α = attenuation factor (unitless);

 C_{IA} = concentration of COPC in indoor air ($\mu g/m^3$); and

 C_{SG} = concentration of COPC in soil gas ($\mu g/m^3$).

An attenuation factor is calculated with the Johnson and Ettinger model for each COPC in soil gas (38 total), at a depth of 5.0 feet bgs. These attenuation factors are presented in Table 16.

Chemical-specific risk-based soil gas cleanup goals at 5.0 feet bgs are calculated from the attenuation factors developed using the Johnson and Ettinger model and from the target indoor air concentrations estimated using standard USEPA and Cal/EPA inhalation risk-assessment methodology. For each chemical, the risk-based soil gas cleanup goal is calculated from:

$$RBCG = \frac{CA}{\alpha}$$
 (Eq. 8)

where:

RBCG = chemical- and depth-specific risk-based soil gas cleanup goal ($\mu g/m^3$);

CA = risk-based target concentration of COPC in indoor air $(\mu g/m^3)$; and

 α = chemical- and depth-specific attenuation factor (unitless).

Risk-based cleanup goals are presented in Table 16. By definition, each soil gas cleanup goal represents the concentration of that COPC in soil gas at 5.0 feet bgs that would be considered safe and acceptable for commercial use of the existing onsite building. It is important to note that exceedances of these soil gas screening levels should not be interpreted to mean that conditions in the building are unsafe. Rather, the soil gas screening levels are set sufficiently low, and incorporate many levels of conservatism, in order to allow for prudent and proactive additional analyses and actions. The soil gas screening levels developed in this assessment can be used to assess the effectiveness of the cleanup activities.

9.0 UNCERTAINTIES

Many of the assumptions used in the human health screening evaluation – regarding the representativeness of the sampling data, human exposures, fate and transport modeling, and chemical toxicity – are conservative, follow agency guidance, and reflect a 90th or

95th percentile value rather than a typical or average value. The use of several conservative exposure and toxicity assumptions can introduce considerable uncertainty into the human health screening evaluation. By using conservative exposure or toxicity estimates, the evaluation can develop a significant conservative bias that may result in the calculation of significantly higher cancer risks or noncancer hazards than are actually posed by the chemicals present in soil, soil gas, and groundwater. The key uncertainties in the human health screening evaluation are discussed in Appendix B. The uncertainty analysis focuses on the site-specific assumptions contributing most to uncertainty in the risk and hazard calculations, and does not assess the validity of default assumptions used in the health screening evaluation. Assumptions/data evaluated in the uncertainty analysis are: representative concentrations in soil gas, soil properties, and building height.

As noted above and discussed in Appendix B, uncertainties exist in the human health risk evaluation regarding representative concentrations of COPCs in soil gas, soil properties, and building air exchange rate. The baseline evaluation, the results of which are summarized above in Section 7, is based on a combination of assumptions regarding these three parameters, based primarily on DTSC vapor intrusion guidance (Cal/EPA, 2005b), and represents a relatively conservative estimate of potential risk and hazard. Discussed in Appendix B and documented in Appendices C and D are two sensitivity analyses which bound the range of potential risks and hazards associated with the uncertainties in these three input parameters. The first sensitivity analysis combines the most conservative, but likely least representative, options of the three parameters, to produce a high-end estimate of potential risk and hazard. The second sensitivity analysis combines the least conservative, but likely most representative, options of the three parameters, to produce a low-end estimate of potential risks and hazards. The conceptual differences between the three evaluations are summarized in the following table.

Summary of Health Risk Evaluations

Evaluation	Where Documented	Soil Gas Concentrations	Soil Properties	Building Ventilation Rate
High-end estimate	Appendix C	Maxima	Default	Default
Baseline estimate	Main report	Maxima	Site-specific	Default
Low-end estimate	Appendix D	Averages	Site-specific	Site-specific

Based on the results of these analyses, the following conclusions may be drawn regarding the sensitivity of the results of the health risk evaluation to the uncertainties regarding representative concentrations of COPCs in soil gas, soil properties, and building air exchange rate.

• Estimated potential risk and hazard for onsite indoor commercial/industrial workers range over approximately 2-1/2 orders of magnitude. The high-end estimates are approximately 8 times greater than the baseline estimates presented above in Section 7. The low-end estimates are less than the baseline estimates by a factor of approximately 47.

- Estimated potential risk and hazard for onsite outdoor intrusive construction workers are not sensitive to these uncertainties, as the estimated potential health effects for this receptor are driven by dermal contact with groundwater.
- Estimated potential risk and hazard for offsite residents range over approximately 2 orders of magnitude. The high-end estimates are approximately 17 times greater than the baseline estimates presented above in Section 7. The low-end estimates are less than the baseline estimates by a factor of approximately 11.
- The high-end estimates of potential risk are the within the 1×10^{-6} to 1×10^{-4} risk management range for all three receptors (onsite indoor commercial/industrial workers, onsite outdoor intrusive construction workers, and offsite residents). The high-end estimates of potential hazard are below the threshold hazard level of 1 for onsite indoor commercial/industrial workers and offsite residents. The high-end estimate of potential hazard exceeds the threshold hazard level of 1 for onsite outdoor intrusive construction workers; as noted above, this hazard is driven by dermal contact with groundwater.
- The low-end estimates of potential risk are below the 1×10⁻⁶ to 1×10⁻⁴ risk management range for onsite indoor commercial/industrial workers and offsite residents, and within the risk management range for onsite outdoor intrusive construction workers. The low-end estimates of potential hazard are below the threshold hazard level of 1 for onsite indoor commercial/industrial workers and offsite residents. The low-end estimate of potential hazard exceeds the threshold hazard level of 1 for onsite outdoor intrusive construction workers; as noted above, this hazard is driven by dermal contact with groundwater.

10.0 CONCLUSIONS

This report describes the methodology and results of a screening level human health risk evaluation for the northwestern portion of the commercial property located at 1310 14th Street in Oakland, California. The potential health impacts to onsite and offsite populations, associated with exposures to site-related chemicals, have been quantified. Of note, the site deed restriction (COFS, 2000) effectively breaks certain transport pathways, including transport from site soil to other media via surface water runoff or dust emissions, and extraction of groundwater for use as potable water supply; and physically prevents direct contact between onsite commercial workers and impacted site soils.

Potentially exposed populations which have been considered in this evaluation are onsite indoor commercial/industrial workers who are assumed to work full-time in the onsite commercial building for 25 years, onsite outdoor intrusive construction workers who are assumed to work onsite for 4 weeks, and offsite residents who are assumed to live near the site for 30 years. Estimated potential cancer risks for residential and commercial receptor populations are compared to the typical points of departure, with respect to risk management, of 1×10^{-6} and 1×10^{-5} , respectively. Estimated potential noncancer hazard indices for all receptor populations are compared to the threshold noncancer hazard index of 1.

The main conclusions of the screening health risk evaluation are as follows.

- The estimated potential cancer risk for onsite indoor commercial workers is 8.0×10^{-6} . The estimated potential noncancer hazard index for onsite indoor commercial workers is 0.051. Both the estimated cancer risk and the noncancer hazard index are below levels of concern.
- The estimated potential cancer risk for onsite outdoor intrusive construction workers is 9.8×10⁻⁵. The estimated potential noncancer hazard index for onsite outdoor intrusive construction workers is 21. Both the estimated cancer risk and the noncancer hazard index are above levels of concern. However, this cancer risk and noncancer hazard are attributable entirely to assumed dermal contact with COPCs in groundwater at the bottom of a construction trench, and do not account for personal protective equipment that intrusive construction workers would be required to use. Actual exposures after implementation of a site-specific Environmental Health and Safety Plan are highly likely to much lower than estimated here, and the actual cancer risk and hazard are likely to be below levels of concern.
- The estimated potential cancer risk for offsite residents is 4.1×10^{-7} . The estimated noncancer hazard index for offsite residents is 0.0040. Both the estimated cancer risk and the noncancer hazard index are below levels of concern.

The human health risk evaluation presented in this report is a screening-level evaluation that is based on a combination of conservative assumptions – regarding vapor intrusion modeling assumptions, exposure assumptions, toxicological data, summation of health effects across chemicals and exposure routes – and therefore it is likely that actual health risks to exposed populations could be lower, or significantly lower, than those estimated in this analysis.

11.0 REFERENCES

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Table 1. Representative Concentrations of Chemicals of Potential Concern in Soil Gas

Chemical of Potential Concern	Concentration
	$(\mu g/m^3)$
Acetone	620
Benzene	40,000
1,3-Butadiene	310
2-Butanone (methyl ethyl ketone)	420
Carbon disulfide	440
Chlorobenzene	160
Chloroform	170
Chloromethane (methyl chloride)	75
Cyclohexane	480
1,2-Dichlorobenzene	2,900
1,3-Dichlorobenzene	210
1,4-Dichlorobenzene	460
Dichlorodifluoromethane (Freon 12)	10,000
1,1-Dichloroethane	140
1,2-Dichloroethane	140
1,1-Dichloroethene (1,1-DCE)	140
cis-1,2-Dichloroethene (cis-1,2-DCE)	140
1,4-Dioxane	500
Ethanol	2,600
Ethylbenzene	7,700
4-Ethyltoluene	3,700
Heptane	550
Hexane	63,000
Methyl tertiary butyl ether (MTBE)	500
Methylene chloride	1,200
4-Methyl-2-pentanone (methyl isobutyl ketone)	550
2-Propanol	350
Styrene	150
Tetrachloroethene (PCE)	1,100
Tetrahydrofuran	420
Toluene	32,000
1,1,1-Trichloroethane	190

Table 1. Representative Concentrations of Chemicals of Potential Concern in Soil Gas

Chemical of Potential Concern	Concentration $(\mu g/m^3)$
Trichloroethene (TCE)	190
Trichlorofluoromethane (Freon 11)	200
1,1,2-Trichloro-1,2,2-trifluoroethane (Freon 113)	270
1,2,4-Trimethylbenzene	2,900
1,3,5-Trimethylbenzene	3,600
Xylenes	19,000

⁽¹⁾ Soil gas samples were collected at depths of 3 and 5 feet below ground surface (bgs) in 1999 and 2008, respectively.

⁽²⁾ Concentration units are micrograms per cubic meter (µg/m³).

⁽³⁾ Representative concentration is the maximum of all samples collected in 1999 and 2008.

Table 2. Physicochemical Properties of Chemicals of Potential Concern

Chemical of Potential Concern	Diffusivit (D _i	-	Diffusivity (D _w)		Henry's Law at Refere Tempera (H)	ence ature	Henry's Law Referen Tempera (T _R)	ice ture	Henry's Law Unitless		Enthalp Vaporizatio Normal Boil (DH _v	on at The ling Point	Normal Boi (T _B	-	Critical Ter	•	Organic (Partition Co (K _{oo}	efficient,	Pure Com Water Sol (S)	ubility,	Vap Press (VI	ure,	Dermal Per Const (K ₁	ant,		orption Factor ABS)
	Value (cm ² /s)	Source	Value (cm ² /s)	Source	Value (atm-m ³ /mol)	Source	Value (°C)	Source	Value (unitless)	Source	Value (cal/mol)	Source	Value (°K)	Source	Value (°K)	Source	Value (cm ³ /g)	Source	Value (mg/L)	Source	Value (mmHg)	Source	Value (cm/hr)	Source	Value (unitless)	Source
Acetone	1.24E-01	1	1.14E-05	1	3.87E-05	1	2.50E+01	1	1.59E-03	1	6.96E+03	1	3.29E+02	1	5.08E+02	1	5.75E-01	1	1.00E+06	1	5.06E+02	1	NA	NA	0.1	7
Benzene	8.80E-02	1	9.80E-06	1	5.54E-03	1	2.50E+01	1	2.27E-01	1	7.34E+03	1	3.53E+02	1	5.62E+02	1	5.89E+01	1	1.79E+03	1	9.64E+01	1	1.49E-02	3	0.1	7
Bromodichloromethane	2.98E-02	1	1.06E-05	1	1.60E-03	1	2.50E+01	1	6.54E-02	1	7.80E+03	1	3.63E+02	1	5.86E+02	1	5.50E+01	1	6.74E+03	1	4.99E+01	1	4.62E-03	3	0.1	7
1,3-Butadiene	2.49E-01	1	1.08E-05	1	7.34E-02	1	2.50E+01	1	3.01E+00	1	5.37E+03	1	2.69E+02	1	4.25E+02	1	1.91E+01	1	7.35E+02	1	7.58E+02	1	1.64E-02	3	0.1	7
2-Butanone (methyl ethyl ketone)	8.08E-02	1	9.80E-06	1	5.58E-05	1	2.50E+01	1	2.29E-03	1	7.48E+03	1	3.53E+02	1	5.37E+02	1	2.30E+00	1	2.23E+05	1	1.31E+02	1	9.63E-04	3	0.1	7
Carbon disulfide	1.04E-01	1	1.00E-05	1	3.02E-02	1	2.50E+01	1	1.24E+00	1	6.39E+03	1	3.19E+02	1	5.52E+02	1	4.57E+01	1	1.19E+03	1	3.58E+02	1	1.72E-02	3	0.1	7
Chlorobenzene	7.30E-02	1	8.70E-06	1	3.69E-03	1	2.50E+01	1	1.51E-01	1	8.41E+03	1	4.05E+02	1	6.32E+02	1	2.19E+02	1	4.72E+02	1	1.18E+01	1	2.82E-02	3	0.1	7
Chloroethane (ethyl chloride)	2.71E-01	1	1.15E-05	1	8.80E-03	1	2.50E+01	1	3.61E-01	1	5.88E+03	1	2.85E+02	1	4.60E+02	1	4.40E+00	1	5.68E+03	1	5.89E+02	1	6.07E-03	3	0.1	7
Chloroform	1.04E-01	1	1.00E-05	1	3.66E-03	1	2.50E+01	1	1.50E-01	1	6.99E+03	1	3.34E+02	1	5.36E+02	1	3.98E+01	1	7.92E+03	1	1.85E+02	1	6.83E-03	3	0.1	7
Chloromethane (methyl chloride)	1.26E-01	1	6.50E-06	1	8.80E-03	1	2.50E+01	1	3.61E-01	1	5.11E+03	1	2.49E+02	1	4.16E+02	1	2.12E+00	1	5.33E+03	1	7.05E+02	1	3.28E-03	3	0.1	7
Cyclohexane	8.00E-02	2	9.10E-06	2	1.50E-01	4	2.50E+01	4	6.10E+00	2	NA	NA	3.54E+02	4	NA	NA	1.66E+02	2	5.50E+01	2	9.69E+01	4	NA	NA	0.1	7
1,2-Dichlorobenzene	6.90E-02	1	7.90E-06	1	1.90E-03	1	2.50E+01	1	7.77E-02	1	9.70E+03	1	4.54E+02	1	7.05E+02	1	6.17E+02	1	1.56E+02	1	1.53E+00	1	4.13E-02	3	0.1	7
1,3-Dichlorobenzene	6.92E-02	1	7.86E-06	1	3.09E-03	1	2.50E+01	1	1.27E-01	1	9.23E+03	1	4.46E+02	1	6.84E+02	1	1.98E+03	1	1.34E+02	1	2.14E+00	1	5.79E-02	3	0.1	7
1,4-Dichlorobenzene	6.90E-02	1	7.90E-06	1	2.39E-03	1	2.50E+01	1	9.82E-02	1	9.27E+03	1	4.47E+02	1	6.85E+02	1	6.17E+02	1	7.90E+01	1	9.78E-01	1	4.20E-02	3	0.1	7
Dichlorodifluoromethane (Freon 12)	6.65E-02	1	9.92E-06	1	3.42E-01	1	2.50E+01	1	1.40E+01	1	9.42E+03	1	2.43E+02	1	3.85E+02	1	4.57E+02	1	2.80E+02	1	6.02E+02	1	8.95E-03	3	0.1	7
1,1-Dichloroethane	7.42E-02	1	1.05E-05	1	5.61E-03	1	2.50E+01	1	2.30E-01	1	6.90E+03	1	3.31E+02	1	5.23E+02	1	3.16E+01	1	5.06E+03	1	2.18E+02	1	6.74E-03	3	0.1	7
1.2-Dichloroethane	1.04E-01	1	9.90E-06	1	9.77E-04	1	2.50E+01 2.50E+01	1	4.00E-02	1	7.64E+03	1	3.57E+02	1	5.61E+02	1	1.74E+01	1	8.52E+03	1	6.39E+01	1	4.20E-03	3	0.1	7
1,1-Dichloroethene (1,1-DCE)	9.00E-02	1	1.04E-05	1	2.60E-02	1	2.50E+01 2.50E+01	1	1.07E+00	1	6.25E+03	1	3.05E+02	1	5.76E+02	1	5.89E+01	1	2.25E+03	1	4.59E+02	1	1.17E-02	3	0.1	7
		1		1		1		1		1		1		1		1		1		1		1				7
cis-1,2-Dichloroethene (cis-1,2-DCE)	7.36E-02	1	1.13E-05	2	4.07E-03	1	2.50E+01	1	1.67E-01	1	7.19E+03	I NIA	3.34E+02	1	5.44E+02	I NA	3.55E+01	2	3.50E+03	2	1.12E+02	1	NA	NA 2	0.1	7
1,4-Dioxane	8.70E-02	2	1.10E-05	2	4.80E-06	4	2.50E+01	4	2.00E-04	2	NA	NA	3.75E+02	4	NA	NA	1.00E+00	4	1.00E+06	4	3.81E+01	4	3.32E-04	3	0.1	7
Ethanol	1.23E-01	4	1.30E-05	6	5.00E-06	4	2.50E+01	4	2.05E-04	4	NA	NA	3.51E+02	4	NA	NA	1.85E-01	4	1.00E+06	4	5.93E+01	4	5.38E-04	3	0.1	7
Ethylbenzene	7.50E-02	1	7.80E-06	1	7.86E-03	1	2.50E+01	1	3.22E-01	1	8.50E+03	1	4.09E+02	1	6.17E+02	1	3.63E+02	1	1.69E+02	1	9.51E+00	1	4.93E-02	3	0.1	7
4-Ethyltoluene	6.81E-02	4	7.84E-06	6	5.01E-03	4	2.50E+01	4	2.05E-01	4	NA	NA	4.33E+02	4	NA	NA	1.77E+03	4	9.49E+01	4	3.01E+00	4	NA	NA	0.1	7
Heptane	7.04E-02	4	7.59E-06	6	2.00E+00	4	2.50E+01	4	8.20E+01	4	NA	NA	3.72E+02	4	NA	NA	1.94E+04	4	3.40E+00	4	5.16E+01	4	NA	NA	0.1	7
Hexane	2.00E-01	1	7.77E-06	1	1.66E+00	1	2.50E+01	1	6.82E+01	1	6.90E+03	1	3.42E+02	1	5.08E+02	1	4.34E+01	1	1.24E+01	1	1.82E+02	1	NA	NA	0.1	7
Methyl tertiary butyl ether (MTBE)	1.02E-01	1	1.05E-05	1	6.23E-04	1	2.50E+01	1	2.56E-02	1	6.68E+03	1	3.28E+02	1	4.97E+02	1	7.26E+00	1	5.10E+04	1	2.74E+02	1	NA	NA	0.1	7
Methylene chloride	1.01E-01	1	1.17E-05	1	2.18E-03	1	2.50E+01	1	8.96E-02	1	6.71E+03	1	3.13E+02	1	5.10E+02	1	1.17E+01	1	1.30E+04	1	2.55E+02	1	3.54E-03	3	0.1	7
4-Methyl-2-pentanone (methyl isobutyl ketone)	7.50E-02	1	7.80E-06	1	1.38E-04	1	2.50E+01	1	5.64E-03	1	8.24E+03	1	3.90E+02	1	5.71E+02	1	9.06E+00	1	1.90E+04	1	1.98E+01	1	2.66E-03	3	0.1	7
2-Propanol	1.02E-01	4	1.04E-05	6	8.10E-06	4	2.50E+01	4	3.32E-04	4	NA	NA	3.55E+02	4	NA	NA	4.28E-01	4	1.00E+06	4	4.54E+01	4	NA	NA	0.1	7
Styrene	7.10E-02	1	8.00E-06	1	2.74E-03	1	2.50E+01	1	1.12E-01	1	8.74E+03	1	4.18E+02	1	6.36E+02	1	7.76E+02	1	3.10E+02	1	6.21E+00	1	3.73E-02	3	0.1	7
1,1,2,2-Tetrachloroethane	7.10E-02	1	7.90E-06	1	3.44E-04	1	2.50E+01	1	1.41E-02	1	9.00E+03	1	4.20E+02	1	6.61E+02	1	9.33E+01	1	2.96E+03	1	4.62E+00	1	6.94E-03	3	0.1	7
Tetrachloroethene (PCE)	7.20E-02	1	8.20E-06	1	1.84E-02	1	2.50E+01	1	7.53E-01	1	8.29E+03	1	3.94E+02	1	6.20E+02	1	1.55E+02	1	2.00E+02	1	1.68E+01	1	3.34E-02	3	0.1	7
Tetrahydrofuran	1.02E-01	4	1.05E-05	6	7.05E-05	4	2.50E+01	4	2.89E-03	4	NA	NA	3.38E+02	4	NA	NA	1.11E+00	4	1.00E+06	4	7.43E+02	4	NA	NA	0.1	7
Toluene	8.70E-02	1	8.60E-06	1	6.62E-03	1	2.50E+01	1	2.72E-01	1	7.93E+03	1	3.84E+02	1	5.92E+02	1	1.82E+02	1	5.26E+02	1	2.87E+01	1	3.11E-02	3	0.1	7
1,1,1-Trichloroethane	7.80E-02	1	8.80E-06	1	1.72E-02	1	2.50E+01	1	7.03E-01	1	7.14E+03	1	3.47E+02	1	5.45E+02	1	1.10E+02	1	1.33E+03	1	1.30E+02	1	1.26E-02	3	0.1	7
Trichloroethene (TCE)	7.90E-02	1	9.10E-06	1	1.03E-02	1	2.50E+01	1	4.21E-01	1	7.51E+03	1	3.60E+02	1	5.44E+02	1	1.66E+02	1	1.47E+03	1	8.75E+01	1	1.16E-02	3	0.1	7
Trichlorofluoromethane (Freon 11)	8.70E-02	1	9.70E-06	1	9.68E-02	1	2.50E+01	1	3.97E+00	1	6.00E+03	1	2.97E+02	1	4.71E+02	1	4.97E+02	1	1.10E+03	1	5.89E+02	1	1.27E-02	3	0.1	7
1,1,2-Trichloro-1,2,2-trifluoroethane (Freon 113)	7.80E-02	1	8.20E-06	1	4.80E-01	1	2.50E+01	1	1.97E+01	1	6.46E+03	1	3.21E+02	1	4.87E+02	1	1.11E+04	1	1.70E+02	1	3.31E+02	1	NA	NA	0.1	7
1,2,4-Trimethylbenzene	6.06E-02	1	7.92E-06	1	6.14E-03	1	2.50E+01	1	2.52E-01	1	9.37E+03	1	4.42E+02	1	6.49E+02	1	1.35E+03	1	5.70E+01	1	2.21E+00	1	NA	NA	0.1	7
1,3,5-Trimethylbenzene	6.02E-02	1	8.67E-06	1	5.87E-03	1	2.50E+01	1	2.41E-01	1	9.32E+03	1	4.38E+02	1	6.37E+02	1	1.35E+03	1	2.00E+00	1	7.42E-02	1	NA	NA	0.1	7
Xylenes	8.50E-02	2	9.90E-06	2	7.30E-03	5	2.50E+01	1,8	2.70E-01	2	8.66E+03	1,8	4.18E+02	1,8	6.30E+02	1,8	4.43E+02	2	1.06E+02	2	6.60E+00	1,8	NA	NA	0.1	7

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Table 2. Physicochemical Properties of Chemicals of Potential Concern

Notes:

- (a) Sources of chemical properties are as follows:
- 1 USEPA. 2003. User's Guide for Evaluating Subsurface Vapor Intrusion into Buildings. Office of Emergency and Remedial Response. June 19.
- 2 USEPA. 2008. Regional Screening Levels (RSLs) for Chemical Contaminants at Superfund Sites. URL: http://www.epa.gov/region09/waste/sfund/prg/. September 12.
- 3 USEPA. 2004. Risk Assessment Guidance for Superfund (RAGS), Volume I: Human Health Evaluation Manual (Part E, Supplemental Guidance for Dermal Risk Assessment) Final. July.
- 4 SRC PhysProp Database. 2003. Found at http://esc.syrres.com/interkow/physdemo.htm. And methods from Schwarzenback R. P. et al. 1993. Environmental Organic Chemistry. John Wiley and Sons, Inc., New York, NY.
- 5 Regional Water Quality Control Board (RWQCB). 2000. Risked Based Screening Levels. Table J. Physio-chemical and Toxicity Constants used in Models. August.
- 6 USEPA. 2006. Water9, Version 3. June 29. URL: http://www.epa.gov/ttn/chief/software/water/water9_3
- 7 Cal/EPA, 2004. Preliminary Endangerment Assessment Guidance Manual. Department of Toxic Substances Control. January.
- 8 For xylenes, o-xylene physicochemical properties are used.
- (b) "NA" indicates not available.

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Table 3. Johnson and Ettinger Model Inputs

Parameter	Symbol	Value	Units	Reference
Building Properties				
Depth below grade to bottom of enclosed space floor	L_{F}	15	cm	DTSC/HERD default (Cal/EPA, 2005a; 2005b)
Area of enclosed space below grade	$A_{b,sg}$	2.05E+07	cm^2	Site-specific
Building air exchange rate	AXR_b	1	hr ⁻¹	DTSC default (Cal/EPA, 2005b)
Building height	Bh	503	cm	Site-specific
Building ventilation rate	Q_b	2.86E+06	cm^3/s	Calculated: $A_{b,sg} x AXR_b x B_h$
Vapor flow rate into building	Q_{soil}	102	L/min	Calculated (Cal/EPA, 2005b)
Vapor flow rate into building	Q_{soil}	1707	cm ³ /s	Calculated via units conversion
Soil Properties				
Average soil temperature	T_s	17	°C	Region-specific (USEPA, 2004)
SCS soil type	Site-specific	Site-specific	_	Site-specific
Dry bulk density	$ ho_{b}$	1.79	g/cm ³	Site-specific average
Total porosity	n	0.339	cm ³ /cm ³	Site-specific average
Water-filled porosity	θ_{w}	0.236	cm ³ /cm ³	Site-specific average
Air-filled porosity	θ_a	0.103	cm ³ /cm ³	Site-specific average

Table 4. Exposure Point Concentrations of Chemicals of Potential Concern in Indoor Air of the Onsite Building

	Modeled Sou		Results of Vapor Intrusion Modeling			
Analyte	C_{SG} ($\mu g/m^3$)	Depth (cm)	α	C_{IA} $(\mu g/m^3)$		
Acetone	620	91	9.9E-05	6.1E-02		
Benzene	40,000	91	3.5E-05	1.4E+00		
1,3-Butadiene	310	91	8.8E-05	2.7E-02		
2-Butanone (methyl ethyl ketone)	420	91	6.7E-05	2.8E-02		
Carbon disulfide	440	91	4.0E-05	1.8E-02		
Chlorobenzene	160	91	2.9E-05	4.7E-03		
Chloroform	170	91	4.1E-05	7.0E-03		
Chloromethane (methyl chloride)	75	91	4.8E-05	3.6E-03		
Cyclohexane	480	91	3.2E-05	1.5E-02		
1,2-Dichlorobenzene	2,900	91	2.8E-05	8.3E-02		
1,3-Dichlorobenzene	210	91	2.8E-05	5.9E-03		
1,4-Dichlorobenzene	460	91	2.8E-05	1.3E-02		
Dichlorodifluoromethane (Freon 12)	10,000	91	2.7E-05	2.7E-01		
1,1-Dichloroethane	140	91	3.0E-05	4.2E-03		
1,2-Dichloroethane	140	91	4.3E-05	6.0E-03		
1,1-Dichloroethene (1,1-DCE)	140	91	3.5E-05	5.0E-03		
cis-1,2-Dichloroethene (cis-1,2-DCE)	140	91	3.0E-05	4.2E-03		
1,4-Dioxane	500	91	2.4E-04	1.2E-01		
Ethanol	2,600	91	2.6E-04	6.8E-01		
Ethylbenzene	7,700	91	3.0E-05	2.3E-01		
4-Ethyltoluene	3,700	91	2.7E-05	1.0E-01		
Heptane	550	91	2.8E-05	1.5E-02		
Hexane	63,000	91	7.3E-05	4.6E+00		
Methyl tertiary butyl ether (MTBE)	500	91	4.3E-05	2.2E-02		
Methylene chloride	1,200	91	4.0E-05	4.8E-02		
4-Methyl-2-pentanone (methyl isobutyl ketone)	550	91	4.2E-05	2.3E-02		
2-Propanol	350	91	1.7E-04	6.1E-02		
Styrene	150	91	2.9E-05	4.3E-03		
Tetrachloroethene (PCE)	1,100	91	2.9E-05	3.2E-02		
Tetrahydrofuran	420	91	6.0E-05	2.5E-02		

Table 4. Exposure Point Concentrations of Chemicals of Potential Concern in Indoor Air of the Onsite Building

	Modeled Sou		Results of Vapor Intrusion Modeling				
Analyte	C_{SG} (µg/m ³)	Depth (cm)	α	C_{IA} $(\mu g/m^3)$			
Toluene	32,000	91	3.4E-05	1.1E+00			
1,1,1-Trichloroethane	190	91	3.1E-05	5.9E-03			
Trichloroethene (TCE)	190	91	3.1E-05	6.0E-03			
Trichlorofluoromethane (Freon 11)	200	91	3.4E-05	6.8E-03			
1,1,2-Trichloro-1,2,2-trifluoroethane (Freon 113)	270	91	3.1E-05	8.3E-03			
1,2,4-Trimethylbenzene	2,900	91	2.5E-05	7.1E-02			
1,3,5-Trimethylbenzene	3,600	91	2.4E-05	8.8E-02			
Xylenes	19,000	91	3.4E-05	6.4E-01			

- (1) This vapor intrusion transport analysis is based on maximum concentrations of volatile organic compounds (VOCs) in soil gas from the August 1999 and May 2008 site investigations (see Table 1). Chemicals of potential concern (COPCs) with respect to vapor intrusion are those VOCs detected above reporting limits in at least one soil gas sample.
- (2) The shallower sampling depth between the 1999 and 2008 sampling events, 3 feet below ground surface, is used in the model because it is a more conservative assumption, i.e., it produces higher indoor air concentrations.
- (3) Non source-related inputs to the Johnson and Ettinger Model are documented in Table 3. Shown here are the results of the Johnson and Ettinger Model, consisting of, for each chemical of potential concern, the predicted attenuation factor (α) and the predicted concentration of the chemical in indoor air (CIA).

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Table 5. Dispersion Factor Calculation

Parameter	Name	Value	Units	Reference
A_{site}	Area of site	1.3	acre	Area of deed-restricted portion of the site
A	Dispersion coefficient	13.81	_	Zone 2 / San Francisco (USEPA, 2002)
В	Dispersion coefficient	20.16	-	Zone 2 / San Francisco (USEPA, 2002)
C	Dispersion coefficient	234.29	_	Zone 2 / San Francisco (USEPA, 2002)
$Q/C_{\rm vol}$	Dispersion factor	74.89	g/m ² /s per kg/m ³	Equation D-1 (USEPA, 2002)

Table 6. Exposure Point Concentrations of Chemicals of Potential Concern in Onsite Outdoor Air and in Offsite Air

Chemical of Potential Concern	Concentration
	$(\mu g/m^3)$
Acetone	9.6E-04
Benzene	2.3E-02
1,3-Butadiene	5.0E-04
2-Butanone (methyl ethyl ketone)	4.0E-04
Carbon disulfide	2.9E-04
Chlorobenzene	7.6E-05
Chloroform	1.1E-04
Chloromethane (methyl chloride)	6.1E-05
Cyclohexane	2.5E-04
1,2-Dichlorobenzene	1.3E-03
1,3-Dichlorobenzene	9.5E-05
1,4-Dichlorobenzene	2.1E-04
Dichlorodifluoromethane (Freon 12)	4.3E-03
1,1-Dichloroethane	6.7E-05
1,2-Dichloroethane	9.7E-05
1,1-Dichloroethene (1,1-DCE)	8.1E-05
cis-1,2-Dichloroethene (cis-1,2-DCE)	6.7E-05
1,4-Dioxane	3.1E-03
Ethanol	1.9E-02
Ethylbenzene	3.7E-03
4-Ethyltoluene	1.6E-03
Heptane	2.5E-04
Hexane	8.1E-02
Methyl tertiary butyl ether (MTBE)	3.5E-04
Methylene chloride	8.0E-04
4-Methyl-2-pentanone (methyl isobutyl ketone)	3.4E-04
2-Propanol	1.4E-03
Styrene	7.0E-05
Tetrachloroethene (PCE)	5.1E-04
Tetrahydrofuran	4.3E-04
Toluene	1.8E-02
1,1,1-Trichloroethane	9.6E-05

Table 6. Exposure Point Concentrations of Chemicals of Potential Concern in Onsite Outdoor Air and in Offsite Air

Chemical of Potential Concern	Concentration (µg/m³)
Trichloroethene (TCE)	9.7E-05
Trichlorofluoromethane (Freon 11)	1.1E-04
1,1,2-Trichloro-1,2,2-trifluoroethane (Freon 113)	1.4E-04
1,2,4-Trimethylbenzene	1.1E-03
1,3,5-Trimethylbenzene	1.4E-03
Xylenes	1.0E-02

Table 7. Exposure Point Concentrations of Chemicals of Potential Concern in Onsite Soil

Chemical of Potential Concern	Concentration (mg/kg)
Benzene	140
Chlorobenzene	0.0017
1,2-Dichlorobenzene	3.1
1,3-Dichlorobenzene	0.038
1,4-Dichlorobenzene	0.33
1,2-Dichloroethane	0.43
Ethylbenzene	170
Methyl tertiary butyl ether (MTBE)	0.084
Toluene	580
Xylenes	990

⁽¹⁾ Concentration units are milligrams per kilograms (mg/kg).

⁽²⁾ Exposure point concentrations (EPCs) in soil are historical maximum detected concentrations.

Table 8. Exposure Point Concentrations of Chemicals of Potential Concern in Onsite Groundwater

Chemical of Potential Concern	Concentration
	(μg/L)
Benzene	99,000
Bromodichloromethane	0.84
Chlorobenzene	0.90
Chloroethane (ethyl chloride)	46
Chloroform	4.7
Chloromethane (methyl chloride)	13
1,2-Dichlorobenzene	0.50
Dichlorodifluoromethane (Freon 12)	3.8
1,1-Dichloroethane	240
1,2-Dichloroethane	2,200
1,1-Dichloroethene (1,1-DCE)	31
cis-1,2-Dichloroethene (cis-1,2-DCE)	8.9
Ethylbenzene	80,000
Methyl tertiary butyl ether (MTBE)	2,800
Methylene chloride	7.9
1,1,2,2-Tetrachloroethane	0.50
Toluene	110,000
1,1,1-Trichloroethane	1.0
Trichloroethene (TCE)	3.8
Xylenes	640,000

⁽¹⁾ Concentration units are micrograms per liter ($\mu g/L$).

⁽²⁾ Exposure point concentrations (EPCs) in groundwater are historical maximum detected concentrations.

Table 9. Chronic Daily Intake (CDI) Equations

Inhalation of Vapors

Noncancer

$$CDI_{inhv,worker} = \frac{C_a \times BR_{worker} \times EF_{worker} \times ED_{worker}}{BW_{worker} \times AT_{nc,worker}}$$

$$CDI_{inhv,child} = \frac{C_a \times BR_{child} \times EF_{child} \times ED_{child}}{BW_{child} \times AT_{nc,child}}$$

Canaar

$$CDI_{inhv,worker} = \frac{C_a \times BR_{worker} \times EF_{worker} \times ED_{worker}}{BW_{worker} \times AT_c}$$

$$CDI_{inhv,age-adjusted} = \frac{C_a \times BR_{child} \times EF_{child} \times ED_{child}}{BW_{child} \times AT_c} + \frac{C_a \times BR_{adult} \times EF_{adult} \times ED_{adult, age-adjusted}}{BW_{adult} \times AT_c}$$

Soil Ingestion

Noncancer

$$\mathbf{CDI}_{\mathrm{ing,worker}} = \frac{\mathbf{C}_{\mathrm{s}} \times \mathbf{IR}_{\mathrm{worker}} \times \mathbf{EF}_{\mathrm{worker}} \times \mathbf{ED}_{\mathrm{worker}} \times \mathbf{CF}}{\mathbf{BW}_{\mathrm{worker}} \times \mathbf{AT}_{\mathrm{nc,worker}}}$$

Cancar

$$CDI_{ing,worker} = \frac{C_s \times IR_{worker} \times EF_{worker} \times ED_{worker} \times CF}{BW_{worker} \times AT_c}$$

Soil Dermal Contact

Noncancer

$$CDI_{dem_s, worker} = \frac{C_s \times SA_{worker} \times AF_{worker} \times ABS \times EF_{worker} \times ED_{worker} \times CF}{BW_{worker} \times AT_{nc, worker}}$$

Cancer

$$CDI_{derm_s, worker} = \frac{C_s \times SA_{worker} \times AF_{worker} \times ABS \times EF_{worker} \times ED_{worker} \times CF}{BW_{worker} \times AT_c}$$

Groundwater Dermal Contact

Noncancer

$$CDI_{derm_gw,worker} = \frac{C_w \times SA_{worker} \times K_p \times ET_{worker} \times EF_{worker} \times ED_{worker} \times CF}{BW_{worker} \times AT_{nc,worker}}$$

Cancer

$$CDI_{derm_gw,worker} = \frac{C_{w} \times SA_{worker} \times K_{p} \times ET_{worker} \times EF_{worker} \times ED_{worker} \times CF}{BW_{worker} \times AT_{c}}$$

- (1) Definitions and values of symbols used are given in Tables 2 and 10.
- (2) Worker indicates commercial and construction worker scenario; adult and child indicate residential scenario.

Table 10. Exposure Assumptions

Parameter	Symbol	Units	Offsite Resident Adult	Offsite Resident Age-Adjusted	Offsite Resident Child	Indoor Commercial Worker	Onsite Outdoor Intrusive Construction Worker
Dermal Contact with Soil and Ground	lwater						
Surface area	SA	cm ² /d	NA	NA	NA	NA	5,700
Adherence factor (soil only)	AF	mg/cm ²	NA	NA	NA	NA	0.8
Conversion Factor	CF	kg/mg	NA	NA	NA	NA	1.0E-06
Ingestion of Soil							
Ingestion Rate	IR	mg/d	NA	NA	NA	NA	330
Conversion Factor	CF	kg/mg	NA	NA	NA	NA	0.000001
Inhalation of Volatiles							
Breathing Rate	BR	m^3/d	20	20	10	14	20
General Intake Parameters							
Exposure Time		hrs/d	NA	NA	NA	8	8
Exposure Frequency	EF	d/y	350	350	350	250	20
Exposure Duration	ED	y	30	24	6	25	1
Body Weight	BW	kg	70	70	15	70	70
Averaging Time-Carcinogens	AT_c	d	25,550	25,550	25,550	25,550	25,550
Averaging Time-Noncarcinogens	$AT_{nc} \\$	d	10,950	NA	2,190	9,125	365

⁽¹⁾ Exposure assumptions are derived from default values for the commercial scenario established in Cal/EPA's Recommended DTSC Default Exposure Factors for Use in Risk Assessment at California Military Facilities (Cal/EPA, 2005).

⁽²⁾ For the outdoor instrusive construction worker, it is assumed that the duration of instrusive construction work is 4 weeks (20 work days).

Table 11. Chronic Daily Intakes – Cancer

	Onsite Commercial Worker	Onsite Intrusive Construction Worker				Age-adjusted Offsite Resident
	Indoor Air	Outdoor Air Inhalation	Soil		Groundwater	Indoor/Outdoor Air
Chemical of Potential Concern	Inhalation		Ingestion	Dermal Contact	Dermal Contact	Inhalation
Acetone	3.0E-06	2.1E-10	-	-	_	1.4E-07
Benzene	6.8E-05	5.1E-09	5.2E-07	7.1E-07	7.5E-04	3.4E-06
Bromodichloromethane	-	_	_	_	2.0E-09	-
1,3-Butadiene	1.3E-06	1.1E-10	_	_	_	7.4E-08
2-Butanone (methyl ethyl ketone)	1.4E-06	9.1E-11	_	_	_	6.0E-08
Carbon disulfide	8.7E-07	6.6E-11	_	_	_	4.4E-08
Chlorobenzene	2.3E-07	1.7E-11	6.3E-12	8.7E-12	1.3E-08	1.1E-08
Chloroethane (ethyl chloride)	_	_	_	_	1.4E-07	_
Chloroform	3.4E-07	2.6E-11	_	_	1.6E-08	1.7E-08
Chloromethane (methyl chloride)	1.8E-07	1.4E-11	_	_	2.2E-08	9.1E-09
Cyclohexane	7.4E-07	5.5E-11	_	_	_	3.7E-08
1,2-Dichlorobenzene	4.0E-06	2.9E-10	1.1E-08	1.6E-08	1.1E-08	2.0E-07
1,3-Dichlorobenzene	2.9E-07	2.1E-11	1.4E-10	1.9E-10	_	1.4E-08
1,4-Dichlorobenzene	6.4E-07	4.7E-11	1.2E-09	1.7E-09	_	3.1E-08
Dichlorodifluoromethane (Freon 12)	1.3E-05	9.6E-10	_	_	1.7E-08	6.4E-07
1,1-Dichloroethane	2.0E-07	1.5E-11	_	_	8.3E-07	1.0E-08
1,2-Dichloroethane	2.9E-07	2.2E-11	1.6E-09	2.2E-09	4.7E-06	1.4E-08
1,1-Dichloroethene (1,1-DCE)	2.4E-07	1.8E-11	_	_	1.8E-07	1.2E-08
cis-1,2-Dichloroethene (cis-1,2-DCE)	2.0E-07	1.5E-11	_	_	2.9E-08	1.0E-08
1,4-Dioxane	5.9E-06	7.0E-10	_	_	_	4.7E-07
Ethanol	3.3E-05	4.3E-09	_	_	_	2.8E-06
Ethylbenzene	1.1E-05	8.4E-10	6.3E-07	8.7E-07	2.0E-03	5.6E-07
4-Ethyltoluene	4.9E-06	3.7E-10	_	_	_	2.4E-07
Heptane	7.5E-07	5.6E-11	_	_	_	3.7E-08
Hexane	2.3E-04	1.8E-08	_	_	_	1.2E-05

Table 11. Chronic Daily Intakes - Cancer

	Onsite Commercial Worker	(Onsite Intrusive (Age-adjusted Offsite Resident			
	Indoor Air	Outdoor Air		Soil		Indoor/Outdoor Air	
Chemical of Potential Concern	Inhalation	Inhalation	Ingestion	Dermal Contact	Dermal Contact	Inhalation	
Methyl tertiary butyl ether (MTBE)	1.1E-06	7.8E-11	3.1E-10	4.3E-10	5.0E-06	5.2E-08	
Methylene chloride	2.4E-06	1.8E-10	_	_	1.4E-08	1.2E-07	
4-Methyl-2-pentanone (methyl isobutyl ketone)	1.1E-06	7.7E-11	_	_	_	5.1E-08	
2-Propanol	3.0E-06	3.1E-10	_	_	_	2.0E-07	
Styrene	2.1E-07	1.6E-11	_	_	_	1.0E-08	
1,1,2,2-Tetrachloroethane	_	_	_	_	1.8E-09	_	
Tetrachloroethene (PCE)	1.5E-06	1.1E-10	_	_	_	7.6E-08	
Tetrahydrofuran	1.2E-06	9.7E-11	_	_	_	6.5E-08	
Toluene	5.4E-05	4.0E-09	2.1E-06	3.0E-06	1.7E-03	2.7E-06	
1,1,1-Trichloroethane	2.9E-07	2.1E-11	_	_	6.4E-09	1.4E-08	
Trichloroethene (TCE)	2.9E-07	2.2E-11	_	_	2.3E-08	1.4E-08	
Trichlorofluoromethane (Freon 11)	3.3E-07	2.5E-11	_	_	_	1.7E-08	
1,1,2-Trichloro-1,2,2-trifluoroethane (Freon 113)	4.1E-07	3.0E-11	_	_	_	2.0E-08	
1,2,4-Trimethylbenzene	3.5E-06	2.5E-10	_	_	_	1.7E-07	
1,3,5-Trimethylbenzene	4.3E-06	3.1E-10	_	_	_	2.1E-07	
Xylenes	3.1E-05	2.3E-09	3.7E-06	5.0E-06	1.3E-02	1.6E-06	

^{(1) &}quot;-" indicates chemical was not determined to be a COPC for the respective pathway.

Table 12. Chronic Daily Intakes – Noncancer

	Onsite Commercial Worker		Onsite Intrusive	Construction Worker	r	Child Offsite Resident
	Indoor Air	Outdoor Air		Soil	Groundwater	Indoor/Outdoor Air
Chemical of Potential Concern	Inhalation	Inhalation	Ingestion	Dermal Contact	Dermal Contact	Inhalation
Acetone	8.4E-06	1.7E-09	_	_	_	6.9E-08
Benzene	1.9E-04	4.2E-05	9.0E-03	1.2E-02	1.3E+01	1.7E-03
Bromodichloromethane	_	_	_	_	6.9E-06	_
1,3-Butadiene	3.8E-06	1.4E-05	_	_	_	5.6E-04
2-Butanone (methyl ethyl ketone)	3.8E-06	4.4E-09	_	_	_	1.8E-07
Carbon disulfide	2.4E-06	2.3E-08	_	_	_	9.4E-07
Chlorobenzene	6.5E-07	4.2E-09	2.2E-08	3.0E-08	4.5E-05	1.7E-07
Chloroethane (ethyl chloride)	-	_	_	_	2.5E-05	_
Chloroform	9.5E-07	2.1E-08	_	_	1.1E-04	8.6E-07
Chloromethane (methyl chloride)	5.0E-07	3.7E-08	_	_	5.9E-05	1.5E-06
Cyclohexane	2.1E-06	2.3E-09	_	_	_	9.2E-08
1,2-Dichlorobenzene	1.1E-05	3.6E-07	8.9E-06	1.2E-05	8.2E-06	1.5E-05
1,3-Dichlorobenzene	8.1E-07	4.9E-08	3.3E-07	4.5E-07	_	2.0E-06
1,4-Dichlorobenzene	1.8E-06	1.4E-08	2.8E-06	3.9E-06	_	5.8E-07
Dichlorodifluoromethane (Freon 12)	3.6E-05	1.2E-06	_	_	6.1E-06	4.8E-05
1,1-Dichloroethane	5.7E-07	7.5E-09	_	_	2.9E-04	3.1E-07
1,2-Dichloroethane	8.2E-07	1.3E-08	5.6E-06	7.7E-06	1.6E-02	5.4E-07
1,1-Dichloroethene (1,1-DCE)	6.8E-07	6.4E-08	_	_	2.6E-04	2.6E-06
cis-1,2-Dichloroethene (cis-1,2-DCE)	5.7E-07	1.1E-07	_	_	2.0E-04	4.3E-06
1,4-Dioxane	1.6E-05	5.7E-08	_	_	_	2.3E-06
Ethanol	9.3E-05	1.0E-06	-	_	_	4.1E-05
Ethylbenzene	3.2E-05	2.0E-07	4.4E-04	6.1E-04	1.4E+00	8.4E-06
4-Ethyltoluene	1.4E-05	9.0E-07	-	_	_	3.7E-05
Heptane	2.1E-06	1.9E-08	_	_	_	8.0E-07

Table 12. Chronic Daily Intakes – Noncancer

	Onsite Commercial Worker	(Onsite Intrusive	Construction Worker	<u>r</u>	Child Offsite Resident
	Indoor Air	Outdoor Air		Soil	Groundwater	Indoor/Outdoor Air
Chemical of Potential Concern	Inhalation	Inhalation	Ingestion	Dermal Contact	Dermal Contact	Inhalation
Hexane	6.3E-04	6.3E-06	_	_	-	2.6E-04
Methyl tertiary butyl ether (MTBE)	3.0E-06	6.4E-09	2.5E-08	3.5E-08	4.1E-04	2.6E-07
Methylene chloride	6.6E-06	1.1E-07	_	_	1.7E-05	4.5E-06
4-Methyl-2-pentanone (methyl isobutyl ketone)	3.2E-06	6.3E-09	_	_	_	2.6E-07
2-Propanol	8.4E-06	1.1E-08	_	_	_	4.4E-07
Styrene	5.9E-07	4.2E-09	-	_	-	1.7E-07
1,1,2,2-Tetrachloroethane	_	_	_	_	3.1E-05	_
Tetrachloroethene (PCE)	4.3E-06	8.0E-07	_	_	_	3.3E-05
Tetrahydrofuran	3.4E-06	7.9E-09	_	_	_	3.2E-07
Toluene	1.5E-04	3.3E-06	1.9E-03	2.6E-03	1.5E+00	1.3E-04
1,1,1-Trichloroethane	8.1E-07	5.2E-09	_	_	2.3E-07	2.1E-07
Trichloroethene (TCE)	8.2E-07	8.9E-09	_	_	5.3E-03	3.6E-07
Trichlorofluoromethane (Freon 11)	9.4E-07	8.8E-09	_	_	_	3.6E-07
1,1,2-Trichloro-1,2,2-trifluoroethane (Freon 113)	1.1E-06	2.5E-10	_	_	_	1.0E-08
1,2,4-Trimethylbenzene	9.8E-06	8.9E-06	_	_	-	3.6E-04
1,3,5-Trimethylbenzene	1.2E-05	1.3E-05	_	_	_	5.2E-04
Xylenes	8.8E-05	5.7E-06	1.3E-03	1.8E-03	4.7E+00	2.3E-04

^{(1) &}quot;-" indicates chemical was not determined to be a COPC for the respective pathway.

Table 13. Carcinogenic and Noncarcinogenic Toxicity Values

-	C	ancer Slope F	Factor (CSF)	Chronic Noncancer Reference Dose (RfD)							
Chemical of Potential Concern	Inhalation	Source	Oral	Source	Inhalation	Source	Oral	Source			
	(per mg/kg/d)		(per mg/kg/d)		(mg/kg/d)		(mg/kg/d)				
Acetone	NC	NA	NC	NA	8.9E+00	3A	9.0E-01	2			
Benzene	1.0E-01	1	1.0E-01	1	8.6E-03	2	4.0E-03	2			
Bromodichloromethane	1.3E-01	1	1.3E-01	1	2.0E-02	R	2.0E-02	2			
1,3-Butadiene	6.0E-01	1	3.4E+00	1	5.7E-04	2	5.7E-04	R			
2-Butanone (methyl ethyl ketone)	NC	NA	NC	NA	1.4E+00	2	6.0E-01	2			
Carbon disulfide	NC	NA	NC	NA	2.0E-01	2	1.0E-01	2			
Chlorobenzene	NC	NA	NC	NA	2.9E-01	1	2.0E-02	2			
Chloroethane (ethyl chloride)	NC	NA	NC	NA	2.9E+00	2	4.0E-01	4N			
Chloroform	1.9E-02	1	3.1E-02	1	8.6E-02	1	1.0E-02	2			
Chloromethane (methyl chloride)	6.3E-03	3H	1.3E-02	3H	2.6E-02	2	2.6E-02	R			
Cyclohexane	NC	NA	NC	NA	1.7E+00	2	1.7E+00	R			
1,2-Dichlorobenzene	NC	NA	NC	NA	5.7E-02	3H	9.0E-02	2			
1,3-Dichlorobenzene	NC	NA	NC	NA	3.0E-02	R	3.0E-02	4N			
1,4-Dichlorobenzene	4.0E-02	1	5.4E-03	1	2.3E-01	1	3.0E-02	4N			
Dichlorodifluoromethane (Freon 12)	NC	NA	NC	NA	5.7E-02	3Н	2.0E-01	2			
1,1-Dichloroethane	5.7E-03	1	5.7E-03	1	1.4E-01	4H	2.0E-01	3P			
1,2-Dichloroethane	7.2E-02	1	4.7E-02	1	1.1E-01	1	2.0E-02	3P			
1,1-Dichloroethene (1,1-DCE)	NC	NA	NC	NA	2.0E-02	1	5.0E-02	2			
cis-1,2-Dichloroethene (cis-1,2-DCE)	NC	NA	NC	NA	1.0E-02	R	1.0E-02	3P			
1,4-Dioxane	2.7E-02	1	2.7E-02	1	8.6E-01	1	8.6E-01	R			
Ethanol	NC	NA	NC	NA	3.0E-01	R	3.0E-01	2			
Ethylbenzene	8.7E-03	1	1.1E-02	1	2.9E-01	2	1.0E-01	2			
4-Ethyltoluene	NC	NA	NC	NA	2.9E-02	2	2.0E-01	2			
Heptane	NC	NA	NC	NA	2.0E-01	2	6.0E-02	3H			
Hexane	NC	NA	NC	NA	2.0E-01	2	6.0E-02	3H			
Methyl tertiary butyl ether (MTBE)	9.1E-04	1	1.8E-03	1	8.6E-01	2	8.6E-01	R			
Methylene chloride	3.5E-03	1	1.4E-02	1	1.1E-01	1	6.0E-02	2			
4-Methyl-2-pentanone (methyl isobutyl ketone)	NC	NA	NC	NA	8.6E-01	2	8.0E-02	3H			
2-Propanol	NC	NA	NC	NA	2.0E+00	1	2.0E+00	R			
Styrene	NC	NA	NC	NA	2.6E-01	1	2.0E-01	2			
1,1,2,2-Tetrachloroethane	2.0E-01	1	2.7E-01	1	4.0E-03	R	4.0E-03	3P			
Tetrachloroethene (PCE)	2.1E-02	1	5.4E-01	1	1.0E-02	1	1.0E-02	2			
Tetrahydrofuran	7.0E-03	N	7.6E-03	4N	8.6E-01	N	3.0E-01	N			
Toluene	NC	NA	NC	NA	8.6E-02	1	8.0E-02	2			
1,1,1-Trichloroethane	NC	NA	NC	NA	2.9E-01	1	2.0E+00	2			
Trichloroethene (TCE)	7.0E-03	1	1.3E-02	1	1.7E-01	1	3.0E-04	4N			
Trichlorofluoromethane (Freon 11)	NC	NA	NC	NA	2.0E-01	3H	3.0E-01	2			
1,1,2-Trichloro-1,2,2-trifluoroethane (Freon 113)	NC	NA	NC	NA	8.6E+00	3H	3.0E+00	2			
1,2,4-Trimethylbenzene	NC	NA	NC	NA	2.0E-03	3P	5.0E-02	4P			
1,3,5-Trimethylbenzene	NC	NA	NC	NA	1.7E-03	3P	5.0E-02	3P			
Xylenes	NC	NA	NC	NA	2.9E-02	2	2.0E-01	2			

Table 13. Carcinogenic and Noncarcinogenic Toxicity Values

- (a) Sources of toxicity data are as follows.
 - 1 OEHHA Toxicity Criteria Database
 - 2 USEPA Integrated Risk Information System (IRIS)
 - 3 USEPA Region 9 Regional Screening Levels Table (2008)
 - 4 USEPA Region 9 PRG Table (2004)
 - A Agency for Toxic Substances & Disease Registry (ATSDR)
 - P USEPA Provisional Peer Reviewed Toxicity Values (PPRTVs) Database
 - N USEPA National Center for Environmental Assessment
 - H USEPA HEAST
 - R Route-to-route extrapolation
- (b) Isobutanol was used as the surrogate for ethanol's oral and inhalation noncancer reference dose.
- (c) Xylene was used as the surrogate for 4-ethyltoluene's oral and inhalation noncancer reference dose.
- (d) Hexane was used as the surrogate for this heptane's oral and inhalation noncancer reference dose.
- (e) Tetrahydrofuran toxicological values were derived from Draft Toxicological Review of Tetrahydrofuran (USEPA, 2007).
- (f) "NC" indicates that the chemical is classified as a noncarcinogen for the inhalation pathway.
- (g) "NA" means not-applicable (see note [e]).

Table 14. Summary of Estimated Cancer Risks – Baseline

	Onsite Commercial Worker		Onsite	Intrusive Constructi	on Worker		Age-adjusted Offsite Resident
	Indoor Air	Outdoor Air		Soil	Groundwater	Total	Indoor/Outdoor Air
Chemical of Potential Concern	Inhalation	Inhalation	Ingestion	Dermal Contact	Dermal Contact		Inhalation
Acetone	NC	NC	-	-	=	-	NC
Benzene	6.8E-06	5.1E-10	5.2E-08	7.1E-08	7.5E-05	7.5E-05	3.4E-07
Bromodichloromethane	-	-	_	-	2.6E-10	2.6E-10	-
1,3-Butadiene	8.0E-07	6.7E-11	-	_	_	6.7E-11	4.4E-08
2-Butanone (methyl ethyl ketone)	NC	NC	-		-	-	NC
Carbon disulfide	NC	NC	_		-	-	NC
Chlorobenzene	NC	NC	NC	NC	NC	_	NC
Chloroethane (ethyl chloride)	_	_	-	_	NC	_	-
Chloroform	6.5E-09	4.9E-13	=	_	5.1E-10	5.1E-10	3.2E-10
Chloromethane (methyl chloride)	1.1E-09	8.6E-14	=	_	2.8E-10	2.8E-10	5.7E-11
Cyclohexane	NC	NC	_	_	_	_	NC
1,2-Dichlorobenzene	NC	NC	NC	NC	NC	=	NC
1,3-Dichlorobenzene	NC	NC	NC	NC	_	=	NC
1,4-Dichlorobenzene	2.5E-08	1.9E-12	6.6E-12	9.1E-12	_	1.8E-11	1.2E-09
Dichlorodifluoromethane (Freon 12)	NC	NC	-	_	NC	=	NC
1,1-Dichloroethane	1.2E-09	8.6E-14	=	_	4.7E-09	4.7E-09	5.7E-11
1,2-Dichloroethane	2.1E-08	1.6E-12	7.5E-11	1.0E-10	2.2E-07	2.2E-07	1.0E-09
1,1-Dichloroethene (1,1-DCE)	NC	NC	=	_	NC	=	NC
cis-1,2-Dichloroethene (cis-1,2-DCE)	NC	NC	-	_	NC	=	NC
1,4-Dioxane	1.6E-07	1.9E-11	-	_	_	1.9E-11	1.3E-08
Ethanol	NC	NC	_	_	_	_	NC
Ethylbenzene	9.8E-08	7.3E-12	6.9E-09	9.5E-09	2.2E-05	2.2E-05	4.8E-09
4-Ethyltoluene	NC	NC	_	_	=	_	NC
Heptane	NC	NC	_	_	_	=	NC
Hexane	NC	NC	-	_	_	=	NC
Methyl tertiary butyl ether (MTBE)	9.6E-10	7.1E-14	5.6E-13	7.7E-13	9.0E-09	9.0E-09	4.7E-11
Methylene chloride	8.3E-09	6.2E-13	=	_	2.0E-10	2.0E-10	4.1E-10
4-Methyl-2-pentanone (methyl isobutyl ketone)	NC	NC	-	_	_	-	NC

Table 14. Summary of Estimated Cancer Risks – Baseline

	Onsite Commercial Worker		Onsite		Age-adjusted Offsite Resident		
	Indoor Air	Outdoor Air		Soil	Groundwater	Total	Indoor/Outdoor Air
Chemical of Potential Concern	Inhalation	Inhalation	Ingestion	Dermal Contact	Dermal Contact		Inhalation
2-Propanol	NC	NC	-	=	_	=	NC
Styrene	NC	NC	_	_	-	-	NC
1,1,2,2-Tetrachloroethane	-	_	_	_	4.8E-10	4.8E-10	-
Tetrachloroethene (PCE)	3.2E-08	2.4E-12	_	_	-	2.4E-12	1.6E-09
Tetrahydrofuran	8.6E-09	6.8E-13	_	_	-	6.8E-13	4.5E-10
Toluene	NC	NC	NC	NC	NC	-	NC
1,1,1-Trichloroethane	NC	NC	_	_	NC	_	NC
Trichloroethene (TCE)	2.0E-09	1.5E-13	-	_	2.9E-10	2.9E-10	1.0E-10
Trichlorofluoromethane (Freon 11)	NC	NC	_	_	_	_	NC
1,1,2-Trichloro-1,2,2-trifluoroethane (Freon 113)	NC	NC	-	_	-	-	NC
1,2,4-Trimethylbenzene	NC	NC	=	_	_	=	NC
1,3,5-Trimethylbenzene	NC	NC	_	_	-	-	NC
Xylenes	NC	NC	NC	NC	NC	=	NC
Cumulative Risk	8.0E-06	6.1E-10	5.9E-08	8.1E-08	9.7E-05	9.8E-05	4.1E-07

^{(1) &}quot;-" indicates chemical was not determined to be a COPC for the respective pathway.

^{(2) &}quot;NC" indicates chemical was determined to be a noncarcinogen.

Table 15. Summary of Estimated Noncancer Hazard Indices – Baseline

	Onsite Commercial Worker		Onsite		Child Offsite Resident		
	Indoor Air	Outdoor Air		Soil	Groundwater	Total	Indoor/Outdoor Air
Chemical of Potential Concern	Inhalation	Inhalation	Ingestion	Dermal Contact	Dermal Contact		Inhalation
Acetone	9.5E-07	1.7E-09	_	-	-	1.7E-09	6.9E-08
Benzene	2.2E-02	4.2E-05	9.0E-03	1.2E-02	1.3E+01	1.3E+01	1.7E-03
Bromodichloromethane	=	_	-	-	6.9E-06	6.9E-06	=
1,3-Butadiene	6.6E-03	1.4E-05	-	-	_	1.4E-05	5.6E-04
2-Butanone (methyl ethyl ketone)	2.7E-06	4.4E-09	-	-	_	4.4E-09	1.8E-07
Carbon disulfide	1.2E-05	2.3E-08	-	-	_	2.3E-08	9.4E-07
Chlorobenzene	2.3E-06	4.2E-09	2.2E-08	3.0E-08	4.5E-05	4.5E-05	1.7E-07
Chloroethane (ethyl chloride)	=	-	_	-	2.5E-05	2.5E-05	-
Chloroform	1.1E-05	2.1E-08	-	-	1.1E-04	1.1E-04	8.6E-07
Chloromethane (methyl chloride)	1.9E-05	3.7E-08	=	_	5.9E-05	5.9E-05	1.5E-06
Cyclohexane	1.2E-06	2.3E-09	=	_	_	2.3E-09	9.2E-08
1,2-Dichlorobenzene	2.0E-04	3.6E-07	8.9E-06	1.2E-05	8.2E-06	3.0E-05	1.5E-05
1,3-Dichlorobenzene	2.7E-05	4.9E-08	3.3E-07	4.5E-07	_	8.3E-07	2.0E-06
1,4-Dichlorobenzene	7.8E-06	1.4E-08	2.8E-06	3.9E-06	_	6.8E-06	5.8E-07
Dichlorodifluoromethane (Freon 12)	6.4E-04	1.2E-06	-	_	6.1E-06	7.2E-06	4.8E-05
1,1-Dichloroethane	4.1E-06	7.5E-09	=	_	2.9E-04	2.9E-04	3.1E-07
1,2-Dichloroethane	7.1E-06	1.3E-08	5.6E-06	7.7E-06	1.6E-02	1.6E-02	5.4E-07
1,1-Dichloroethene (1,1-DCE)	3.4E-05	6.4E-08	=	_	2.6E-04	2.6E-04	2.6E-06
cis-1,2-Dichloroethene (cis-1,2-DCE)	5.7E-05	1.1E-07	-	_	2.0E-04	2.0E-04	4.3E-06
1,4-Dioxane	1.9E-05	5.7E-08	=	_	_	5.7E-08	2.3E-06
Ethanol	3.1E-04	1.0E-06	=	_	_	1.0E-06	4.1E-05
Ethylbenzene	1.1E-04	2.0E-07	4.4E-04	6.1E-04	1.4E+00	1.4E+00	8.4E-06
4-Ethyltoluene	4.8E-04	9.0E-07	-	_	_	9.0E-07	3.7E-05
Heptane	1.1E-05	1.9E-08	=	_	_	1.9E-08	8.0E-07
Hexane	3.2E-03	6.3E-06	_	_	=	6.3E-06	2.6E-04
Methyl tertiary butyl ether (MTBE)	3.4E-06	6.4E-09	2.5E-08	3.5E-08	4.1E-04	4.1E-04	2.6E-07
Methylene chloride	5.8E-05	1.1E-07	_	_	1.7E-05	1.7E-05	4.5E-06
4-Methyl-2-pentanone (methyl isobutyl ketone)	3.7E-06	6.3E-09	-	_	=	6.3E-09	2.6E-07
2-Propanol	4.2E-06	1.1E-08	_	-	_	1.1E-08	4.4E-07

Table 15. Summary of Estimated Noncancer Hazard Indices – Baseline

	Onsite Commercial Worker		Onsite	Intrusive Construction	on Worker		Child Offsite Resident
	Indoor Air	Outdoor Air		Soil	Groundwater	Total	Indoor/Outdoor Air
Chemical of Potential Concern	Inhalation	Inhalation	Ingestion	Dermal Contact	Dermal Contact		Inhalation
Styrene	2.3E-06	4.2E-09	=	=	-	4.2E-09	1.7E-07
1,1,2,2-Tetrachloroethane	=	-	-	-	3.1E-05	3.1E-05	=
Tetrachloroethene (PCE)	4.3E-04	8.0E-07	-	_	-	8.0E-07	3.3E-05
Геtrahydrofuran	4.0E-06	7.9E-09	-	-	-	7.9E-09	3.2E-07
Toluene	1.8E-03	3.3E-06	1.9E-03	2.6E-03	1.5E+00	1.5E+00	1.3E-04
,1,1-Trichloroethane	2.8E-06	5.2E-09	_	_	2.3E-07	2.3E-07	2.1E-07
Trichloroethene (TCE)	4.8E-06	8.9E-09	-	-	5.3E-03	5.3E-03	3.6E-07
Trichlorofluoromethane (Freon 11)	4.7E-06	8.8E-09	_	_	_	8.8E-09	3.6E-07
,1,2-Trichloro-1,2,2-trifluoroethane (Freon 113)	1.3E-07	2.5E-10	-	-	-	2.5E-10	1.0E-08
,2,4-Trimethylbenzene	4.9E-03	8.9E-06	-	_	-	8.9E-06	3.6E-04
,3,5-Trimethylbenzene	7.0E-03	1.3E-05	_	_	_	1.3E-05	5.2E-04
Kylenes	3.1E-03	5.7E-06	1.3E-03	1.8E-03	4.7E+00	4.7E+00	2.3E-04
Cumulative Hazard	5.1E-02	9.7E-05	1.3E-02	1.7E-02	2.1E+01	2.1E+01	4.0E-03

^{(1) &}quot;-" indicates chemical was not determined to be a COPC for the respective pathway.

Table 16. Risk-based Soil Gas Cleanup Goals

Chemical of Potential Concern	Cancer-based Indoor Air Target Concentration (CA_c) $(\mu g/m^3)$	Noncancer-based Indoor Air Target Concentration (CA_{nc}) $(\mu g/m^3)$	Controlling Indoor Air Target Concentration (CA) (µg/m³)	Attenuation Factor at 5.0 feet bgs	Risk-based Cleanup Goal at 5.0 ft bgs (µg/m³)
Acetone	NC	6.5E+04	6.5E+04	5.9E-05	1.1E+09
Benzene	2.0E+00	6.3E+01	2.0E+00	2.0E-05	1.0E+05
1,3-Butadiene	3.4E-01	4.2E+00	3.4E-01	5.3E-05	6.5E+03
2-Butanone (methyl ethyl ketone)	NC	1.0E+04	1.0E+04	3.9E-05	2.7E+08
Carbon disulfide	NC	1.5E+03	1.5E+03	2.3E-05	6.3E+07
Chlorobenzene	NC	2.1E+03	2.1E+03	1.7E-05	1.2E+08
Chloroform	1.1E+01	6.3E+02	1.1E+01	2.3E-05	4.6E+05
Chloromethane (methyl chloride)	3.2E+01	1.9E+02	3.2E+01	2.8E-05	1.2E+06
Cyclohexane	NC	1.3E+04	1.3E+04	1.8E-05	7.0E+08
1,2-Dichlorobenzene	NC	4.2E+02	4.2E+02	1.6E-05	2.6E+07
1,3-Dichlorobenzene	NC	2.2E+02	2.2E+02	1.6E-05	1.4E+07
1,4-Dichlorobenzene	5.1E+00	1.7E+03	5.1E+00	1.6E-05	3.2E+05
Dichlorodifluoromethane (Freon 12)	NC	4.2E+02	4.2E+02	1.5E-05	2.8E+07
1,1-Dichloroethane	3.6E+01	1.0E+03	3.6E+01	1.7E-05	2.1E+06
1,2-Dichloroethane	2.8E+00	8.3E+02	2.8E+00	2.4E-05	1.2E+05
1,1-Dichloroethene (1,1-DCE)	NC	1.5E+02	1.5E+02	2.0E-05	7.2E+06
cis-1,2-Dichloroethene (cis-1,2-DCE)	NC	7.3E+01	7.3E+01	1.7E-05	4.3E+06
1,4-Dioxane	7.6E+00	6.3E+03	7.6E+00	1.6E-04	4.7E+04
Ethanol	NC	2.2E+03	2.2E+03	1.8E-04	1.2E+07

Table 16. Risk-based Soil Gas Cleanup Goals

Chemical of Potential Concern	Cancer-based Indoor Air Target Concentration (CA _c) (µg/m³)	Noncancer-based Indoor Air Target Concentration (CA _{nc}) (µg/m³)	Controlling Indoor Air Target Concentration (CA) (µg/m³)	Attenuation Factor at 5.0 feet bgs	Risk-based Cleanup Goal at 5.0 ft bgs (µg/m³)
Ethylbenzene	2.3E+01	2.1E+03	2.3E+01	1.7E-05	1.4E+06
4-Ethyltoluene	NC	2.1E+02	2.1E+02	1.5E-05	1.3E+07
Heptane	NC	1.5E+03	1.5E+03	1.6E-05	9.2E+07
Hexane	NC	1.5E+03	1.5E+03	4.3E-05	3.4E+07
Methyl tertiary butyl ether (MTBE)	2.2E+02	6.3E+03	2.2E+02	2.5E-05	9.1E+06
Methylene chloride	5.8E+01	8.3E+02	5.8E+01	2.3E-05	2.5E+06
4-Methyl-2-pentanone (methyl isobutyl ketone)	NC	6.3E+03	6.3E+03	2.4E-05	2.6E+08
2-Propanol	NC	1.5E+04	1.5E+04	1.1E-04	1.3E+08
Styrene	NC	1.9E+03	1.9E+03	1.6E-05	1.1E+08
Tetrachloroethene (PCE)	9.7E+00	7.3E+01	9.7E+00	1.6E-05	6.0E+05
Tetrahydrofuran	2.9E+01	6.3E+03	2.9E+01	3.5E-05	8.4E+05
Toluene	NC	6.3E+02	6.3E+02	2.0E-05	3.2E+07
1,1,1-Trichloroethane	NC	2.1E+03	2.1E+03	1.8E-05	1.2E+08
Trichloroethene (TCE)	2.9E+01	1.3E+03	2.9E+01	1.8E-05	1.6E+06
Trichlorofluoromethane (Freon 11)	NC	1.5E+03	1.5E+03	2.0E-05	7.5E+07
1,1,2-Trichloro-1,2,2-trifluoroethane (Freon 113)	NC	6.3E+04	6.3E+04	1.8E-05	3.6E+09
1,2,4-Trimethylbenzene	NC	1.5E+01	1.5E+01	1.4E-05	1.1E+06
1,3,5-Trimethylbenzene	NC	1.3E+01	1.3E+01	1.4E-05	9.0E+05
Xylenes	NC	2.1E+02	2.1E+02	1.9E-05	1.1E+07

Table 16. Risk-based Soil Gas Cleanup Goals

Chemical of Potential Concern	Cancer-based Indoor Air Target Concentration (CA _c) (µg/m³)	Noncancer-based Indoor Air Target Concentration (CA _{nc}) (µg/m³)	Controlling Indoor Air Target Concentration (CA) (µg/m³)	Attenuation Factor at 5.0 feet bgs	Risk-based Cleanup Goal at 5.0 ft bgs (µg/m³)
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- (1) Cancer-based indoor air target concentrations assume a target risk of 1×10^{-5} . Noncancer-based indoor air target concentrations assume a target hazard quotient of 1.
- (2) Soil gas screening levels are calculated from the chemical-specific risk-based indoor air target concentration and attenuation factor:

$$C_{SG} = C_{IA} / \alpha$$

(3) NC = Noncarcinogenic

Date: 11/12/08

Contract Number: 07-557-A

I:\CAD\07\07-557-A\Site location map.dwg

Drafter: EC



Total and Effective Porosity Report

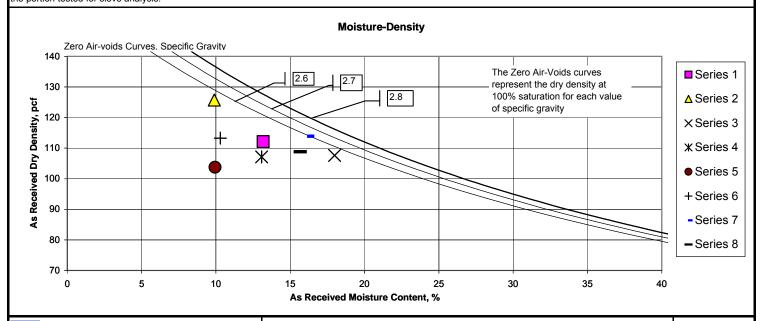
(API RP40 and ASTM D6836m)

 Job No:
 634-013
 Project No.:
 720-17784-1

 Client:
 TestAmerica
 Date:
 2/17/09

Project Name:	Nestle-Oakla	and		By:	PJ							
Boring:	GT-A-4.5	GT-A-5.0	GT-B-4.5	GT-B-5.0	GT-C-4.5	GT-C-5.0	GT-D-4.5	GT-D-5.0				
Sample:	720-17784-1	720-17784-2	720-17784-3	720-17784-4	720-17784-5	720-17784-6	720-17784-7	720-17784-8				
Depth, ft:												
Visual	Grayish	Grayish	Mottled	Mottled	Grayish	Grayish	Mottled	Brown SANI				
Description:	Brown	Brown	Reddish	Reddish	Brown SAND	Brown	Brown	w/ Clay*				
	Clayey	Clayey	Brown	Brown SAND	w/ Clay*	Clayey	Clayey					
	SAND w/	GRAVEL w/	Clayey	w/ Clay*		SAND	SAND					
	Gravel	Sand	SAND									
Total Porosity,												
%	33.5	25.7	36.4	36.5	38.3	32.4	32.5	35.7				
Effective												
Porosity, %	12.9	7.0	7.3	16.1	23.7	4.2	4.9	10.3				
Air-filled												
Porosity, %	9.9	5.8	5.5	14.1	21.8	13.7	3.0	8.4				
Water-filled												
Porosity, %	23.7	19.9	31.0	22.4	16.5	18.7	29.5	27.3				
Saturation, %	70.6	77.3	85.0	61.5	43.1	57.7	90.9	76.4				
Moisture, %	13.2	9.9	18.0	13.1	9.9	10.3	16.2	15.7				
Wet Unit wt, pcf	126.9	138.2	127.0	121.1	114.1	124.9	132.3	125.9				
Dry Unit wt, pcf	112.1	125.7	107.6	107.1	103.7	113.2	113.9	108.8				
Series	1	2	3	4	5	6	7	8				

Note: All reported values above are for the "as received" condition except for the effective porosity which is measured at a tension of 1/3 Bar. Both GT-A samples required significant patching due to gravel. This could have a significant impact on the reported values. * The material tested for effective porosity was slightly more coarsely grained than the portion tested for sieve analysis.



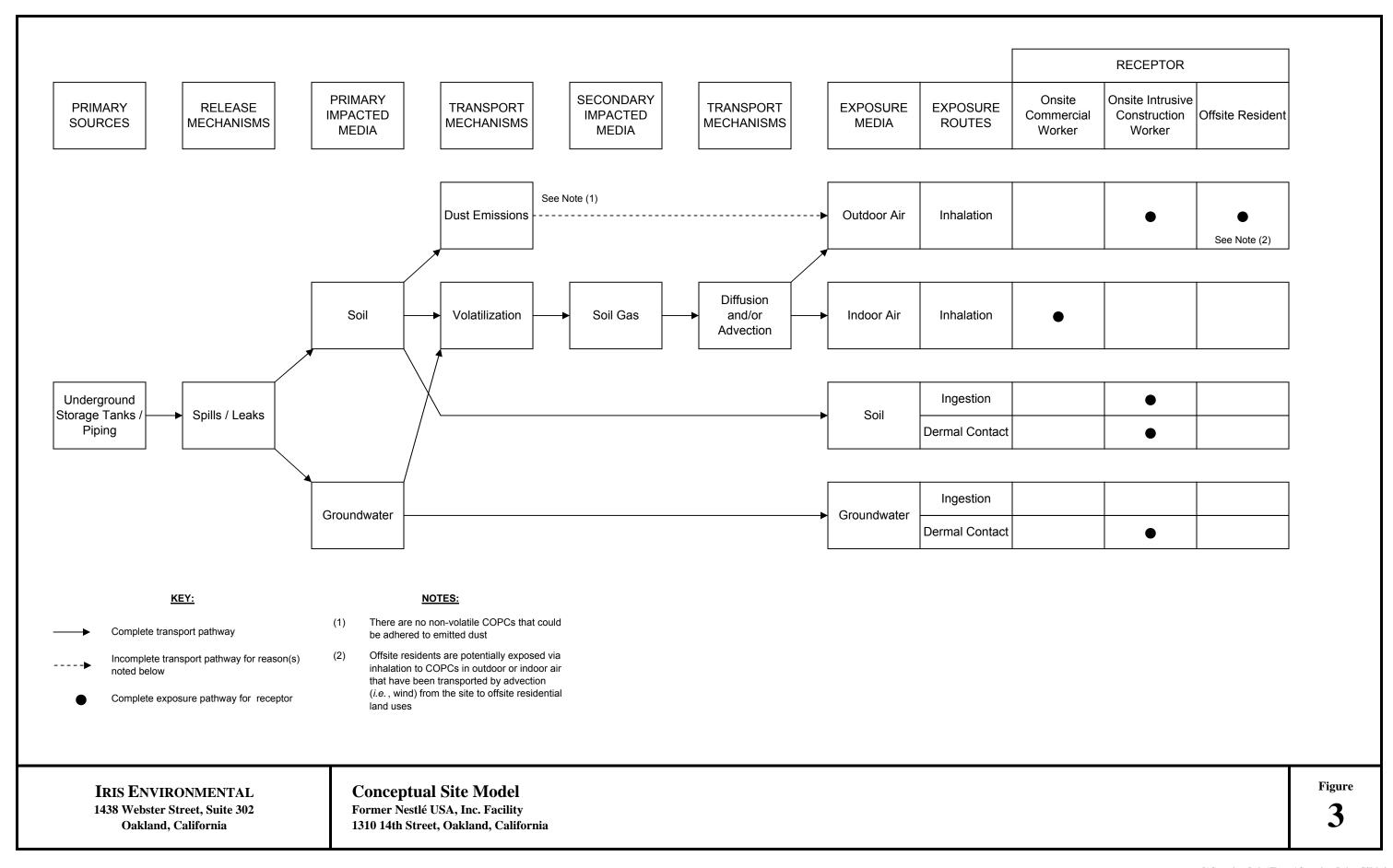
IRIS ENVIRONMENTAL 1438 Webster Street, Suite 302 Oakland, California 94612

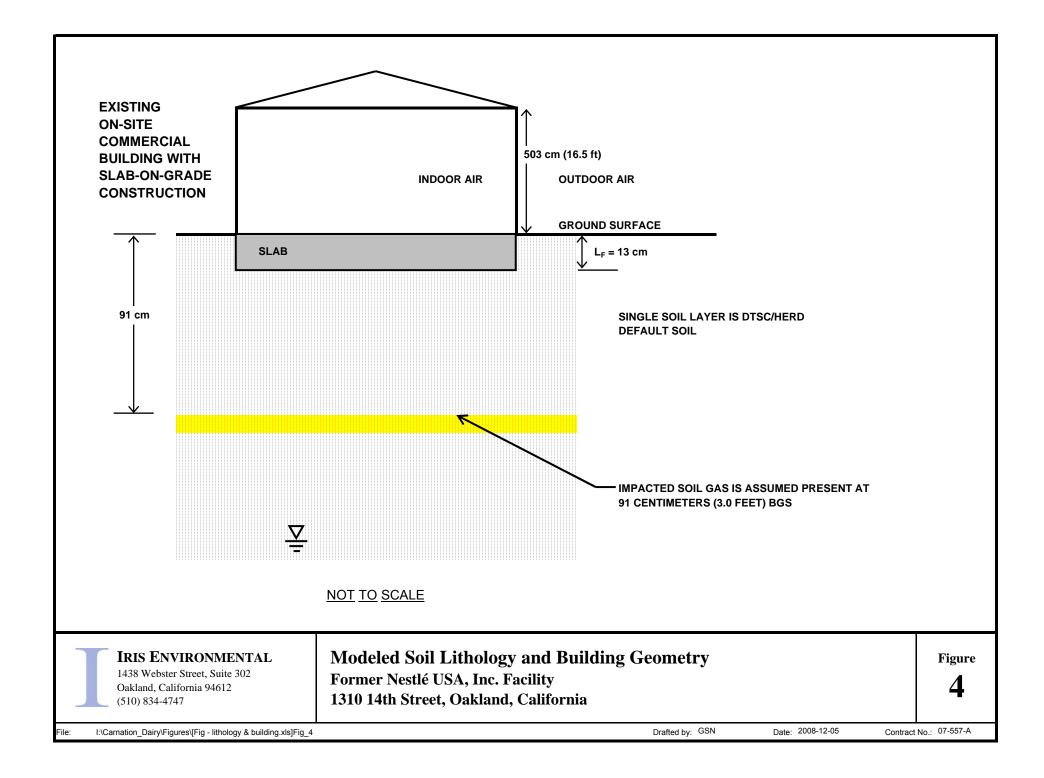
(510) 834-4747

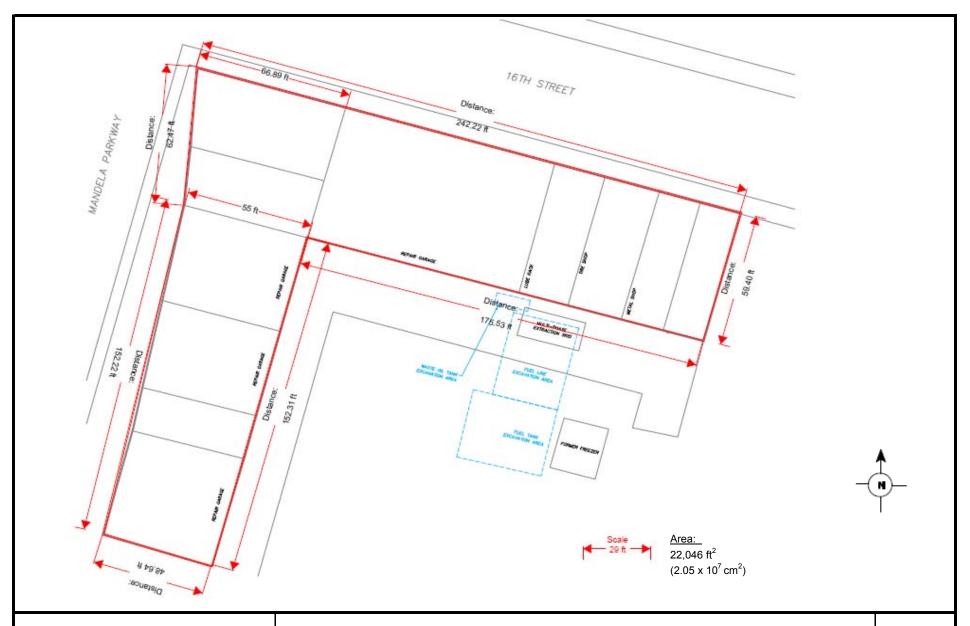
Total and Effective Porosity Report (Cooper, 2009) Former Nestlé USA, Inc. Facility 1310 14th Street, Oakland, California Figure

2

Drafted by: DT Date: 2009-03-18







IRIS ENVIRONMENTAL
1438 Webster Street, Suite 302
Oakland, California 94612
(510) 834-4747

Building Dimensions and Area Former Nestlé USA, Inc. Facility 1310 14th Street, Oakland, California

Figure 5

APPENDIX A SUMMARY OF PREVIOUS SITE INVESTIGATIONS

Table 1a: Soil Gas Sampling Results Vapors in Soil - August 99

																			Conce	ntration (ppbv)																			
															1,2-Di-	1,3-Di-	1,4-Di-	1,1-Di-	1,2-Di-	1,1-Di-	cis-1,2-									4-Methyl-		Methyl			Tetra-	Tetra-	1,1,1-Tri-	Tri-	1,2,4-Tri-	1,3,5-Tri-
Sample			Ethyl-	Total				1,3-Bu-	2-Bu-	Carbon	Chloro-	Chloro-	Chloro-	Cyclo-	chloro-	chloro-	chloro-	chloro-	chloro-	chloro-	Dichloro-	1,4-Di-		4-Ethyl-	Freon	Freon	Freon	Нер-	Hex-	2-penta- N	lethylene	t-butyl	2-Pro-	Sty-	chloro-	hydro-	chloro-	chloro-	methyl-	methyl-
ID	Benzene	Toluene	benzene	Xylenes	TPH-g	TPH-d	Acetone	tadiene	tanone	Disulfide	benzene	form	methane	hexane	benzene	benzene	benzene	ethane	ethane	ethene	ethene	oxane	Ethanol	toluene	11	12	113	tane	ane	none	Chloride	ether	panol	rene	ethene	furan	ethane	ethene	benzene	benzene
SB1, 3'	4.3	3.1	<0.65	2.74	800	NA	77 a	2.8	13	6.2	<0.65	<0.65	<0.65	<2.6	<0.65	<0.65	0.77	<0.65	<0.65	<0.65	<0.65	<2.6	63	<2.6	0.74	0.93	27	<2.6	4.4	3.8	3.7	<2.6	5.6	<0.65	1.2	<2.6	<0.65	<0.65	1.1	<0.65
SB2, 3'	7.5	12	3.6	17.6	1,100	NA	260 a	<2.7	24	9.0	<0.67	3.9	<0.67	12	<0.67	<0.67	1.8	<0.67	<0.67	<0.67	<0.67	<2.7	110	<2.7	1.2	200	<0.67	3.3	5.3	8.1	2.2	<2.7	<2.7	3.0	<0.67	<2.7	<0.67	<0.67	2.0	0.77
SB3, 3' SB3, 3' du	9,900 9,500	230 240	68 <140	67 <140	,	NA NA	<190 <580	<190 <580	<190 <580	<190 <580	<48 <140	<48 <140	<48 <140	<190 <580	<48 <140	<190 <580	<190 <580	<190 <580	<48 <140	180 160	<48 <140	<190 <580	590 580	<190 <580	<48 <140	<190 <580	<190 <580	<48 <140	<48 <140	<190 <580	<48 <140	<48 <140	<48 <140	<48 <140						
SB4, 3'	1,200	76	8.1	18.7	4,600	NA	200 a	19	<14	<14	<3.5	<3.5	<3.5	32	<3.5	<3.5	<3.5	<3.5	<3.5	<3.5	<3.5	<14	1,400	<14	<3.5	100	<3.5	<14	19	15	340	<14	22	<3.5	160	<14	21	<3.5	<3.5	<3.5
SB5, 3'	7.6	5.6	0.80	1.9	1,900	NA	45 a	61	12	18	<0.71	<0.71	0.77	8.2	<0.71	<0.71	<0.71	<0.71	<0.71	<0.71	<0.71	3.3	55	<2.8	4.4	1.2	3.4	<2.8	<2.8	<2.8	<0.71	<2.8	<2.8	<0.71	<0.71	<2.8	<0.71	<0.71	<0.71	<0.71
SB6, 3'	3.0	4.2	<0.68	2.52	560	NA	11 a	<2.7	4.0	<2.7	<0.68	<0.68	<0.68	<2.7	<0.68	<0.68	<0.68	<0.68	<0.68	<0.68	<0.68	<2.7	35	<2.7	<0.68	<0.68	<0.68	<2.7	<2.7	<2.7	<0.68	<2.7	<2.7	<0.68	<0.68	<2.7	<0.68	<0.68	1.1	<0.68
SB7, 3'	5.9	6.2	0.87	4.3	780	NA	43 a	3.4	7.9	3.3	<0.73	<0.73	<0.73	5.1	<0.73	<0.73	2.0	<0.73	<0.73	<0.73	<0.73	8.2	94	<2.9	0.74	1.1	<0.73	<2.9	6.8	4.4	<0.73	<2.9	3.8	1.0	2.0	<2.9	<0.73	<0.73	1.8	<0.73
SB8, 3'	10	12	3.8	15.7	1,300	NA	42 a	<11	<11	<11	<2.8	<2.8	<2.8	<11	<2.8	<2.8	<2.8	<2.8	<2.8	<2.8	<2.8	<11	62	<11	6.5	630	<2.8	<11	<11	<11	<2.8	<11	<11	<2.8	<2.8	<11	<2.8	<2.8	5.3	<2.8
SB9, 3'	12	18	1.7	9.9	690	NA	19 a	<2.7	6.0	<2.7	<0.68	1.1	<0.68	4.9	<0.68	<0.68	<0.68	<0.68	<0.68	<0.68	<0.68	<2.7	47	<2.7	1.5	20	<0.68	<2.7	4.3	<2.7	<0.68	<2.7	<2.7	<0.68	<0.68	<2.7	<0.68	<0.68	2.3	0.77
SB10, 3'	3.5	2.8	<0.80	1.7	610	NA	39 a	<3.2	9.7	<3.2	<0.80	1.6	<0.80	<3.2	<0.80	<0.80	<0.80	<0.80	<0.80	<0.80	<0.80	<3.2	40	<3.2	<0.80	1.4	<0.80	<3.2	3.9	<3.2	<0.80	<3.2	<3.2	<0.80	<0.80	<3.2	<0.80	<0.80	1.2	<0.80
SB11, 3'	2.7	1.9	<0.82	0.91	520	NA	38 a	<3.3	9.9	<3.3	<0.82	<0.82	3.7	<3.3	<0.82	<0.82	<0.82	<0.82	<0.82	<0.82	<0.82	22	23	<3.3	4.6	<0.82	<0.82	<3.3	<3.3	<3.3	1.2	<3.3	<3.3	<0.82	<0.82	<3.3	<0.82	<0.82	0.85	<0.82
SB12, 3'	250	<70	<70	610	750,000	NA	<280	<280	<280	<280	<70	<70	<70	<280	480	<70	76	<70	<70	<70	<70	<280	<280	760	<70	<70	<70	<280	18,000	<280	<70	<280	<280	<70	<70	<280	<70	<70	580	740
SB13, 3'	0.91	8.5	<0.67	1.3	550	NA	49 a	<2.7	5.5	6.4	<0.67	<0.67	<0.67	<2.7	<0.67	<0.67	<0.67	<0.67	<0.67	<0.67	<0.67	4.3	410 b	<2.7	<0.67	<0.67	<0.67	3.4	<2.7	<2.7	5.6	<2.7	26	<0.67	<0.67	58	<0.67	<0.67	1.1	<0.67
SB14, 3'	2.7	5.3	0.87	4.7	620	NA	10 a	<2.8	3.5	<2.8	<0.70	<0.70	<0.70	<2.8	<0.70	<0.70	1.6	<0.70	<0.70	<0.70	<0.70	<2.8	67	<2.8	<0.70	<0.70	<0.70	<2.8	<2.8	2.8	1.3	2.9	<2.8	0.82	<0.70	<2.8	<0.70	<0.70	2.0	0.81
SB15, 3'	42	12	1.6	6.7	2,100	NA	51 a	13	13	<5.8	<1.4	<1.4	<1.4	<5.8	<1.4	<1.4	<1.4	<1.4	<1.4	<1.4	<1.4	<5.8	190	<5.8	<1.4	46	<1.4	<5.8	50	<5.8	4.8	<5.8	<5.8	<1.4	2.1	<5.8	<1.4	<1.4	1.8	<1.4

Notes:

Parts per billion volumetric.

Compound present in laboratory blank greater than reporting limit (background subtraction not performed). Exceeds instrument calibration range.

NA Not analyzed.

Total Petroleum Hydrocarbons as gasoline. Total Petroleum Hydrocarbons as diesel. TPH-g

Revised Site Conceptual Model Former Nestlé USA, Inc. Facility-Oakland, CA 1310 14th Street, Oakland, CA

Table 1b: Soil Gas Sampling Results
Vapors in Soil - May 08

Boring Location	Sample Depth (feet	Date of Sample	e Analytical results (ug/L) of Vapor												
	bgs)	Collection	TPH g	TPH d	Benzene	Ethylbenzene	Toluene	Xylenes, Tot	1,2-DCA	Others					
SB-16	5	19-May-08	<10	<50	<0.10	<0.10	<0.20	<0.30	<0.10						
SB-17	5	19-May-08	<10	<50	<0.10	<0.10	<0.20	< 0.30	<0.10						
SB-18	5	19-May-08	630	<50	2.2	<0.10	0.44	< 0.30	<0.10						
SB-19	5	19-May-08	<10	<50	<0.10	<0.10	<0.20	< 0.30	<0.10						
SB-20/ PCB-7	5	19-May-08	19	<50	<0.10	<0.10	<0.20	< 0.30	<0.10						
SB-21/ PCB-8	5	19-May-08	25	<50	<0.10	<0.10	<0.20	< 0.30	<0.10						
SB-22	5	19-May-08	2,600	<50	40	7.7	32	19.1	<0.10	Dichlorodifluoromethane: 0.39					
SB-23	5	19-May-08	<10	<50	<0.10	<0.10	<0.20	< 0.30	<0.10						
SB-24/ PCB-1	5	19-May-08	<10	<50	<0.10	<0.10	0.22	< 0.30	<0.10						
SB-25/ PCB-2	5	19-May-08	<10	<50	<0.10	<0.10	<0.20	< 0.30	<0.10						
SB-26	5	19-May-08	<10	<50	<0.10	<0.10	<0.20	< 0.30	<0.10	Dichlorodifluoromethane: 10					
SB-27/ PCB-3	5	19-May-08	<10	<50	<0.10	<0.10	<0.20	< 0.30	<0.10						
SB-22 dup	5	19-May-08	2,600	<50	40	7.5	32	18.0	<0.10	Dichlorodifluoromethane: 0.38					
Probe Blank	NA	19-May-08	<10	<50	<0.10	<0.10	<0.20	< 0.30	<0.10						

Notes:

EPA Method 8260B for VOC Analyses of soil vapor EPA Mentod 8015m for TPH-g and TPH-d analyses of soil vapor

Table 2: Historical Soil Sample Results (1999 - 2008)

Boring Location	Sample Depth	Date of Sample					Analytica	ıl results (mç	ı/Kg)		
	(feet bgs)	Collection	TPH g	TPH d	TPH mo	Benzene	Toluene	Ethylbenzene	Xylenes, Tot	1,2-DCA	Others
SB-1	3.5-4.0	08/12/99	<0.13	1,200	NA	< 0.0013	<0.0013	< 0.0013	< 0.0013	< 0.0011	
SB-1	6.5-7.0	08/12/99	<0.10	<5.9	NA	<0.001	<0.001	< 0.001	< 0.001	<0.0008	
SB-2	3.5-4.0	08/12/99	< 0.09	<5.6	NA	< 0.0009	< 0.0009	< 0.0009	< 0.0009	< 0.001	
SB-2	6.5-7.0	08/12/99	<0.10	<5.9	NA	< 0.001	<0.001	< 0.001	< 0.001	0.001	
SB-3	3.5-4.0	08/12/99	<0.10	<5.6	NA	< 0.001	<0.001	< 0.001	< 0.001	0.0007	
SB-3	6.5-7.0	08/12/99	6,160	<5.7	NA	11	190	100	460	0.0018	MTBE: 0.073
SB-4	3.5-4.0	08/12/99	<0.10	<5.5	NA	< 0.001	< 0.001	< 0.001	< 0.001	< 0.0007	
SB-4	6.5-7.0	08/12/99	1	94	NA	0.082	0.0085	0.0073	0.013	0.001	
SB-5	3.5-4.0	08/12/99	< 0.09	<5.5	NA	< 0.0009	< 0.0009	< 0.0009	< 0.0009	0.0006	
SB-5	6.5-7.0	08/12/99	<0.08	<5.9	NA	<0.0008	<0.0008	<0.0008	<0.0008	0.0009	
SB-6	3.5-4.0	08/13/99	<0.10	<5.5	NA	< 0.001	<0.001	< 0.001	< 0.001	<0.0008	
SB-6	6.5-7.0	08/13/99	10,100	1,100	NA	76	490	170	990	0.43	
SB-7	3.5-4.0	08/12/99	<0.10	<5.4	NA	<0.001	<0.001	<0.001	<0.001	<0.0008	
SB-7	6.5-7.0	08/12/99	<0.11	<5.8	NA	<0.0011	<0.0011	<0.0011	<0.0011	<0.0009	
SB-8	3.5-4.0	08/12/99	<0.10	<5.6	NA NA	<0.001	<0.001	<0.001	<0.001	<0.0007	1
SB-8	6.5-7.0	08/12/99	13	<5.8	NA NA	0.43	0.36	0.12	0.83	0.0012	MTBE: 0.022
SB-9	3.5-4.0	08/13/99	<0.09	<5.6	NA	<0.0009	<0.0009	<0.0009	<0.0009	<0.001	
SB-9	6.5-7.0	08/13/99	<0.61	<5.8	NA NA	0.024	<0.0061	<0.0061	<0.0061	<0.0011	
SB-10	3.5-4.0	08/13/99	<0.09	<5.6	NA NA	<0.0009	<0.0009	<0.0009	<0.0009	<0.0008	
SB-10	6.5-7.0	08/13/99	<0.13	<6.4	NA.	<0.0013	<0.0013	< 0.0013	<0.0013	<0.001	
SB-11	3.5-4.0	08/13/99	<0.20	<5.5	NA NA	<0.002	<0.002	<0.002	<0.002	<0.0011	
SB-11	6.5-7.0	08/13/99	<0.11	<5.7	NA NA	<0.002	<0.0011	<0.0011	<0.0011	<0.0011	
SB-12	3.5-4.0	08/12/99	<0.10	<5.5	NA NA	<0.001	<0.001	<0.001	<0.001	<0.0006	
SB-12	4.5-5.0	08/12/99	496	2,900	NA	0.07	0.032	4	6.7	<0.0009	Chlorobenzene: 0.0017 1,2-DCB: 3.1 1,3-DCB: 0.038 1,4-DCB: 0.33 MTBE:
SB-12	6.5-7.0	08/12/99	2	60		< 0.001	<0.001	0.023	0.0098	< 0.0011	MTBE: 0.001
SB-13	3.5-4.0	08/13/99	1	390	NA	<0.0012	0.002	0.0027	0.0027	0.0025	
SB-13	6.5-7.0	08/13/99	12	65	NA	0.25	0.048	0.15	0.49	0.0014	
SB-14	3.5-4.0	08/12/99	<0.08	<5.5	NA	<0.0008	<0.0008	<0.0008	<0.0008	<0.0008	MTBE: 0.084
SB-14	6.5-7.0	08/12/99	29	450	NA	0.56	0.29	0.33	1.7	0.0097	
SB-15	3.5-4.0	08/12/99	<0.51	140	NA	< 0.0054	< 0.0054	< 0.0054	<0.0054	<0.0091	
SB-15	6.5-7.0	08/12/99	<0.57	81	NA	<0.0061	0.012	<0.0061	0.0085	<0.0098	
SB-16	6-6.5	05/19/08	<0.22	30	<50	<0.0043	< 0.0043	< 0.0043	<0.0087	< 0.0043	
SB-17	8-8.5	05/22/08	2,500	3,600	2,900	30	130	27	120	ND	
SB-17	10-10.5	05/22/08	12.000	17.000	13.000	140	580	120	620	<8.3	
SB-17	15-15.5	05/22/08	64	1,400	1,300	<0.89	< 0.89	<0.89	<1.8	<0.89	
SB-17	20-20.5	05/22/08	<0.21	<0.99	<49	< 0.0042	< 0.0042	< 0.0042	<0.0084	< 0.0042	
SB-18	8-8.5	05/21/08	1,900	67	<49	41	110	28	130	<19	
SB-19	8-8.5	05/21/08	<0.25	< 0.99	<49	< 0.0050	< 0.0050	< 0.0050	<0.010	< 0.0050	
SB-20/ PCB-7	8-8.5	05/22/08	5,600	390	51	86	280	54	280	<8.3	
SB-21/ PCB-8	8-8.5	05/21/08	3,800	2,500	<49	40	210	69	360	<19	
SB-22	8-8.5	05/21/08	3,200	1,100	<500	<47	140	<47	190	<47	
SB-23	11.5-12	05/22/08	<0.21	1.2	<49	< 0.0041	<0.0041	< 0.0041	<0.0082	<0.0041	
SB-24/ PCB-1	9-9.5	05/20/08	<0.19	1.6	<50	< 0.0039	< 0.0039	< 0.0039	<0.0078	< 0.0039	
SB-25/ PCB-2	8-8.5	05/20/08	<0.19	1.1	<50	< 0.0037	< 0.0037	< 0.0037	< 0.0075	< 0.0037	
SB-26	8.5-9	05/21/08	< 0.23	10	<50	< 0.0047	< 0.0047	< 0.0047	< 0.0093	< 0.0047	
SB-27/ PCB-3	8.5-9	05/20/08	<0.27	< 0.99	<49	< 0.0054	<0.0054	< 0.0054	<0.011	< 0.0054	
SB-20/ PCB-7 Dup	8-8.5	05/22/08	4,900	610	<250	99	300	64	340	<21	
SB-25/ PCB-2 Dup	8-8.5	05/20/08	NA	<1.0	<50	NA	NA	NA	NA	NA	

NA = Not Analyzed
EPA Method 8260 for BTEX and 1,2-DCA analyses of soil
EPA Mentod 8015m for TPH-g, TPH-d, and TPM-mo analyses of soil

Table 3: Historical Groundwater Sample Results (1993 - 2008)

Well Number	Date Sampled	Benzene µg/L	Toluene μg/L	Ethyl- Benzene µg/L	Xylenes μg/L	TPH-G μg/L	TPH-D μg/L	1,1- DCA µg/L	1,2- DCA µg/L	1,1,1- TCA µg/L	TCE µg/L	MTBE µg/L	Notes
MW-2	03/23/93	ND	ND	ND	ND	ND	ND						
	07/27/93 11/05/93	ND 	ND 	ND 	ND 	ND 	ND 						
	02/25/94	<1	<1	<1	<1	<100	<1,000						
	06/03/94	<0.5	<0.5	<0.5	<0.5	<50	<20,000						
	08/31/94	<0.3	<0.3	<0.3	<0.6	<500	<500						
	12/22/94 03/13/95	<0.5 0.8	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<50 <50	<50 <400						Non-diesel peak reported.
	06/09/95	<0.5	<0.5	<0.5	<0.5	<100	<50						
	09/21/95	0.7	<0.5	<0.5	<0.5	<50	<50						
	12/12/95	<0.5	<0.5	<0.5	<1.0	<100	<50						
	03/12/96 06/21/96	<0.5	<0.5 	<0.5	<0.5	<100 	<50 						
	08/29/96	<0.5	<0.5	<0.5	<0.5	<50	<150						
	01/16/97	<0.5	<0.5	<0.5	<0.5	<50	<150	0.7	<0.5	<0.5	<0.5		
	07/07/97	<0.5	<0.5	<0.5	<0.5	<50	<150					<0.5	
	01/27/98 07/22/98	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	100 <50	<150 					<0.5 <0.5	
	07/22/98	<0.5	<0.5	<0.5	<0.5	<50 <50	<200	<0.5	<0.5	<0.5	<0.5	<0.5	
MW-3		35	2.9			300	ND						
C-AA IAI	03/23/93 07/27/93	35 97	2.9	2 4	3.2 1.1	220	ND ND						
	11/05/93	4.9	ND	ND	1.2	170	ND						
	02/25/94	42	<1	<1	<1	100	<1,000						
	06/03/94	120	8.2	8.4	4.5	320	<20,000						
	08/31/94 12/22/94	83 1,460	1.1 18	5.3 100	2.9 50	<500 3,800	<500 270						
	03/13/95	3,600	260	270	280	14,000	1,700						
	06/09/95	4,700	58	140	71	3,700	120						
	09/21/95	9,800	58	600	95	14,000	300						
	12/12/95 03/12/96	330 350	2.1 4.6	47 23	5.3 8.7	700 600	<50 <50						
	06/21/96	940	76	98	57	1,900	<50						
	08/29/96	420	29	44	28	900	<150						
	01/16/97	1,600	270	120	194	3,600	700	<0.5	9.2	<0.5	<0.5		
	04/15/97	1,300	300	180	160	4,300	800	<0.5	16	<0.5	1.1	6.9	
	07/07/97 10/27/97	100 1,030	84 60	100 54	67 40	1,900 2,200	350 	 <0.5	2.4	<0.5	 <0.5	3.8 3.1	
	01/27/98	1,070	98	73	69	3,200						3.9	
	04/22/98	610	56	49	54	1,800		<0.5	3.0	<0.5	<0.5	1.1	
	07/22/98	1,800	230	160	180	3,600	370					5.0	
	10/21/98 07/23/99	78 1,500	1.0 140	3.8 76.0	0.6 260	110 4,000	<250 790	<0.5 <0.5	0.6 1.0	<0.5 <0.5	<0.5 <0.5	<0.5 5.60	
	10/28/99	1,100	43	58	102	3,000	600	<0.5	0.9		<0.5		
	02/10/00	690	22	36	49	1,400	520	<0.5	<0.5	<0.5	<0.5	2.20	
	04/27/00	1,100	140	73	163	2,400	250	<0.5	0.6	<0.5	<0.5	<0.5	
	08/03/00 10/23/00	520 2,000	7.7 16	21 22	27 46	1,100 3,800	750 760	<0.5 <0.5	0.6 0.7	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	
	01/31/01	360	8.6	14	28	860	300	<0.5	0.6	<0.5	<0.5	<0.5	
	04/26/01	808	60.6	46.8	115	1,530	280	<0.5	8.0	<0.5	<0.5	<0.5	
	07/30/01	788	23.3	44.6	80.7	1,400	350	<0.5	0.6	<0.5	<0.5	<0.5	
	10/29/01 01/29/02	852 1,250	14.3 85.3	24.5 64.7	38.6 95.7	1,730 4,240	500 490	<0.5 <0.5	0.5 1.4	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	
	04/29/02	1,120	51.5	84.4	117	5,710	700	<0.5	1.1	<0.5	<0.5	<0.5	
MW-5	02/05/99	<0.5	<0.5	<0.5	<0.5	<50	<150	<0.5	<0.5	<0.5	<0.5	<0.5	
MW-6	03/23/93	ND	ND	ND	ND	ND	ND						
	07/27/93	ND	ND	ND	ND	ND	ND						
	11/05/93	ND	ND	ND	ND	ND	ND						
	02/25/94 06/03/94	<1 2.7	<1 <0.5	<1 <0.5	3.5 <0.5	<100 69	<1,000 <20,000						
	08/31/94	<0.3	8.7	1.6	3.5	<500	<500						
	12/22/94	<0.5	<0.5	<0.5	<0.5	<50	<50						Non-diesel peak reported.
	03/13/95	1.2	<0.5	<0.5	<0.5	<50 <100	<400						
	06/09/95 09/21/95	0.6 <0.5	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<100 <50	<50 <50						
	12/12/95	<0.5	<0.5	<0.5	<1.0	<100	<50						
	03/12/96	<0.5	<0.5	<0.5	<0.5	<100	<50						
	06/21/96												
	08/29/96 01/16/97	<0.5 5.5	<0.5 16	<0.5 2.9	<0.5 16	<50 140	<150 220	 <0.5	6.3	<0.5	<0.5		
	07/07/97	5.5 <0.5	<0.5	<0.5	<0.5	<50	<150	<0.5 		<0.5 	<0.5 	<0.5	
	07/22/98	<0.5	<0.5	<0.5	<0.5	<50	<250					<0.5	
	10/24/00	<0.5	<0.5	<0.5	<0.5	<50	<250	<0.5	7.7	<0.5	<0.5	<0.5	
	01/31/01	<0.5	<0.5	<0.5	<0.5	<50	<250	<0.5	6.9	<0.5	<0.5	<0.5	

Table 3: Historical Groundwater Sample Results (1993 - 2008)

Well Number	Date Sampled	Benzene µg/L	Toluene μg/L	Ethyl- Benzene µg/L	Xylenes μg/L	TPH-G μg/L	TPH-D μg/L	1,1- DCA µg/L	1,2- DCA µg/L	1,1,1- ΤCA μg/L	TCE µg/L	MTBE μg/L	Notes
MW-6	04/27/01	<0.5	<0.5	<0.5	<0.5	<200	<250	<0.5	6.6	<0.5	<0.5	<0.5	
(cont.)	07/30/01	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<0.5	<200 <200	<250 <500	<0.5 <0.5	9.2 10	<0.5	<0.5 <0.5	<0.5	
	10/30/01 01/29/02	0.54	<0.5	<0.5 <0.5	<1.0 <1.0	<200	<500 <250	<0.5	10	<0.5 <0.5	<0.5	<0.5 <0.5	
	04/30/02	<0.5	<0.5	<0.5	<1.0	<200	<250	<0.5	14	<0.5	<0.5	<0.5	
MW-11	02/05/99	<0.5	<0.5	<0.5	<0.5	<50	<150					<0.5	
MW-12	02/05/99	<0.5	<0.5	<0.5	<0.5	<50	<150					<0.5	
MW-13	02/05/99	<0.5	<0.5	<0.5	<0.5	<50	<150		-		-	<0.5	
MW-15	02/05/99	<0.5	<0.5	<0.5	<0.5	<50	430	<0.5	<0.5	<0.5	<0.5	<0.5	
	07/22/99	<0.5	<0.5	<0.5	<0.5	<50	<200	<0.5	<0.5	<0.5	<0.5	<0.5	
MW-25	03/23/93	ND ND	ND ND	ND	ND	ND ND	ND ND		-				
	07/27/93 11/05/93	4.2	4.4	ND 2.5	ND 20	170	ND ND						
	02/25/94	2.1	<1	<1	<1	<100	<1,000						
	06/03/94	2.4	14	<0.5	3.4	97	<20,000						
	08/31/94	0.5	<0.3	<0.3	<0.6	<500	<500						
	12/22/94	0.5	<0.5	<0.5	<0.5	<50	<50						Non-diesel peak reported.
	03/13/95	0.58	<0.5	<0.5	<0.5	150	950						
	06/09/95	0.8	<0.5	<0.5	<0.5	<100	60						
	09/21/95	<0.5	<0.5	<0.5	<0.5	50	<50						
	12/12/95	<0.5	<0.5	<0.5	<1.0	<100	<50						
	03/12/96	<0.5	<0.5	<0.5	<0.5	120	<50						
	06/21/96							-					
	08/29/96	<0.5	<0.5	<0.5	< 0.5	90	<150						
	01/16/97 07/07/97	0.6	<0.5	<0.5	<0.5 <0.5	80	<150	25	41	<0.5	<0.5		
	01/07/97	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	140 <100	<150 					11 10	
	07/22/98	<0.5	<0.5	<0.5	<0.5	<50	<250					24	
	02/05/99	<0.5	<0.5	<0.5	<0.5	<50	340	28	59	<0.5	<0.5	28	1,1-DCE detected, 0.9 µg/L.
	04/07/99	<0.5	<0.5	<0.5	<0.5	<50	<250	27	72	<0.5	<0.5	27	1,1-DCE detected, 1.6 µg/L.
	07/23/99	1.80	<0.5	<0.5	<0.5	<50	<200	30	58	<0.5	<0.5	23.0	,,
	10/27/99	< 0.5	1.4	<0.5	1.0	<100	<200	35	47		<0.5		
	02/08/00	<0.5	<0.5	<0.5	<0.5	100	<250	39	41	<0.5	<0.5	29.0	1,1-Dichloroethene detected at 3.1 µg/L.
	04/26/00	<0.5	<0.5	<0.5	<0.5	<100	<250	51	38	<0.5	<0.5	18	1,1-Dichloroethene detected at 4.2 µg/L.
	08/03/00	<0.5	<0.5	<0.5	<0.5	<50	<250	40	57	<0.5	<0.5	27	1,1-Dichloroethene detected at 2.6 μg/L.
	10/23/00	<0.5	<0.5	<0.5	<0.5	<50	<250	54	68	<0.5	<0.5	38	1,1-Dichloroethene detected at 3.5 μg/L.
	01/31/01	<0.5	<0.5	<0.5	<0.5	90	<250	52	46	<0.5	<0.5	22	1,1-Dichloroethene detected at 6.5 µg/L.
	04/26/01 07/30/01	<0.5 <0.5	0.62 <0.5	<0.5 <0.5	<0.5 <0.5	<200 <200	<250 <250	49 33	37 36	<0.5 <0.5	<0.5 <0.5	15.8 10.9	1,1-Dichloroethene detected at 6.0 µg/L. Chloromethane detected at 0.8 µg/L;
	10/29/01	<0.5	<0.5	<0.5	<1.0	<200	<500	22	38	<0.5	<0.5	10.5	1,1-Dichloroethene detected at 4.6 μg/L. Chloromethane detected at 0.5 μg/L;
	01/28/02	<0.5	<0.5	<0.5	<1.0	<200	<250	25	56	<0.5	<0.5	8.90	1,1-Dichloroethene detected at 1.8 µg/L. 1,1-Dichloroethene detected at 2.8 µg/L.
	04/29/02	<0.5	<0.5	<0.5	<1.0	<200	<250	14	44	<0.5	<0.5	6.92	1,1-Dichloroethene detected at 1.7 μg/L; 1,1,2,2-Tetrachloroethane detected at 0.5 μg/L.
	10/22/02	7.64	248	133	843	4,790	1,240	9.6	34	<0.5	<0.5	1,410	1,1-Dichloroethene detected at 0.9 µg/L.
	11/15/02 05/06/03	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<1.0 <1.0	<200 <200	<250 <250	11 8.5	35 34	<0.5 <0.5	<0.5 <0.5	7.3 5.7	Chloroethane detected at 22 µg/L.
	10/14/03	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<1.0 <1.0	<200 <200	<250 <250	7.6	34 27	<0.5	<0.5 <0.5	5.7 6.3	1,1-Dichloroethene detected at 0.8 μg/L.
	04/27/04	<0.5	<0.5	<0.5	<1.0	<200	<250	5.1	18	<0.5	<0.5	5.2	
	11/17/04	<0.50	<0.50	<0.50	<0.50	<50	190	6.7	25	<0.50	<0.50	6.1	1,1-Dichloroethene detected at 0.51 µg/L.
MW-26	03/23/93	180	190	55	330	7,000	1,300	ND	ND	ND	ND		
	07/27/93	470	96	30	80	1,800	ND	ND	140	ND	ND		
	11/05/93	4,700	1,300	9	1,400	19,000	ND	ND	120	ND	ND		
	02/25/94	4,800	570	200	860	14,000	<1,000	<1	28	<1	<1		
	06/03/94	4,100	300	120	230	12,000	<20,000	1.7	140	<0.5	<0.5		Bromodichloromethane detected, 0.84 µg/L.
	08/31/94	4,100	360	170	450	93,000	1,400	<4.0	<4.0	<4.0	<4.0		0 -45
	12/22/94	1,030	170	85	290	5,000	560	<2.0	<2.0	<2.0	<2.0		8 other volatiles detected by 8260.
	03/13/95 06/09/95	320 14,000	19 64	23 31	66 230	3,000 10,800	810 310	53 240	5.8 3.1	<0.5 1	<0.5 <0.5		
	09/21/95	1,900	160	160	330	8,000	200	1.3	120	<0.5	<0.5		
	12/12/95	13,000	38	36	120	25,000	0.6	1.4	180	<0.5	<0.5		No diesel pattern detected; result due to high gasoline concentration.
	03/12/96	9,000	33	30	65	4,400	<50	<0.5	180	<0.5	<0.5		
	06/21/96	14,000	27	16	66	5,400	<50	3.2	170	<0.5	<0.5		
	08/29/96	8,500	26	28	74	19,000	<150	<0.5	160	<0.5	<0.5		
	01/16/97	6,500	21	31	47	4,600		4.3	>50	<0.5	<0.5	26	
	04/15/97	16,000	33	40	160	26,000	2,200	3.5	97	<0.5	2.4	40	cis-1,2-DCE detected, 0.7 μg/L.
	07/07/97	22,000	44	170	200	28,000	1,100	<5.0	<5.0	<5.0	<5.0	95	
	10/27/97	16,000	26 <5.0	100	37	30,000	420	3.6	92 100	<0.5	<0.5	38	
	01/27/98 04/22/98	23,600 5,000	<5.0 4.3	<5.0 9.2	<5.0 16	26,000 14,000	420	8.3 13	100	<0.5 <0.5	<0.5 <0.5	100 27	
	04/22/98	5,000	4.3	9.2	16	14,000		13	130	<0.5	<0.5	21	

Table 3: Historical Groundwater Sample Results (1993 - 2008)

Well Number	Date Sampled	Benzene µg/L	Toluene μg/L	Ethyl- Benzene µg/L	Xylenes μg/L	TPH-G μg/L	TPH-D μg/L	1,1- DCA µg/L	1,2- DCA µg/L	1,1,1- ΤCA μg/L	TCE µg/L	MTBE μg/L	Notes
MW-26	07/22/98	3,800	5.7	6.9	11	5,200	750	10	110		<1.0	33	
(cont.)	10/21/98	420 20	<0.5 <0.5	2.1 0.60	2.7 0.80	820 230	<250 230	24 10	82 51	<0.5 <0.5	<0.5 <0.5	31 29	
	02/05/99 04/07/99	<0.5	<0.5	<0.5	<0.5	80	<250 <250	15	54	<0.5	<0.5	25	
	07/23/99	7.10	<0.5	<0.5	0.80	180	<200	12	32	<0.5	<0.5	12.0	
	10/27/99	14	1.4	2.9	7.8	400	<200	13	30		<0.5		
	02/08/00	<0.5	<0.5	<0.5	<0.5	80	<250	13	32	<0.5	<0.5	28.0	
	04/26/00	0.7	<0.5	0.6	<0.5	200	340	7.5	39	<0.5	<0.5	22	
	08/03/00 10/23/00	6.8 10	<0.5 0.8	0.6 1.7	1.4 1.7	<50 80	<250 <250	7.4 5.1	19 37	<0.5 <0.5	<0.5 <0.5	19 26	
	01/31/01	26	0.70	2.4	2.2	390	320	5.7	51	<0.5	<0.5	33	
	04/26/01	10.6	<0.5	0.70	1.04	400	350	16	39	<0.5	<0.5	28.5	
	07/30/01	107	<0.5	1.42	1.06	1,920	380	22	44	<0.5	<0.5	31.4	
	10/29/01	31.6	<0.5	<0.5	<1.0	2,020	500	26	25	<0.5	<0.5	27	
	01/28/02 04/29/02	30.0	<0.5 <0.5	0.70	<1.0	450	380 550	43 50	<0.5 23	<0.5 <0.5	<0.5 <0.5	14.5 8.62	1,1-Dichloroethene detected at 1.8 µg/L.
	10/22/02	394 1,440	25.7	<0.5 6.60	<1.0 20.4	1,870 4,440	890	53	26	<0.5	<0.5	168	1,1-Dichloroethene detected at 2.5 µg/L. 1,1-Dichloroethene detected at 3.7 µg/L.
	11/15/02	1,630	0.56	3.22	3.86	5,590	780	18	33	<0.5	<0.5	49.2	1,1-dichloroethene detected at 3.7 µg/L.
	05/06/03	1,250	<0.5	2.42	<1.0	3,730	380	46	24	<0.5	<0.5	13.1	1,1-Dichloroethene detected at 3.1 µg/L.
	10/14/03	51	<0.5	1.38	<1.0	3,100	<250	83	28	<0.5	<0.5	23.8	1,1-Dichloroethene detected at 3.3 µg/L.
	04/27/04	467	<0.5	1.24	<1.0	1,380	<250	82	33	<0.5	<0.5	<0.5	1,1-Dichloroethene detected at 5.2 μg/L.
	11/17/04	120	<1.0	2.50	1.3	740	820	31	44	<0.50	<0.50	120	1,1-Dichloroethene detected at 1.1 μg/L.
MW-27	06/21/96	<0.5	<0.5	<0.5	<0.5	<50	<50	<0.5	6.8	<0.5	<0.5		
	08/29/96 01/16/97	 12	 5.0	 <0.5	 2.6	 70	 <150	 <0.5	5.7	<0.5	 <0.5		
	07/22/98	<0.5	<0.5	<0.5	<0.5	<50	<250	<1.0	1.4	<0.5	<1.0	<0.5	
	02/05/99	<0.5	<0.5	<0.5	<0.5	<50	<150	<0.5	0.7	<0.5	<0.5	<0.5	
	07/23/99	<0.5	<0.5	<0.5	<0.5	<50	<200	<0.5	0.7	<0.5	<0.5	<0.5	
	10/27/99	<0.5	<0.5	<0.5	<0.5	<100	<200	<0.5	<0.5		<0.5		
	02/08/00	<0.5	<0.5	<0.5	<0.5	<50	<250	<0.5	<0.5	<0.5	<0.5	<0.5	
	04/27/00 08/16/00	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<100 <50	250	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	
	10/23/00	<0.5	<0.5	<0.5	<0.5	<50 <50	<250	<0.5	<0.5	<0.5	<0.5	<0.5	
	01/31/01	<0.5	<0.5	<0.5	<0.5	<50	<250	<0.5	<0.5	<0.5	<0.5	<0.5	
	04/26/01	<0.5	<0.5	<0.5	<0.5	<200	<250	<0.5	<0.5	<0.5	<0.5	<0.5	
	07/30/01	<0.5	<0.5	<0.5	<0.5	<200	<250	<0.5	<0.5	<0.5	<0.5	<0.5	
	10/29/01	<0.5	<0.5	<0.5	<1.0	<200	<500	<0.5	<0.5	<0.5	<0.5	<0.5	
	01/28/02 04/29/02	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<1.0 <1.0	<200 <200	<250 <250	<0.5 <0.5	0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	
	10/22/02	8.56	56.2	9.37	59.3	650	600	<0.5	<0.5	<0.5	<0.5	331	
	11/15/02	<0.5	<0.5	<0.5	<1.0	<200	<250	<0.5	<0.5	<0.5	<0.5	<0.5	
	05/06/03	<0.5	<0.5	<0.5	<1.0	<200	<250	<0.5	<0.5	<0.5	<0.5	<0.5	
	10/14/03	<0.5	<0.5	<0.5	<1.0	<200	<250	<0.5	< 0.5	<0.5	<0.5	<0.5	
	04/27/04 11/17/04	<0.5 <0.50	<0.5 <0.50	<0.5 <0.50	<1.0 <0.50	<200 <50	<250 64	<0.5 < 0.50	<0.5 < 0.50	<0.5 < 0.50	<0.5 < 0.50	<0.5 <5.0	
MIN 20													
MW-28	03/23/93 07/27/93	ND ND	ND ND	ND ND	ND ND	110 ND	ND ND						
	11/05/93	ND	ND	ND	2.1	ND	ND						
	02/25/94	<1	<1	<1	<1	<100	<1						
	06/03/94	3.1	<0.5	<0.5	<0.5	<50	<20,000						
	08/31/94	1.4	<0.3	<0.3	<0.6	<500	<500						
	12/22/94 03/13/95	<0.5 0.91	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<50 <50	<50 <400						Non-diesel peak reported.
	06/09/95	<0.5	<0.5	<0.5	<0.5	<100	<50						
	09/21/95	<0.5	<0.5	<0.5	<0.5	<50	<50						
	12/12/95	<0.5	<0.5	<0.5	<1.0	<100	<50						
	03/12/96	<0.5	<0.5	<0.5	< 0.5	<100	<50						
	06/21/96 08/29/96	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<100 <50	<50 <150						
	08/29/96	<0.5 18	<0.5 20	<0.5 2.2	<0.5 13	<50 220	<150 <150	5.1	85	<0.5	<0.5	8.2	
	04/15/97	<0.5	<0.5	<0.5	<0.5	120	<150	1.1	150	<0.5	<0.5	7.1	
	07/07/97	<0.5	<0.5	<0.5	<0.5	110	<150	<5.0	170	<5.0	<5.0	7.2	
	10/27/97	3.6	<0.5	<0.5	<0.5	300		6.2	120	<0.5	<0.5	36	
	01/27/98	7.6	<0.5	<0.5	<0.5	500 <50	<150	1.0			 -0.5	56	
	04/22/98 07/22/98	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<50 <50		1.0 <1.0	89 85	<0.5	<0.5 <1.0	8.6 18	
	10/21/98	<0.5	<0.5	<0.5 <0.5	<0.5 <0.5	<50 <50	<250	0.5	80	<0.5	<0.5	12	
	02/05/99	<0.5	<0.5	<0.5	<0.5	<50	<150	32	29	<0.5	<0.5	5.0	1,1-DCE detected, 0.9 µg/L.
	04/07/99	<0.5	<0.5	<0.5	<0.5	<50	<250	<0.5	62	<0.5	<0.5	4.5	
	07/23/99	<0.5	<0.5	<0.5	<0.5	<50	<200	<0.5	50	<0.5	<0.5	1.80	
	10/27/99	0.7	 -0.5	 -0 F	 -0.5	 -100	<200	 <0.5	 32		 <0.5		
	11/02/99 02/08/00	0.7 <0.5	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<100 <50	 <250	<0.5 <0.5	32	<0.5	<0.5 <0.5	4.30	
	04/26/00	<0.5	<0.5	<0.5	<0.5	<100	<250	<0.5	50	<0.5	<0.5	1.5	
	08/03/00	<0.5	<0.5	<0.5	<0.5	<50	<250	<0.5	47	<0.5	<0.5	3.7	

Table 3: Historical Groundwater Sample Results (1993 - 2008)

Well Number	Date	Benzene	Toluene	Ethyl- Benzene	Xylenes	TPH-G	TPH-D	1,1- DCA	1,2- DCA	1,1,1- TCA	TCE	MTBE	Notes
	Sampled	μg/L	μg/L	μg/L	μg/L	μg/L	μg/L	μg/L	μg/L	μg/L	μg/L	μg/L	
MW-28	10/23/00	<0.5	<0.5	<0.5	<0.5	<50	<250	<0.5	57	<0.5	<0.5	4.7	
(cont.)	01/31/01	<0.5	<0.5	<0.5	<0.5	<50	<250	<0.5	46	<0.5	<0.5	4.4	
	04/26/01	<0.5	<0.5 <0.5	<0.5	<0.5	<200 <200	<250	<0.5 <0.5	26 38	<0.5	<0.5 <0.5	1.98	Obligation of the standard of 2.2 mg/l
	07/30/01 10/29/01	0.5 <0.5	<0.5	0.64 <0.5	2.58 <1.0	<200 <200	<250 <500	<0.5	29	<0.5 <0.5	<0.5	3.0 3.74	Chloromethane detected at 3.3 µg/L.
	01/28/02	6.20	<0.5	<0.5	<1.0	<200	<250	2.8	50	<0.5	<0.5	6.00	
	04/29/02	1.64	<0.5	<0.5	<1.0	<200	<250	3.7	44	<0.5	<0.5	4.81	
	10/22/02	25.0	<0.5	<0.5	<1.0	750	<250	2.0	59	<0.5	<0.5	<0.5	
	11/15/02	13.4	<0.5	1.29	<1.0	610	<250	1.3	54	<0.5	<0.5	<0.5	Chloromethane detected at 1.0 μg/L.
	05/06/03 10/14/03	3.1 <0.5	<0.5 <0.5	<0.5 <0.5	<1.0 <1.0	390 <200	<250 <250	0.8 <0.5	70 38	<0.5 <0.5	<0.5 <0.5	9.29 6.44	Chloroethane detected at 0.8 µg/L.
	04/27/04	<0.5	<0.5	<0.5	<1.0	<200	<250	<0.5	< 0.5	<0.5	<0.5	9.29	
	11/17/04	<0.50	<0.50	<0.50	<0.50	<50	<50	<0.50	4.7	<0.50	<0.50	<5.0	
MW-29	03/23/93	ND	ND	ND	ND	ND	ND						
	07/27/93	ND	ND	ND	ND	ND	ND						
	11/05/93	ND	ND	2.1	11	ND	ND						
	02/25/94	<1	<1	<1	<1	<100	<1,000						
	06/03/94	<0.5	<0.5	<0.5	<0.5	<50	<20,000						
	08/31/94 12/22/94	<0.3 <0.5	<0.3 <0.5	<0.3 <0.5	<0.6 <0.5	<500 <50	<500 <50						Non-diesel peak reported.
	03/13/95	0.59	<0.5	<0.5	<0.5	<50	<400						Non dieser peak reported.
	06/09/95	<0.5	<0.5	<0.5	<0.5	<100	<50						
	09/21/95	<0.5	<0.5	<0.5	<0.5	<50	<50						
	12/12/95	< 0.5	<0.5	<0.5	<1.0	<100	<50						
	03/12/96 06/21/96	<0.5 	<0.5 	<0.5 	<1.0 	<100 	<50 						
	08/29/96	<0.5	<0.5	<0.5	<0.5	<50	<150						
	01/16/97	6.6	8.9	0.6	9.3	120	<150	47	24	<0.5	<0.5	1.8	
	07/07/97	<0.5	<0.5	<0.5	<0.5	<50	<150	52	21	<5.0	<5.0	1.2	
	01/27/98	<0.5	<0.5	<0.5	<0.5	100	<150					8.0	
	07/22/98	<0.5	<0.5	<0.5	<0.5	<50	<250	12	29	 -0.5	<1.0	7.8	
	02/05/99 04/07/99	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<50 <50	<150 <250	<0.5 30	68 38	<0.5 <0.5	<0.5 <0.5	8.5 4.9	1,1-DCE detected, 1.4 µg/L.
	07/23/99	<0.5	<0.5	<0.5	<0.5	<50	<200	44	33	<0.5	1.9	4.70	1,1-Dichloroethene detected at 2.3 µg/L; cis-1,2-Dichloroethene detected at 2.3 µg/L.
	10/27/99	<0.5	<0.5	<0.5	<0.5	<100	<200	36	23		<0.5		
	02/08/00	< 0.5	<0.5	<0.5	<0.5	<50	<250	87	25	<0.5	<0.5	18.0	1,1-Dichloroethene detected at 9.6 µg/L.
	04/26/00 08/16/00	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<100 <50	<250 	61 49	38 21	<0.5 <0.5	<0.5 <0.5	12 17	1,1-Dichloroethene detected at 5.2 µg/L. 1,1-Dichloroethene detected at 6.0 µg/L.
	10/23/00	<0.5	<0.5	<0.5	<0.5	<50	<250	94	40	<0.5	<0.5	34	1,1-Dichloroethene detected at 0.0 µg/L.
	01/31/01	<0.5	<0.5	<0.5	<0.5	60	<250	100	35	<0.5	<0.5	26	1,1-Dichloroethene detected at 13 µg/L.
	04/26/01	<0.5	<0.5	<0.5	<0.5	<200	270	87	38	<0.5	<0.5	39.1	1,1-Dichloroethene detected at 12 μg/L.
	07/30/01	1.25	1.28	1.1	5.99	220	<250	120	42	<0.5	<0.5	42.3	1,1-Dichloroethene detected at 13 µg/L.
	10/29/01 01/28/02	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<1.0 <1.0	<200 <200	<500 <250	120 120	34 44	<0.5 <0.5	<0.5 <0.5	28.0 28.9	1,1-Dichloroethene detected at 14 µg/L. 1,1-Dichloroethene detected at 26 µg/L.
	04/29/02	4.95	<0.5	<0.5	<1.0	<200	<250	130	29	<0.5	<0.5	20.9	1,1-Dichloroethene detected at 23 µg/L.
	10/22/02	<0.5	<0.5	<0.5	<1.0	<200	<250	140	26	<0.5	<0.5	18.1	1,1-Dichloroethene detected at 19 µg/L.
	11/15/02	<0.5	<0.5	<0.5	<1.0	<200	<250	120	26	<0.5	<0.5	13.9	1,1-dichloroethene detected at 15 μg/L.
	05/06/03	<0.5	<0.5	<0.5	<1.0	<200	<250	140	31	<0.5	<0.5	13.1	1,1-Dichloroethene detected at 24 μg/L.
	10/14/03	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<1.0	<200 <200	<250 <250	110 160	22 28	<0.5 <0.5	<0.5 <0.5	11.9	Chloromethane detected at 0.9 µg/L.
	04/27/04 11/17/04	<1.0	<1.0	<1.0	<1.0 <1.0	120	< 50	33	6.5	<0.50	<0.50	15.3 120	1,1-Dichloroethene detected at 31 µg/L. 1,1-Dichloroethene detected at 5.5 µg/L.
MW-30	03/23/93	ND	ND	ND	ND	ND	ND						
ĺ	07/27/93	ND	ND	ND	ND	ND	ND						
ĺ	11/05/93 02/25/94	ND 13	ND <1	ND <1	2.8 <1	ND <100	ND <1.000						
	02/25/94 06/03/94	1.3 1.1	<1 <0.5	<1 <0.5	<1 <0.5	<100 <50	<1,000 <20,000						
	08/31/94	0.8	<0.3	<0.3	<0.6	<500	<500						
	12/22/94	0.6	<0.5	<0.5	<0.5	<50	<50						Non-diesel peak reported.
	03/13/95	0.98	<0.5	<0.5	<0.5	<50	<400						
	06/09/95	<0.5	<0.5	<0.5	<0.5	<100	<50						
	09/21/95 12/12/95	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<0.5 <1.0	<50 <100	<50 <50						
ĺ	03/12/96	<0.5 <0.5	<0.5	<0.5	<0.5	<100	<50 <50						
	06/21/96												
ĺ	08/29/96	<0.5	<0.5	<0.5	<0.5	<50	<150						
ĺ	01/16/97	<0.5	<0.5	<0.5	0.6	80	<150	<0.5	<0.5	<0.5	0.9		
ĺ	07/07/97	<0.5	<0.5	<0.5	<0.5	<50	<150					<0.5	
ĺ	01/27/98 07/22/98	5.4 <0.5	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	100 <50						<0.5 <0.5	
	04/07/99	<0.5	<0.5	<0.5	<0.5	<50 <50	<250					<0.5	
	07/22/99	<0.5	<0.5	<0.5	<0.5	<50		<0.5	<0.5	<0.5	<0.5	<0.5	
	10/28/99	<0.5	<0.5	<0.5	<0.5	<100	<200	<0.5	<0.5		<0.5		
ĺ	02/08/00	<0.5	<0.5	<0.5	<0.5	<50	<250	<0.5	<0.5	<0.5	<0.5	<0.5	
	04/27/00	<0.5	<0.5	<0.5	<0.5	<100	250	<0.5	<0.5	<0.5	<0.5	<0.5	

Table 3: Historical Groundwater Sample Results (1993 - 2008)

Well Number	Date Sampled	Benzene µg/L	Toluene µg/L	Ethyl- Benzene	Xylenes μg/L	TPH-G μg/L	TPH-D μg/L	1,1- DCA	1,2- DCA	1,1,1- TCA	TCE µg/L	MTBE µg/L	Notes
	•			μg/L				μg/L	μg/L	μg/L			
MW-30	08/04/00	<0.5	<0.5	<0.5	<0.5	<50	<250	<0.5	<0.5	<0.5	<0.5	<0.5	
(cont.)	10/24/00	5.4	<0.5	<0.5 <0.5	<0.5	<50	<250	<0.5	<0.5	<0.5	<0.5	<0.5	
	01/31/01 04/27/01	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<50 <200	<250 <250	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	
	04/27/01	<0.5	<0.5	<0.5	<0.5	<200	<250	<0.5	<0.5	<0.5	<0.5	<0.5	
	10/29/01	<0.5	<0.5	<0.5	<1.0	<200	<500	<0.5	<0.5	<0.5	<0.5	<0.5	Chloroethane detected at 1.3 µg/L.
	01/29/02	<0.5	<0.5	<0.5	<1.0	<200	<250	<0.5	<0.5	<0.5	<0.5	<0.5	
	04/30/02	<0.5	<0.5	<0.5	<1.0	<200	<250	<0.5	<0.5	<0.5	<0.5	<0.5	
	10/22/02	<0.5	<0.5	<0.5	<1.0	<200	<250	<0.5	<0.5	<0.5	<0.5	<0.5	
	05/06/03	<0.5	<0.5	<0.5	<1.0	<200	<250	<0.5	<0.5	<0.5	<0.5	<0.5	
	10/14/03	<0.5	<0.5	<0.5	<1.0	<200	<250	<0.5	<0.5	<0.5	<0.5	<0.5	
	04/27/04	<0.5	<0.5	<0.5	<1.0	<200	<250	<0.5	<0.5	<0.5	<0.5	<0.5	
	11/17/04	<0.50	<0.50	<0.50	<0.50	<50	140	<0.50	<0.50	<0.50	<0.50	<5.0	
MW-32	03/23/93	391 ND	6.2 ND	3.1 ND	9 6	440 ND	ND	ND	60	ND	ND		
	07/27/93 11/05/93	ND 20	ND ND	ND 1.8	ND 2.1	ND 170	ND ND	ND ND	14 7.9	ND ND	ND ND		
	02/25/94	5.6	(1)	<1	<1	<100	<1,000	<1	7.9 <1	<1	<1		
	06/03/94	120	1.3	<0.5	1.4	350	<20,000	<0.5	11	<0.5	<0.5		
	08/31/94	39	0.5	2.2	1.2	<500	<500	<4.0	10	<4.0	<4.0		
	12/22/94	4.8	<0.5	<0.5	<0.5	<50	<50	<2.0	4.6	<2.0	<2.0		Non-diesel peak reported.
	03/13/95	220	3.6	6.5	5.8	1,100	<400	<0.5	16	<0.5	<0.5		
	06/09/95	1,500	7.9	43	14	2,200	180	0.7	<0.5	0.5	<0.5		
	09/21/95	1,200	2.4	72	4.5	2,300	60	<0.5	6.7	<0.5	1.4		
	12/12/95	230	<0.5	8.9	<1.0	500	<50	<0.5	28	<0.5	<0.5		
I	03/12/96 06/21/96	40	<0.5	1.7	<0.5	110 	<50 	<0.5	6.8	<0.5	<0.5		
	08/29/96	150	<0.5	49	<0.5	700	<150	<0.5	27	<0.5	<0.5		
	01/16/97	14	<0.5	1.9	<0.5	150	<150	<0.5	10	<0.5	0.7		cis-1,2-DCE detected, 0.8 µg/L.
	07/07/97	370	11	110	21	1,600	190					11	old 1,2 DOE detected, 0.0 pg/2.
	01/27/98	13	<0.5	1.0	<0.5	300		<0.5	7.5	<0.5	<0.5	2.5	
	07/22/98	700	55	88	66	2,300						14	
	07/22/99	59.0	0.80	1.80	<0.5	900	220	<0.5	5.9	<0.5	<0.5	8.70	
	10/28/99	95	2.5	2.1	1.6	500	<200	<0.5	12		<0.5		
	02/10/00	7.0	<0.5	<0.5	<0.5	120	<250	<0.5	4.3	<0.5	<0.5	1.10	
	04/27/00	240	7.0	12	18.8	800	250	<0.5	9.8	<0.5	<0.5	<0.5	
	08/03/00	620 430	3.0 4.30	14 5.50	4.1	1,300	<250	<0.5	3.0	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	
	10/23/00 01/31/01	430	1.5	0.90	8.80 2.8	1,200 280	260 <250	<0.5 <0.5	7.8 5.7	<0.5	<0.5	3.6	
	04/26/01	268	13.0	22.1	22.0	780	<250	<0.5	6.3	<0.5	<0.5	<0.5	
	07/30/01	29.4	<0.5	0.52	0.51	320	<250	<0.5	6.6	<0.5	<0.5	<0.5	
	10/29/01	16.1	2.01	1.14	3.96	<200	<500	<0.5	5.4	<0.5	<0.5	<0.5	
	01/29/02	12.0	<0.5	0.70	<1.0	<200	<250	<0.5	4.9	<0.5	2.0	<0.5	cis 1,2-Dichloroethene detected at 1.3 µg/L.
	04/29/02	188	5.52	9.70	13.0	680	<250	<0.5	6.0	<0.5	<0.5	<0.5	
	10/22/02	4.84	<0.5	<0.5	<1.0	<200	<250	<0.5	4.8	<0.5	<0.5	<0.5	
	05/06/03	20.72	0.76	0.86	2.08	<200	<250	<0.5	5.8	<0.5	<0.5	<0.5	
	10/14/03 04/27/04	6.02 23.60	<0.5 1.68	<0.5 0.67	<1.0 3.91	<200 <200	<250 <250	<0.5 <0.5	3.2 3.0	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	
	11/17/04	2.0	<0.50	<0.50	<0.50	< 50	< 50	<0.50	2.1	<0.50	<0.50	< 5.0	
MW-33	04/07/99	0.60	<0.5	0.90	<0.5	<50	<250					<0.5	
IVIVV-33	04/07/99	8.90	<0.5	1.00	<0.5	<50 <50	<200	0.6	0.7	<0.5	<0.5	<0.5	
	10/28/99	40	0.9	21	3.8	200	<200	0.8	1.3		<0.5		
I	02/10/00	20	0.7	12	10.0	380	<250	0.9	0.6	<0.5	<0.5	1.30	
	04/27/00	6.9	<0.5	6.4	<0.5	<100	250	4.3	0.9	<0.5	<0.5	<0.5	
	08/03/00	31	0.5	20	1.0	150	550	<0.5	0.6	<0.5	<0.5	<0.5	
	10/23/00	89	1.5	36	3.9	350	<250	<0.5	2.1	<0.5	<0.5	<0.5	
 	01/31/01	6.8	<0.5	2.0	<0.5	<50	<250	1.9	0.6	<0.5	<0.5	0.7	
 	04/26/01 07/30/01	6.61	0.56 2.61	1.63	0.61 6.6	<200 <200	<250 <250	2.6 2.2	<0.5 0.5	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	Diabloradifluoromethane detected at 0.6
 	10/29/01	4.43 14.2	<0.5	1.34 0.63	6.6 <1.0	<200 <200	<250 <500	1.3	0.5	<0.5	<0.5	<0.5 <0.5	Dichlorodifluoromethane detected at 0.6 μg/L.
I	01/28/02	<0.5	<0.5	<0.5	<1.0	<200	<250	1.1	0.7	<0.5	3.8	<0.5	Dichlorodiflouromethane detected at 1.9 μg/L.;
[O.O_	3.0	3.0	3.0		_00			0	3.0	0	5.0	cis 1,2-Dichloroethene detected at 8.9 µg/L.
	04/29/02	14.6	<0.5	1.41	<1.0	<200	<250	8.0	0.9	<0.5	<0.5	<0.5	Dichlorodiflouromethane detected at 1.9 μg/L.
MW-100	07/06/01	<0.5	<0.5	<0.5	<0.5	<200	<250	<0.5	<0.5	<0.5	<0.5	<0.5	
 	07/30/01	<0.5	<0.5	<0.5	<0.5	<200	<250	<0.5	<0.5	<0.5	<0.5	<0.5	Chloromethane detected at 1.8 µg/L.
	10/30/01	<0.5	<0.5	<0.5	<1.0	<200	<500	<0.5	<0.5	<0.5	<0.5	<0.5	
	01/28/02 04/29/02	<0.5 <0.5	<0.5 <0.5	<0.5	<1.0	<200 <200	<250 <250	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	
	10/22/02	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<1.0 <1.0	<200 <200	<250 <250	<0.5	<0.5	<0.5	<0.5	<0.5	
1	10/22/02				<1.0	<200	<250	<0.5	<0.5	<0.5	<0.5	<0.5	
	05/06/03	< 0.5	<0.5	<0.5									
	05/06/03 10/14/03	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<1.0	<200	<250	<0.5	<0.5	<0.5	<0.5	<0.5	

Table 3: Historical Groundwater Sample Results (1993 - 2008)

Well Number	Date Sampled	Benzene µg/L	Toluene μg/L	Ethyl- Benzene µg/L	Xylenes μg/L	TPH-G μg/L	TPH-D μg/L	1,1- DCA µg/L	1,2- DCA µg/L	1,1,1- ΤCA μg/L	TCE µg/L	MTBE μg/L	Notes
MW-?	02/05/99	<0.5	<0.5	<0.5	<0.5	<50	430					<0.5	
PR-26	07/26/99 10/26/99	20,000 28,000	15,000 25,000	1,100 2,300	7,250 8,400	82,500 110,000	11,000 60,000	 <0.5	 24		 <0.5	33.0	
PR-45	07/26/99 10/28/99 02/09/00 04/27/00 08/04/00	13,200 12,000 24,000 17,000 20,000	8,200 8,200 25,000 9,500 8,800	2,600 1,700 10,000 16,000 2,600	15,600 8,500 53,000 92,000 16,000	82,500 45,000 360,000 1,300,000 73,000	39,000 25,000 82,000 20,300 54,500	 <0.5 <0.5 <5.0 <0.5	 <0.5 4.0 <5.0 1.0	 <0.5 <5.0 <0.5	 <0.5 <0.5 <5.0 <0.5	35.0 1,000 <5.0 <0.5	
	10/23/00 04/27/01 07/30/01 10/29/01 01/29/02	26,000 16,200 14,500 12,600 8,930	12,000 8,600 8,900 6,650 4,860	4,000 3,220 4,400 2,260 2,640	20,000 19,000 24,700 12,400 12,700	96,000 178,000 132,000 86,100 114,000	36,000 22,700 29,700 50,000 19,400	<0.5 <0.5 <0.5 <0.5	1.2 14 11 7.8 30	<0.5 <0.5 <0.5 <0.5 <0.5	<0.5 <0.5 <0.5 <0.5	<5.0 <25 <50 <25 <0.5	Chloroethane detected at 6.0 µg/L. Chloroethane detected at 4.6 µg/L. Chloromethane detected at 0.6 µg/L; Chloroethane detected at 11 µg/L; Methylene chloride detected at 0.5 µg/L. Chloroethane detected at 6.0 µg/L. Chloroethane detected at 7.5 µg/L.
	05/16/02	14,300	2,630	1,580	7,780	125,000	15,600	<0.5	1.0	<0.5	<0.5	<0.5	Chloroethane detected at 7.3 µg/L.
PR-52	07/26/99 10/28/99 02/09/00 04/28/00 08/04/00 10/24/00 01/31/01	12,000 19,000 22,000 20,000 26,000 52,000 81,000	1,720 530 1,600 2,200 1,600 13,000 840	750 1,800 4,100 4,700 2,900 41,000 57,000	12,400 5,800 15,800 18,600 15,000 180,000 210,000	172,000 40,000 200,000 270,000 150,000 650,000 5,300,000	40,000 450,000 140,000 88,000 110,000 280,000 276,000	<0.5 <0.5 <0.5 <1.0 <0.5 <5.0 <0.5	1.8 <0.5 1.3 <1.0 2.3 <5.0 1.0	<0.5 <0.5 <1.0 <0.5 <5.0 <0.5	<0.5 <0.5 <0.5 <1.0 <0.5 <5.0 <0.5	217 430 <5.0 <0.5 <5.0 500	Methylene chloride detected at 7.9 µg/L. Chloroethane detected at 2.4 µg/L;
	04/27/01 07/30/01 10/29/01	25,000 31,100 22,700	16,300 2,480 1,630	14,700 13,500 3,070	55,000 51,700 11,500	886,000 340,000	134,000 185,000 140,000	<0.5 <0.5 <0.5	<0.5 1.3	<0.5 <0.5 <0.5	<0.5 <0.5 <0.5	1,040 2,510	Methylene chloride detected at 0.6 μg/L. Chloroethane detected at 1.5 μg/L. Chloroethane detected at 1.3 μg/L; Chloroethane detected at 46 μg/L; Methylene chloride detected at 0.6 μg/L. Chloromethane detected at 0.6 μg/L; Chloroethane detected at 0.6 μg/L;
	01/29/02 05/16/02	21,500 31,600	1,840 53,600	4,540 43,800	16,800 216,000	517,000 2,020,000	272,000 75,000	<0.5 <5.0	<0.5 <5.0	<0.5 <5.0	<0.5 <5.0	44.1 63.5	Methylene chloride detected at 0.7 μg/L. Chloroethane detected at 1.5 μg/L. Chloroethane detected at 8.3 μg/L.
PR-53	07/26/99 10/27/99 02/09/00 04/28/00 08/04/00	31,000 17,000 21,000 34,000 35,000	12,000 3,900 5,000 30,000 17,000	1,900 890 1,200 9,300 3,800	8,800 3,320 5,300 51,000 24,000	110,000 54,000 65,000 730,000 180,000	98,000 16,000 9,400 104,000 69,500	<0.5 <0.5 0.6 <1.0 <0.5	43 18 20 <1.0 1.7	<0.5 <0.5 <1.0 <0.5	<0.5 <0.5 <0.5 <1.0 <0.5	43.0 67.0 340 110	Methylene chloride detected at 6.2 μg/L. Methylene chloride detected at 0.8 μg/L.
	10/24/00 01/31/01 04/27/01	99,000 66,000 55,500	110,000 15,000 10,000	80,000 28,000 23,700	640,000 140,000 137,000	580,000 2,400,000 4,240,000	380,000 960,000 806,000	<5.0 <0.5 <0.5	5.0 1.5 <0.5	<5.0 <0.5 <0.5	<5.0 <0.5 <0.5	380 660 <5,000	Chloroethane detected at 1.7 µg/L; Methylene chloride detected at 0.9 µg/L. Chloroethane detected at 1.7 µg/L; Methylene chloride detected at 1.1 µg/L.
	10/29/01 01/29/02 05/16/02	46,500 33,000 35,800	9,520 7,340 10,500	12,900 10,300 18,700	74,000 41,800 130,000	1,630,000 495,000 3,280,000	130,000 462,000 113,000	<0.5 <0.5 <5.0	0.8 1.8 <5.0	<0.5 <0.5 <5.0	<0.5 <0.5 <5.0	<500 122 242	Chloroethane detected at 3.0 µg/L; Methylene chloride detected at 0.9 µg/L. Chloroethane detected at 3.2 µg/L.
PR-54	07/26/99 10/26/99 02/09/00 04/28/00 08/04/00	32,000 27,000 27,000 24,000 27,000	22,000 10,000 23,000 14,000 7,600	1,500 3,700 9,900 1,200 1,400	21,800 19,500 50,000 9,000 11,000	170,000 190,000 960,000 76,000 120,000	28,000 350,000 110,000 80,000 54,500	<0.5 <0.5 <0.5 <1.0 <0.5	3.0 <0.5 3.9 1.6 2.0	<0.5 <0.5 <1.0 <0.5	<0.5 <0.5 <0.5 <1.0 <0.5	56.0 1,000 300 200	Methylene chloride detected at 2.5 μg/L.
	10/24/00 01/31/01	23,000	4,400 8,300	2,000 3,300	13,000 21,000	140,000 220,000	96,000 236,000	<0.5 <0.5	2.3	<0.5 <0.5	<0.5 <0.5	<100 480	Chloroethane detected at 5.3 µg/L; Methylene chloride detected at 2.3 µg/L. Chloroethane detected at 2.8 µg/L; Methylene chloride detected at 1.7 µg/L.
	04/27/01 07/30/01	26,100 31,700	8,650 18,000	2,120 9,880	15,900 58,400	51,300 320,000	108,000 71,200	<0.5 <0.5	<0.5 3.9	<0.5 <0.5	<0.5 <0.5	<500 2,750	Chloroethane detected at 3.0 µg/L. Chloromethane detected at 2.2 µg/L; Chloroethane detected at 2.2 µg/L; Methylene chloride detected at 2.6 µg/L.
	10/30/01 01/29/02 05/16/02	25,400 13,300 27,900	11,300 9,850 34,500	3,500 4,240 5,630	18,800 33,100 36,400	222,000 108,000 324,000	530,000 48,000 172,000	<0.5 <0.5 <5.0	7.5 43	<0.5 <0.5 <5.0	<0.5 <0.5 <5.0	276 51.3 251	Chloroethane detected at 7.4 µg/L; Methylene chloride detected at 2.0 µg/L. Chloroethane detected at 6.2 µg/L. Chloroethane detected at 9.8 µg/L.
PR-64	07/26/99 10/27/99 02/09/00 04/28/00 05/16/02	22,000 11,000 22,000 19,000 18,300	18,000 7,400 20,000 16,000 40,100	1,700 1,200 6,000 1,800 10,400	10,300 3,900 17,000 13,900 104,000	110,000 66,000 120,000 130,000 30,600,000	50,000 40,000 78,000 419,000	<0.5 <0.5 <0.5 <1.0 <5.0	130 110 >50 67 <5.0	<0.5 <0.5 <1.0 <5.0	<0.5 <0.5 <0.5 <1.0 <5.0	35.0 110 300 <500	Methylene chloride detected at 1.4 μg/L.
PR-65	07/26/99 10/26/99	12,000 14,000	1,400 2,300	1,300 1,800	13,000 11,000	68,000 65,000	16,500 50,000	<0.5 <0.5	2.6 <0.5	<0.5	<0.5 <0.5	20.0	
PR-68	07/26/99 10/26/99	1,900 2,800	24.0 36	27.0 86	62.0 62	4,900 8,000	11,000 2,800	<0.5 <0.5	1.2 <0.5	<0.5	<0.5 <0.5	4.40	

Table 3: Historical Groundwater Sample Results (1993 - 2008)

Well Number	Date Sampled	Benzene µg/L	Toluene μg/L	Ethyl- Benzene µg/L	Xylenes μg/L	TPH-G μg/L	TPH-D μg/L	1,1- DCA µg/L	1,2- DCA µg/L	1,1,1- ΤCA μg/L	TCE µg/L	MTBE μg/L	Notes
PR-76	04/07/99	<0.5	<0.5	<0.5	<0.5	<50	<250					<0.5	
	10/22/02 05/06/03	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<1.0 <1.0	<200 <200	<250 <250	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	
	10/14/03	<0.5	<0.5	<0.5	<1.0	<200	<250	<0.5	<0.5	<0.5	<0.5	<0.5	
	04/27/04	<0.5	<0.5	<0.5	<1.0	<200	<250	<0.5	<0.5	<0.5	<0.5	<0.5	
	11/17/04	<0.50	<0.50	<0.50	<0.50	<50	85	<0.50	<0.50	<0.50	<0.50	<5.0	
V-24	04/07/99	<0.5	<0.5	<0.5	<0.5	120	<250				-	0.5	
V-31	07/26/99 10/26/99	7,000 7,000	600 120	550 850	1,370 950	17,500 18,000	5,350 3,000	 <0.5	 <0.5		 <0.5	19.0	
V-46	02/05/99	<0.5	<0.5	<0.5	<0.5	<50	270	<0.5	<0.5	<0.5	<0.5	<0.5	
V-55	07/22/99	8,000	480	740	2,880	30,000	2,100	<0.5	<0.5	<0.5	<0.5	13.0	
	10/28/99	11,000	59	1,200	317	28,000	38,000	<0.5	<0.5		<0.5		
	02/09/00	2,200	59 540	760	350	7,900	10,000	<0.5	<0.5	<0.5	<0.5	9.70	
	04/28/00 08/03/00	2,900 9,400	510 380	440 720	2,340 2,200	14,000 28,000	26,500 70,000	<5.0 <0.5	<5.0 <0.5	<5.0 <0.5	<5.0 <0.5	<5.0 <0.5	
	10/23/00	11,000	140	900	1,300	30,000	51,000	<0.5	<0.5	<0.5	<0.5	<12	
	01/31/01	4,600	57	550	1,200	34,000	88,500	<0.5	<0.5	<0.5	<0.5	44	
	04/26/01	6,400	61.5	250	336	34,200	227,000	<0.5	<0.5	<0.5	<0.5	<25	
	10/30/01 01/29/02	5,360 1,660	70.0 140	1,090 492	1,450 818	32,700 12,000	78,000 4,100	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<25 <0.5	
	04/29/02	5,170	95.1	572	523	30,600	35,100	<0.5	<0.5	<0.5	<0.5	1.06	
V-72	07/26/99	13,500	6.80	1.10	3.90	3,900	12,900	<0.5	11	<0.5	<0.5	<0.5	
1 2 2	10/28/99	2,900	58	21	3.90 47.7	6,000	48,000	<0.5	3.4	<0.5	<0.5	<0.5	
	02/09/00	670	8.2	<0.5	17.8	890	6,100	<0.5	3.0	<0.5	<0.5	<0.5	
	04/28/00	130	<0.5	<0.5	<0.5	200	5,950	<0.5	0.7	<0.5	<0.5	<0.5	
	08/04/00	460	0.8	<0.5	0.6	440 3.500	4,120	<0.5	2.8	<0.5	<0.5 <0.5	<0.5 <0.5	
	10/24/00 04/27/01	2,700 1,240	3.2 2.05	0.5 <0.5	2.3 2.78	3,500 1,310	17,000 6,290	<0.5 <0.5	4.0 5.1	<0.5 <0.5	<0.5	<0.5	Dichlorodifluoromethane detected at 0.8 μg/L.
	07/30/01	1,790	69.8	1.22	2.50	1,490	4,290	<0.5	6.2	<0.5	<0.5	<0.5	Chloromethane detected at 1.5 μg/L.
	10/29/01	1,330	4.38	0.55	3.32	1,960		<0.5	5.6	<0.5	<0.5	<0.5	Chloromethane detected at 1.1 µg/L.
	01/29/02	655	6.40	<0.5	8.00	1,840	2,250	<0.5	3.9	<0.5	<0.5	<0.5	Chloromethane detected at 1.8 µg/L.
	05/16/02	43.8	1.09	<0.5	4.36	230	5,120	<0.5	<0.5	<0.5	<0.5	<0.5	Chloromethane detected at 1.8 μg/L.
V-84	07/26/99 10/26/99	2,400 1,100	440 130	80.0 46	340 108	8,700 4,000	2,350 700	<0.5 <0.5	2.4 <0.5	<0.5 	<0.5 <0.5	6.40	
	02/09/00	300	30	8.9	53	2,300	1,100	<0.5	1.2	<0.5	<0.5	<0.5	
	04/28/00	30	1.9	<0.5	<0.5	100	550	<5.0	<5.0	<5.0	<5.0	<0.5	
	08/04/00	900	110	34	120	2,700	1,380	<0.5	1.0	<0.5	<0.5	<0.5	
	10/24/00	2,000 68	480	24	110	48,000 970	1,900	<0.5	1.0 <0.5	<0.5	<0.5 <0.5	<0.5	
	01/31/01 04/26/01	925	1.3 97.0	5.3 45.4	8.2 59.7	2,360	1,820 1,180	<0.5 <0.5	0.8	<0.5 <0.5	<0.5	<0.5 <0.5	
	07/30/01	1,720	282	50	359	8,100	7,040	<0.5	1.5	<0.5	<0.5	<0.5	
	10/30/01	870	250	27.6	167	8,960		<0.5	1.0	<0.5	<0.5	<0.5	
	01/29/02 04/29/02	197 318	4.90 34.4	1.70 15.4	3.60 18.4	640 1,070	500 400	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	
29 (CC-1)	07/23/99	<0.5	<0.5	<0.5	<0.5	<50	<200	<0.5	<0.5	<0.5	<0.5	<0.5	
29 (00-1)	10/28/99	<0.5	<0.5	<0.5	<0.5	<100	<200	<0.5	<0.5		<0.5		
	02/08/00	<0.5	<0.5	<0.5	<0.5	<50	<250	<0.5	<0.5	<0.5	<0.5	<0.5	
	04/26/00	<0.5	<0.5	<0.5	< 0.5	<100	<250	<0.5	<0.5	<0.5	<0.5	<0.5	
	08/03/00 10/23/00	1.4 <0.5	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<50 <50	<250 <250	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	
	01/31/01	<0.5	<0.5	<0.5	<0.5	<50	<250	<0.5	<0.5	<0.5	<0.5	<0.5	
	04/26/01	<0.5	<0.5	<0.5	<0.5	<200	<250	<0.5	<0.5	<0.5	<0.5	<0.5	
	07/30/01	<0.5	<0.5	<0.5	<0.5	<200	<250	<0.5	<0.5	<0.5	<0.5	<0.5	
	10/30/01 01/28/02	1.12 <0.5	0.56 <0.5	<0.5 <0.5	<0.5 <1.0	<200 <200	<500 <250	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	
	04/29/02	<0.5	<0.5	<0.5	<0.5	<200	<250	<0.5	<0.5	<0.5	<0.5	<0.5	
	10/22/02	1 20	14.6	2.44	16.4	220	-250	-0 F	-0 E	-0 F	-0 F	02.0	Chloromethane detected at 1.3 µg/L, Chloroform
	10/22/02 11/15/02	1.38 <0.50	14.6 <0.50	2.44 <0.50	16.4 <1.0	220 <200	<250 <250	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	92.0 <0.5	detected at 4.7 µg/L. Chloroform detected at 2.6 µg/L.
	05/06/03	<0.50	<0.50	<0.50	<1.0	<200	<250	<0.5	<0.5	<0.5	<0.5	<0.5	
	10/14/03	<0.50	<0.50	<0.50	<1.0	<200	<250	<0.5	<0.5	<0.5	<0.5	<0.5	Chloroform detected at 0.7 μg/L.
	04/27/04	< 0.50	<0.50	<0.50	<1.0	<200	<250	<0.5	<0.5	<0.5	<0.5	<0.5	
00 (00 S;	11/17/04	<0.50	<0.50	<0.50	<0.50	<50	<50	<0.50	<0.50	<0.50	<0.50	<5.0	
30 (CC-2)	07/22/99 10/28/99	0.90 <0.5	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<50 <100	<200 <200	<0.5 <0.5	<0.5 <0.5	<0.5	<0.5 <0.5	<0.5	
	02/08/00	<0.5 <0.5	<0.5	<0.5	<0.5 <0.5	<50	<250	<0.5	<0.5	<0.5	<0.5	<0.5	
	04/26/00	<0.5	<0.5	<0.5	<0.5	<100	<250	<0.5	0.7	<0.5	<0.5	<0.5	
	08/03/00	<0.5	<0.5	<0.5	<0.5	<50	<250	<0.5	<0.5	<0.5	<0.5	<0.5	
	10/23/00 01/31/01	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<50 <50	340 <250	<0.5 <0.5	0.9 <0.5	<0.5 <0.5	<0.5 <0.5	<2.5 <0.5	
	04/26/01	<0.5 <0.5	<0.5	<0.5	<0.5 <0.5	<200	<250	<0.5	<0.5	<0.5	<0.5	<0.5	
	07/30/01	<0.5	1.43	<0.5	1.63	<200	<250	<0.5	1.6	<0.5	<0.5		Dichlorodifluoromethane detected at 2.8 μg/L.

Table 3: Historical Groundwater Sample Results (1993 - 2008)

Well Number	Date Sampled	Benzene µg/L	Toluene μg/L	Ethyl- Benzene	Xylenes μg/L	TPH-G μg/L	TPH-D μg/L	1,1- DCA	1,2- DCA	1,1,1- TCA	TCE µg/L	MTBE μg/L	Notes
	·	. 0		μg/L				μg/L	μg/L	μg/L			
30 (CC-2)	10/29/01	<0.5	<0.5	<1.0	<0.5	<200	<500	<0.5	<0.5	<0.5	<0.5	<0.5	
(cont.)	01/28/02	<0.5	<0.5	<0.5	<1.0	<200	<250	<0.5	1.9	<0.5	<0.5	<0.5	Dichlorodifluoromethane detected at 3.8 μg/L.
	04/29/02	<0.5	<0.5	<0.5	<0.5	<200	<250	<0.5	2.5	<0.5	<0.5	0.86	Dichlorodifluoromethane detected at 3.6 μg/L.
	10/10/02	<0.5	<0.5	<0.5	<1.0	<200	<250	<0.5	<0.5	<0.5	<0.5	<0.5	Chloroform detected at 0.6 µg/L.
	11/15/02	<0.5	<0.5	<0.5	<1.0	<200	<250	<0.5	<0.5	<0.5	<0.5	<0.5	Chloroform detected at 0.5 μg/L.
	05/06/03	<0.5	<0.5	<0.5	<1.0	<200	<250	<0.5	<0.5	<0.5	<0.5	<0.5	
81	02/05/99 07/22/99	<0.5 0.70	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<50 <50	<150 <200	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	
94	02/05/99	<0.5	<0.5	<0.5	<0.5	<50	170					<0.5	
	07/22/99	<0.5	<0.5	<0.5	<0.5	<50	<200	<0.5	<0.5	<0.5	<0.5	<0.5	
210	02/05/99	<0.5	<0.5	<0.5	<0.5	<50	960					<0.5	
223	10/26/99	<0.5	<0.5	<0.5	<0.5	<100	<200	<0.5	<0.5		<0.5		
	02/10/00	<0.5	<0.5	<0.5	<0.5	<50	640	<0.5	<0.5	<0.5	<0.5	<0.5	
ĺ	04/27/00	<0.5	<0.5	<0.5	<0.5	<100	250	<0.5	<0.5	<0.5	<0.5	<0.5	
	08/03/00	<0.5	<0.5	<0.5	<0.5	<50	680	<0.5	<0.5	<0.5	<0.5	<0.5	
	10/23/00	1.30	<0.5	<0.5	<0.5	<50	<250	<0.5	<0.5	<0.5	<0.5	<0.5	Chlorobenzene detected at 0.9 µg/L.
	01/31/01	<0.5	<0.5	<0.5	<0.5	<50	<250	<0.5	<0.5	<0.5	<0.5	<0.5	
	04/26/01	<0.5	<0.5	<0.5	<0.5	<200	390	<0.5	<0.5	<0.5	<0.5	<0.5	1,2-Dichlorobenzene detected at 0.5 µg/L.
	07/30/01	<0.5	<0.5	<0.5	<0.5	<200	<250	<0.5	<0.5	<0.5	<0.5	<0.5	Dichlorodifluoromethane detected at 0.5 μg/L.
	10/30/01	<0.5	<0.5	<0.5	<1.0	<200	<500	<0.5	<0.5	<0.5	<0.5	<0.5	Chloromethane detected at 0.8 µg/L.
	01/29/02	<0.5	<0.5	<0.5	<1.0	<200	<250	<0.5	<0.5	<0.5	<0.5	<0.5	
	04/29/02	<0.5	<0.5	<0.5	<1.0	<200	<250	<0.5	<0.5	<0.5	<0.5	<0.5	
224	07/26/99	<0.5	<0.5	<0.5	<0.5	<50	640	<0.5	<0.5	<0.5	<0.5	<0.5	
239	07/26/99	55,000	85.0	1,500	190	30,000		<0.5	<0.5	<0.5	<0.5	5.30	
	10/26/99	23,000	53	1,500	103.2	28,000	10,000	<0.5	<0.5		<0.5		
	02/10/00	40,000	48	1,900	52	44,000	21,000	<0.5	1.0	<0.5	<0.5	14.0	
	04/28/00	25,000	540	2,000	710	36,000	12,500	<5.0	<5.0	<5.0	<5.0	<5.0	
	08/04/00	25,000	220	1,900	920	45,000	32,500	<0.5	0.6	<0.5	<0.5	<0.5	
	10/24/00	24,000	100	1,500	390	50,000	50,000	<0.5	<0.5	<0.5	<0.5	<5.0	
	01/31/01	23,000	84	1,900	200	52,000	112,000	<0.5	0.9	<0.5	<0.5	<0.5	
	04/26/01	23,900	113 384	1,990	590 966	298,000	143,000	<0.5	<0.5	<0.5	<0.5	<25	
	07/30/01 10/30/01	30,200 41,200	273	2,000 1,470	215	66,500 54,300	19,100 120,000	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<0.5 <0.5	<0.5 <50	
	01/28/02	24,500	273	1,470	352	112,000	6,900	<0.5	<0.5	<0.5	<0.5	<0.5	Chloroethane detected at 0.6 μg/L.
	04/29/02	25,900	280	1,380	491	71,600	9,400	<0.5	<0.5	<0.5	<0.5	<0.5	Chloroethane detected at 0.0 µg/L.
241	04/07/99	<0.5	<0.5	<0.5	<0.5	<50	<250					<0.5	
249	07/22/99	<0.5	<0.5	<0.5	<0.5	<50	<200	<0.5	<0.5	<0.5	<0.5 NA	<0.5 NA	
SB-16	05/20/08	< 0.50	< 0.50	< 0.50	530	<50	530	NA NA	<0.50	NA NA	NA NA	NA NA	
SB-17 SB-18	05/22/08 05/22/08	12,000 50.000	3,200	17,000	560,000	120,000	560,000 23.000	NA NA	<0.50 2,200	NA NA	NA NA	NA NA	
SB-18 SB-19	05/22/08	50,000 <12	2,300 220	46,000 <12	23,000 1,600	190,000 8,200	1,600	NA NA	2,200 <12	NA NA	NA NA	NA NA	
SB-19 SB-20/ PCB-7	05/22/08	41,000	3,000	30,000	47,000	8,200 170,000	47,000	NA NA	930	NA	NA NA	NA NA	
SB-20/ PCB-7 SB-21/ PCB-8	05/22/08	12,000	2,600	20,000	3,500	110,000	3,500	NA	<250	NA	NA	NA	
SB-21/ PCB-8 SB-22	05/23/08	27,000		39,000	73,000	870,000	73,000	NA NA	<2,500	NA	NA NA	NA NA	
SB-24/ PCB-1	05/22/08		13,000 <0.50	<0.50	360	<50	360	NA	<0.50	NA	NA	NA	
SB-25/ PCB-1 SB-25/ PCB-2		1.1						NA NA		NA	NA NA	NA NA	
SB-25/ PCB-2 SB-26	05/21/08 05/22/08	<0.50 <0.50	<0.50 <0.50	<0.50 <0.50	140 270	<50 <50	140 270	NA	<0.50 <0.50	NA	NA	NA	
						<50 NA	_	NA		NA	NA	NA NA	
SB-27/ PCB-3	05/20/08	<0.50	<0.50	< 0.50	NA	NΑ	NA	IAM	<0.50	INA	IAM	INA	l

Not detected.

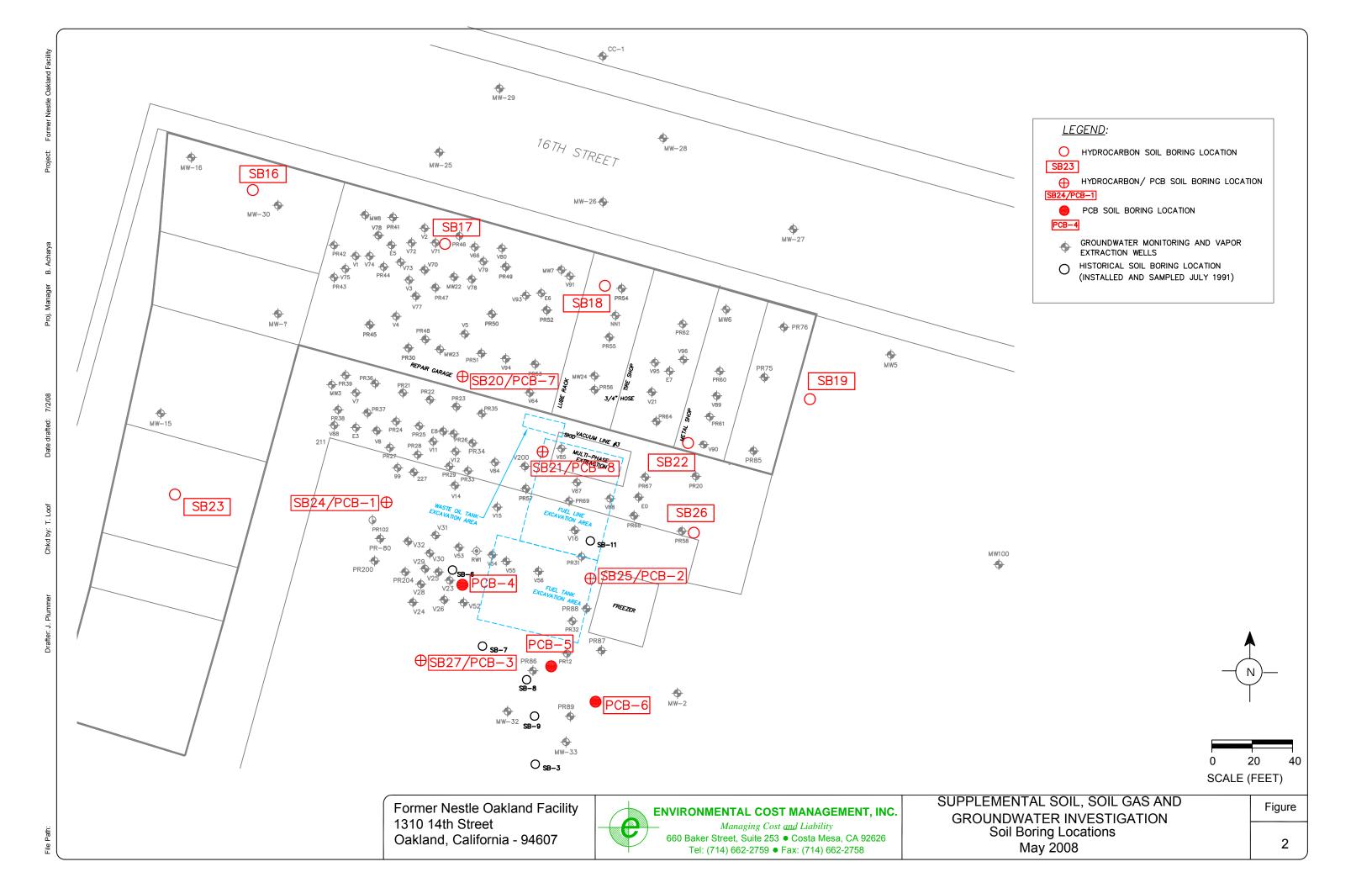
ND NA Not analyzed or not sampled.

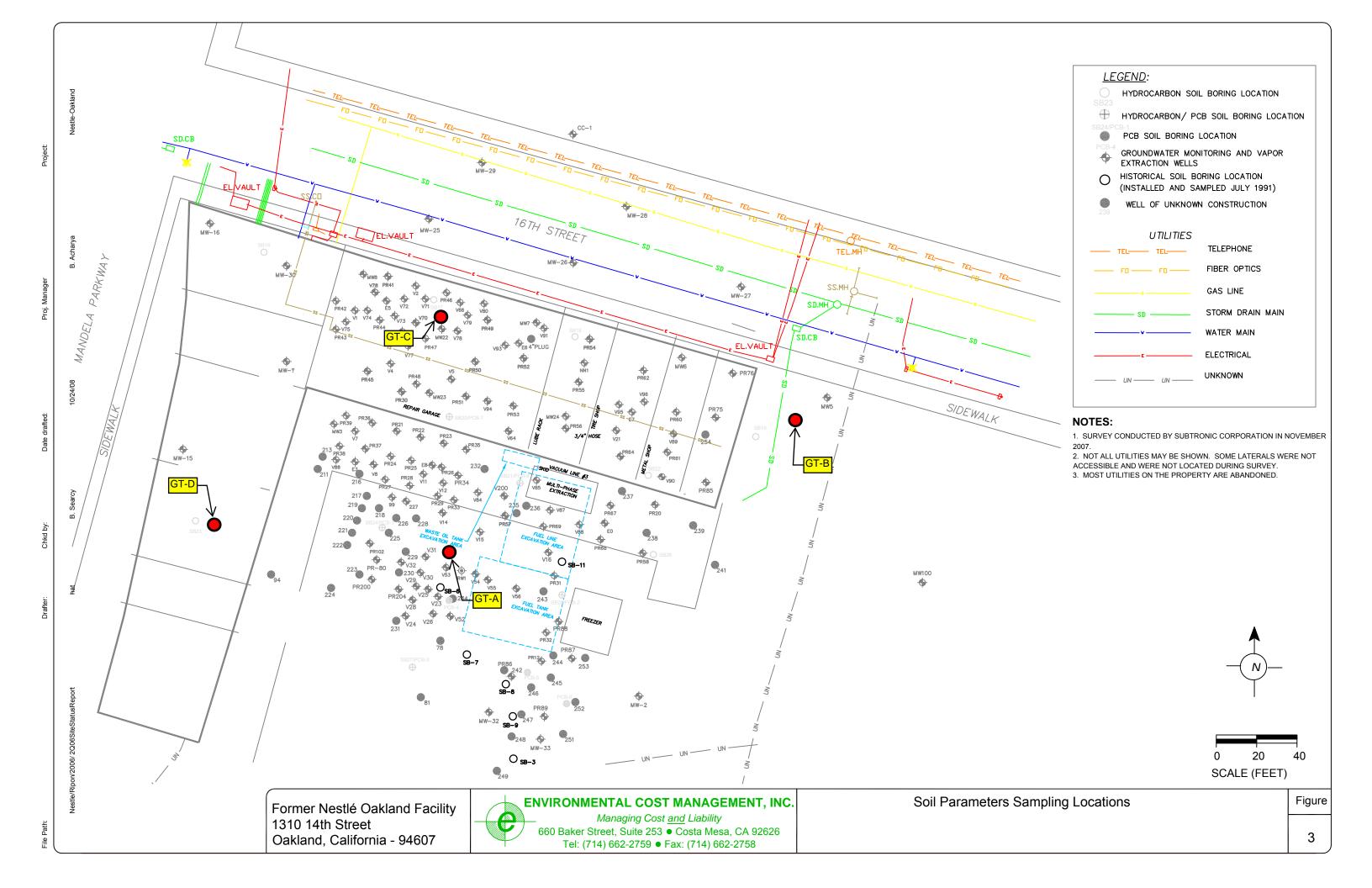
Micrograms per liter.

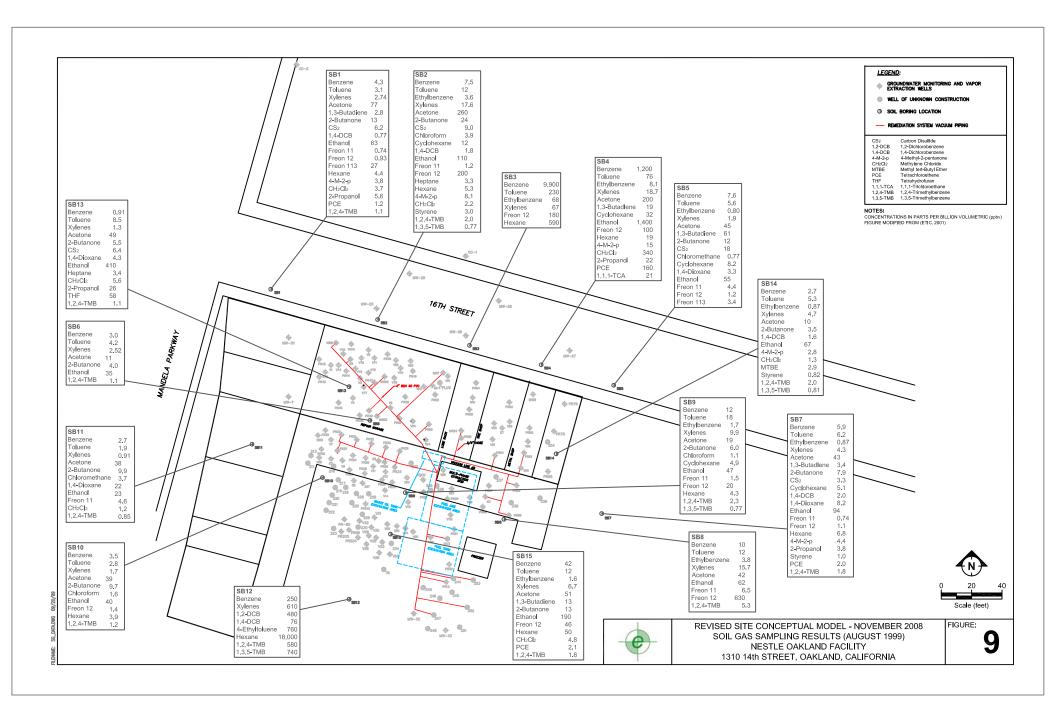
μg/L TPH-G Total Petroleum Hydrocarbons as gasoline. TPH-D Total Petroleum Hydrocarbons as diesel.

1,1-DCA 1,1-Dichloroethane. 1,1-DCA cis-1,1-DCE 1,1,1-TCA 1,2-Dichloroethane. 1,1-Dichloroethene. 1,1,1-Trichloroethane. 1,2-DCE TCE cis 1,2-Dichloroethylene. Trichloroethene. MTBE Methyl tertiary butyl ether.

10/22/02 Data was confirmed anomalous by resampling on 11/15/02.







APPENDIX B UNCERTAINTIES ANALYSIS

1.0 INTRODUCTION

Many of the assumptions used in the screening level human health risk evaluation — regarding the representativeness of the sampling data, human exposures, fate and transport modeling, and chemical toxicity — are conservative, follow agency guidance, and reflect a 90th or 95th percentile value rather than a typical or average value. The use of several conservative exposure and toxicity assumptions can introduce considerable uncertainty into the human health screening evaluation. By using conservative exposure or toxicity estimates, the evaluation can develop a significant conservative bias that may result in the calculation of significantly higher cancer risks or noncancer hazards than are actually posed by the chemicals present in soil, soil gas, and groundwater. The key uncertainties in the human health screening evaluation are discussed below. This uncertainty analysis focuses on the site-specific assumptions contributing most to uncertainty in the risk and hazard calculations, and does not assess the validity of default assumptions used in the health screening evaluation. The parameters evaluated in the uncertainty analysis are: representative concentrations in soil gas, soil properties, and building air exchange rate.

The uncertainties associated with representative concentrations in soil gas, soil properties, and building air exchange rate are discussed below. Two sensitivity analyses have been performed, to bound the range of potential risks and hazards associated with the uncertainties in these three input parameters. The first sensitivity analysis combines the most conservative options of the three parameters, to produce a high-end estimate of potential risk and hazard. The second sensitivity analysis combines the least conservative options of the three parameters, to produce a low-end estimate of potential risks and hazards. The baseline health risk evaluation, the results of which are presented in Section 7 of the main report, is based on a combination of assumptions regarding these three parameters, based primarily on DTSC vapor intrusion guidance (Cal/EPA, 2005b), and represents a relatively conservative estimate of potential risk and hazard. The conceptual differences between the three evaluations are summarized in the following table.

Summary of Health Risk Evaluations

Evaluation	Where Documented	Soil Gas Concentrations	Soil Properties	Building Ventilation Rate
High-end estimate	Appendix C	Maxima	Default	Default
Baseline estimate	Main report	Maxima	Site-specific	Default
Low-end estimate	Appendix D	Averages	Site-specific	Site-specific

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2.0 Uncertainty in Representative Concentrations in Soil Gas

As discussed in Sections 4.5.1, and 4.5.2 of the main report, the baseline analyses of vapor transport from soil gas to indoor air and outdoor air are based on the historical maximum detected concentration of each COPC in soil gas, from the combined 1999 and 2008 datasets. This assumption is consistent with current DTSC vapor intrusion guidance (Cal/EPA, 2005b), which recommends the use of maximum detected soil gas concentrations in vapor intrusion screening risk evaluations. It should be noted that this recommendation is based on an assumed scenario of residential development on quarteracre lots, with soil gas data collected at this same density (quarter-acre density is equivalent to 100-foot spacing between sampling locations), so that the worst-case residence is evaluated. At this site, however, soil gas sampling has been performed at a much greater density, including numerous samples collected beneath the one on-site commercial building (see ECM Figures 2 and 9 in Appendix A). Furthermore, the 1999 results are generally higher than the 2008 results and so provide most of the maximum concentrations, but are likely to be less representative of current site conditions than the 2008 data. Therefore, the use of average, rather than historical maximum, soil gas concentrations is arguably more appropriate for estimating the long-term average indoor air concentrations associated with vapor intrusion into the onsite commercial building. The sensitivity of the results of the health risk evaluation to the use of maximum versus average concentrations of COPCs in soil gas is discussed below in Section 5.0.

3.0 Uncertainty in Soil Properties

As discussed in Sections 4.5.1, and 4.5.2 of the main report, the baseline analyses of vapor transport from soil gas to indoor air and outdoor air are based on site-specific soil properties (total porosity, water-filled porosity, and bulk density) as recommended in the DTSC vapor intrusion guidance (Cal/EPA, 2005b). These properties were measured during the January 2009 site investigation. While measured site-specific soil properties are likely to be most representative of actual site soil conditions, these site-specific properties are less conservative than the DTSC/HERD default soil properties (Cal/EPA, 2005a). The sensitivity of the results of the health risk evaluation to the use of default versus site-specific soil properties is discussed below in Section 5.0.

4.0 Uncertainty in Building Air Exchange Rate

The predicted vapor intrusion transport of COPCs from soil gas into the onsite building is dependent upon the building air exchange rate; specifically, predicted concentrations of COPCs in indoor air are inversely proportional to the air exchange rate. As noted in Section 4.5.1 of the main report, the baseline vapor intrusion transport analysis assumes the DTSC/HERD default air exchange rate of 1 building volume per hour (Cal/EPA, 2005a). This default air exchange rate is conservative and very likely underestimates the actual air exchange rate of the existing onsite commercial building. The existing onsite building is an old, warehouse-type structure with several rollup doors; a reasonable estimate of the actual air exchange rate of this structure, based on engineering judgment, is 4 building volumes per hour. The sensitivity of the results of the health risk evaluation

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to the use of a default versus site-specific building air exchange rate is discussed below in Section 5.0.

5.0 High-end and Low-end Estimates of Potential Risk and Hazard

As discussed above, uncertainties exist in the human health risk evaluation regarding representative concentrations of COPCs in soil gas, soil properties, and building air exchange rate. The baseline health risk evaluation, the results of which are summarized in Section 7 of the main report, is based on a combination of assumptions regarding these three parameters; these assumptions are consistent with DTSC vapor intrusion guidance (Cal/EPA, 2005b) and produce a reasonably conservative estimate of potential risk and hazard.

Discussed here are two sensitivity analyses which bound the range of potential risks and hazards associated with the uncertainties in these three input parameters. The first sensitivity analysis combines the most conservative, but likely least representative, options of the three parameters, to produce a high-end estimate of potential risk and hazard. The second sensitivity analysis combines the least conservative, but likely most representative, options of the three parameters, to produce a low-end estimate of potential risks and hazards

The high-end estimate of potential risk and hazard is based on a combination of historical maximum concentrations of COPCs in soil gas, DTSC/HERD default soil properties, and the DTSC/HERD default building air exchange rate. This worst-case estimate is likely the least representative of actual exposures and associated health effects, of the three evaluations. This analysis is documented in Tables C-1 through C-5 of Appendix C. The results of this high-end estimate may be summarized as follows.

- Using default soil properties, the estimated cancer risk for onsite indoor commercial/industrial workers, associated with vapor intrusion (the only complete exposure pathway for this receptor), is 6.7×10⁻⁵, which is within the 1×10⁻⁶ to 1×10⁻⁴ risk management range (see Table C-4). The estimated noncancer hazard index is 0.44, which is below the threshold hazard index of 1 (see Table C-5).
- Using default soil properties, the estimated cancer risk for onsite outdoor intrusive construction workers, summed across the four complete exposure pathways, is 9.8×10⁻⁵, which is within the 1×10⁻⁶ to 1×10⁻⁴ risk management range (see Table C-4). The estimated noncancer hazard index is 21, which is above the threshold hazard index of 1 (see Table C-5).
- Using default soil properties, the estimated cancer risk for offsite residents, associated with inhalation of volatile COPCs in indoor or outdoor air that have migrated downwind from the site (the only complete exposure pathway for this receptor), is 6.9×10⁻⁶, which is within the 1×10⁻⁶ to 1×10⁻⁴ risk management range (see Table C-4). The estimated noncancer hazard index is 0.069, which is below the threshold hazard index of 1 (see Table C-5).

The low-end estimate of potential risk and hazard is based on a combination of average concentrations of COPCs in soil gas, site-specific soil properties, and site-specific building air exchange rate. This estimate is likely the most representative of actual exposures and associated health effects, of the three evaluations. This analysis is documented in Tables D-1 through D-6 of Appendix C. The results of this low-end estimate may be summarized as follows.

- Using average concentrations of COPCs in soil gas and a site-specific building air exchange rate, the estimated cancer risk for onsite indoor commercial/industrial workers, associated with vapor intrusion (the only complete exposure pathway for this receptor), is 1.7×10^{-7} , which is below the 1×10^{-6} to 1×10^{-4} risk management range (see Table D-5). The estimated noncancer hazard index is 0.0011, which is below the threshold hazard index of 1 (see Table D-6).
- Using average concentrations of COPCs in soil gas and a site-specific building air exchange rate, the estimated cancer risk for onsite outdoor intrusive construction workers, summed across the four complete exposure pathways, is 9.8×10^{-5} , which is within the 1×10^{-6} to 1×10^{-4} risk management range (see Table D-5). The estimated noncancer hazard index is 21, which is above the threshold hazard index of 1 (see Table D-6).
- Using average concentrations of COPCs in soil gas and a site-specific building air exchange rate, the estimated cancer risk for offsite residents, associated with inhalation of volatile COPCs in indoor or outdoor air that have migrated downwind from the site (the only complete exposure pathway for this receptor), is 3.5×10^{-8} , which is below the 1×10^{-6} to 1×10^{-4} risk management range (see Table D-5). The estimated noncancer hazard index is 0.00033, which is below the threshold hazard index of 1 (see Table D-6).

The following conclusions may be drawn regarding the sensitivity of the results of the human health risk evaluation to the uncertainties regarding representative concentrations of COPCs in soil gas, soil properties, and building air exchange rate.

- Estimated potential risk and hazard for onsite indoor commercial/industrial workers range over approximately 2-1/2 orders of magnitude. The high-end estimates are approximately 8 times greater than the baseline estimates presented in Section 7 of the main report. The low-end estimates are less than the baseline estimates by a factor of approximately 47.
- Estimated potential risk and hazard for onsite outdoor intrusive construction workers are not sensitive to these uncertainties, as the estimated potential health effects for this receptor are driven by dermal contact with groundwater.
- Estimated potential risk and hazard for offsite residents range over approximately 2 orders of magnitude. The high-end estimates are approximately 17 times greater than the baseline estimates presented above in Section 7 of the main report. The low-end estimates are less than the baseline estimates by a factor of approximately 11.

APPENDIX C HIGH-END ESTIMATE OF RISK AND HAZARD

Table C-1. Johnson and Ettinger Model Inputs – High-end Estimate

Parameter	Symbol	Value	Units	Reference
Building Properties				
Depth below grade to bottom of enclosed space floor	L_{F}	15	cm	DTSC/HERD default (Cal/EPA, 2005a; 2005b)
Area of enclosed space below grade	$A_{b,sg}$	2.05E+07	cm^2	Site-specific
Building air exchange rate	AXR_b	1	hr ⁻¹	DTSC default (Cal/EPA, 2005b)
Building height	Bh	503	cm	Site-specific
Building ventilation rate	Q_b	2.86E+06	cm^3/s	Calculated: $A_{b,sg} x AXR_b x B_h$
Vapor flow rate into building	Q_{soil}	102	L/min	Calculated (Cal/EPA, 2005b)
Vapor flow rate into building	Q_{soil}	1707	cm ³ /s	Calculated via units conversion
Soil Properties				
Average soil temperature	T_s	17	°C	Site-specific (USEPA, 2004)
SCS soil type	_	Herd Defau	lt —	Site-specific
Dry bulk density	ρ_{b}	1.50	g/cm ³	DTSC default (Cal/EPA, 2005b)
Total porosity	n	0.430	cm ³ /cm ³	DTSC default (Cal/EPA, 2005b)
Water-filled porosity	θ_{w}	0.150	cm ³ /cm ³	DTSC default (Cal/EPA, 2005b)
Air-filled porosity	θ_a	0.280	cm ³ /cm ³	DTSC default (Cal/EPA, 2005b)

Table C-2. Exposure Point Concentrations of Chemicals of Potential Concern in the Indoor Air of the Onsite Building – High-end Estimate

	Modeled Sou			apor Intrusion eling
Chemical of Potential Concern	C_{SG} ($\mu g/m^3$)	Depth (cm)	α	C_{IA} ($\mu g/m^3$)
Acetone	620	91	3.6E-04	2.2E-01
Benzene	40,000	91	3.1E-04	1.2E+01
1,3-Butadiene	310	91	4.5E-04	1.4E-01
2-Butanone (methyl ethyl ketone)	420	91	3.0E-04	1.3E-01
Carbon disulfide	440	91	3.3E-04	1.5E-01
Chlorobenzene	160	91	2.8E-04	4.5E-02
Chloroform	170	91	3.3E-04	5.7E-02
Chloromethane (methyl chloride)	75	91	3.6E-04	2.7E-02
Cyclohexane	480	91	3.0E-04	1.4E-01
1,2-Dichlorobenzene	2,900	91	2.7E-04	7.9E-01
1,3-Dichlorobenzene	210	91	2.7E-04	5.8E-02
1,4-Dichlorobenzene	460	91	2.7E-04	1.3E-01
Dichlorodifluoromethane (Freon 12)	10,000	91	2.7E-04	2.7E+00
1,1-Dichloroethane	140	91	2.8E-04	4.0E-02
1,2-Dichloroethane	140	91	3.3E-04	4.7E-02
1,1-Dichloroethene (1,1-DCE)	140	91	3.1E-04	4.4E-02
cis-1,2-Dichloroethene (cis-1,2-DCE)	140	91	2.8E-04	4.0E-02
1,4-Dioxane	500	91	3.2E-04	1.6E-01
Ethanol	2,600	91	3.7E-04	9.6E-01
Ethylbenzene	7,700	91	2.9E-04	2.2E+00
4-Ethyltoluene	3,700	91	2.7E-04	1.0E+00
Heptane	550	91	2.8E-04	1.5E-01
Hexane	63,000	91	4.2E-04	2.7E+01
Methyl tertiary butyl ether (MTBE)	500	91	3.3E-04	1.7E-01
Methylene chloride	1,200	91	3.3E-04	4.0E-01
4-Methyl-2-pentanone (methyl isobutyl ketone)	550	91	2.9E-04	1.6E-01
2-Propanol	350	91	3.4E-04	1.2E-01
Styrene	150	91	2.8E-04	4.2E-02
Tetrachloroethene (PCE)	1,100	91	2.8E-04	3.1E-01
Tetrahydrofuran	420	91	3.3E-04	1.4E-01

Table C-2. Exposure Point Concentrations of Chemicals of Potential Concern in the Indoor Air of the Onsite Building – High-end Estimate

	Modeled Sou		Results of Vapor Intrusion Modeling		
Chemical of Potential Concern	C_{SG} (µg/m ³)	Depth (cm)	α	C_{IA} (µg/m ³)	
Toluene	32,000	91	3.1E-04	9.9E+00	
1,1,1-Trichloroethane	190	91	2.9E-04	5.5E-02	
Trichloroethene (TCE)	190	91	2.9E-04	5.6E-02	
Trichlorofluoromethane (Freon 11)	200	91	3.1E-04	6.2E-02	
1,1,2-Trichloro-1,2,2-trifluoroethane (Freon 113)	270	91	2.9E-04	7.9E-02	
1,2,4-Trimethylbenzene	2,900	91	2.5E-04	7.4E-01	
1,3,5-Trimethylbenzene	3,600	91	2.5E-04	9.1E-01	
Xylenes	19,000	91	3.0E-04	5.8E+00	

- (1) This vapor intrusion transport analysis is based on maximum concentrations of volatile organic compounds (VOCs) in soil gas from the August 1999 and May 2008 site investigations (see Table 1). Chemicals of potential concern (COPCs) with respect to vapor intrusion are those VOCs detected above reporting limits in at least one soil gas sample.
- (2) The shallower sampling depth between the 1999 and 2008 sampling events, 3 feet below ground surface, is used in the model because it is a more conservative assumption, *i.e.*, it produces higher indoor air concentrations.
- (3) Non source-related inputs to the Johnson and Ettinger Model are documented in Table C-1. Shown here are the results of the Johnson and Ettinger Model, consisting of, for each chemical of potential concern, the predicted attenuation factor (α) and the predicted concentration of the chemical in indoor air (C_{IA}).

Table C-3. Exposure Point Concentrations of Chemicals of Potential Concern in Onsite Outdoor Air and in Offsite Air – High-end Estimate

Chemical of Potential Concern	Concentration
	$(\mu g/m^3)$
Acetone	8.8E-03
Benzene	4.0E-01
1,3-Butadiene	8.8E-03
2-Butanone (methyl ethyl ketone)	3.9E-03
Carbon disulfide	5.2E-03
Chlorobenzene	1.3E-03
Chloroform	2.0E-03
Chloromethane (methyl chloride)	1.1E-03
Cyclohexane	4.4E-03
1,2-Dichlorobenzene	2.3E-02
1,3-Dichlorobenzene	1.6E-03
1,4-Dichlorobenzene	3.6E-03
Dichlorodifluoromethane (Freon 12)	7.5E-02
1,1-Dichloroethane	1.2E-03
1,2-Dichloroethane	1.7E-03
1,1-Dichloroethene (1,1-DCE)	1.4E-03
cis-1,2-Dichloroethene (cis-1,2-DCE)	1.2E-03
1,4-Dioxane	5.3E-03
Ethanol	3.9E-02
Ethylbenzene	6.6E-02
4-Ethyltoluene	2.9E-02
Heptane	4.4E-03
Hexane	1.4E+00
Methyl tertiary butyl ether (MTBE)	5.8E-03
Methylene chloride	1.4E-02
4-Methyl-2-pentanone (methyl isobutyl ketone)	4.7E-03
2-Propanol	4.2E-03
Styrene	1.2E-03
Tetrachloroethene (PCE)	9.0E-03
Tetrahydrofuran	4.9E-03
Toluene	3.2E-01
1,1,1-Trichloroethane	1.7E-03

Table C-3. Exposure Point Concentrations of Chemicals of Potential Concern in Onsite Outdoor Air and in Offsite Air – High-end Estimate

Concentration (μg/m³)
1.7E-03
2.0E-03
2.4E-03
2.0E-02
2.5E-02
1.8E-01

Table C-4. Summary of Estimated Cancer Risks – High-end Estimate

	Onsite Commercial Worker		Onsite Intrusive Construction Worker					
	Indoor Air	Outdoor Air		Soil	Groundwater	Total	Indoor/Outdoor Air	
Chemical of Potential Concern	Inhalation	Inhalation	Ingestion	Dermal Contact	Dermal Contact		Inhalation	
Acetone	NC	NC	-	-	_	=	NC	
Benzene	6.1E-05	8.9E-09	5.2E-08	7.1E-08	7.5E-05	7.5E-05	5.9E-06	
Bromodichloromethane	_	_	_	_	2.6E-10	2.6E-10	-	
1,3-Butadiene	4.1E-06	1.2E-09	_	_	_	1.2E-09	7.8E-07	
2-Butanone (methyl ethyl ketone)	NC	NC	_	_	_	_	NC	
Carbon disulfide	NC	NC	-	-	-	_	NC	
Chlorobenzene	NC	NC	NC	NC	NC	_	NC	
Chloroethane (ethyl chloride)	_	_	_	_	NC	_	-	
Chloroform	5.3E-08	8.5E-12	_	_	5.1E-10	5.2E-10	5.7E-09	
Chloromethane (methyl chloride)	8.4E-09	1.5E-12	=	_	2.8E-10	2.8E-10	1.0E-09	
Cyclohexane	NC	NC	_	_	_	_	NC	
1,2-Dichlorobenzene	NC	NC	NC	NC	NC	_	NC	
1,3-Dichlorobenzene	NC	NC	NC	NC	_	_	NC	
1,4-Dichlorobenzene	2.5E-07	3.2E-11	6.6E-12	9.1E-12	_	4.8E-11	2.1E-08	
Dichlorodifluoromethane (Freon 12)	NC	NC	=	_	NC	=	NC	
1,1-Dichloroethane	1.1E-08	1.5E-12	_	_	4.7E-09	4.7E-09	1.0E-09	
1,2-Dichloroethane	1.7E-07	2.7E-11	7.5E-11	1.0E-10	2.2E-07	2.2E-07	1.8E-08	
1,1-Dichloroethene (1,1-DCE)	NC	NC	_	_	NC	_	NC	
cis-1,2-Dichloroethene (cis-1,2-DCE)	NC	NC	_	_	NC	_	NC	
1,4-Dioxane	2.1E-07	3.2E-11	=	_	_	3.2E-11	2.1E-08	
Ethanol	NC	NC	_	_	_	_	NC	
Ethylbenzene	9.4E-07	1.3E-10	6.9E-09	9.5E-09	2.2E-05	2.2E-05	8.5E-08	
4-Ethyltoluene	NC	NC	_	_	_	_	NC	
Heptane	NC	NC	_	_	_	_	NC	
Hexane	NC	NC	=	_	_	=	NC	
Methyl tertiary butyl ether (MTBE)	7.4E-09	1.2E-12	5.6E-13	7.7E-13	9.0E-09	9.0E-09	7.9E-10	
Methylene chloride	6.8E-08	1.1E-11	_	-	2.0E-10	2.1E-10	7.2E-09	
4-Methyl-2-pentanone (methyl isobutyl ketone)	NC	NC	_	_	=		NC	

Table C-4. Summary of Estimated Cancer Risks – High-end Estimate

	Onsite Commercial Worker		Onsite	Intrusive Construction	Age-adjusted Offsite Resident		
Chemical of Potential Concern	Indoor Air	Outdoor Air	Soil		Groundwater	Total	Indoor/Outdoor Air
	Inhalation	Inhalation	Ingestion	Dermal Contact	Dermal Contact		Inhalation
2-Propanol	NC	NC	-	=	-	=	NC
Styrene	NC	NC	-	-	-	_	NC
1,1,2,2-Tetrachloroethane	_	-	_		4.8E-10	4.8E-10	-
Tetrachloroethene (PCE)	3.2E-07	4.2E-11	_		-	4.2E-11	2.8E-08
Tetrahydrofuran	4.8E-08	7.6E-12	_	-	-	7.6E-12	5.1E-09
Toluene	NC	NC	NC	NC	NC	-	NC
1,1,1-Trichloroethane	NC	NC	_		NC	-	NC
Trichloroethene (TCE)	1.9E-08	2.7E-12	_	-	2.9E-10	3.0E-10	1.8E-09
Trichlorofluoromethane (Freon 11)	NC	NC	_	_	_	_	NC
1,1,2-Trichloro-1,2,2-trifluoroethane (Freon 113)	NC	NC	-	_	_	-	NC
1,2,4-Trimethylbenzene	NC	NC	_	_	_	_	NC
1,3,5-Trimethylbenzene	NC	NC	_	_	_	_	NC
Xylenes	NC	NC	NC	NC	NC	_	NC
Cumulative Risk	6.7E-05	1.0E-08	5.9E-08	8.1E-08	9.7E-05	9.8E-05	6.9E-06

^{(1) &}quot;-" indicates chemical was not determined to be a COPC for the respective pathway.

^{(2) &}quot;NC" indicates chemical is classified as a noncarcinogen.

Table C-5. Summary of Estimated Noncancer Hazard Indices – High-end Estimate

	Onsite Commercial Worker		Onsite	Intrusive Construction	on Worker		Child Offsite Resident
	Indoor Air	Outdoor Air		Soil	Groundwater	Total	Indoor/Outdoor Air
Chemical of Potential Concern	Inhalation	Inhalation	Ingestion	Dermal Contact	Dermal Contact		Inhalation
Acetone	3.5E-06	1.6E-08	_	_	-	1.6E-08	6.3E-07
Benzene	2.0E-01	7.3E-04	9.0E-03	1.2E-02	1.3E+01	1.3E+01	3.0E-02
Bromodichloromethane	-	-	_	_	6.9E-06	6.9E-06	-
1,3-Butadiene	3.3E-02	2.4E-04	-	_	-	2.4E-04	9.8E-03
2-Butanone (methyl ethyl ketone)	1.2E-05	4.2E-08	_	-	-	4.2E-08	1.7E-06
Carbon disulfide	1.0E-04	4.1E-07	_	-	-	4.1E-07	1.7E-05
Chlorobenzene	2.2E-05	7.3E-08	2.2E-08	3.0E-08	4.5E-05	4.5E-05	3.0E-06
Chloroethane (ethyl chloride)	-	_	_	_	2.5E-05	2.5E-05	-
Chloroform	9.1E-05	3.7E-07	_	_	1.1E-04	1.1E-04	1.5E-05
Chloromethane (methyl chloride)	1.4E-04	6.5E-07	_	_	5.9E-05	6.0E-05	2.7E-05
Cyclohexane	1.1E-05	4.0E-08	_	_	_	4.0E-08	1.6E-06
1,2-Dichlorobenzene	1.9E-03	6.2E-06	8.9E-06	1.2E-05	8.2E-06	3.6E-05	2.5E-04
1,3-Dichlorobenzene	2.6E-04	8.6E-07	3.3E-07	4.5E-07	_	1.6E-06	3.5E-05
1,4-Dichlorobenzene	7.5E-05	2.5E-07	2.8E-06	3.9E-06	-	7.0E-06	1.0E-05
Dichlorodifluoromethane (Freon 12)	6.4E-03	2.1E-05	_	_	6.1E-06	2.7E-05	8.4E-04
1,1-Dichloroethane	3.9E-05	1.3E-07	_	_	2.9E-04	2.9E-04	5.4E-06
1,2-Dichloroethane	5.6E-05	2.3E-07	5.6E-06	7.7E-06	1.6E-02	1.6E-02	9.2E-06
1,1-Dichloroethene (1,1-DCE)	3.0E-04	1.1E-06	_	_	2.6E-04	2.6E-04	4.6E-05
cis-1,2-Dichloroethene (cis-1,2-DCE)	5.4E-04	1.8E-06	_	_	2.0E-04	2.0E-04	7.5E-05
1,4-Dioxane	2.6E-05	9.7E-08	_	_	_	9.7E-08	4.0E-06
Ethanol	4.4E-04	2.0E-06	_	_	_	2.0E-06	8.2E-05
Ethylbenzene	1.1E-03	3.6E-06	4.4E-04	6.1E-04	1.4E+00	1.4E+00	1.5E-04
4-Ethyltoluene	4.8E-03	1.6E-05	_	_	-	1.6E-05	6.4E-04
Heptane	1.0E-04	3.4E-07	_	_	_	3.4E-07	1.4E-05
Hexane	1.8E-02	1.1E-04	_	_	_	1.1E-04	4.6E-03
Methyl tertiary butyl ether (MTBE)	2.7E-05	1.1E-07	2.5E-08	3.5E-08	4.1E-04	4.1E-04	4.3E-06
Methylene chloride	4.8E-04	1.9E-06	_	_	1.7E-05	1.9E-05	7.7E-05

Table C-5. Summary of Estimated Noncancer Hazard Indices – High-end Estimate

	Onsite Commercial Worker		Onsite	Intrusive Construction	on Worker		Child Offsite Resident
	Indoor Air	Outdoor Air		Soil	Groundwater	Total	Indoor/Outdoor Air
Chemical of Potential Concern	Inhalation	Inhalation	Ingestion	Dermal Contact	Dermal Contact		Inhalation
4-Methyl-2-pentanone (methyl isobutyl ketone)	2.5E-05	8.6E-08	-	-	_	8.6E-08	3.5E-06
2-Propanol	8.1E-06	3.3E-08	-	-	-	3.3E-08	1.4E-06
Styrene	2.2E-05	7.4E-08	-	-	-	7.4E-08	3.0E-06
1,1,2,2-Tetrachloroethane	_	_	_		3.1E-05	3.1E-05	_
Tetrachloroethene (PCE)	4.2E-03	1.4E-05	_	-	_	1.4E-05	5.7E-04
Tetrahydrofuran	2.2E-05	8.9E-08	_		_	8.9E-08	3.6E-06
Toluene	1.6E-02	5.8E-05	1.9E-03	2.6E-03	1.5E+00	1.5E+00	2.4E-03
1,1,1-Trichloroethane	2.7E-05	9.2E-08	_		2.3E-07	3.2E-07	3.8E-06
Trichloroethene (TCE)	4.5E-05	1.6E-07	_		5.3E-03	5.3E-03	6.3E-06
Trichlorofluoromethane (Freon 11)	4.2E-05	1.5E-07	_	-	_	1.5E-07	6.3E-06
1,1,2-Trichloro-1,2,2-trifluoroethane (Freon 113)	1.3E-06	4.4E-09	_		_	4.4E-09	1.8E-07
1,2,4-Trimethylbenzene	5.1E-02	1.6E-04	_	_	_	1.6E-04	6.4E-03
1,3,5-Trimethylbenzene	7.3E-02	2.2E-04	_	-	_	2.2E-04	9.2E-03
Xylenes	2.8E-02	1.0E-04	1.3E-03	1.8E-03	4.7E+00	4.7E+00	4.1E-03
Cumulative Hazard	4.4E-01	1.7E-03	1.3E-02	1.7E-02	2.1E+01	2.1E+01	6.9E-02

^{(1) &}quot;-" indicates chemical was not determined to be a COPC for the respective pathway.

APPENDIX D LOW-END ESTIMATE OF RISK AND HAZARD

Table D-1. Average Concentrations of Chemicals of Potential Concern in Soil Gas

Chemical of Potential Concern	Concentration (μg/m³)
Acetone	180
Benzene	2,900
1,3-Butadiene	51
2-Butanone (methyl ethyl ketone)	70
Carbon disulfide	62
Chlorobenzene	20
Chloroform	35
Chloromethane (methyl chloride)	9.9
Cyclohexane	72
1,2-Dichlorobenzene	210
1,3-Dichlorobenzene	27
1,4-Dichlorobenzene	45
Dichlorodifluoromethane (Freon 12)	940
1,1-Dichloroethane	32
1,2-Dichloroethane	32
1,1-Dichloroethene (1,1-DCE)	32
cis-1,2-Dichloroethene (cis-1,2-DCE)	32
1,4-Dioxane	71
Ethanol	350
Ethylbenzene	330
4-Ethyltoluene	290
Heptane	72
Hexane	4,400
Methyl tertiary butyl ether (MTBE)	64
Methylene chloride	77
4-Methyl-2-pentanone (methyl isobutyl ketone)	77
2-Propanol	52
Styrene	20
Tetrachloroethene (PCE)	80
Tetrahydrofuran	63
Toluene	1,300
1,1,1-Trichloroethane	39
Trichloroethene (TCE)	36

Table D-1. Average Concentrations of Chemicals of Potential Concern in Soil Gas

Chemical of Potential Concern	Concentration (µg/m³)
Trichlorofluoromethane (Freon 11)	39
1,1,2-Trichloro-1,2,2-trifluoroethane (Freon 113)	50
1,2,4-Trimethylbenzene	210
1,3,5-Trimethylbenzene	250
Xylenes	890

⁽¹⁾ Soil gas samples were collected at depths of 3 and 5 feet below ground surface (bgs) in 1999 and 2008, respectively.

⁽²⁾ Concentration units are micrograms per cubic meter (µg/m³).

⁽³⁾ Representative concentration is the mean of all samples collected in 1999 and 2008. For purposes of this averaging, non-detect results are assumed equal to one-half the reporting limit.

Table D-2. Johnson and Ettinger Model Inputs – Low-end Estimate

Parameter	Symbol	Value	Units	Reference
Building Properties				
Depth below grade to bottom of enclosed space floor	L_{F}	15	cm	DTSC/HERD default (Cal/EPA, 2005a; 2005b)
Area of enclosed space below grade	$A_{b,sg}$	1.00E+06	cm ²	DTSC default (Cal/EPA, 2005b)
Building air exchange rate	AXR_b	4	hr ⁻¹	DTSC default (Cal/EPA, 2005b)
Building height	Bh	503	cm	DTSC default (Cal/EPA, 2005b)
Building ventilation rate	Q_b	5.59E+05	cm ³ /s	Calculated: $A_{b,sg} x AXR_b x B_h$
Vapor flow rate into building	Q_{soil}	5	L/min	Calculated (Cal/EPA, 2005b)
Vapor flow rate into building	Q_{soil}	83	cm^3/s	Calculated via units conversion
Soil Properties				
Average soil temperature	T_s	17	°C	Site-specific (USEPA, 2004)
SCS soil type	_	Site-specific	_	Site-specific
Dry bulk density	$ ho_b$	1.79	g/cm ³	Site-specific average
Total porosity	n	0.339	cm ³ /cm ³	Site-specific average
Water-filled porosity	θ_{w}	0.236	cm ³ /cm ³	Site-specific average
Air-filled porosity	θ_a	0.103	cm ³ /cm ³	Site-specific average

Table D-3. Exposure Point Concentrations of Chemicals of Potential Concern in the Indoor Air of the Onsite Building – Low-end Estimate

	Modeled Sou		Results of Vapor Intrusion Modeling		
Chemical of Potential Concern	C_{SG} ($\mu g/m^3$)	Depth (cm)	α	C_{IA} $(\mu g/m^3)$	
Acetone	180	91	2.5E-05	4.4E-03	
Benzene	2,900	91	8.7E-06	2.5E-02	
1,3-Butadiene	51	91	2.2E-05	1.1E-03	
2-Butanone (methyl ethyl ketone)	70	91	1.7E-05	1.2E-03	
Carbon disulfide	62	91	1.0E-05	6.3E-04	
Chlorobenzene	20	91	7.4E-06	1.5E-04	
Chloroform	35	91	1.0E-05	3.6E-04	
Chloromethane (methyl chloride)	9.9	91	1.2E-05	1.2E-04	
Cyclohexane	72	91	7.9E-06	5.7E-04	
1,2-Dichlorobenzene	210	91	7.1E-06	1.5E-03	
1,3-Dichlorobenzene	27	91	7.0E-06	1.9E-04	
1,4-Dichlorobenzene	45	91	7.1E-06	3.2E-04	
Dichlorodifluoromethane (Freon 12)	940	91	6.6E-06	6.2E-03	
1,1-Dichloroethane	32	91	7.4E-06	2.4E-04	
1,2-Dichloroethane	32	91	1.1E-05	3.4E-04	
1,1-Dichloroethene (1,1-DCE)	32	91	8.8E-06	2.8E-04	
cis-1,2-Dichloroethene (cis-1,2-DCE)	32	91	7.4E-06	2.4E-04	
1,4-Dioxane	71	91	6.0E-05	4.3E-03	
Ethanol	350	91	6.5E-05	2.3E-02	
Ethylbenzene	330	91	7.5E-06	2.5E-03	
4-Ethyltoluene	290	91	6.8E-06	2.0E-03	
Heptane	72	91	7.0E-06	5.0E-04	
Hexane	4,400	91	1.8E-05	8.0E-02	
Methyl tertiary butyl ether (MTBE)	64	91	1.1E-05	6.9E-04	
Methylene chloride	77	91	1.0E-05	7.8E-04	
4-Methyl-2-pentanone (methyl isobutyl ketone)	77	91	1.1E-05	8.1E-04	
2-Propanol	52	91	4.4E-05	2.3E-03	
Styrene	20	91	7.2E-06	1.4E-04	
Tetrachloroethene (PCE)	80	91	7.2E-06	5.7E-04	
Tetrahydrofuran	63	91	1.5E-05	9.4E-04	

Table D-3. Exposure Point Concentrations of Chemicals of Potential Concern in the Indoor Air of the Onsite Building – Low-end Estimate

	Modeled Sou		Results of Va Mod	por Intrusion eling
Chemical of Potential Concern	C_{SG} (μ g/m ³)	Depth (cm)	α	C_{IA} ($\mu g/m^3$)
Toluene	1,300	91	8.6E-06	1.1E-02
1,1,1-Trichloroethane	39	91	7.7E-06	3.0E-04
Trichloroethene (TCE)	36	91	7.9E-06	2.8E-04
Trichlorofluoromethane (Freon 11)	39	91	8.6E-06	3.3E-04
1,1,2-Trichloro-1,2,2-trifluoroethane (Freon 113)	50	91	7.7E-06	3.9E-04
1,2,4-Trimethylbenzene	210	91	6.1E-06	1.3E-03
1,3,5-Trimethylbenzene	250	91	6.1E-06	1.5E-03
Xylenes	890	91	8.4E-06	7.5E-03

- (1) This vapor intrusion transport analysis is based on average concentrations of volatile organic compounds (VOCs) in soil gas from the August 1999 and May 2008 site investigations (see Table 1). Chemicals of potential concern (COPCs) with respect to vapor intrusion are those VOCs detected above reporting limits in at least one soil gas sample. For the purpose of calculating average concentrations, non-detect results are assumed equal to one-half the laboratory reporting limit.
- (2) The shallower sampling depth between the 1999 and 2008 sampling events, 3 feet below ground surface, is used in the model because it is a more conservative assumption, *i.e.*, it produces higher indoor air concentrations.
- (3) Non source-related inputs to the Johnson and Ettinger Model are documented in Table D-1. Shown here are the results of the Johnson and Ettinger Model, consisting of, for each chemical of potential concern, the predicted attenuation factor (α) and the predicted concentration of the chemical in indoor air (C_{IA}).

Table D-4. Exposure Point Concentrations of Chemicals of Potential Concern in Onsite Outdoor Air and in Offsite Air – Low-end Estimate

Chemical of Potential Concern	Concentration
	$(\mu g/m^3)$
Acetone	2.8E-04
Benzene	1.7E-03
1,3-Butadiene	8.2E-05
2-Butanone (methyl ethyl ketone)	6.7E-05
Carbon disulfide	4.2E-05
Chlorobenzene	9.5E-06
Chloroform	2.4E-05
Chloromethane (methyl chloride)	8.0E-06
Cyclohexane	3.7E-05
1,2-Dichlorobenzene	9.5E-05
1,3-Dichlorobenzene	1.2E-05
1,4-Dichlorobenzene	2.0E-05
Dichlorodifluoromethane (Freon 12)	4.0E-04
1,1-Dichloroethane	1.5E-05
1,2-Dichloroethane	2.2E-05
1,1-Dichloroethene (1,1-DCE)	1.9E-05
cis-1,2-Dichloroethene (cis-1,2-DCE)	1.5E-05
1,4-Dioxane	4.4E-04
Ethanol	2.6E-03
Ethylbenzene	1.6E-04
4-Ethyltoluene	1.3E-04
Heptane	3.3E-05
Hexane	5.7E-03
Methyl tertiary butyl ether (MTBE)	4.5E-05
Methylene chloride	5.1E-05
4-Methyl-2-pentanone (methyl isobutyl ketone)	4.8E-05
2-Propanol	2.0E-04
Styrene	9.3E-06
Tetrachloroethene (PCE)	3.7E-05
Tetrahydrofuran	6.5E-05
Toluene	7.3E-04
1,1,1-Trichloroethane	2.0E-05

Table D-4. Exposure Point Concentrations of Chemicals of Potential Concern in Onsite Outdoor Air and in Offsite Air – Low-end Estimate

Trichloroethene (TCE) 1.8E-05 Trichlorofluoromethane (Freon 11) 2.2E-05 1,1,2-Trichloro-1,2,2-trifluoroethane (Freon 113) 2.5E-05 1,2,4-Trimethylbenzene 8.3E-05 1,3,5-Trimethylbenzene 9.8E-05 Xylenes 4.9E-04	Chemical of Potential Concern	Concentration $(\mu g/m^3)$
1,1,2-Trichloro-1,2,2-trifluoroethane (Freon 113)2.5E-051,2,4-Trimethylbenzene8.3E-051,3,5-Trimethylbenzene9.8E-05	Trichloroethene (TCE)	1.8E-05
1,2,4-Trimethylbenzene8.3E-051,3,5-Trimethylbenzene9.8E-05	Trichlorofluoromethane (Freon 11)	2.2E-05
1,3,5-Trimethylbenzene 9.8E-05	1,1,2-Trichloro-1,2,2-trifluoroethane (Freon 113)	2.5E-05
	1,2,4-Trimethylbenzene	8.3E-05
Xylenes 4.9E-04	1,3,5-Trimethylbenzene	9.8E-05
	Xylenes	4.9E-04

Table D-5. Summary of Estimated Cancer Risks – Low-end Estimate

	Onsite Commercial Worker		Age-adjusted Offsite Resident				
Chemical of Potential Concern	Indoor Air	Outdoor Air		Soil	Groundwater Dermal Contact	Total	Indoor/Outdoor Air
	Inhalation	Inhalation	Ingestion	Dermal Contact			Inhalation
Acetone	NC	NC	-	=	_	=	NC
Benzene	1.2E-07	3.7E-11	5.2E-08	7.1E-08	7.5E-05	7.5E-05	2.5E-08
Bromodichloromethane	-	-	_	-	2.6E-10	2.6E-10	-
1,3-Butadiene	3.3E-08	1.1E-11	_	_	_	1.1E-11	7.3E-09
2-Butanone (methyl ethyl ketone)	NC	NC	_	_	_	_	NC
Carbon disulfide	NC	NC	_		-	_	NC
Chlorobenzene	NC	NC	NC	NC	NC	_	NC
Chloroethane (ethyl chloride)	-	_	_	-	NC	_	_
Chloroform	3.3E-10	1.0E-13	-	_	5.1E-10	5.1E-10	6.7E-11
Chloromethane (methyl chloride)	3.7E-11	1.1E-14	_	_	2.8E-10	2.8E-10	7.5E-12
Cyclohexane	NC	NC	-	_	_	=	NC
1,2-Dichlorobenzene	NC	NC	NC	NC	NC		NC
1,3-Dichlorobenzene	NC	NC	NC	NC	=		NC
1,4-Dichlorobenzene	6.2E-10	1.8E-13	6.6E-12	9.1E-12	=	1.6E-11	1.2E-10
Dichlorodifluoromethane (Freon 12)	NC	NC	_	_	NC		NC
1,1-Dichloroethane	6.6E-11	2.0E-14	_	_	4.7E-09	4.7E-09	1.3E-11
1,2-Dichloroethane	1.2E-09	3.6E-13	7.5E-11	1.0E-10	2.2E-07	2.2E-07	2.4E-10
1,1-Dichloroethene (1,1-DCE)	NC	NC	_	_	NC		NC
cis-1,2-Dichloroethene (cis-1,2-DCE)	NC	NC	_	-	NC	_	NC
1,4-Dioxane	5.6E-09	2.7E-12	_	-	_	2.7E-12	1.8E-09
Ethanol	NC	NC	_	_	=		NC
Ethylbenzene	1.1E-09	3.1E-13	6.9E-09	9.5E-09	2.2E-05	2.2E-05	2.1E-10
4-Ethyltoluene	NC	NC	_	-	_	_	NC
Heptane	NC	NC	_	_	_	_	NC
Hexane	NC	NC	_	_	_	_	NC
Methyl tertiary butyl ether (MTBE)	3.1E-11	9.1E-15	5.6E-13	7.7E-13	9.0E-09	9.0E-09	6.1E-12
Methylene chloride	1.3E-10	4.0E-14	_	_	2.0E-10	2.0E-10	2.7E-11
4-Methyl-2-pentanone (methyl isobutyl ketone)	NC	NC	_	_	_	_	NC

Table D-5. Summary of Estimated Cancer Risks – Low-end Estimate

	Onsite Commercial Worker		Age-adjusted Offsite Resident				
	Indoor Air	Outdoor Air	Soil		Groundwater	Total	Indoor/Outdoor Air
Chemical of Potential Concern	Inhalation	Inhalation	Ingestion	Dermal Contact	Dermal Contact		Inhalation
2-Propanol	NC	NC	-	=	_	=	NC
Styrene	NC	NC	_	-	-	-	NC
1,1,2,2-Tetrachloroethane	_	-	_	-	4.8E-10	4.8E-10	-
Tetrachloroethene (PCE)	5.9E-10	1.7E-13	_	-	-	1.7E-13	1.2E-10
Tetrahydrofuran	3.2E-10	1.0E-13	_	_	_	1.0E-13	6.8E-11
Toluene	NC	NC	NC	NC	NC	-	NC
1,1,1-Trichloroethane	NC	NC	_	-	NC	-	NC
Trichloroethene (TCE)	9.7E-11	2.9E-14	_	-	2.9E-10	2.9E-10	1.9E-11
Trichlorofluoromethane (Freon 11)	NC	NC	_	_	_	_	NC
1,1,2-Trichloro-1,2,2-trifluoroethane (Freon 113)	NC	NC	_	_	_	_	NC
1,2,4-Trimethylbenzene	NC	NC	_	_	-	_	NC
1,3,5-Trimethylbenzene	NC	NC	_	_	_	_	NC
Xylenes	NC	NC	NC	NC	NC	_	NC
Cumulative Risk	1.7E-07	5.2E-11	5.9E-08	8.1E-08	9.7E-05	9.8E-05	3.5E-08

^{(1) &}quot;-" indicates chemical was not determined to be a COPC for the respective pathway.

^{(2) &}quot;NC" indicates chemical is classified as a noncarcinogen.

Table D-6. Summary of Estimated Noncancer Hazard Indices – Low-end Estimate

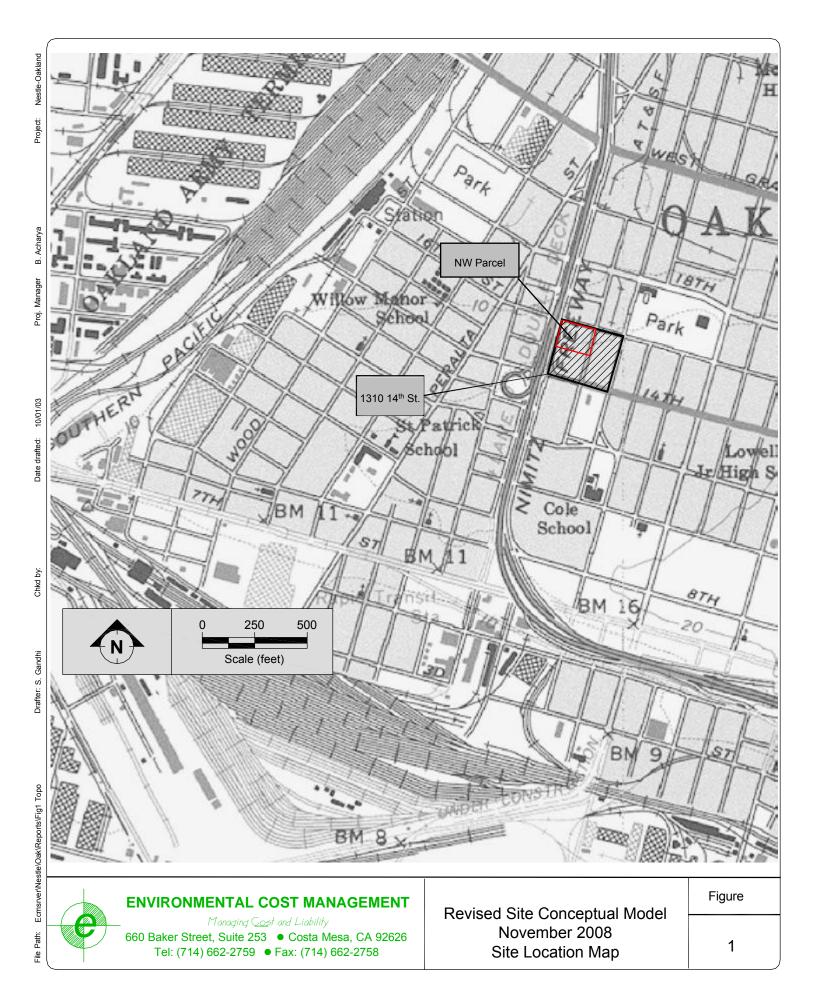
	Onsite Commercial Worker		Onsite	Intrusive Construction	on Worker		Child Offsite Resident Indoor/Outdoor Air
Chemical of Potential Concern	Indoor Air	Outdoor Air		Soil	Groundwater	Total	
	Inhalation	Inhalation	Ingestion	Dermal Contact	Dermal Contact		Inhalation
Acetone	6.9E-08	4.9E-10	_	_	_	4.9E-10	2.0E-08
Benzene	4.0E-04	3.0E-06	9.0E-03	1.2E-02	1.3E+01	1.3E+01	1.2E-04
Bromodichloromethane	_	-	-	_	6.9E-06	6.9E-06	-
1,3-Butadiene	2.7E-04	2.2E-06	-	_	_	2.2E-06	9.1E-05
2-Butanone (methyl ethyl ketone)	1.1E-07	7.4E-10	_		_	7.4E-10	3.0E-08
Carbon disulfide	4.3E-07	3.3E-09	-	_	-	3.3E-09	1.3E-07
Chlorobenzene	7.1E-08	5.2E-10	2.2E-08	3.0E-08	4.5E-05	4.5E-05	2.1E-08
Chloroethane (ethyl chloride)	_	_	_		2.5E-05	2.5E-05	-
Chloroform	5.7E-07	4.3E-09	_		1.1E-04	1.1E-04	1.8E-07
Chloromethane (methyl chloride)	6.4E-07	4.9E-09	_	_	5.9E-05	5.9E-05	2.0E-07
Cyclohexane	4.5E-08	3.4E-10	_		_	3.4E-10	1.4E-08
1,2-Dichlorobenzene	3.6E-06	2.6E-08	8.9E-06	1.2E-05	8.2E-06	2.9E-05	1.1E-06
1,3-Dichlorobenzene	8.7E-07	6.4E-09	3.3E-07	4.5E-07	_	7.9E-07	2.6E-07
1,4-Dichlorobenzene	1.9E-07	1.4E-09	2.8E-06	3.9E-06	_	6.8E-06	5.7E-08
Dichlorodifluoromethane (Freon 12)	1.5E-05	1.1E-07	_	_	6.1E-06	6.2E-06	4.5E-06
1,1-Dichloroethane	2.3E-07	1.7E-09	-	-	2.9E-04	2.9E-04	7.0E-08
1,2-Dichloroethane	4.1E-07	3.0E-09	5.6E-06	7.7E-06	1.6E-02	1.6E-02	1.2E-07
1,1-Dichloroethene (1,1-DCE)	1.9E-06	1.5E-08	_	_	2.6E-04	2.6E-04	5.9E-07
cis-1,2-Dichloroethene (cis-1,2-DCE)	3.3E-06	2.4E-08	-	_	2.0E-04	2.0E-04	9.8E-07
1,4-Dioxane	6.8E-07	8.1E-09	-	_	_	8.1E-09	3.3E-07
Ethanol	1.0E-05	1.3E-07	-	_	_	1.3E-07	5.5E-06
Ethylbenzene	1.2E-06	8.8E-09	4.4E-04	6.1E-04	1.4E+00	1.4E+00	3.6E-07
4-Ethyltoluene	9.5E-06	7.0E-08	_	_	_	7.0E-08	2.9E-06
Heptane	3.4E-07	2.6E-09	-	_	_	2.6E-09	1.0E-07
Hexane	5.5E-05	4.4E-07	_	_	_	4.4E-07	1.8E-05
Methyl tertiary butyl ether (MTBE)	1.1E-07	8.2E-10	2.5E-08	3.5E-08	4.1E-04	4.1E-04	3.3E-08
Methylene chloride	9.3E-07	7.0E-09	_	_	1.7E-05	1.7E-05	2.9E-07

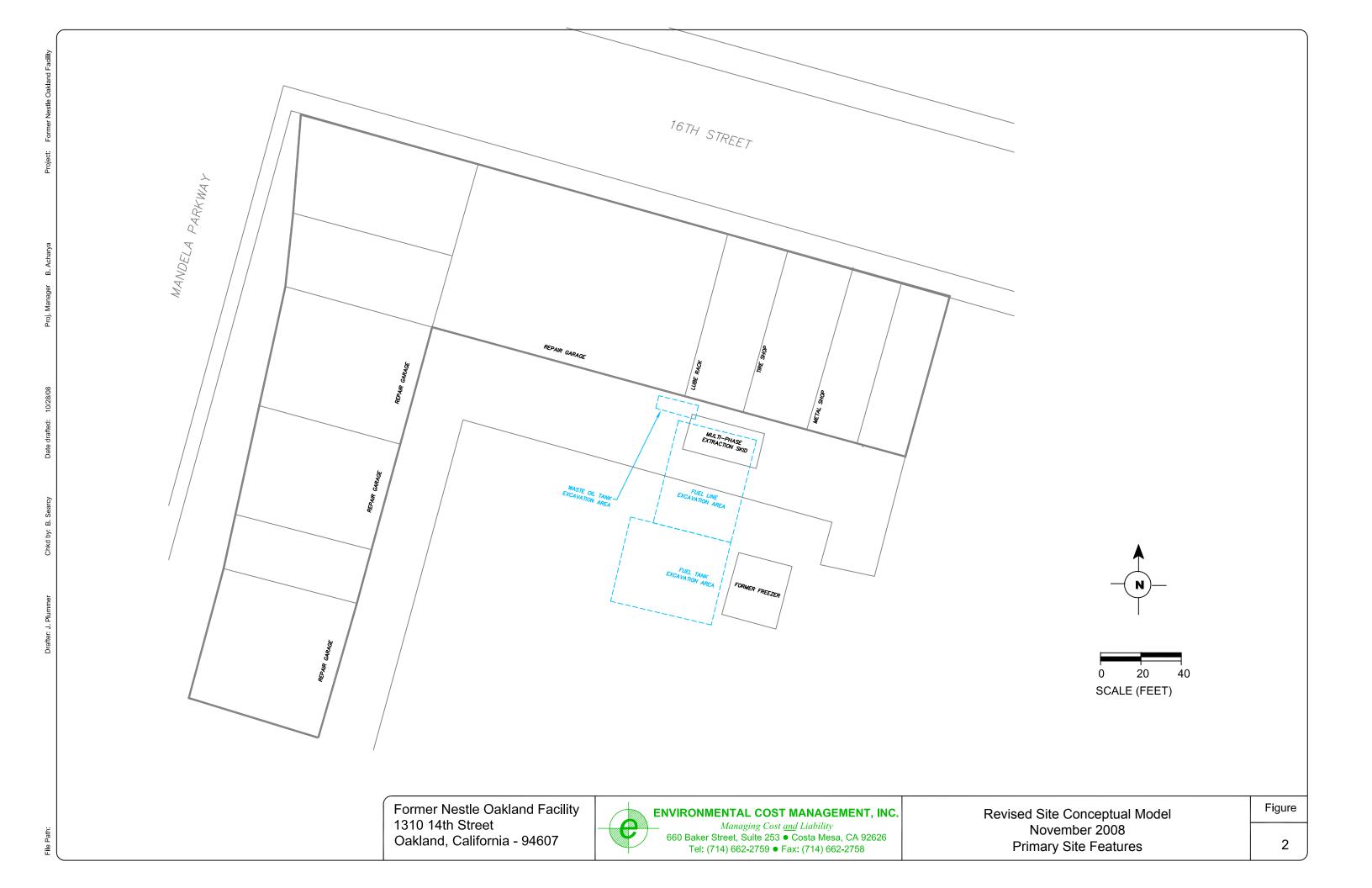
Table D-6. Summary of Estimated Noncancer Hazard Indices – Low-end Estimate

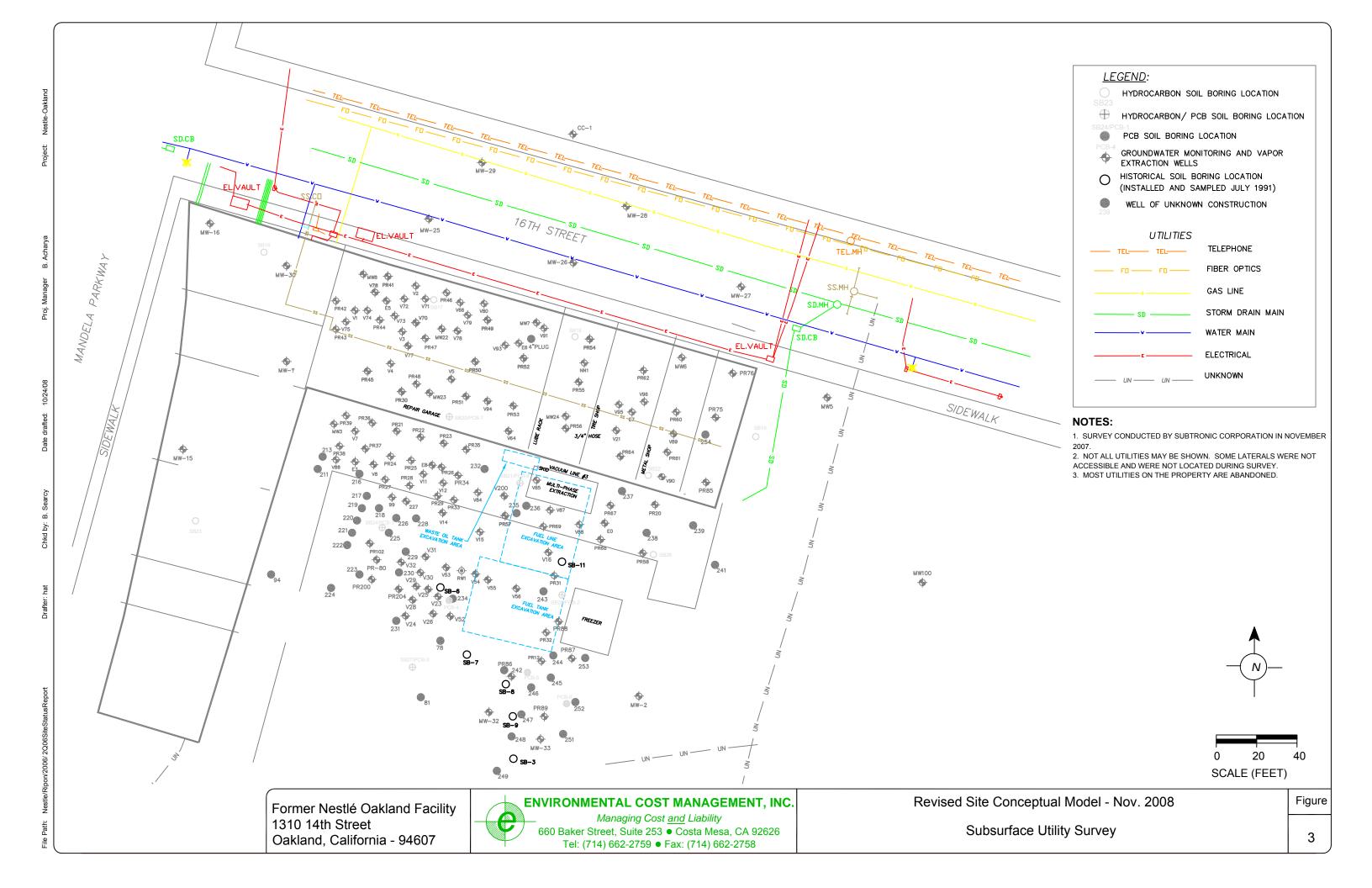
	Onsite Commercial Worker			Child Offsite Resident			
	Indoor Air	Outdoor Air		Soil	Groundwater	Total	Indoor/Outdoor Air
Chemical of Potential Concern	Inhalation	Inhalation	Ingestion	Dermal Contact	Dermal Contact		Inhalation
4-Methyl-2-pentanone (methyl isobutyl ketone)	1.3E-07	8.8E-10	-	-	_	8.8E-10	3.6E-08
2-Propanol	1.6E-07	1.6E-09	-	-	_	1.6E-09	6.5E-08
Styrene	7.7E-08	5.7E-10	_	-	_	5.7E-10	2.3E-08
1,1,2,2-Tetrachloroethane	-	_	-	-	3.1E-05	3.1E-05	-
Tetrachloroethene (PCE)	7.9E-06	5.8E-08	_	-	_	5.8E-08	2.4E-06
Tetrahydrofuran	1.5E-07	1.2E-09	_	-	_	1.2E-09	4.9E-08
Toluene	1.8E-05	1.3E-07	1.9E-03	2.6E-03	1.5E+00	1.5E+00	5.5E-06
1,1,1-Trichloroethane	1.4E-07	1.1E-09	_	-	2.3E-07	2.3E-07	4.4E-08
Trichloroethene (TCE)	2.3E-07	1.7E-09	-	-	5.3E-03	5.3E-03	6.9E-08
Trichlorofluoromethane (Freon 11)	2.3E-07	1.7E-09	-	_	_	1.7E-09	7.0E-08
1,1,2-Trichloro-1,2,2-trifluoroethane (Freon 113)	6.2E-09	4.6E-11	-	_	_	4.6E-11	1.9E-09
1,2,4-Trimethylbenzene	8.8E-05	6.5E-07	-	_	_	6.5E-07	2.6E-05
1,3,5-Trimethylbenzene	1.2E-04	8.9E-07	_	_	_	8.9E-07	3.6E-05
Xylenes	3.6E-05	2.7E-07	1.3E-03	1.8E-03	4.7E+00	4.7E+00	1.1E-05
Cumulative Hazard	1.1E-03	8.1E-06	1.3E-02	1.7E-02	2.1E+01	2.1E+01	3.3E-04

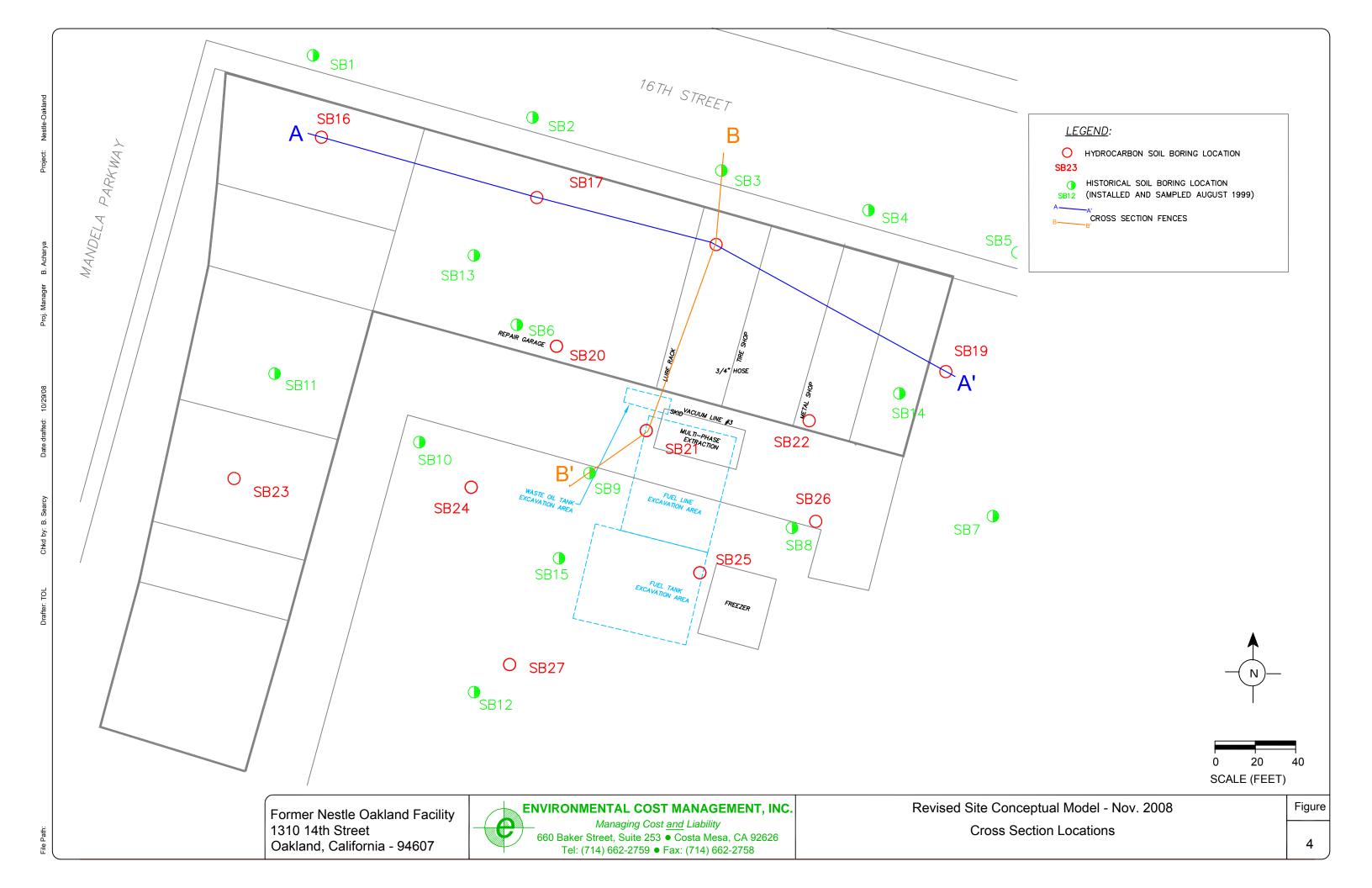
^{(1) &}quot;-" indicates chemical was not determined to be a COPC for the respective pathway.

Appendix B: Site Conceptual Model Figures

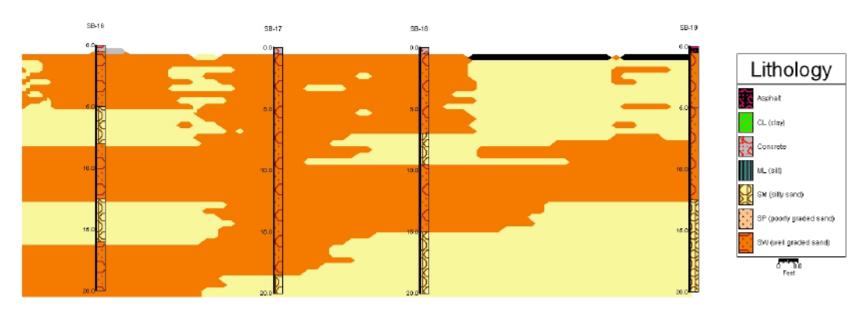








W E



Former Nestle USA, Inc. Facility 1310 14th Street Oakland, California 94607



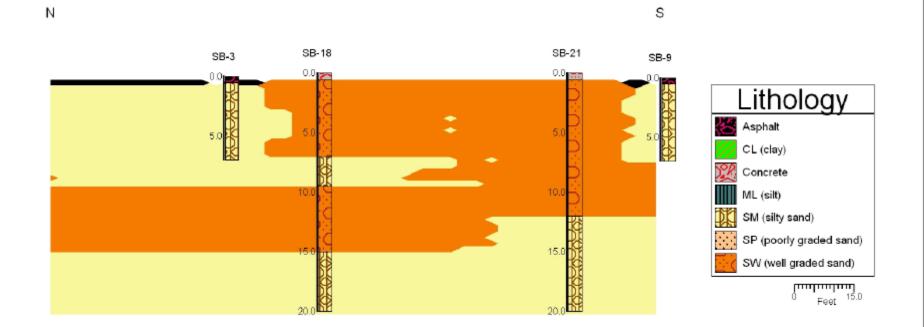
ENVIRONMENTAL COST MANAGEMENT, INC.

Managing Cost <u>and</u> Liability
660 Baker Street, Suite 253 ● Costa Mesa, CA 92626
Tel: (714) 662-2759 ● Fax: (714) 662-2758

Revised Site Conceptual Model – Nov. 2008 E-W Cross Section

Figure

5



Former Nestle USA, Inc. Facility 1310 14th Street Oakland, California 94607

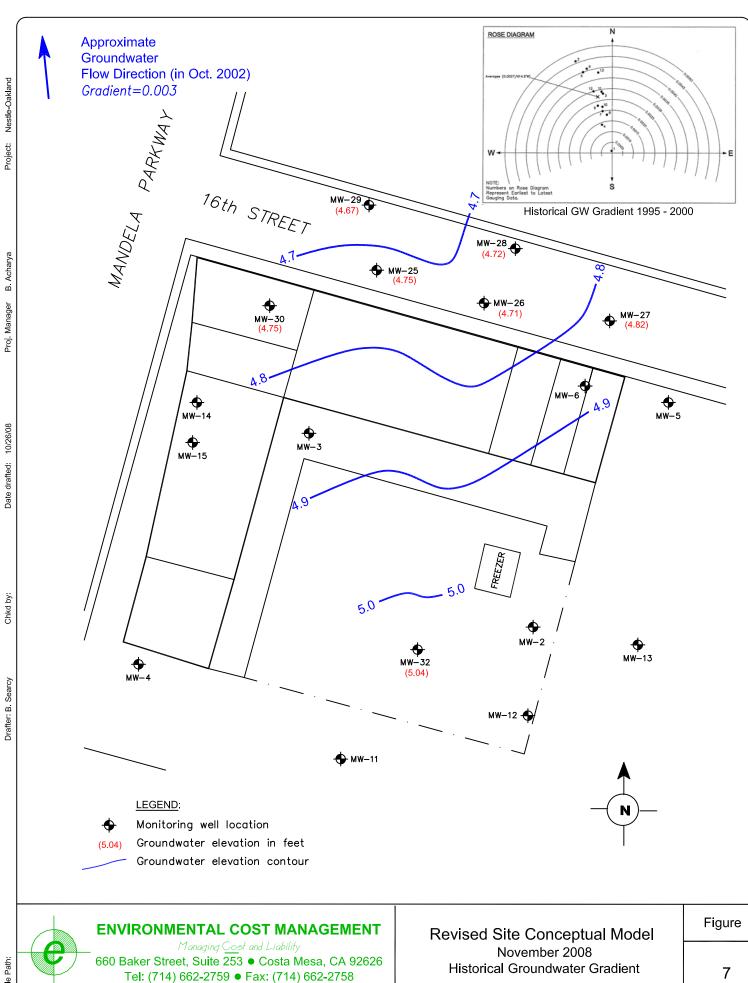


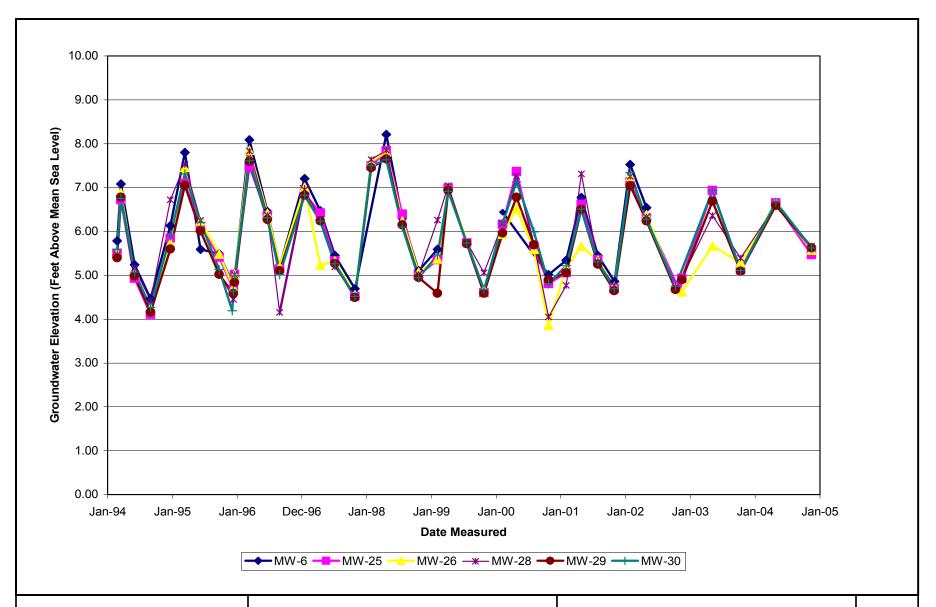
ENVIRONMENTAL COST MANAGEMENT, INC.

- Managing Cost <u>and</u> Liability 660 Baker Street, Suite 253 ● Costa Mesa, CA 92626 Tel: (714) 662-2759 ● Fax: (714) 662-2758 Revised Site Conceptual Model – Nov. 2008 N-S Cross Section

Figure

6





Former Nestlé Oakland Facility 1310 14th Street Oakland, California - 94607

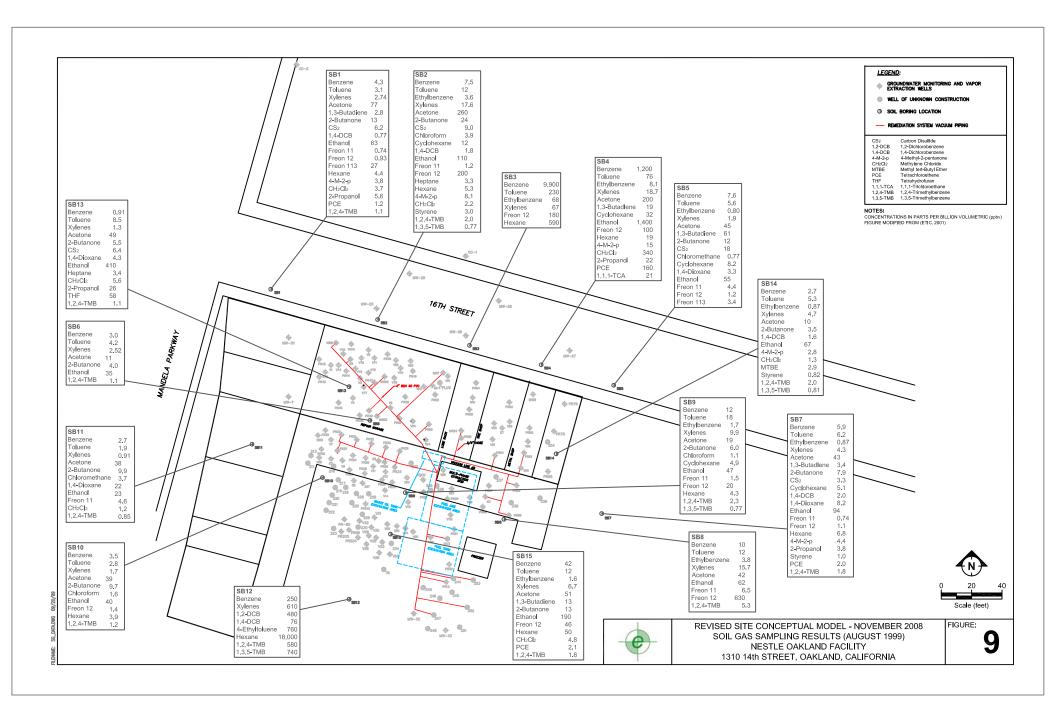


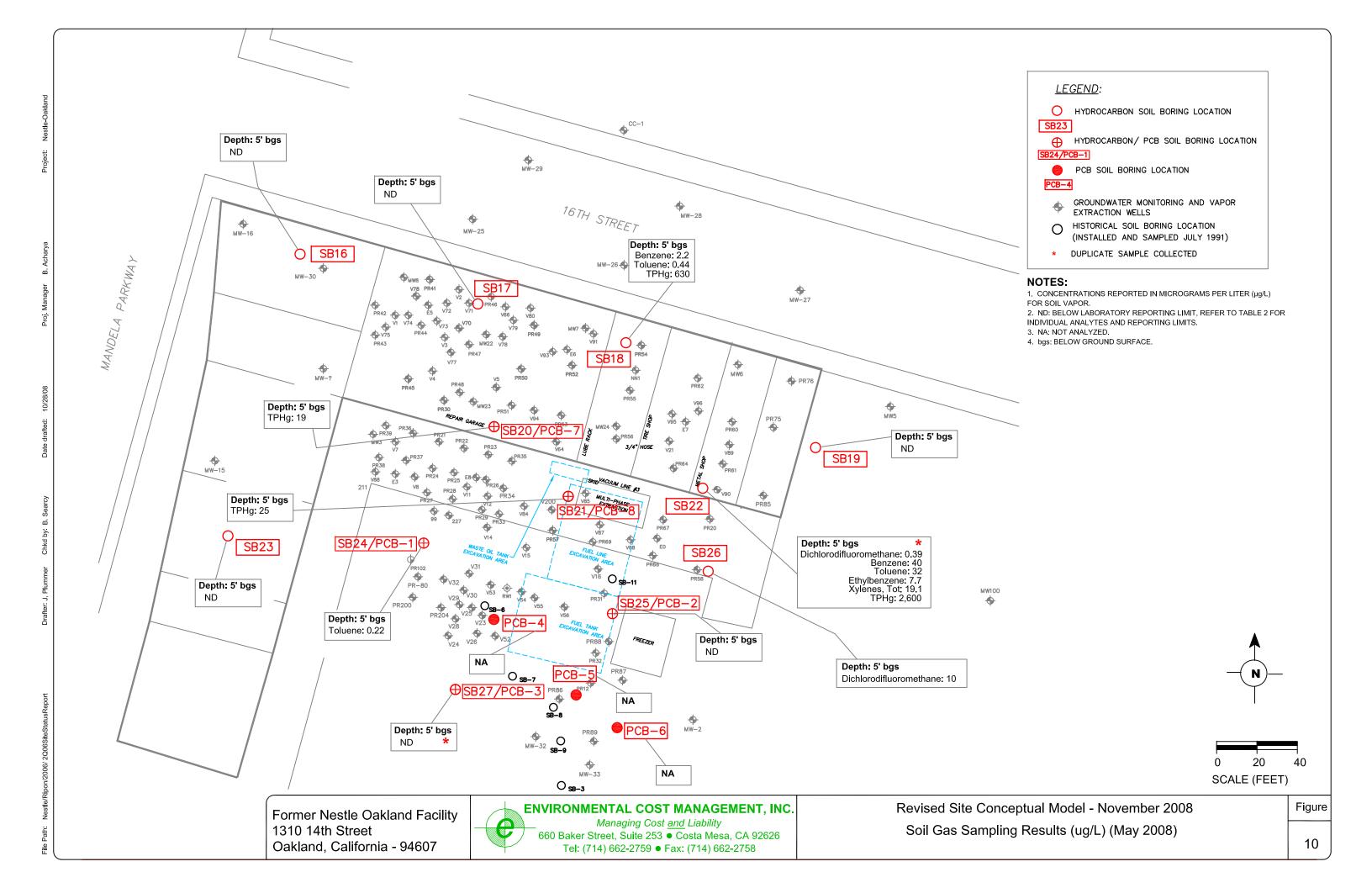
Revised Site Conceptual Model – Nov. 2008

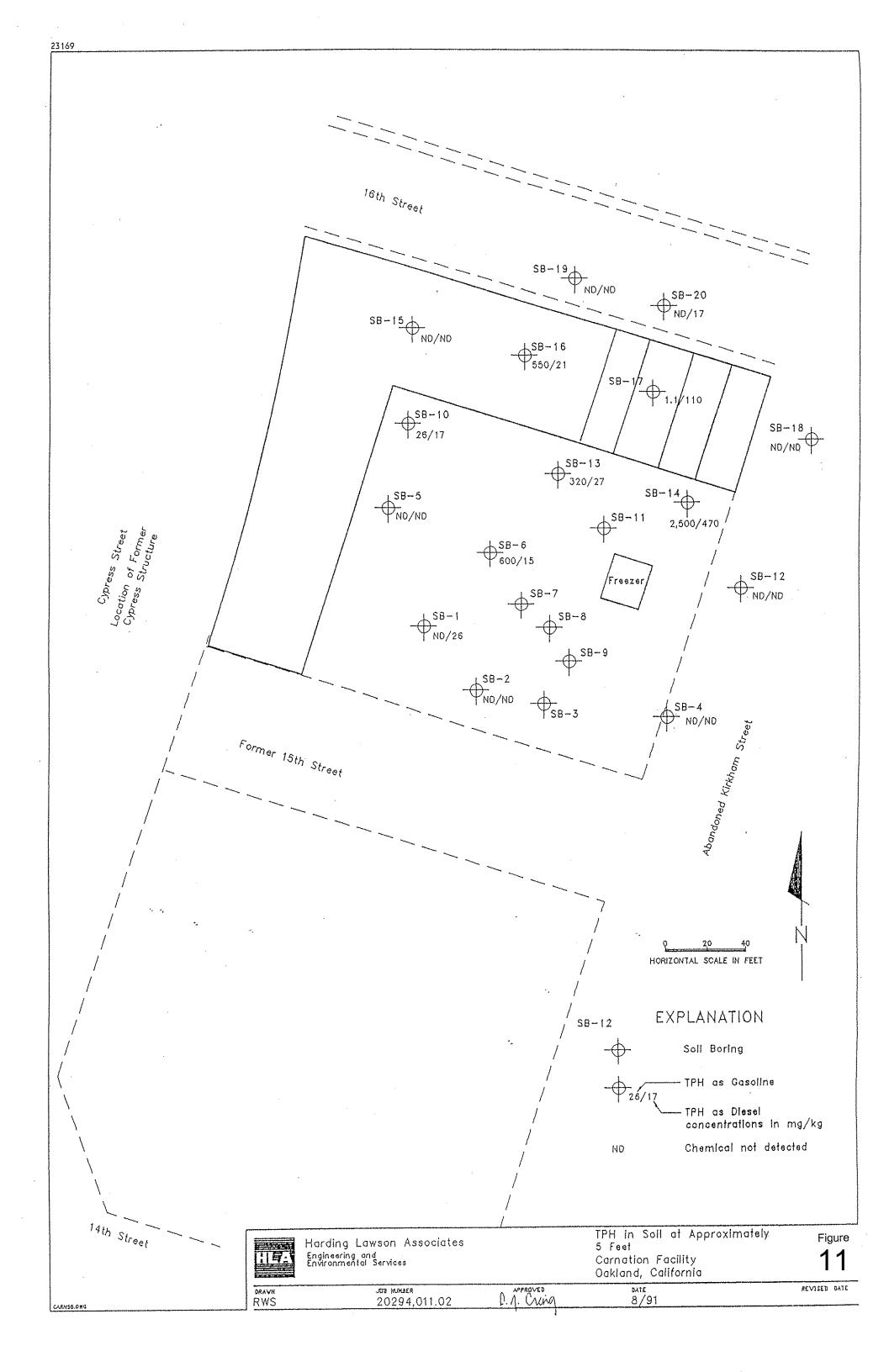
Groundwater Elevation Trends

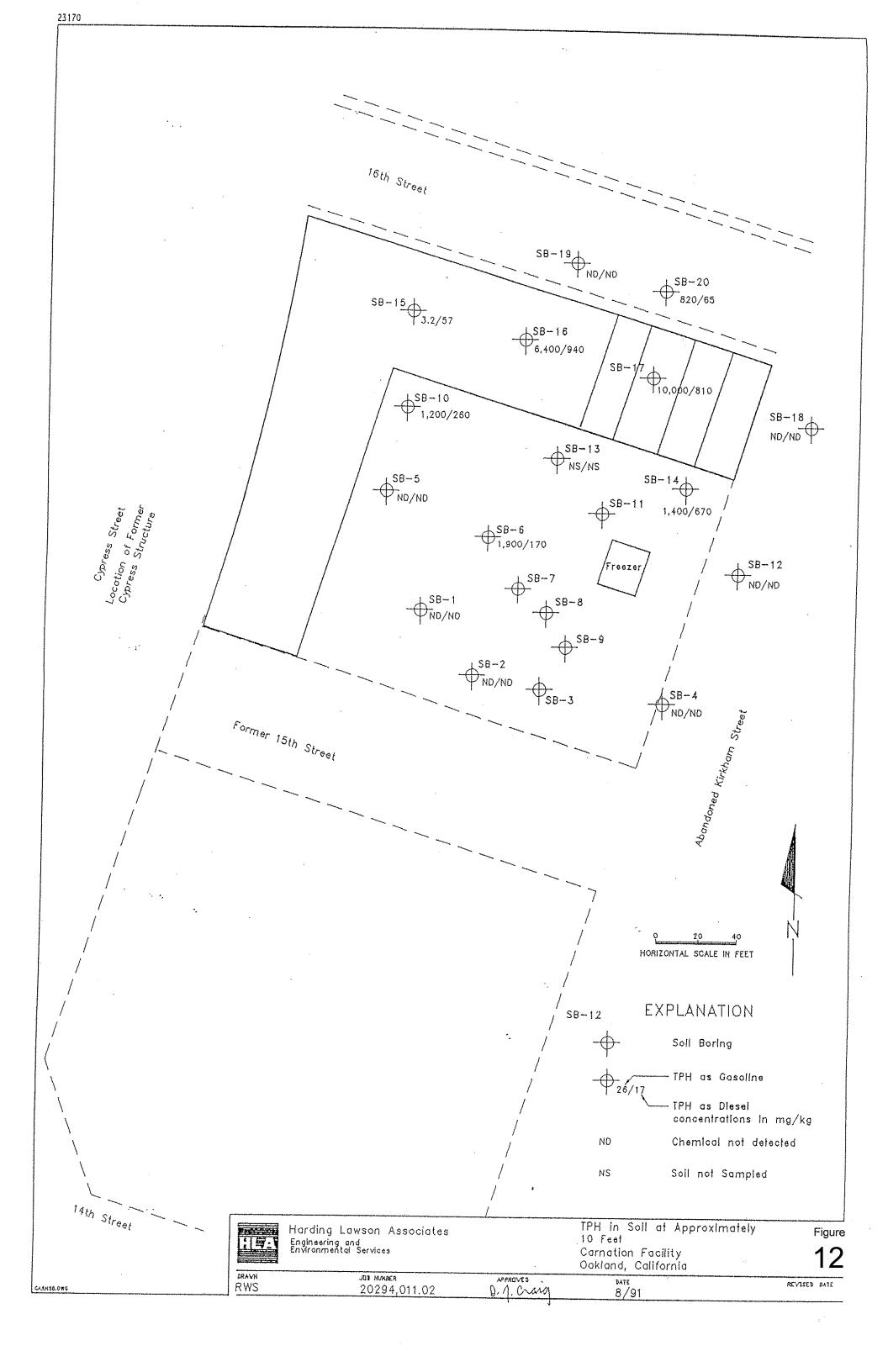
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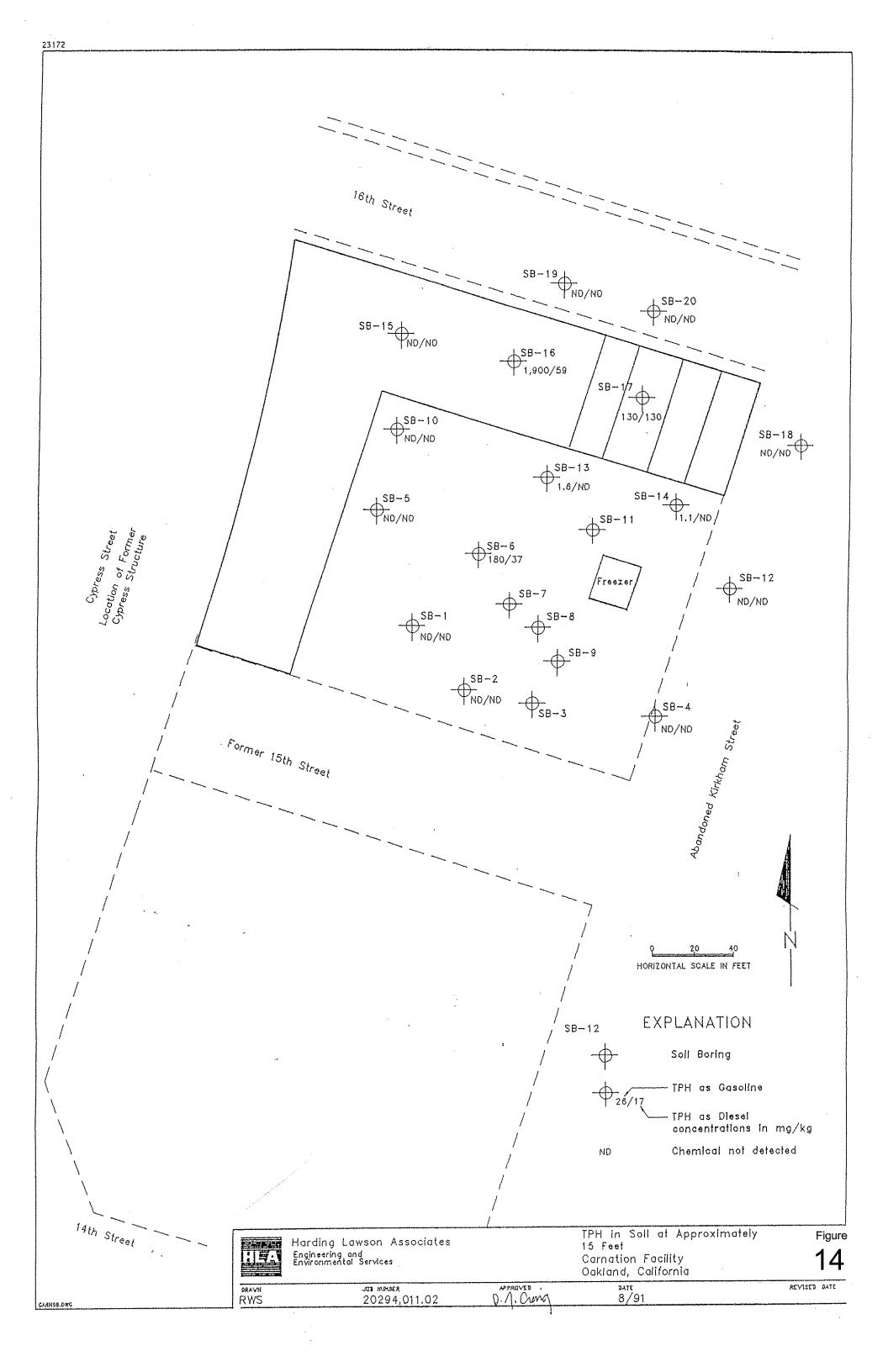
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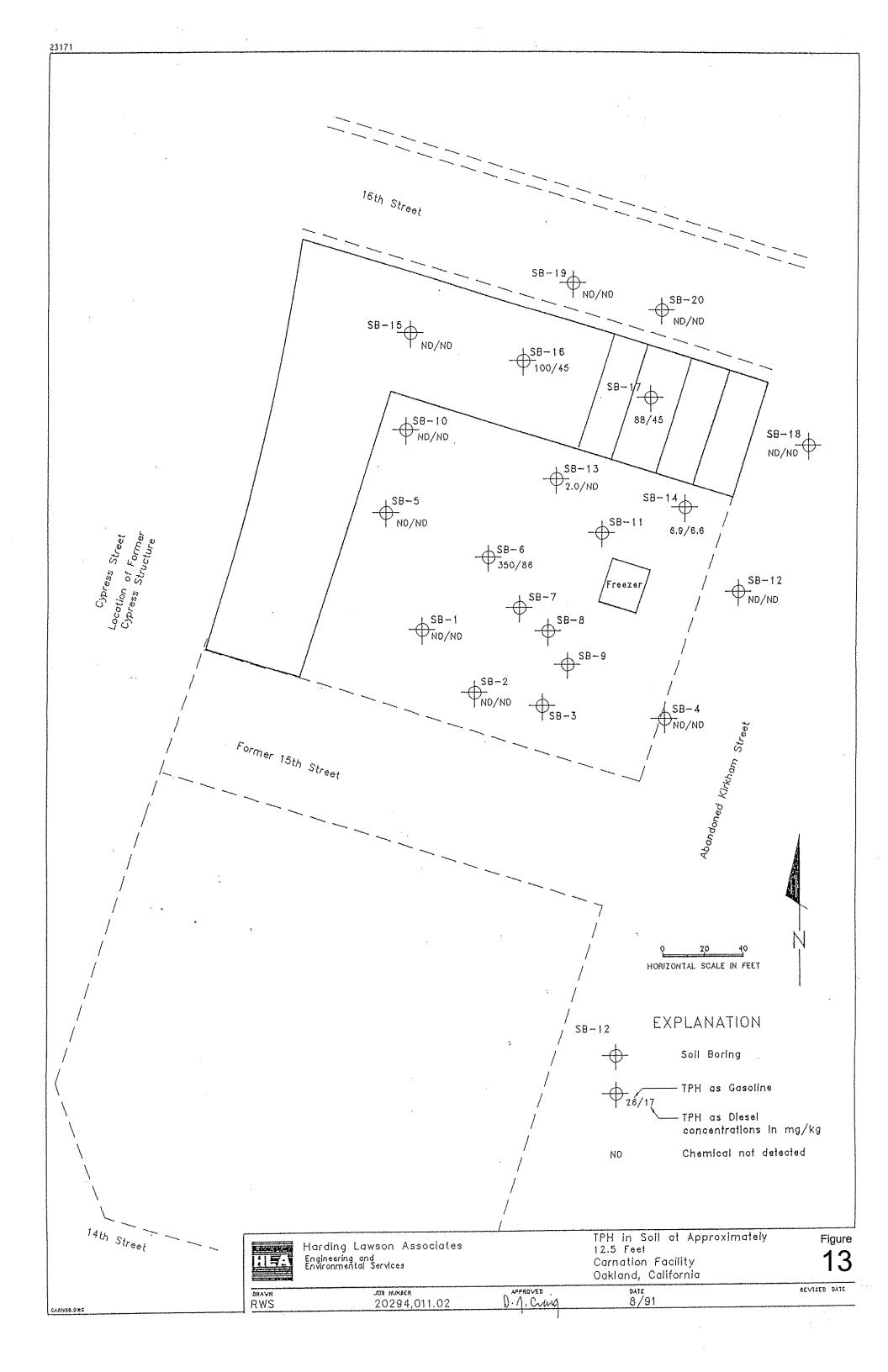


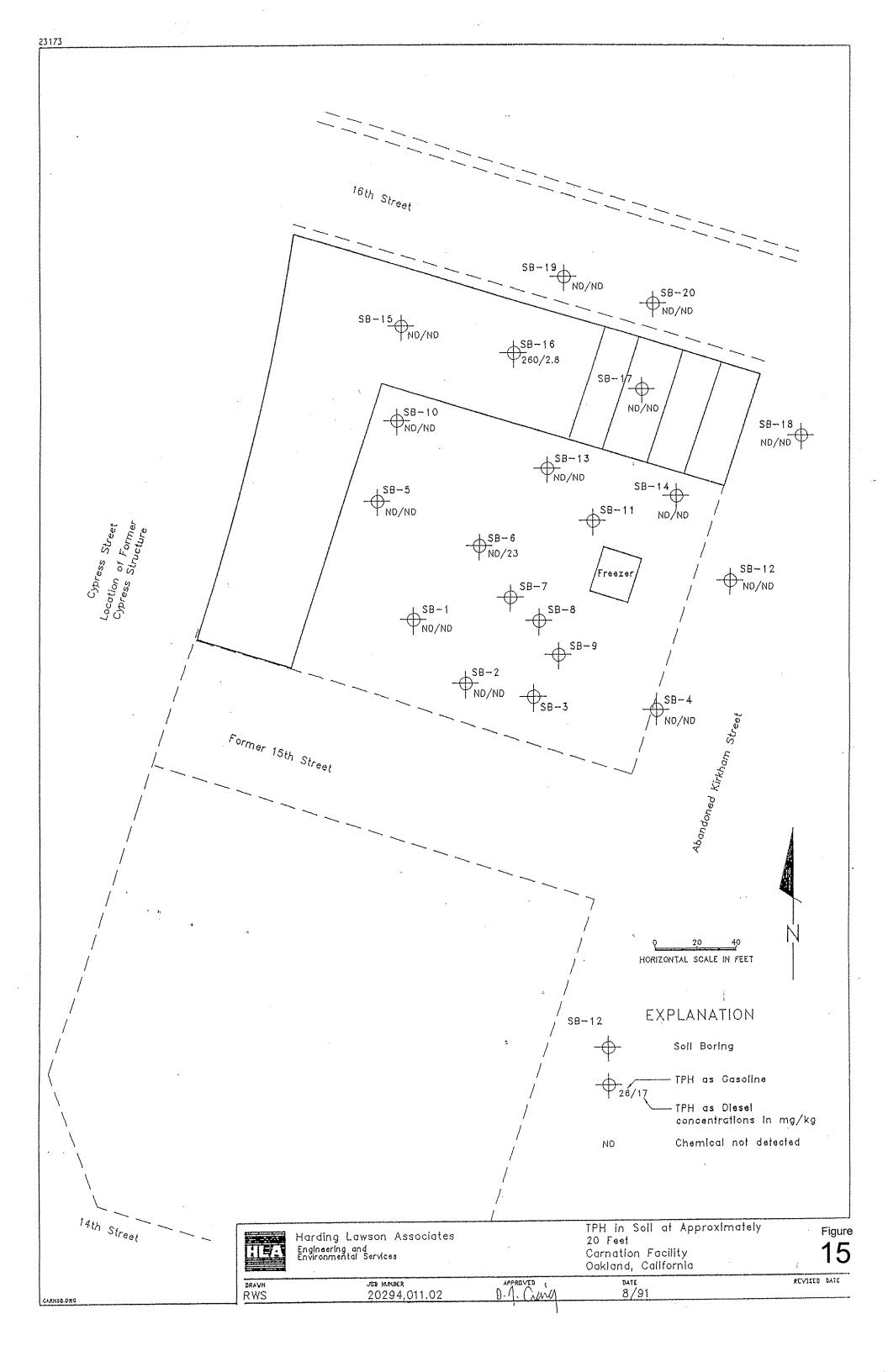


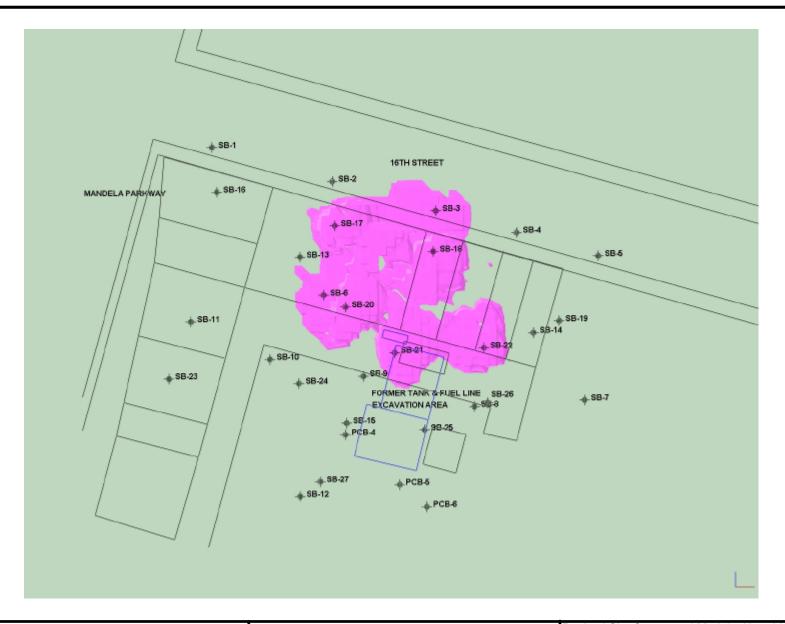












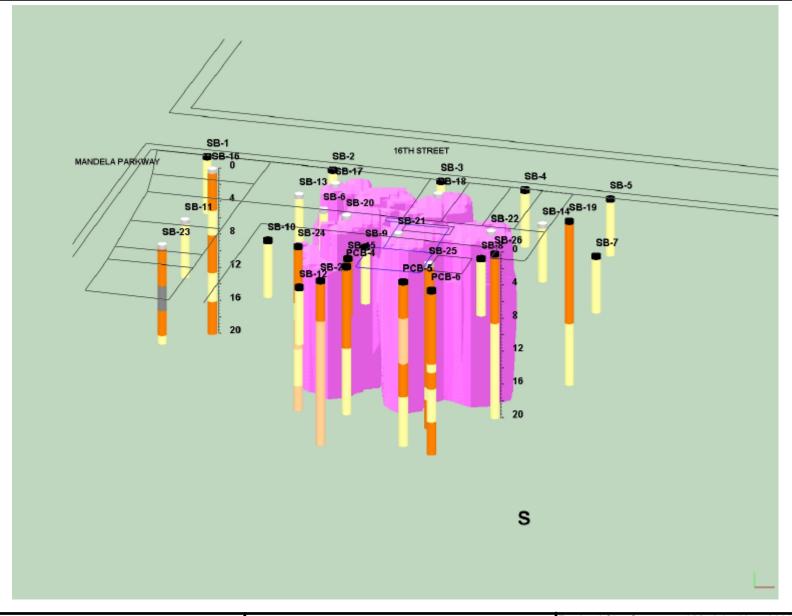


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Revised Site Conceptual Model – Nov. 2008
3D Interpolation of
TPHg Soil Sample Results
> 500 mg/kg – Plan View
(August 1999 and May 2008 Data)

Figure





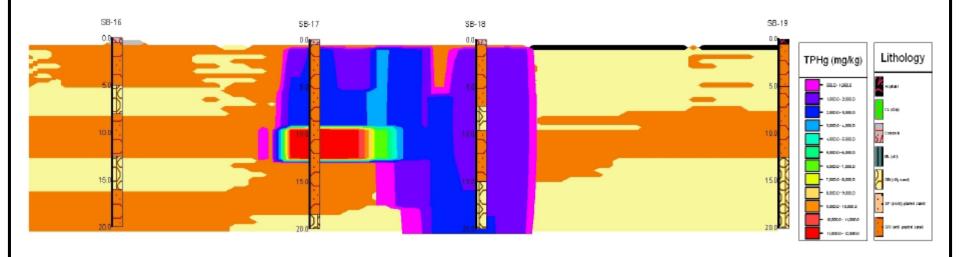
ENVIRONMENTAL COST MANAGEMENT, INC.

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Tel: (714) 662-2759 ● Fax: (714) 662-2758

Revised Site Conceptual Model – Nov. 2008
3D Interpolation of
TPHg Soil Sample Results
> 500 mg/kg – South View
(August 1999 and May 2008 Data)

Figure

W



Former Nestle USA, Inc. Facility 1310 14th Street Oakland, California 94607

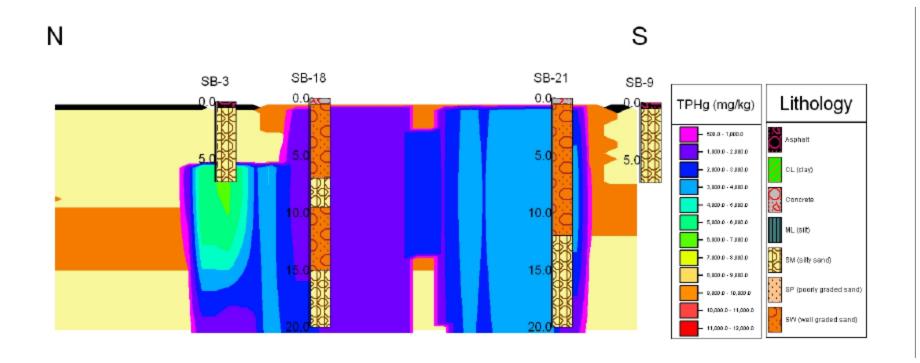


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Managing Cost and Liability
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Revised Site Conceptual Model – Nov. 2008 E-W Cross Section TPHg Soil Sample Results (August 1999 and May 2008 Data)

Figure





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Tel: (714) 662-2759 ● Fax: (714) 662-2758

Revised Site Conceptual Model – Nov. 2008 N-S Cross Section TPHg Soil Sample Results (August 1999 and May 2008 Data)

Figure



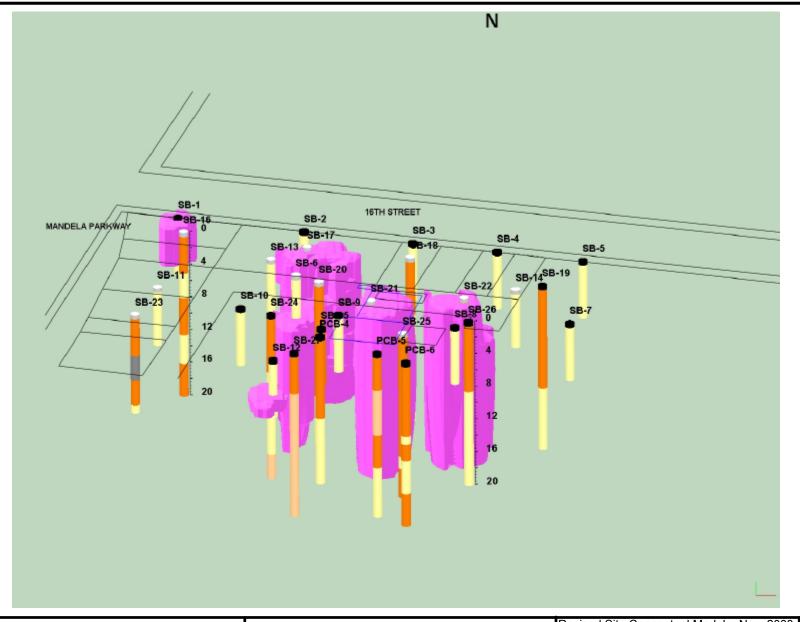


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Tel: (714) 662-2759 ● Fax: (714) 662-2758

Revised Site Conceptual Model – Nov. 2008
3D Interpolation of
TPHd Soil Sample Results
> 500 mg/kg - Plan View
(August 1999 and May 2008 Data)

Figure



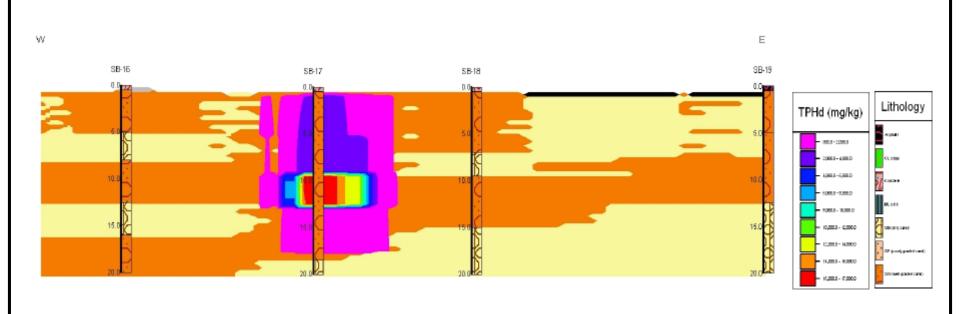


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Revised Site Conceptual Model – Nov. 2008
3D Interpolation of
TPHd Soil Sample Results
> 500 mg/kg – South View
(August 1999 and May 2008 Data)

Figure



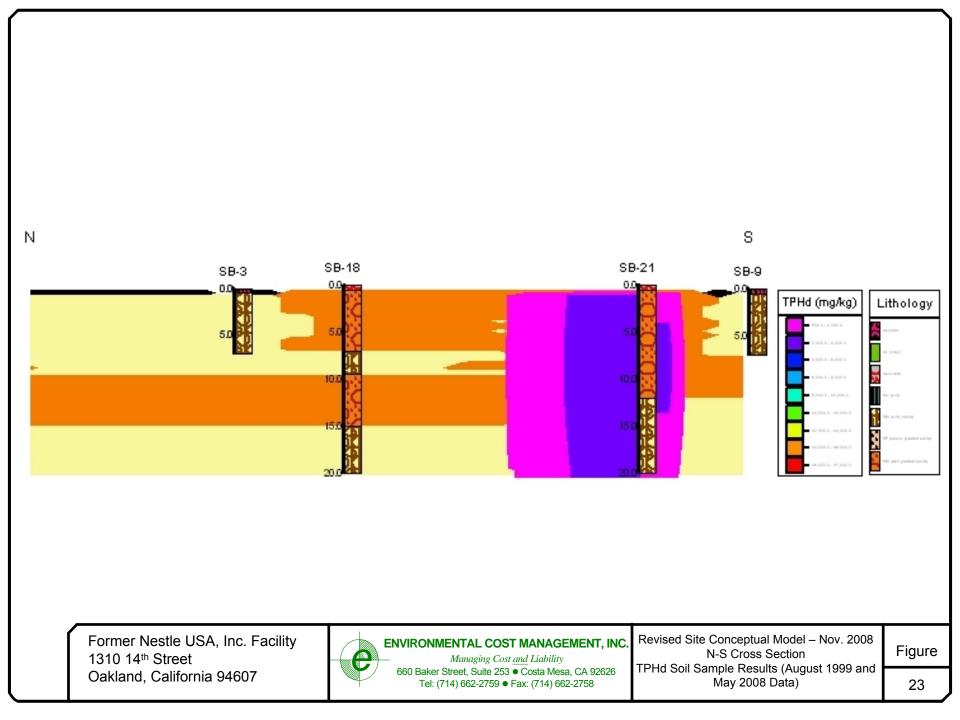


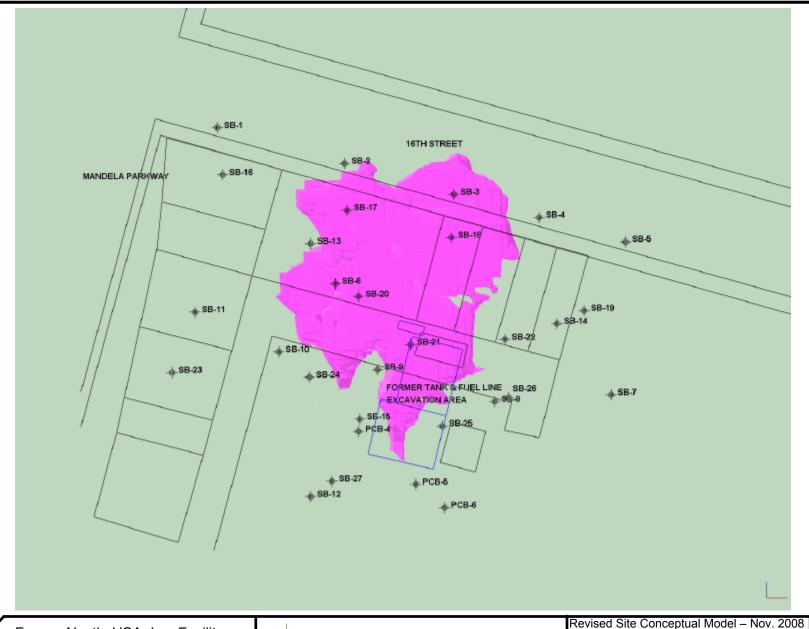
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Revised Site Conceptual Model – Nov. 2008 E-W Cross Section TPHd Soil Sample Results (August 1999 and May 2008 Data)

Figure





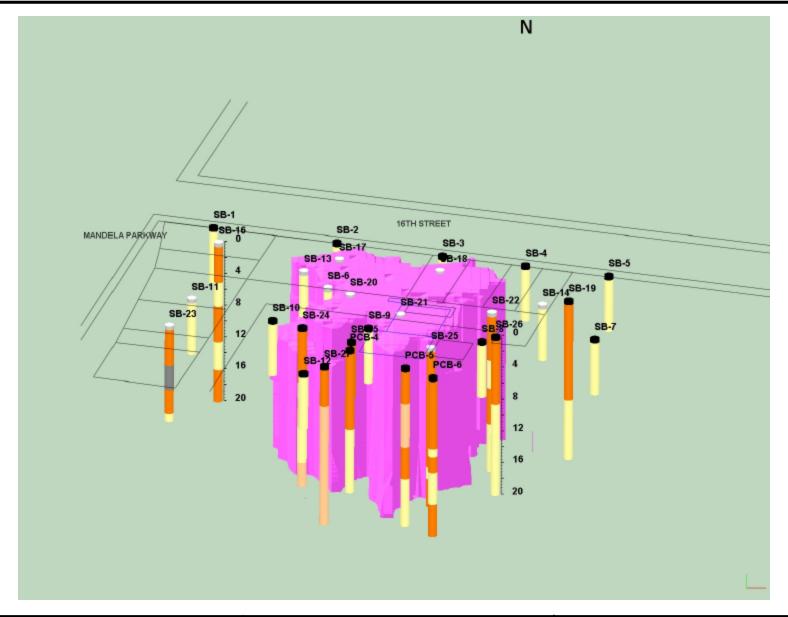


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Revised Site Conceptual Model – Nov. 20 3D Interpolation of Benzene Soil Sample Results > 0.5 mg/kg – Plan View (August 1999 and May 2008 Data)

Figure



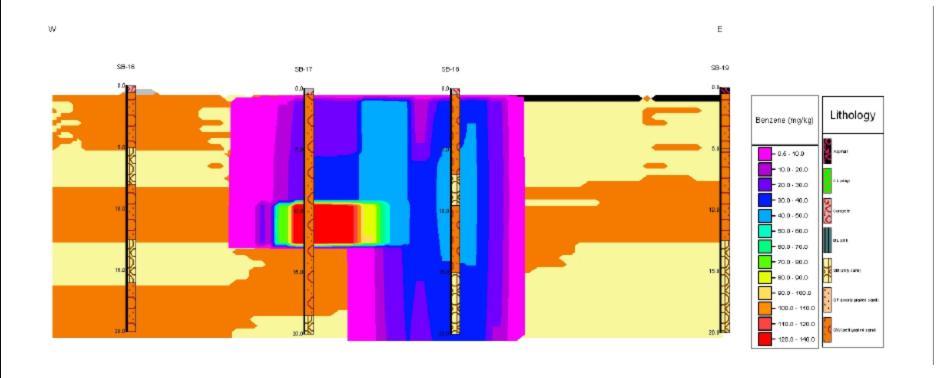


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Revised Site Conceptual Model – Nov. 2008
3D Interpolation of
Benzene Soil Sample Results
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(August 1999 and May 2008 Data)

Figure

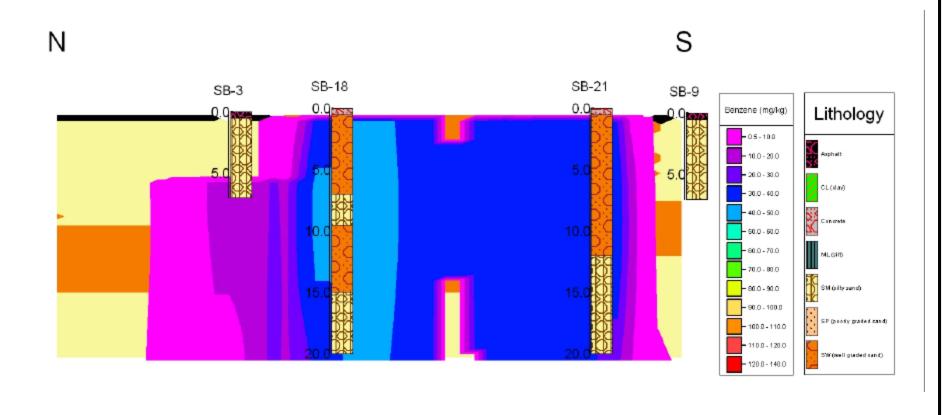




ENVIRONMENTAL COST MANAGEMENT, INC.

Managing Cost <u>and</u> Liability
660 Baker Street, Suite 253 ● Costa Mesa, CA 92626
Tel: (714) 662-2759 ● Fax: (714) 662-2758

Revised Site Conceptual Model – Nov. 2008 E-W Cross Section Benzene Soil Sample Results (August 1999 and May 2008 Data) Figure



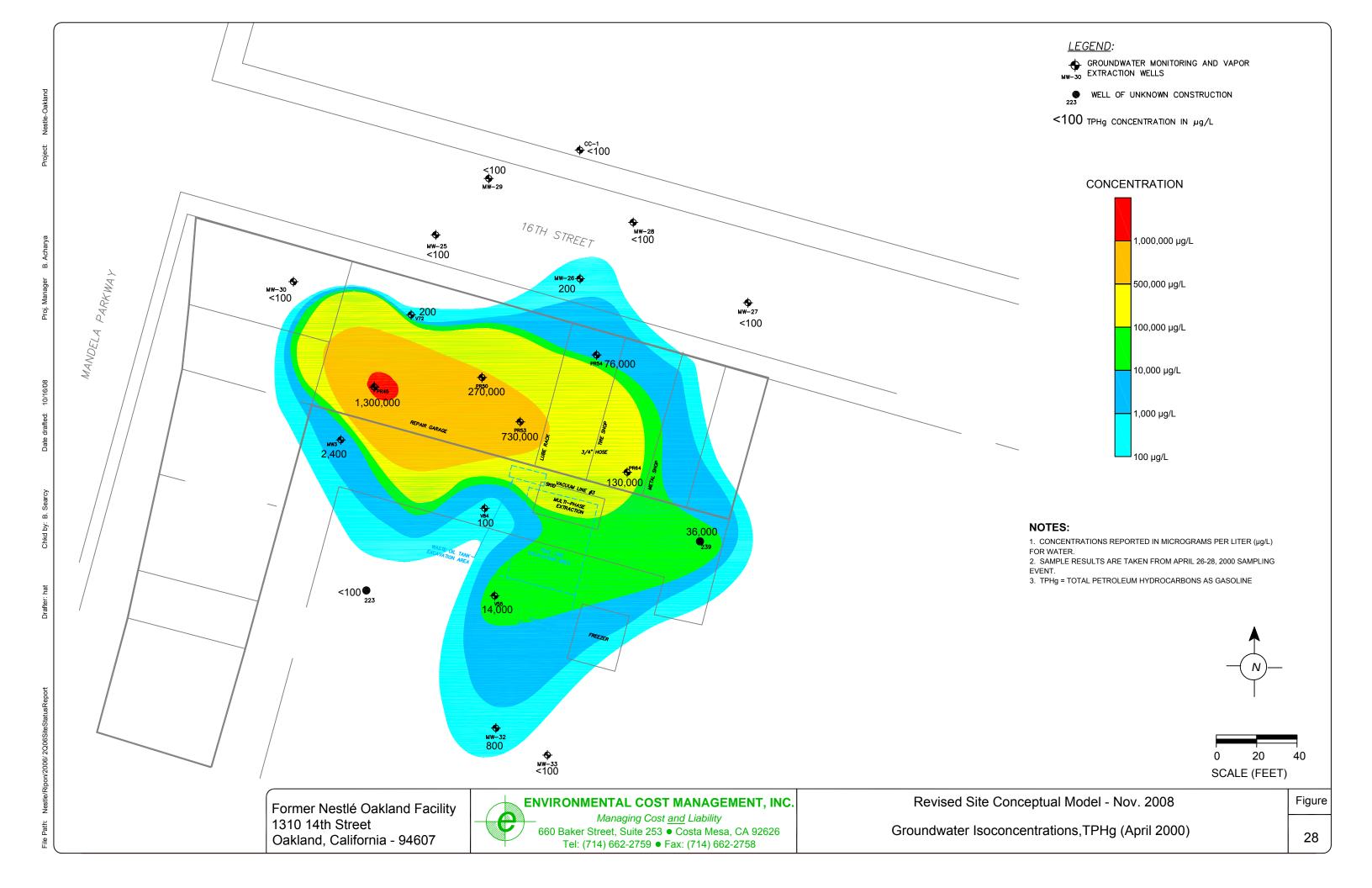


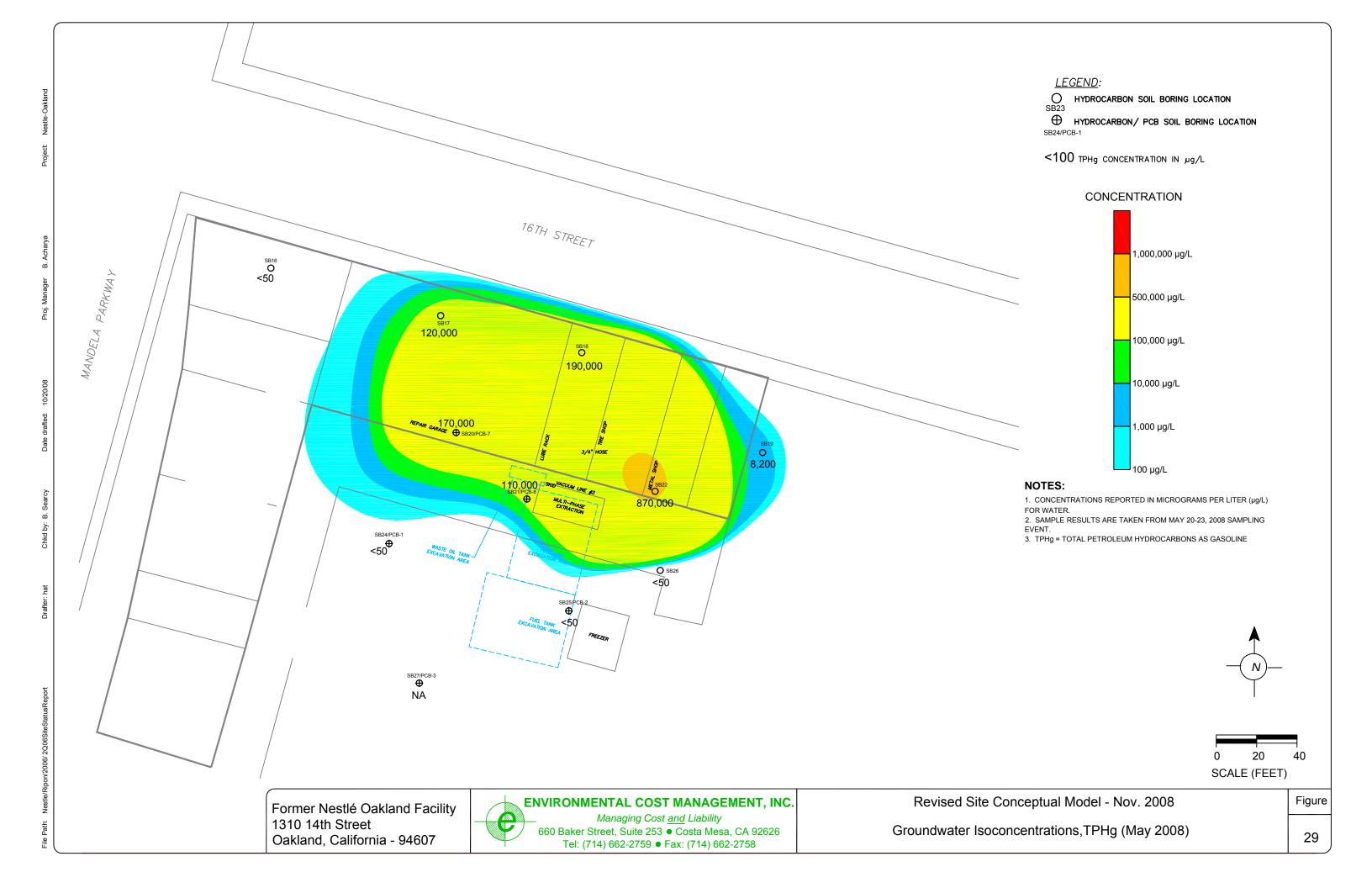
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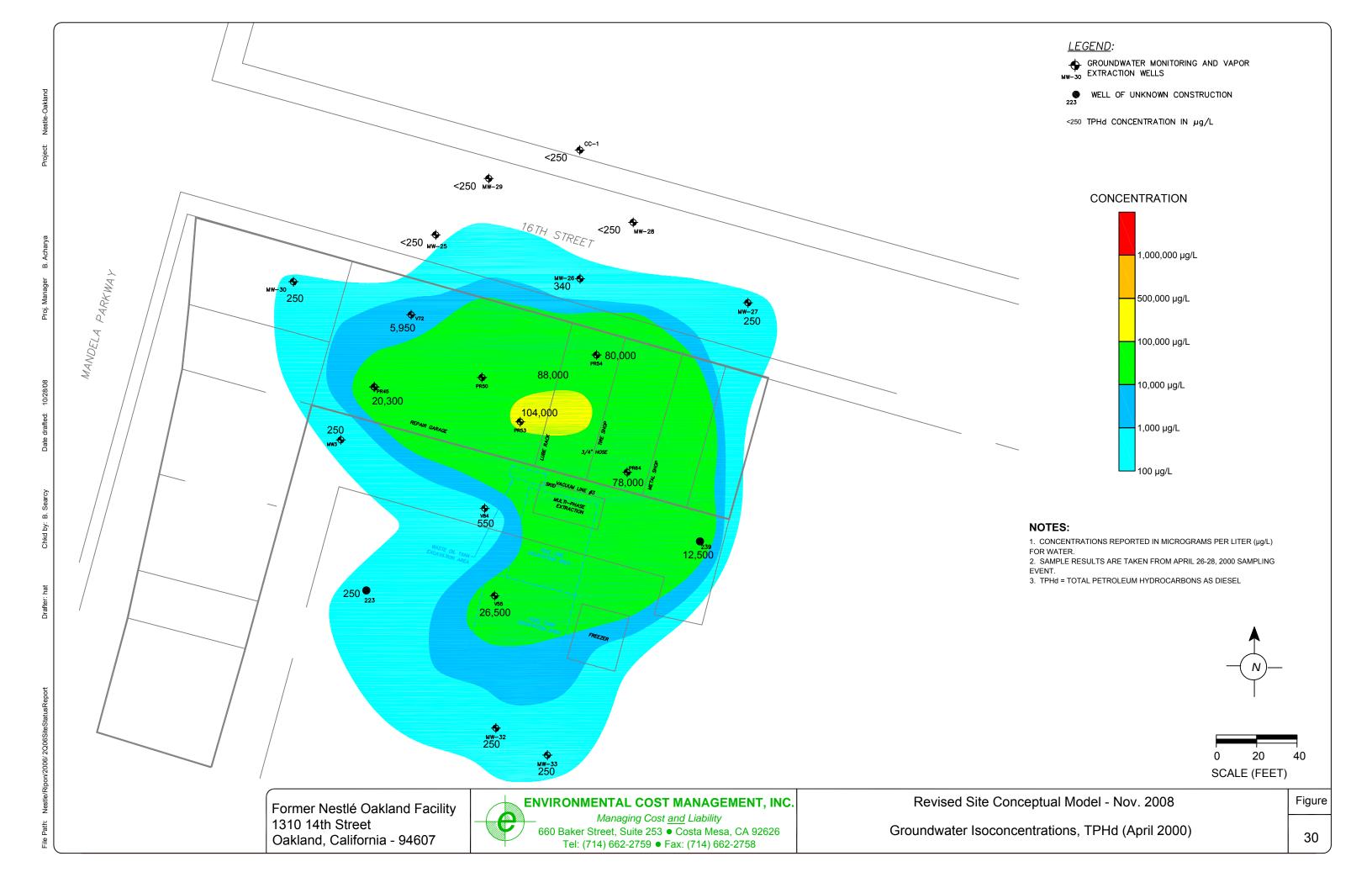
Managing Cost and Liability
660 Baker Street, Suite 253 ● Costa Mesa, CA 92626
Tel: (714) 662-2759 ● Fax: (714) 662-2758

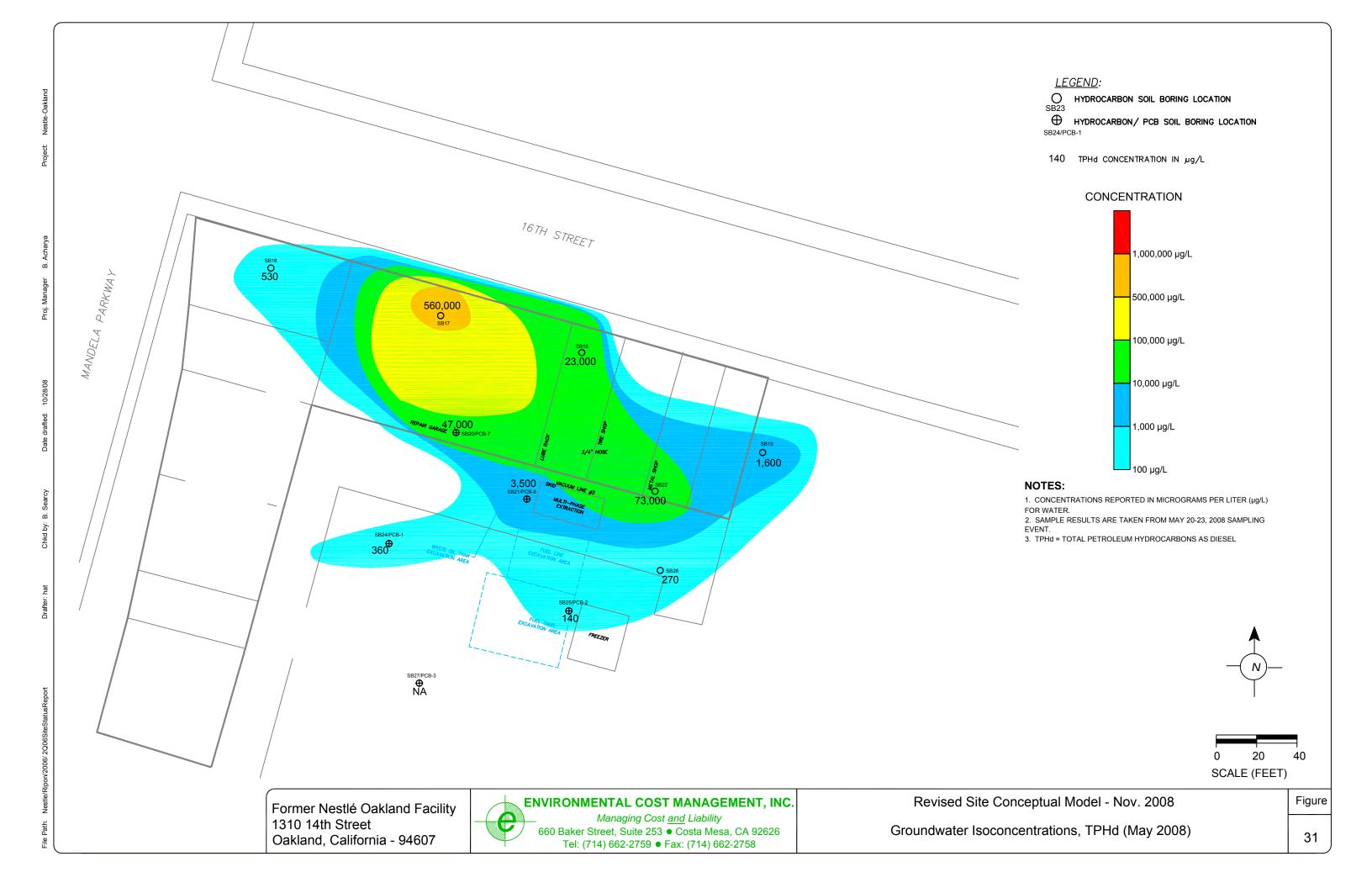
Revised Site Conceptual Model – Nov. 2008 N-S Cross Section Benzene Soil Sample Results (August 1999 and May 2008 Data)

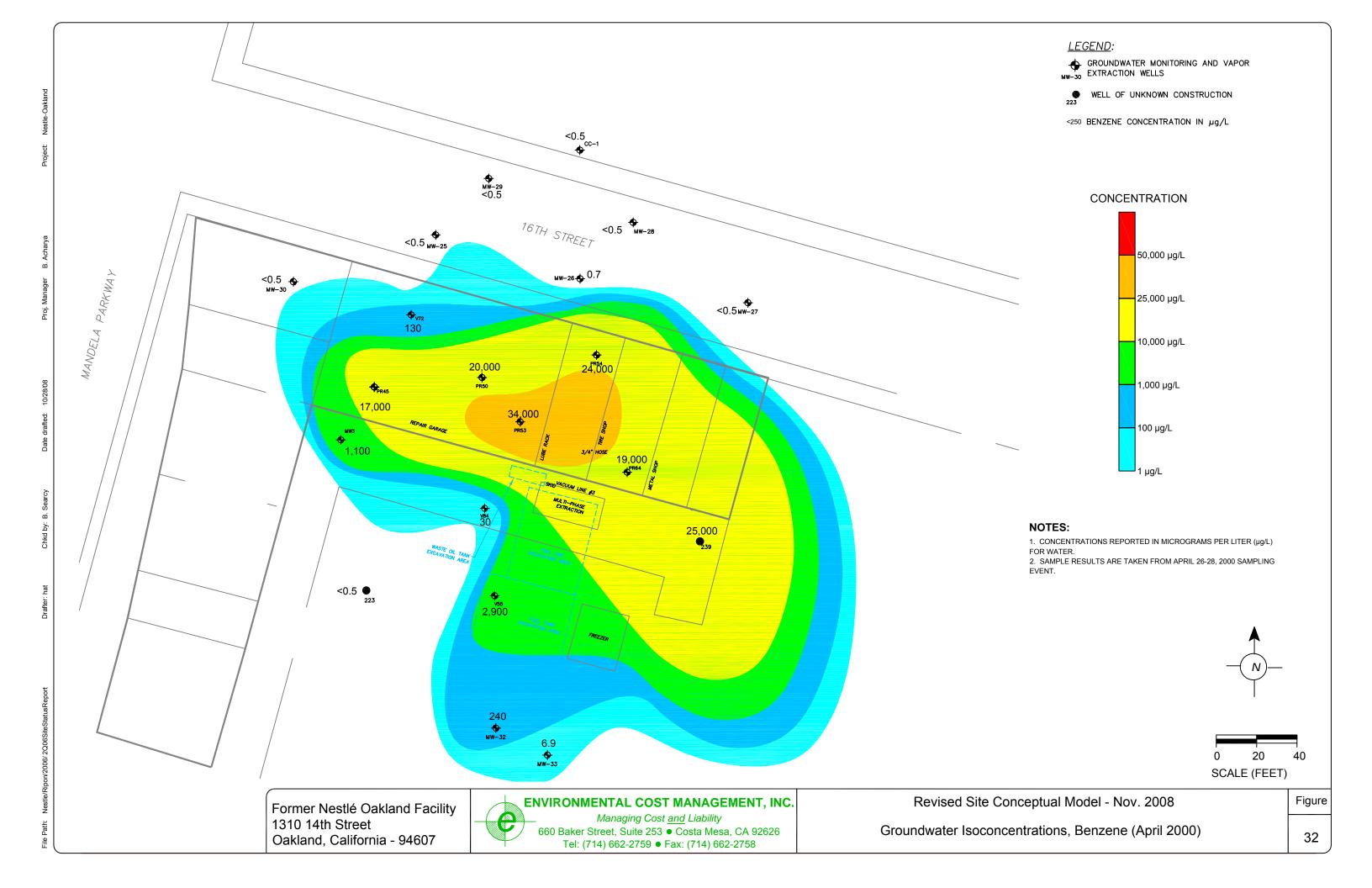
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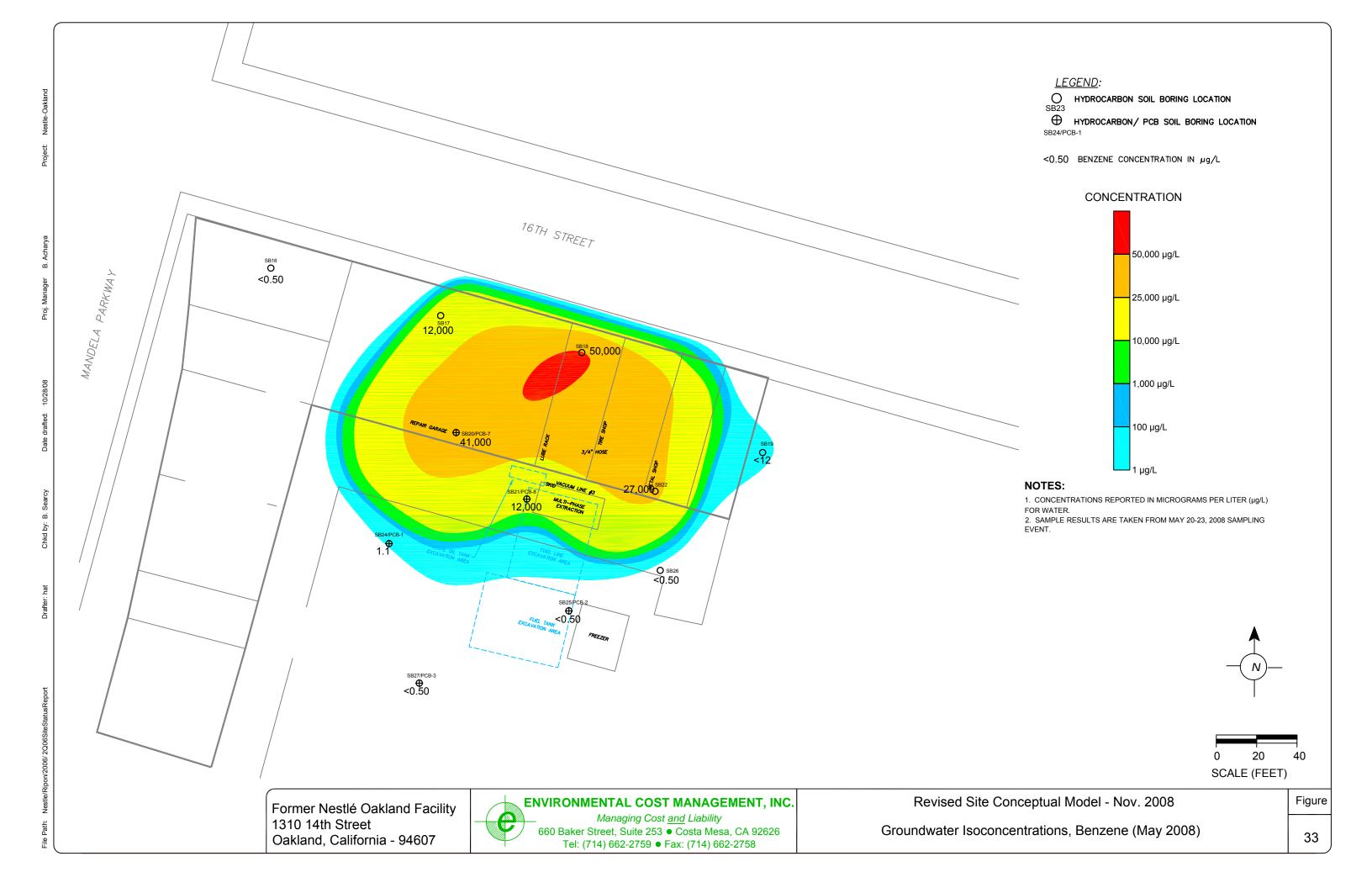




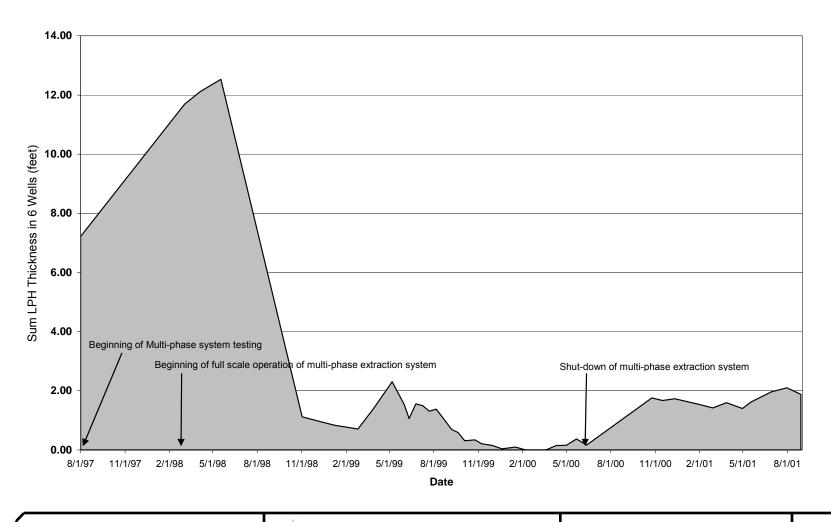








Sum of LPH Thickness in 6 Wells (MW23, MW24, PR48, PR58, PR61, and PR64)



Former Nestle USA, Inc. Facility 1310 14th Street Oakland, California 94607



Revised Site Conceptual Model November 2008 Historical LPH Thickness at 6 wells

Figure

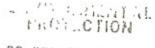
Appendix C: Covenant and Environmental Restriction (Deed Restriction)



Nestlé USA

800 NORTH BRAND BLVD GLENDALE, CA 91203

FAX (818) 543-6000 FAX (818) 543-5840



00 JUN 27 MILL: 19



LEGAL DEPARTMENT

June 23, 2000

Larry Seto Sr. Hazardous Materials Specialist ENVIRONMENTAL HEALTH SERVICES Environmental Protection 1131 Harbor Bay Parkway, Suite 250 Alameda, CA 94502-6577

Re.

Covenant and Environmental Restriction on

Nestlé USA, Inc. - Oakland Property

Dear Mr. Seto:

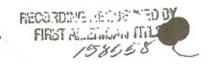
Enclosure

Enclosed please find a copy of the recorded deed restriction on the above property for your records. Please let me know if you need anything else from us at this time. Thanks again for your cooperation on this matter.

Sincerely yours,

Noelia Martí-Colon Senior Counsel

ACCESSOR CONTRACTOR



Recording Requested By:

Nestle USA Inc. 800 North Brand Blvd. Glendale, California 91203

When Recorded, Mail To:

Leroy Griffin
Hazardous Materials Program Supervisor
City of Oakland Fire Services
1605 Martin Luther King Jr. Way
Oakland, California 94612

	(000 total
in the Office of the St	recorded
in the Office of the Re	corder of
ITIO WALL	County
California, as Instrume	ent No
2000 175664	
124	on the
12th day of Line	, 2000
FIRST AMERICAN TITLE G	Designation of the Contraction

· file Clathard

COVENANT AND ENVIRONMENTAL RESTRICTION ON PROPERTY

Northeast Portion of the Former Carnation Dairy Facility which Occupies 1315-1372 14th Street and 1315-1385 16th Street

This Covenant and Environmental Restriction on Property (this "Covenant") is made as of the grad day of Juns , 2000 by Nestle USA ("Covenantor") who is the Owner of record of that certain property situated at 1315-1372 14th Street and 1315-1385 16th Street, in the City of Oakland, County of Alameda, State of California, which contains a contaminated area which is more particularly described in Exhibit A attached hereto and incorporated herein by this reference (such contaminated area hereinafter referred to as the "Burdened Property"), for the benefit of the City of Oakland Fire Services (COFS), with reference to the following facts:

- A. The Burdened Property and groundwater underlying the property contains hazardous materials.
- B. Contamination of the Burdened Property. Soil at the Burdened Property was contaminated by releases from petroleum underground storage tanks. These releases resulted in contamination of soil and groundwater with organic chemicals including benzene, toluene, ethylbenzene, xylenes, and 1,2—dichloroethane, which are hazardous materials as that term is defined in Health & Safety Code Section 25260. Removal of underground storage tanks and remediation of the petroleum hydrocarbons was initiated in January 1988 and is summarized below:

Tank, Line, and Dispenser Removal

Four (4) underground fuel storage tanks and associated piping were removed in December 1988. One (1) 1,000 gallon used-oil tank was removed in January 1989.

Remedial Actions

Soil Excavation: Between January and March 1989, 1,200 cubic yards of soil were removed in the area of the former underground storage tanks and associated piping. This soil was treated on-site and replaced back in the excavated area.

Liquid Petroleum Hydrocarbon Removal: Liquid petroleum hydrocarbons were removed using a product skimming system from the subsurface during January through March 1989. Approximately 1,800 gallons were removed during this time period.

Soil Vapor Extraction: A soil vapor extraction system operated from January 1994 to December 1995 and removed an estimated 5,200 gallons of hydrocarbon.

Multi-phase Extraction: A multi-phase extraction system has been operating at the site since August 1997. Approximately 10,500 pounds of hydrocarbons have been removed using this system. Thickness of petroleum hydrocarbons decreased since August 1997.

- C. Exposure Pathways. The contaminants addressed in this Covenant are present in soil and groundwater on the Burdened Property. Without the mitigation measures which have been performed on the Burdened Property, exposure to these contaminants could take place via the following pathways (onsite workers only):
 - > Ingestion and dermal contact with surface soils;
 - > Inhalation of volatile emissions from subsurface soils and groundwater

The risk of public exposure to the contaminants has been substantially lessened by the remediation and controls described in part B.

- D. Adjacent Land Uses and Population Potentially Affected. The Burdened Property is currently an unused industrial facility and is adjacent to industrial, commercial, and residential land uses.
- E. Full and voluntary disclosure to the COFS of the presence of hazardous materials on the Burdened Property has been made and extensive sampling of the Burdened Property has been conducted.
- F. Covenantor desires and intends that in order to benefit the COFS, and to protect the present and future public health and safety, the Burdened Property shall be used in such a manner as to avoid potential harm to persons or property that may result from hazardous materials that may have been deposited on portions of the Burdened Property.

ARTICLE I GENERAL PROVISIONS

- 1.1 Provisions to Run with the Land. This Covenant sets forth protective provisions, covenants, conditions and restrictions (collectively referred to as "Restrictions") upon and subject to which the Burdened Property and every portion thereof shall be improved, held, used, occupied, leased, sold, hypothecated, encumbered, and/or conveyed. The restrictions set forth in Article III are reasonably necessary to protect present and future human health and safety or the environment as a result of the presence of hazardous materials in the subsurface below the Burdened Property. Each and all of the Restrictions shall run with the land, and pass with each and every portion of the Burdened Property, and shall apply to, inure to the benefit of, and bind the respective successors in interest thereof, for the benefit of the COFS and all Owners and Occupants. Each and all of the Restrictions are imposed upon the entire Burdened Property. Each and all of the Restrictions run with the land pursuant to section 1471 of the Civil Code. Each and all of the Restrictions are enforceable by the California Regional Water Quality Control Board for the San Francisco Bay Region (the "Board").
- 1.2 <u>Concurrence of Owners and Lessees Presumed</u>. All purchasers, lessees, or possessors of any portion of the Burdened Property shall be deemed by their purchase, leasing, or possession of such Burdened Property, to be in accord with the foregoing and to agree for and among themselves, their heirs, successors, and assignees, and the agents, employees, and lessees of such owners, heirs, successors, and assignees, that the Restrictions as herein established must be adhered to for the benefit of the COFS and the Owners and Occupants of the Burdened Property and that the interest of the Owners and Occupants of the Burdened Property shall be subject to the Restrictions contained herein.
- 1.3 Apportionment of Burden Among Multiple Owners. Where ownership of the Burdened Property is held by multiple persons, holding by several titles, the burdens imposed by this Covenant shall be apportioned between them proportionate to the value of the property held by each owner, if such value can be ascertained, and if not, then according to their respective interests in point of quantity. (Cal. Civ. Code, § 1467.)
- 1.4 <u>Incorporation into Deeds and Leases</u>. Covenantor desires and covenants that the Restrictions set out herein shall be incorporated in and attached to each and all deeds and leases of any portion of the Burdened Property. Recordation of this Covenant shall be deemed binding on all successors, assigns, and lessees, regardless of whether a copy of this Covenant and Agreement has been attached to or incorporated into any given deed or lease.
- 1.5 <u>Purpose</u>. It is the purpose of this instrument to convey to the COFS real property rights, which will run with the land, to facilitate the remediation of past environmental contamination and to protect human health and the environment by reducing the risk of exposure to residual hazardous materials.

ARTICLE II DEFINITIONS

- 2.1 <u>COFS</u>. "COFS" shall mean the City of Oakland Fire Services and shall include its successor agencies, if any.
- 2.2 <u>Board</u>. "Board" shall mean the California Regional Water Quality Control Board for the San Francisco Bay Region and shall include its successor agencies, if any.
- 2.3 <u>Improvements</u>. "Improvements" shall mean all buildings, roads, driveways, regradings, and paved parking areas, constructed or placed upon any portion of the Burdened Property.
- 2.4 Occupants. "Occupants" shall mean Owners and those persons entitled by ownership, leasehold, or other legal relationship to the exclusive right to use and/or occupy all or any portion of the Burdened Property.
- 2.5 Owner or Owners. "Owner" or "Owners" shall mean the Covenantor and/or its successors in interest, who hold title to all or any portion of the Burdened Property.

ARTICLE III DEVELOPMENT, USE AND CONVEYANCE OF THE BURDENED PROPERTY

- 3.1 <u>Restrictions on Development and Use</u>. Covenantor promises to restrict the use of the Burdened Property as follows:
- a. Development of the Burdened Property shall be restricted to industrial, commercial or office space;
 - b. No residence for human habitation shall be permitted on the Burdened Property;
 - c. No hospitals shall be permitted on the Burdened Property;
- d. No schools for persons under 21 years of age shall be permitted on the Burdened
 Property;
- e. No day care centers for children or day care centers for Senior Citizens shall be permitted on the Burdened Property;
- f. No Owners or Occupants of the Burdened Property or any portion thereof shall conduct any excavation work on the Burdened Property, unless expressly permitted in writing by the COFS. Any contaminated soils brought to the surface by grading, excavation, trenching, or

backfilling shall be managed by Covenantor or his agent in accordance with all applicable provisions of local, state and federal law;

- g. All uses and development of the Burdened Property shall be consistent with any applicable Board Order or Risk Management Plan, each of which is hereby incorporated by reference including future amendments thereto. All uses and development shall preserve the integrity of any cap, any remedial measures taken or remedial equipment installed, and any groundwater monitoring system installed on the Burdened Property pursuant to the requirements of the COFS, unless otherwise expressly permitted in writing by the COFS. Any development of the Burdened Property will maintain a surface cap of the soil, exclusive of minor landscape areas, by buildings or paved surfaces.
- h. No Owners or Occupants of the Property or any portion thereof shall drill, bore, otherwise construct, or use a well for the purpose of extracting water for any use, including but not limited to, domestic, potable, or industrial uses, unless expressly permitted in writing by the Board.

3.1.1 Notifications/Access/Non Aggravation

- a. The Owner shall notify the COFS of each of the following: (1) The type, cause, location and date of any disturbance to any cap, any remedial measures taken or remedial equipment installed, and of the groundwater monitoring system installed on the Burdened Property pursuant to the requirements of the COFS, which could affect the ability of such cap or remedial measures, remedial equipment, or monitoring system to perform their respective functions and (2) the type and date of repair of such disturbance. Notification to the COFS shall be made by registered mail within ten (10) working days of both the discovery of such disturbance and the completion of repairs;
- b. The Covenantor agrees that the COFS, and/or any persons acting pursuant to COFS orders, shall have reasonable access to the Burdened Property for the purposes of inspection, surveillance, maintenance, or monitoring, as provided for in Division 7 of the Water Code.
- c. No Owner or Occupant of the Burdened Property shall act in any manner that will aggravate or contribute to the existing environmental conditions of the Burdened Property. All use and development of the Burdened Property shall preserve the integrity of any capped areas.
- 3.2 <u>Enforcement</u>. Failure of an Owner or Occupant to comply with any of the restrictions, as set forth in paragraph 3.1, shall be grounds for the COFS, by reason of this Covenant, to have the authority to require that the Owner modify or remove any Improvements constructed in violation of that paragraph. Violation of the Covenant shall be grounds for the COFS to file civil actions against the Owner as provided by law.
- 3.3 Notice in Agreements. After the date of recordation hereof, all Owners and Occupants shall execute a written instrument which shall accompany all purchase agreements or leases relating to the property. Any such instrument shall contain the following statement:

The land described herein contains hazardous materials in soils and in the ground water under the property, and is subject to a deed restriction dated as of <u>June 8</u>, 2000, and recorded an <u>Concurrently herewith 2000</u>, in the Official Records of <u>Alameda</u> County, California, which covenant and Restriction imposes certain covenants, conditions, and restrictions on usage of the property described herein. This statement is not a declaration that a hazard exists.

ARTICLE IV VARIANCE AND TERMINATION

- 4.1 <u>Variance</u>. Any Owner or, with the Owner's consent, any Occupant of the Burdened Property or any portion thereof may apply to the COFS for a written variance from the provisions of this Covenant.
- 4.2 <u>Termination</u>. Any Owner or, with the Owner's consent, any Occupant of the Burdened Property or a portion thereof may apply to the COFS for a termination of the Restrictions as they apply to all or any portion of the Burdened Property which consent to termination shall not be unreasonably withheld.
- 4.3 <u>Term</u>. Unless terminated in accordance with paragraph 4.2 above, by law or otherwise, this Covenant shall continue in effect in perpetuity.

ARTICLE V MISCELLANEOUS

- 5.1 No Dedication Intended. Nothing set forth herein shall be construed to be a gift or dedication, or offer of a gift or dedication, of the Burdened Property or any portion thereof to the general public.
- 5.2 Notices. Whenever any person gives or serves any notice, demand, or other communication with respect to this Covenant, each such notice, demand, or other communication shall be in writing and shall be deemed effective (1) when delivered, if personally delivered to the person being served or official of a government agency being served, or (2) three (3) business days after deposit in the mail if mailed by United States mail, postage paid certified, return receipt requested:

If To: "Covenantor"
Nestlé USA, Inc.
Legal Department
800 North Brand Boulevard
Glendale, Ca. 91203

If To: "COFS"

City of Oakland Fire Services

Attention: Hazardous Materials Program Supervisor

1605 Martin Luther King Jr. Way

Oakland, California 94612

- 5.3 <u>Partial Invalidity</u>. If any portion of the Restrictions or terms set forth herein is determined to be invalid for any reason, the remaining portion shall remain in full force and effect as if such portion had not been included herein.
- 5.4 <u>Article Headings</u>. Headings at the beginning of each numbered article of this Covenant are solely for the convenience of the parties and are not a part of the Covenant.
- 5.5 <u>Recordation</u>. This instrument shall be executed by the Hazardous Materials Program Supervisor of the COFS. This instrument shall be recorded by the Covenantor in the County of Alanceda within ten (10) days of the date of execution.
 - 5.6 References. All references to Code sections include successor provisions.
- 5.7 <u>Construction</u>. Any general rule of construction to the contrary notwithstanding, this instrument shall be liberally construed in favor of the Covenant to effect the purpose of this instrument and the policy and purpose of the Water Code. If any provision of this instrument is found to be ambiguous, an interpretation consistent with the purpose of this instrument that would render the provision valid shall be favored over any interpretation that would render it invalid.

IN WITNESS WHEREOF, the parties execute this Covenant as of the date set forth above.

Covenar	itor: NESTL	E USA,	INC.
By:	Mande	Lo Robert	H. Sanders
Title:	V. P.		٠.
Date:	4.8.00		
Agency:	City of O	akland Fire Ser	vices

By: LeRoy Griff:
Title: Hazardous Materials Program Supervisor

APPENDIX A

LEGAL DESCRIPTION DEED RESTRICTION AREA

ROB BEIN, WILLIAM FROST & ASSOCIATED 1981 N. Broadway, Suite 235 Walnut Creek, California 94596

LEGAL DESCRIPTION DEED RESTRICTION AREA

That certain parcel of land situated in the City of Oakland, County of Alameda, State of California described as follows:

Being a portion of Lots 4 through 23 and a portion Kirkham Street of the Scotchler Tract and Vicinity, Oakland, as shown on a map thereof filed in Book 7 of Maps at Page 21 on December 10, 1874 in the Office of the County Recorder of Alameda County more particularly described as follows:

BEGINNING at the intersection of said Kirkam Street and the northwest corner of lot 17, in block 584, as shown on the map of "Re-division of Blocks 584, 585, 601, 153 and 580-A, City of Oakland, County of Alameda, California", filed May 1, 1885, in Book 4 of Maps, at Page 25, in said office of the County Recorder;

Thence, along the northerly line of said Kirkham Street and said lots 13, 12, 11, 10, 9, 8, 7, 6 and 5, North 72°53'28" West 292.25 feet to the northwest corner of said lot 5, said point also being the northeasterly corner of that certain parcel of land described in the deed to the State of California, recorded May 12, 1955 in Volume 7658, of Official Records at Page 299, in said office of the County Recorder;

Thence, continuing along said northerly line of Kirkham Street, North 72°53'28" West 8.64 feet;

Thence, along said State of California parcel, along a non-tangent 1240 foot radius curve to the right, through a central angle of 2°59'04" to the easterly line of the parcel of land described in the deed to the State of California, recorded August 12, 1955 in Book 7749, of Official Records at Page 447, as Instrument Number AK-86901, in said office of the County Recorder;

Thence, along last said State of California parcel (7749 OR 447), along a non-tangent 1240 foot radius curve to the right from a tangent that bears South 10°54'36" West to the south line of said lot 22, said southerly line also being the north line of 15th Street, as shown on said map of the Scotchler Tract (7 M 21);

Thence, along said northerly line of 15th Street and the easterly prolongation of said north line, South 74°03'30" East 285.05 feet to the easterly line of said Kirkham Street;

Thence, along said easterly line, North 15°56'30" West 209.50 feet to the POINT OF BEGINNING.

EXHIBIT attached and by this reference made a part hereof.

No. 5816
END. 6-30 2000

Patrick J. Tami, L.S. 5816

DEE RESTRICTION AREA

	16TH STREET N74*53'28'W 300.89'							P.O.B.					
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											1	11 .	
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		15TH STREET							9				
		(VACATED)					1		8				
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SCALE: 1"=60"	5	6	7	8	9	10	11	12	13				

17TH STREET

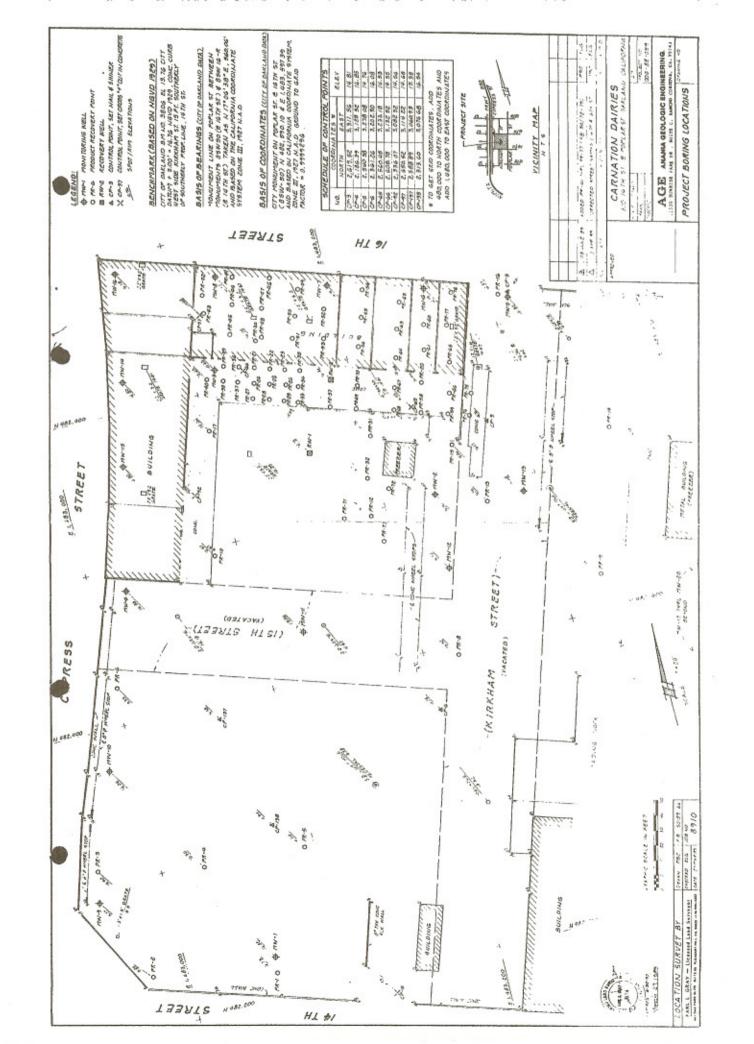


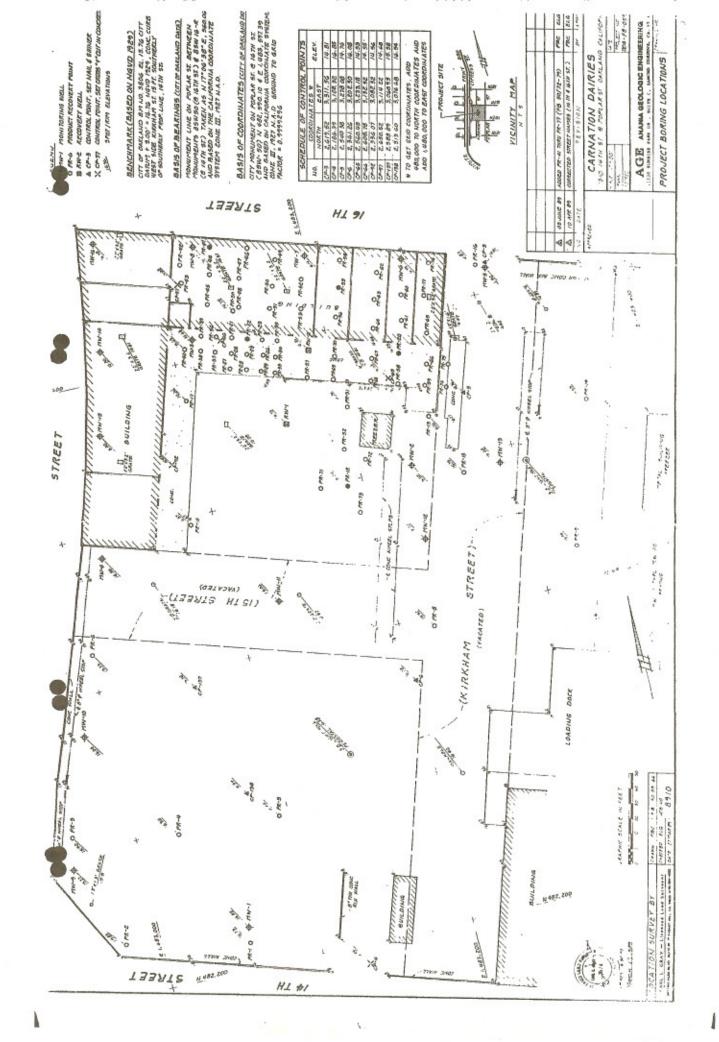
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State of California County of Alameda	ancessoccessocial (1900-1900-1900-1900-1900-1900-1900-1900
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County of Hlameda	
On 6-7-00 before me,	Ellen N. Dolese
personally appeared LeRoy Grittin	Name and Title of Officer (e.g., "Jane Doe, Notary Public")
personally appeared Lekay Grittin	Name(s) of Signer(s)
Noe	rsonally known to me 🛩
	oved to me on the basis of satisfactory evidence
ELLEN N, DOLESE Commission # 1171028 Notary Public - Collifornia Alameda County My Comm. Expires Jan 28, 2002 OPT	the person(s) whose name(s) is/are subscribed to the instrument and acknowledged to me that he/she/they uted the same in his/her/their authorized capacity(ies), hat by his/her/their signature(s) on the instrument the in(s), or the entity upon behalf of which the person(s) is, executed the instrument. IESS my hand and official seal. Signature of Notary Public TONAL To ve valuable to persons relying on the document and could prevent
fraudulent removal and reattachme	ent of this form to another document.
Title or Type of Document: Covenant & Env	Number of Pages: 12
Document Date: 5-25-00	Number of Pages: 12
Signer(s) Other Than Named Above:	
Capacity(ies) Claimed by Signer(s)	
Signer's Name: LERay Griff.:	Signer's Name:
S- Individual	- Individual
☐ Corporate Officer	☐ Corporate Officer
Title(s):	Title(s):
☐ Partner — ☐ Limited ☐ General	□ Partner — □ Limited □ General
☐ Attorney-in-Fact ☐ Trustee	☐ Attorney-in-Fact ☐ Trustee
☐ Guardian or Conservator ☐ Guardian or Conservator ☐ Grandian or Conservator ☐ Grandian or Conservator ☐ Grandian or Conservator ☐ Grandian or Conservator	Guardian or Conservator Guardian or Conservator RIGHT THUMBPRINT OF SIGNER
Other: Top of thumb here	Other: Top of thumb here
Signer Is Representing:	Signer Is Representing:

CALIFORNIA ALL-PURPOSE ACKNOWLEDGMENT

State of California County of		
On JUNE 8, 2000, before me, MARIA HAZEL PERLI NUTRAY PUBLIC Date Personally appeared ROBERT H. SANDERS Name(a) of Signet(a). "Jame Goe, Notary Public") Personally known to me Troved to me on the basis of satisfactory evidence It is Angeles County evidence It is an evidence	State of California)
On JUNE 8, 2000, before me, MARIA HAZEL PERRI NOTALY PUBLIC Personally appeared ROBERT H. SANDERS Name(s) of Signet(s) Personally known to me Though the information below is not required by law, it mey prove valuable to persons relying on the document and condition of this form to another document. Description of Attached Document: Lovenant Date: MAY 25, 2000 Name and Title of Odlice (a.g., Jane Doe, Notary Public) ROBERT H. SANDERS Name(s) of Signet(s) Personally known to me Thought known to me Thought the information below is not required by law, it mey prove valuable to persons relying on the document and could prevent fraudulent removal and reattachment of this form to another document. Description of Attached Document Ittle or Type of Document: Lovenant & Supracial Notary Public Suprace(s) Other Than Named Above: LEVRY CRUENT Signature (s) Other Than Named Above: LEVRY CRUENT Signature (s) Office (a.g., Jane Doe, Notary Public) Named and Title of Office (a.g., Jane Doe, Notary Public) Named of Different (a.g., Jane Doe, Notary Public) Personally known to me Thought known to me Thou	100000	ss.
Place Notery Saal Above Place Notery Saal Above Place Notery Saal Above Place Notery Saal Above ROBERT H. SANDERS Name(s) of Signer(s) Personally known to me proved to me on the basis of satisfactory evidence to be the person(s) whose name(s) is/are subscribed to the within instrument and acknowledged to me that he/she/their authorized capacity(ies), and that by his/her/their signature(s) on the instrument the person(s) acted, executed the instrument. WITNESS my hand and official seal. WITNESS my hand and official seal. WITNESS my hand and official seal. Place Notery Saal Above OPTIONAL Though the information below is not required by lew, it may prove valuable to persons relying on the document and could prevent fraudulent removal and reattachment of this form to another document. Description of Attached Document Title or Type of Document: Document Date: MARIA HAZEL PERRI Document Date: MARIA HAZEL PERRI Document Date: Number of Pages: TWELVE (12) Signellar	County of LOS ANGELES	
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Signature of Notary Public With and Above MARIA HAZEL PERRI Commission # 1113713 Norcry Public — California Los Angeles County My Comm. Expres Oct 13, 2000 My Comm. Expres Oct 14, 2000 My Comm. Expres Oct 14, 2000 My Comm. Expres Oct 14, 20		Proved to me on the basis of satisfactory.
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Though the information below is not required by law, it may prove valuable to persons relying on the document and could prevent fraudulent removal and reattachment of this form to another document. Description of Attached Document Title or Type of Document: LOVINANT & ENVIRONMENTAL RESTRICTION ON PROPER Document Date: MAY 25, 2000 Number of Pages: TWELVE (12) Signer(s) Other Than Named Above: LEVAY GRIFFIA	Piace Notary Seal Above	WITNESS my hand and official seal. Maria Signature of Notary Public
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Signer(s) Other Than Named Above: LEVGY GRIFFIN Capacity(les) Claimed by Signer Signer's Name: Nobert H SANDERS Individual Corporate Officer — Title(s): VICE PRESIDEN Partner — Limited General Attorney in Fact Trustee Guardian or Conservator Other: Signer Is Representing: NESTLE VSA, INC.	Document Date: MAY 25, 2000	Number of Pages: TWELVE (12)
Capacity(les) Claimed by Signer Signer's Name: Nobert H SANDERS Individual Corporate Officer — Title(s): VICE PRESIDEN Partner — Limited General Attorney in Fact Trustee Guardian or Conservator Other: Signer Is Representing: NESTLE VSA, INC.	Signer(s) Other Than Named Above: LEVOY	GRIFFIN
Individual Corporate Officer — Title(s): VICE PRESIDEN Partner — Limited General Attorney in Fact Trustee Guardian or Conservator Other: Signer Is Representing: NESTLE VSA, INC.	Capacity(les) Claimed by Signer Signer's Name: <u>ROBENT HI</u> SANDEN	RIGHT THUMBPRINT
Corporate Officer — Title(s):	Individual	Top of thumb here
□ Partner □ Limited □ General □ Attorney in Fact □ Trustee □ Guardian or Conservator □ Other:	Corporate Officer — Title(s): VICE PICE	ESTOGN
☐ Attorney in Fact ☐ Trustee ☐ Guardian or Conservator ☐ Other: ☐ Signer Is Representing: NESTLE VSA, INC.	☐ Partner — ☐ Limited ☐ General	
☐ Guardian or Conservator ☐ Other: ☐ Signer Is Representing: NESTLE VSA, INC.	☐ Attorney in Fact	
☐ Guardian or Conservator ☐ Other: ☐ Signer Is Representing: NESTLE USA, INC.	☐ Trustee	
Signer Is Representing: NESTLE VSA, INC.	☐ Guardian or Conservator	
Signer Is Representing: NESTLE USA, INC.	U Other:	
•	Signer Is Representing: NETTLE VSA,	INC.
		,





Appendix D: Risk Management Plan



Risk Management Plan for Deed Restricted Portion of Former Nestlé USA Facility

1310 14th Street Oakland, California

Prepared for

Nestlé USA, Inc. 800 North Brand Boulevard Glendale, California 91203

Prepared by

ETIC Engineering, Inc. 2285 Morello Avenue Pleasant Hill, California 94523 (925) 602-4710

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Brent Searcy					Date
Project Engineer					
11/1//			24 Jan	2001	
Douglas E. Oram, Ph.D.		TERED GEOL			Date
Project Manager	N. S.	STERED GEOLOGICAL OF THE STERES			
Hadi SieNerbach-Corle		No. 6793 (R) *	January	24,2000	,
Heidi Dieffenbach-Carle, R.G. #6793	11	Lex03/31/02/	/ 0 0		Date
Senior Geologist	1	ATE OF CALIFORNIA			

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Risk Management Plan

Former Nestle Facility, 1310 14th Street, Oakland, California

Number	<u>Description</u>
1	Site location map.
2	Site map showing northwest portion of property to which environmental restrictions apply.
3	Site plan showing area of hydrocarbon impact to groundwater.
4	Diagram of conceptual site model.

LIST OF APPENDIXES

Risk Management Plan

Former Nestle Facility, 1310 14th Street, Oakland, California

<u>Appendix</u>	<u>Description</u>		
A	Covenant and Environmental Restriction Document		
В	Construction Worker Risk/Hazard Calculation Summary		

1. INTRODUCTION

ETIC Engineering, Inc. (ETIC) was contracted by Nestle USA, Inc. (Nestle) to prepare this Risk Management Plan (RMP) for the former Nestle property in Oakland, California. The property is located at 1310 14th Street, as shown in Figure 1. Figure 2 outlines the area for which a Deed Restriction has been recorded (Appendix A) and to which the restrictions and risk management protocols discussed in this document apply. This area will be referred to as the "subject facility" throughout this RMP. The RMP was prepared to fulfill property transfer requirements for the former Nestle property. The RMP presents the decision framework and the specific protocols for managing potential human health risks associated with the subsurface presence of chemicals and proposed future land use at the "subject facility". Potential health risks associated with daily occupants at the "subject facility" have been documented in a risk-based corrective action (RBCA) analysis for the "subject facility" (JCI 2000, as reported in ETIC 2001). This RBCA analysis was originally submitted to the Alameda County Health Agency (ACHA) and the California Regional Water Quality Control Board - San Francisco Bay Region (RWQCB) on 21 March 2000. Following discussions amongst ETIC, Nestle, ACHA, and RWQCB, comments on the RBCA analysis were addressed in a 27 June 2000 letter from ETIC to ACHA and RWQCB. A copy of the final RBCA analysis for the "subject facility" is included in ETIC Engineering's Comprehensive Site Characterization Report for the site dated January 2001. The risk to construction workers is evaluated in this RMP (see Appendix B).

The RMP delineates the specific risk management measures that will be implemented prior to, during, and after development of the "subject facility". It was prepared solely for use within the "subject facility" and is not intended for management of risks outside of this area. Although this RMP sets forth the requirements to appropriately manage the chemicals in soil and groundwater, the RMP is not intended to catalog all other legal requirements that may apply to the project or to activities conducted within the "subject facility" area.

Current and future owners and lessees, occupants and managers, or contractors delegated or authorized to perform property maintenance or construction are required to comply with the measures identified in the RMP when engaging in the relevant activities discussed. A Deed Restriction for the "subject facility" portion of the former Nestle property was recorded on 12 June 2000 at the Office of the Recorder of Alameda County (see Appendix A). Figure 2 shows the northwestern portion of the site, referred to as the "subject facility" in this RMP report, for which the deed restriction measures apply. The Deed Restriction requires Owner and/or Lessee compliance with the RMP measures. Specifically, the Deed Restriction places responsibility for compliance with the Owner and/or Lessee of the "subject facility" at the time the activity is conducted, even when such Owner or Lessee has contracted with another party to perform those measures. The term "Owner" or "Owners", as used in this RMP, shall mean those persons (whether individuals, corporations, or other legal entities) who, at such time when activities regulated by this RMP are conducted, hold title to the "subject facility". The term "Lessee" or 'Lessees" as used in this RMP shall mean those persons who are entitled by ownership, leasehold, license, permit, or other legal relationship with the Owner, to enter and exclusively occupy the "subject facility" and to engage in activities that are regulated by this RMP. A former Owner or former Lessee, licensee, permittee, or other former holder of a property or contract right who, at such time when activities regulated by this RMP are conducted, no longer holds an interest in title to a parcel or no longer has a property or contract interest in a parcel will not be considered an Owner or Lessee for the purposes of this RMP.

The California Environmental Protection Agency (Cal/EPA) has designated the ACHA as the "Administering Agency" under Assembly Bill (AB) 2061, in December 1998. As the Administering Agency, the ACHA is responsible for overseeing completion of the comprehensive site characterization study, the RBCA analysis, and the closure requirements of the "subject facility". The comprehensive site characterization study and RBCA analysis included the following tasks:

- Compilation and evaluation of historical soil and groundwater quality data;
- Field investigation (including collection of soil vapor, soil matrix, and groundwater quality data); and
- Preparation of comprehensive site characterization and RBCA reports.

The RBCA analysis concluded that the chemicals of potential concern (COPCs) observed at the "subject facility" do not pose a significant risk to daily site occupants following proposed redevelopment and commercial/industrial land use at the site. Accordingly, additional remediation at the site is not warranted, provided that future development will maintain a surface cap of the soil, exclusive of minor landscape areas, by buildings or paved surfaces. In addition, implementation of risk management practices, as described in this RMP, is recommended to address potential health risks associated with direct exposure of construction workers to chemicals beneath the site during redevelopment. To aid in development and implementation of risk management practices, the risk to construction workers was quantitatively evaluated and is summarized in Appendix B.

2. SITE BACKGROUND

2.1 SITE LOCATION

The former Nestle property is located at 1310 14th Street, Oakland, California. The property covers two city blocks and is bounded by 14th Street, 16th Street, Poplar Street, and Mandela Parkway (Figure 1). The entire site is covered with buildings and concrete or asphalt paving. The "subject facility" area of the property to which the environmental restrictions discussed in this document apply is located in the northwest portion of the property (Figure 2). This "subject facility" portion of the site contains an "L" shaped building which formerly housed warehouse and service bay facilities. The Cypress Structure of Interstate 880 (I-880), a former elevated freeway structure, existed west of the "subject facility" until it sustained extensive damage during the October 1989 Loma Prieta earthquake. This portion of I-880 was subsequently demolished and redeveloped as Mandela Parkway.

The topography slopes gently to the west, toward San Francisco Bay. Land use in the immediate area is primarily light industrial, with some commercial property and residences located east and west of the property.

2.1.1 Climate

Climatic conditions in the region are moderate, with mild, wet winters and warm, dry summers. Representative mean high/low temperatures and wind conditions are presented below:

	January	April	July	October	Annual
1990 Average temperature	52.3	61.5	66.0	65.5	60.9
(degrees F)					
Average wind speed	7.4	9.7	9.7	7.3	8.6
(mph, long term average)					
Average wind direction	SE	W	NW	W	W
(long term average)	***************************************				
1990 Rainfall	4.41	0.24	0.00	0.35	14.27
(inches)					

Temperature, rainfall from NOAA for Oakland Museum station (1990) Wind data from California Air Resources Board for Oakland International Airport (1984)

2.2 SITE HISTORY

The former facility was used to manufacture ice cream and packaged milk. The facility was also used for the distribution of ice cream and packaged fresh milk by trucks. A maintenance yard for vehicles used in the distribution of dairy products operated at the facility and included underground fuel and waste oil storage tanks.

Facilities at the property were originally constructed by American Creamery in 1915. Carnation purchased the property in 1929 and made additions and improvements to the buildings between 1946 and 1973 for dairy product processing and distribution. Nestle USA, Inc. assumed

operation of the property following its purchase of Carnation in 1985. Nestle ceased operations at the property in 1991 (HLA 1991).

2.2.1 Adjacent Land Use

Land use surrounding the site is light industrial and residential. Facilities to the north and south of the site are primarily light industrial. Immediately east of the site are light industrial facilities, with residential land use extending from approximately one block east of the site to Interstate 980 (I-980). West of the site is a mixed light industrial and residential area.

ETIC has conducted database searches and door-to-door well surveys for areas surrounding the site. No active water supply wells were identified during these efforts. Documentation of the surrounding area well surveys and database searches is provided in the Comprehensive Site Characterization Report (ETIC 2001).

2.3 SUMMARY OF SITE INVESTIGATIONS AND CURRENT ENVIRONMENTAL CONDITIONS

Previous environmental investigations conducted at the former Nestle property are briefly summarized below. More complete and detailed documentation of previous investigations and remediation activities is provided in the Comprehensive Site Characterization Report (ETIC 2001).

Four fuel underground storage tanks (USTs) were removed from the site in 1989. During removal of the USTs, gasoline and diesel fuel was observed as floating product in the tank cavity. Approximately 1,200 cubic yards of soil was excavated from the tank cavity and stockpiled onsite. Nutrients were applied to the soil stockpile in an attempt to bioremediate the soil. No further information regarding soil removal or disposal was available at the time this report was written.

Anania Geological Engineering (AGE) was retained by Carnation in 1989 to conduct a preliminary site characterization and to implement several interim remedial measures designed to contain and eliminate the presence of petroleum hydrocarbons in the soil and groundwater. A number of interim remedial actions were implemented, including installation of product recovery wells and removal of floating product, installation and operation of groundwater extraction and vapor extraction systems, and ex-situ bioremediation of soil. Thirty-three groundwater monitoring wells and 103 product recovery wells were installed at the site. Approximately 1.5 million gallons of groundwater were pumped and treated by carbon adsorption, resulting in the removal of approximately 5,000 gallons of gasoline and diesel fuel from soil and groundwater (HLA 1991).

In December 1990, Harding Lawson Associates was retained to review the preliminary site characterization and remediation data and to conduct additional site investigations. Between April and August 1991, HLA oversaw the installation of 20 soil borings. A soil vapor extraction (SVE) system was operated from January 1994 to December 1995 and removed an estimated 5,200 gallons of hydrocarbon equivalent (Park 1994; EA 1996).

At the end of 1995 the SVE system had removed most of the hydrocarbons that this technology is capable of removing, but floating product, or liquid-phase hydrocarbons (LPH), was still present in a number of wells. A multi-phase extraction system was installed and operated from August 1997 through June 2000. The system was installed to remove LPH trapped in the soil and floating on the groundwater. A total of 10,875 pounds of hydrocarbons have been removed since August 1997. Product levels have decreased since August 1997, and the hydrocarbon recovery rate has reached an asymptotic level. Figure 3 shows the area beneath which groundwater has historically been impacted by hydrocarbons.

A RBCA analysis for the site is included in the January 2001 Comprehensive Site Characterization Report. The RBCA analysis (JCI 2000) concluded that the chemicals of potential concern (COPCs) observed at the "subject facility" do not pose a significant risk to daily site occupants following proposed redevelopment and commercial/industrial land use at the site. Accordingly, additional remediation at the site is not warranted, provided that future development will maintain a surface cap of the soil, exclusive of minor landscape areas, by buildings or paved surfaces. In order to protect the health and safety of construction workers that may come into direct contact with chemicals beneath the site during future property redevelopment, the implementation of risk management practices, as outlined in Section 5 of this RMP document, is recommended.

2.4 FUTURE SITE DEVELOPMENT

As of January 2001, specific future development and/or construction plans for this site have not been presented. This document, in conjunction with the January 2001 Comprehensive Site Characterization Report for the site, identifies and outlines the risk management procedures which must be followed during any future development of the "subject facility". This RMP provides the specific protocols to be followed in order to mitigate risks to human health and the environment that were identified in the risk assessment portion of the Comprehensive Site Characterization Report.

Sections 5 and 6 of this report provide the risk management protocols which must be followed during and after any future site development activities. Appendix B presents the results of risk analysis efforts performed specifically for the purposes of developing a health and safety plan for protection of construction workers who may be involved in any future development activities at the site.

2.5 DEED RESTRICTION

The entire property was sold by Nestle to Encinal 14th Street, LLC in July 2000. Prior to the sale of the property, Covenants and Environmental Restrictions were developed for the "subject facility" area (northwest portion) of the property. The restrictions were reviewed by the ACHA and the RWQCB, and were signed by the City of Oakland Fire Services (COFS) in June 2000. These restrictions were recorded against the deed for the former Nestle property on 12 June 2000. Figure 2 shows a map of the entire property; the "subject facility" area (northwest portion), to which the environmental restrictions apply, is outlined and identified. A complete copy of the environmental restrictions is included as Appendix A.

3. SUMMARY OF HEALTH RISKS

A RBCA analysis was performed in support of comprehensive site characterization and the low risk designation requirement for the "subject facility" (JCI 2000). The RBCA analysis focused on potential health risks to construction workers and future daily occupants at and in the vicinity of the "subject facility", accounting for potential future development and land use at the "subject facility".

A conceptual site model (CSM) of contaminant occurrence, fate, transport, and potential exposure was developed as the basis for the RBCA analysis. A graphical representation of the CSM is depicted in Figure 4. As indicated on Figure 4, complete exposure pathways associated with daily onsite and offsite occupants include:

- Ingestion, inhalation, and dermal contact with surface soils (onsite industrial/commercial workers);
- Inhalation of volatile emissions and/or particulates from subsurface soils and groundwater to indoor air (onsite industrial/commercial workers);
- Inhalation of volatile emissions and/or particulates from subsurface soils and groundwater to outdoor air (onsite industrial/commercial workers);
- Inhalation of volatile emissions and/or particulates from groundwater to indoor air (offsite residents); and
- Inhalation of volatile emissions and/or particulates from groundwater to outdoor air (offsite residents).

The RBCA analysis did not include an evaluation of health risks to potential intermittent receptors such as site visitors and/or trespassers; however, the risks to daily site occupants may be used as a conservative estimate of risks to intermittent receptors.

Details of the RBCA analysis are documented by JCI (2000), included as an appendix to the Comprehensive Site Characterization report for the property (ETIC 2001). Conclusions of the RBCA analysis for daily onsite and offsite receptors included:

- Risks/hazards associated with direct exposure of daily site (commercial/industrial) occupants
 to observed levels of chemicals in surface soils are protective of USEPA-defined target
 risk/hazard levels;
- Risks/hazards associated with onsite (commercial/industrial) indoor and outdoor air inhalation of volatiles detected in shallow soil vapor samples are protective of USEPAdefined target risk/hazard levels;
- Risks/hazards associated with offsite (residential) indoor and outdoor air inhalation of volatiles detected in groundwater at offsite locations are protective of USEPA-defined target risk/hazard levels; and
- An RMP outlining appropriate risk management practices, health and safety measures, and deed restrictions should be developed prior to initiation of construction activities and redevelopment at the "subject facility".

To aid in development of this RMP and a health and safety plan for protection of construction workers, risks to construction workers were also quantified, as summarized in Appendix B. The construction worker risk analysis indicates that without protective measures, the carcinogenic risk associated with exposure of construction workers to subsurface chemicals is within the target risk range adopted by the USEPA, while the non-carcinogenic hazard marginally exceeds the target hazard level. Therefore, to prevent construction workers from potentially hazardous exposure levels at the "subject facility", the recommendations in this RMP document should be implemented.

4. RISK MANAGEMENT MEASURES PRIOR TO SITE DEVELOPMENT

Potential exposure prior to development of the "subject facility" is limited to intermittent visitors or trespassers. As indicated in Section 3, the risk to intermittent receptors is considered insignificant. Moreover, due to the presence of a fence around the property and a paved surface throughout much of the property, additional risk management measures prior to development of the "subject facility" are not warranted.

5. RISK MANAGEMENT MEASURES DURING SITE DEVELOPMENT

The Deed Restriction for the "subject facility" indicates that no owners or occupants of the "subject facility" or any portion thereof shall conduct any excavation work on the "subject facility", unless expressly permitted in writing by the COFS. Should excavation be permitted as part of redevelopment, the primary exposure to chemicals at the "subject facility" will be limited to that associated with construction workers. As indicated in Section 3, risk management measures are recommended for protection of construction workers. To this end, risk management measures were developed to provide adequate protection to human health for onsite construction workers during development of the "subject facility".

Development activities at the facilities may include various site preparation activities such as, but not limited to, excavation, stockpiling, trenching, site grading, backfilling, and dewatering that may disturb the native soils and/or groundwater beneath the "subject facility". Specifically, potential events or activities associated with development of the "subject facility" that may result in potential health impacts to onsite construction workers during development include:

- Dust generation associated with soil excavation and trenching, grading, loading activities, backfilling, movement of construction and transportation equipment, and fugitive dust generation from winds traversing an exposed soil stockpile; and
- Potential contact with subsurface chemicals during trenching and excavation.

The risk management measures that will control potential impacts associated with each of these activities are described below. Management measures that are recommended to control potential impacts on construction workers, contractors, and short-term intrusive workers who may be engaged in limited excavation activities, such as utility repair, are also described below.

5.1 SITE-SPECIFIC HEALTH AND SAFETY REQUIREMENTS AND SAFETY PLAN

The construction contractor shall assume full responsibility and liability for the compliance with provisions of the Work Hours and Safety Standard Act (40 U.S.C. 327 et seq.). The construction contractor shall comply with all applicable safety regulations and other requirements, including, but not limited to, the following:

- Code of Federal Regulations (CFR), Title 29-Labor
- State of California, California Code of Regulations (CCR), Industrial Relations
- Medical Surveillance Programs (e.g., OSHA, 29 CFR 1200)
- Injury and Illness Prevention Programs (e.g., SB 198, 8 CCR, CAL/OSHA, GISO 3203, Section 5192 and 1509)
- Implementation of mitigation measures under California Environmental Quality Act (CEQA), if any
- The Construction Standard (29 CFR 1926)
- Workers' Right to Know (29 CFR 1910.120)
- Section 6360-99 of the California Labor Code (Hazard Communication)

During construction and site development activities, workers that may directly contact contaminated soil or groundwater at the "subject facility" must perform their activities in accordance with a hazardous operations site-specific health and safety plan (HASP). The construction contractor will be responsible for development and implementation of the HASP in compliance with all applicable federal, state, and local regulations and requirements. The HASP shall be prepared by a Certified Industrial Hygienist. If needed, the construction contractor will submit the HASP to the RWQCB or ACHA for review. Preparation of a HASP will be required for, but not limited to, site preparation work including grading, utility installation, foundation construction, service pit construction, and other activities where workers might directly contact impacted soil or groundwater beneath the "subject facility". The HASP shall include, but not be limited to, the following elements:

- Identification and description of the responsibility of those individuals who control each phase of operations and are responsible for employee and public safety. The plan shall set forth in writing the policies and procedures to be followed by all personnel. This shall include designation of an overall project site safety representative with authority to stop any construction/demolition activity or modify work practices if the site safety plan is being violated, or if such action is necessary to protect workers, property, and the surrounding community during the contract period. This requirement shall apply continuously and not be limited to normal working hours.
- Information identifying and delineating all workplace hazards that have been identified or are generally associated with the proposed work phases, and how this information is communicated to employees (e.g., tailgate safety meetings). Hazardous material communication standards can be found in 29 CFR 1910.120 and 8 OCR 5194. Hazardous waste information can be found in 29 CFR 1910.1200 and 8 CCR 5192.
- Engineering controls, specific work practices, and measures to be used to monitor and control worker and general public exposure to any identified hazard with special emphasis to demolition debris, dust, petroleum impacted soils, LPH, and other hazardous materials. The monitoring of site personnel for contaminant exposure shall be conducted so as to maintain the proper level of personal protection, including action level of protection.
- Level of training required for all specified contractor(s) or subcontractor personnel, possibly, but not limited to, asbestos, lead and hazardous materials awareness training; and the 40-hour Hazardous Waste Operations and Emergency Response Training Program and the associated 8-hour refresher training in accordance with Title 29, Code of Federal Regulations 1910.120, and 8 CCR 5192 for all personnel who will come in direct contact with surface and subsurface contaminated materials when performing their work. Contractors shall maintain and provide all training records to the Resident Engineer.
- Provision of sufficient personnel properly trained to handle, excavate, and dispose of hazardous waste and other contaminated waste that is expected in this project. The training shall be in accordance with 29 CFR 1910.120, 29 CFR 1910.134, 8 CCR 5144, and 8 CCR 5192.

- Requirements of contractors and subcontractors for any applicable medical surveillance programs and Injury and Illness Prevention Program (IIPP) (e.g., SB 198, 8 CCR and CAL/OSHA, GISO 3203, Sections 5192 and 1509); implementation of mitigation measures under CEQA (AB 3180); the Construction Standard (29 CFR 1926); Workers Right to Know (29 CFR 1910.120); Section 6360-99 of the California Labor Code (Hazard Communication); the San Francisco Health Code, Article 21 addressing Hazardous Materials, and the Americans with Disabilities Act (ADA).
- Methods to be used to decontaminate equipment.
- Sanitation facilities to be provided for personal hygiene. Portable toilets and discharge of their waste products into sanitary sewers shall comply with local codes.
- Contingency Plan for emergency including fire, spillage of hazardous/toxic wastes and liquids (with special emphasis to clean up spillage due to fuel/oil from contractors' equipment), traffic accident, personal accident, power failure, or any event that may require modification or abridgment of site control and decontamination procedures. This plan shall also include procedures to be followed in the event of a large-scale spill of contaminated material on a public roadway in accordance with the hazardous Substance Highway Spill Containment and Abatement Act (California Vehicle Code, Section 2450 et seq.), and the Emergency Service Act (California Government Code Section 8571.4 et seq.)

5.2 CONSTRUCTION IMPACT MITIGATION MEASURES

Measures must also be implemented to mitigate potential health impacts on construction workers, should they be exposed directly to chemicals in soil and groundwater underlying the "subject facility". Potential exposure pathways associated with onsite construction workers include inhalation, incidental ingestion, and dermal contact with chemicals in soils and groundwater.

Specifically, measures that must be implemented to mitigate potential impacts during construction include the following:

- Each contractor will prepare and implement a site-specific HASP to address the potential exposure to contaminated soils and groundwater during construction;
- Dust control through spraying of water and other techniques to minimize mobility of impacted soils toward offsite locations;
- Minimize soil and groundwater contact by onsite construction worker.

Details of these mitigation measures, except the site-specific health and safety plan, are described below.

5.2.1 Dust Control

Dust controls must be implemented to prevent offsite dispersion and accumulation of impacted soils and to comply with applicable regulations pertaining to air quality and nuisance control. Potential construction activities that could generate dust and warrant risk management measures include: (1) excavation and stockpile control; (2) onsite construction vehicle traffic, and (3) windblown soil.

Alameda County may require monitoring of dust generation during site construction at the "subject facility". Results of the monitoring will be used by the construction contractor for determining the needs and appropriate dust control practices in accordance with the regulations for excavating and restoring streets in Alameda County.

Dust generation will be minimized by all appropriate measures, which may include, but not be limited to, the following:

- Wetting of surface soils and spoil piles during excavation, trenching, compaction, and site grading and paving;
- Control of excavation techniques to minimize dust generation such as minimizing drop distances; and
- Covering of stockpiles, if present, with visqueen or other suitable membrane covers.

Additional measures, if required, may be utilized at the discretion of the construction contractor.

5.2.2 Minimizing Soil and Groundwater Contact by Construction Worker

Existing data indicate the subsurface presence of chemicals in both unsaturated soils, saturated soils, and groundwater beneath the "subject facility". Shallow groundwater beneath the site occurs at depths ranging from 5.0 to 10.0 feet below ground surface. Details of the hydrogeological characterization are presented in the Comprehensive Site Characterization Report (ETIC 2001).

Future construction work at the site may involve excavation and/or direct contact with chemicals above and below the water table. To mitigate risks associated with this exposure, the construction contractor shall develop and implement a site-specific HASP. Examples of health and safety measures are the use of protective clothing, protective gloves and boots, and suitable respirators with cartridges during construction activities.

6. RISK MANAGEMENT MEASURES AFTER SITE DEVELOPMENT

The post-construction portion of this RMP addresses the precautions that must be undertaken to mitigate the long-term health risks associated with the "subject facility" after all redevelopment activities are complete. Any future reuse of the "subject facility" involving disturbance of soil, pavements, or building foundations must be accomplished in a manner consistent with the objectives of this RMP.

Components of the post-construction portion of this RMP include the following:

- Prevention of the exposure of daily site occupants or visitors to impacted soil by maintaining cover materials in appropriate conditions;
- Establishment of protocols to protect onsite workers engaged in subsurface excavation activities such as buried utility repair, work on buried foundations, or pavement requiring exposure to soil and/or groundwater;
- Prevention of use of groundwater beneath the facility;
- Agency (COFS, ACHA, and RWQCB) notification on change in property use.

6.1 COVERING OF THE SITE

As indicated in the Deed Restriction for the former Nestle property, all uses and development of the "subject facility" shall maintain a surface cap of the soil, exclusive of minor landscape areas, by buildings or paved surfaces. The Maintenance and Operations Facility Manager or their designated representative must annually conduct a visual inspection of the cover to ensure that the cover materials remain in adequate shape. Damage to the integrity of the cover materials, such as major cracks, must be promptly repaired.

Upon completion of the inspection and any necessary repairs, the Maintenance and Operations Facility Manager or their designated representative will prepare a report documenting the inspection and repairs. The report will contain, at a minimum, the following information:

- Date of inspection
- Personnel conducting the inspection
- Results of the inspection
- Repairs completed to maintain the integrity of the cover

Reports must be signed by the Maintenance and Operations Facility Manager or their designated representative. Reports must be filed by the site occupant at the Maintenance and Operations Facility. The reports will be available for review by the COFS, ACHA, and RWQCB.

6.2 PROTOCOLS FOR FUTURE SUBSURFACE DEVELOPMENT

If excavation is permitted by the COFS, health and safety procedures must be followed, as previously described, for all individuals engaged in subsurface excavation activities in which covered soil and groundwater may be exposed. The likely scenarios are buried utility repairs, work on buried foundations, or repairs and alterations to pavements. At a minimum, a site-specific HASP must be prepared and employed in concert with any such work.

If minor soil disturbance is undertaken in the future, the work must follow the guidelines presented herein. Any impacted soil subject to excavation and brought to the surface by grading, excavation, trenching, or backfilling shall be managed in accordance with all applicable provisions of local, state, and federal laws. Excavated soil may be reused as backfill in the excavation area, provided that the excavation will be properly covered with asphalt, concrete, or clean material. Excess material must be disposed of offsite at an appropriate waste facility.

If future activities at the "subject facility" are planned involving a significant reduction in the extent or effectiveness of the cap over the soil, then an addendum to this RMP must be prepared and submitted to the ACHA and RWQCB.

6.3 USE OF GROUNDWATER

As indicated in the Deed Restriction for the former Nestle property, no owner or occupants of the "subject facility" shall drill, bore, otherwise construct, or use a well for the purpose of extracting groundwater for any use, including, but not limited to, domestic, potable, or industrial uses, unless expressly permitted in writing by the ACHA and RWQCB.

6.4 AGENCY NOTIFICATION ON CHANGE OF PROPERTY USE

As indicated in the Deed Restriction for the former Nestle property, land use at the "subject facility" will be restricted to industrial, commercial, or office space. Use of the "subject facility" as a residence for human habitation, hospital, school for persons under 21 years of age, and/or day care center is also prohibited by the Deed Restriction.

REFERENCES

EA (EA Engineering, Science, and Technology). 1996. Product Recoverability and Vapor Extraction/Air Sparging Pilot Test Report for the Nestle USA Former Carnation Dairy Facility, 1310 14th Street, Oakland, California. EA, Lafayette, California. July.

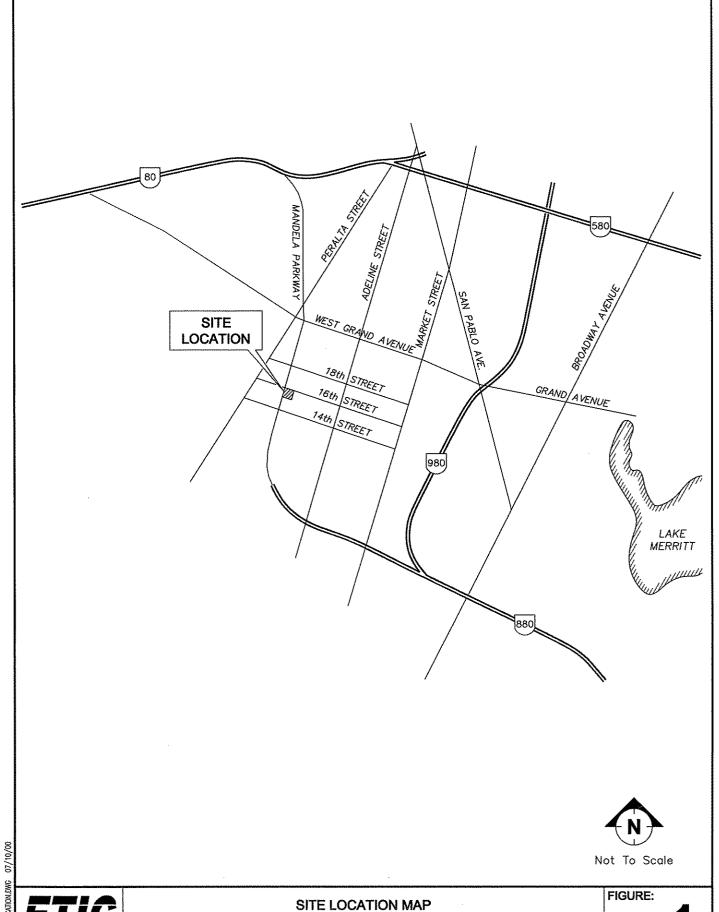
ETIC (ETIC Engineering, Inc.). 2001. Comprehensive Site Characterization Report, Support for the Site as a Low-Risk Soil and Groundwater Case, Former Nestle USA, Inc. Facility, Oakland, California. ETIC, Pleasant Hill, California. January.

HLA (Harding Lawson Associates). 1991. Site Characterization Report, Carnation Facility, Oakland, California. HLA, Novato, California. 17 September.

JCI (Javaherian Consulting, Inc.). 2000. Technical Memorandum: Risk-Based Corrective Action Analysis, Nestle USA, Inc. Facility, 1310 14th Street, Oakland, California. JCI, San Francisco, California. 22 August.

Park (Park Environmental). 1994. Vapor Extraction Remediation Update, October 1993 through April 1994, Carnation Company Facility, 1310 14th Street, Oakland, California. Park, Rocklin, California. 19 May.

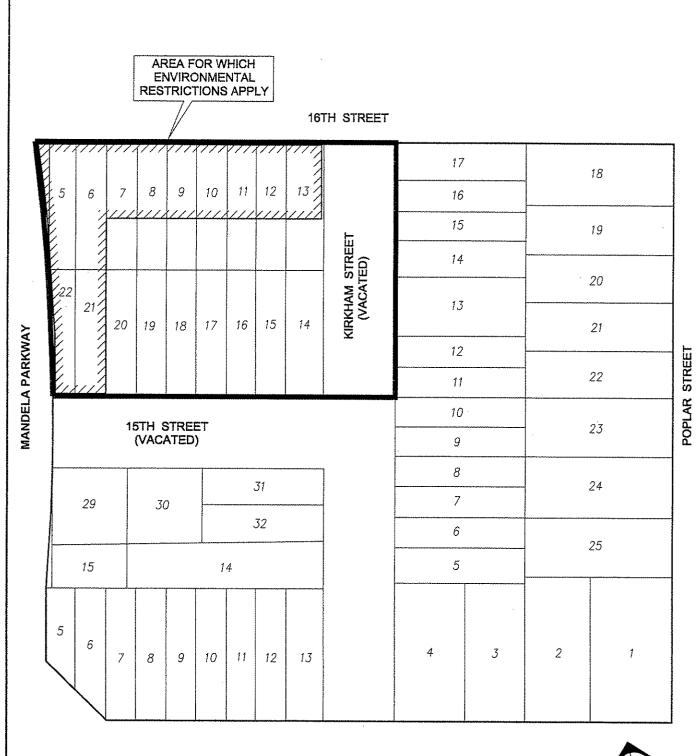




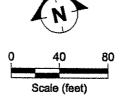
ETIC Engineering, Inc.

SITE LOCATION MAP NESTLE OAKLAND FACILITY 1310 14th STREET, OAKLAND, CALIFORNIA

1



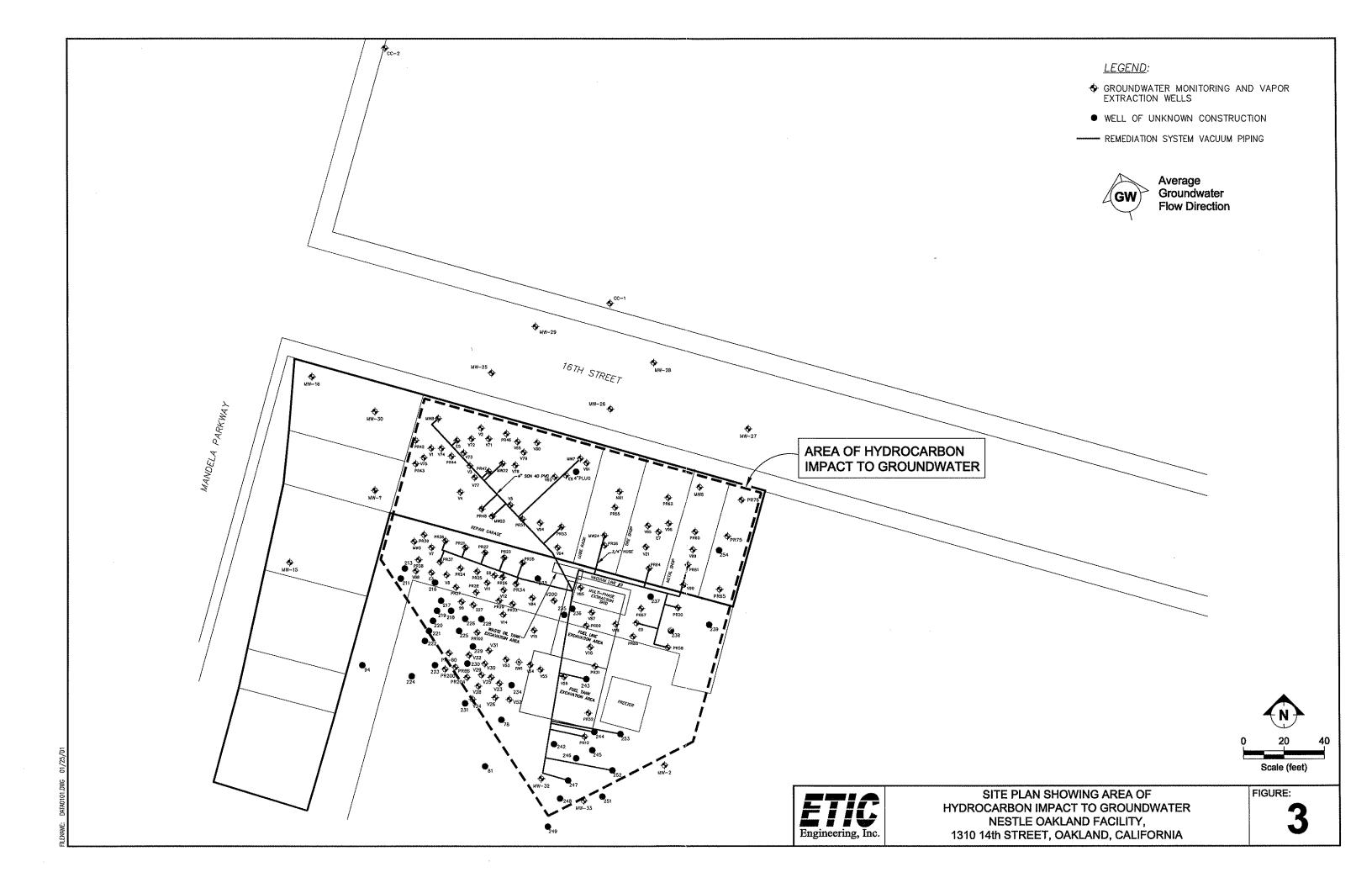
14TH STREET

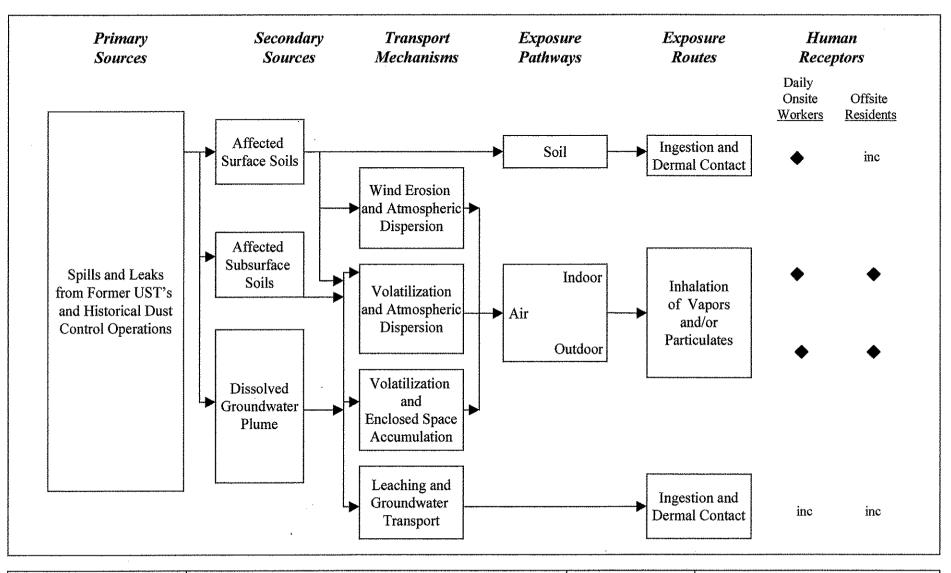


ETIC Engineering, Inc. FORMER NESTLE FACILITY (CARNATION DAIRY FACILITY) SHOWING NORTHWEST SECTION FOR WHICH ENVIRONMENTAL RESTRICTIONS APPLY, NESTLE OAKLAND FACILITY, 1310 14th STREET, OAKLAND, CALIFORNIA

FIGURE:

2





Complete pathwayinc Incomplete pathway

Conceptual Site Model
Nestle Oakland Facility
1310 14th Street, Oakland, California

Figure
4
August 2000



Appendix A

Covenant and Environmental Restriction Document

FIRST ALERNO AT TITLE

/58458

Recording Requested By:

Nestle USA Inc. 800 North Brand Blvd. Glendale, California 91203

When Recorded, Mail To:

Leroy Griffin
Hazardous Materials Program Supervisor
City of Oakland Fire Services
1605 Martin Luther King Jr. Way
Oakland, California 94612

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COVENANT AND ENVIRONMENTAL RESTRICTION ON PROPERTY

Northeast Portion of the Former Carnation Dairy Facility which Occupies 1315-1372 14th Street and 1315-1385 16th Street

This Covenant and Environmental Restriction on Property (this "Covenant") is made as of the 8th day of Juliano, 2000 by Nestle USA ("Covenantor") who is the Owner of record of that certain property situated at 1315-1372 14th Street and 1315-1385 16th Street, in the City of Oakland, County of Alameda, State of California, which contains a contaminated area which is more particularly described in Exhibit A attached hereto and incorporated herein by this reference (such contaminated area hereinafter referred to as the "Burdened Property"), for the benefit of the City of Oakland Fire Services (COFS), with reference to the following facts:

- A. The Burdened Property and groundwater underlying the property contains hazardous materials.
- B. Contamination of the Burdened Property. Soil at the Burdened Property was contaminated by releases from petroleum underground storage tanks. These releases resulted in contamination of soil and groundwater with organic chemicals including benzene, toluene, ethylbenzene, xylenes, and 1,2—dichloroethane, which are hazardous materials as that term is defined in Health & Safety Code Section 25260. Removal of underground storage tanks and remediation of the petroleum hydrocarbons was initiated in January 1988 and is summarized below:

Tank, Line, and Dispenser Removal

Four (4) underground fuel storage tanks and associated piping were removed in December 1988. One (1) 1,000 gallon used-oil tank was removed in January 1989.

Remedial Actions

Soil Excavation: Between January and March 1989, 1,200 cubic yards of soil were removed in the area of the former underground storage tanks and associated piping. This soil was treated on-site and replaced back in the excavated area.

Liquid Petroleum Hydrocarbon Removal: Liquid petroleum hydrocarbons were removed using a product skimming system from the subsurface during January through March 1989. Approximately 1,800 gallons were removed during this time period.

Soil Vapor Extraction: A soil vapor extraction system operated from January 1994 to December 1995 and removed an estimated 5,200 gallons of hydrocarbon.

Multi-phase Extraction: A multi-phase extraction system has been operating at the site since August 1997. Approximately 10,500 pounds of hydrocarbons have been removed using this system. Thickness of petroleum hydrocarbons decreased since August 1997.

- C. <u>Exposure Pathways</u>. The contaminants addressed in this Covenant are present in soil and groundwater on the Burdened Property. Without the mitigation measures which have been performed on the Burdened Property, exposure to these contaminants could take place via the following pathways (onsite workers only):
 - > Ingestion and dermal contact with surface soils;
 - > Inhalation of volatile emissions from subsurface soils and groundwater

The risk of public exposure to the contaminants has been substantially lessened by the remediation and controls described in part B.

- D. Adjacent Land Uses and Population Potentially Affected. The Burdened Property is currently an unused industrial facility and is adjacent to industrial, commercial, and residential land uses.
- E. Full and voluntary disclosure to the COFS of the presence of hazardous materials on the Burdened Property has been made and extensive sampling of the Burdened Property has been conducted.
- F. Covenantor desires and intends that in order to benefit the COFS, and to protect the present and future public health and safety, the Burdened Property shall be used in such a manner as to avoid potential harm to persons or property that may result from hazardous materials that may have been deposited on portions of the Burdened Property.

ARTICLE I GENERAL PROVISIONS

- 1.1 Provisions to Run with the Land. This Covenant sets forth protective provisions, covenants, conditions and restrictions (collectively referred to as "Restrictions") upon and subject to which the Burdened Property and every portion thereof shall be improved, held, used, occupied, leased, sold, hypothecated, encumbered, and/or conveyed. The restrictions set forth in Article III are reasonably necessary to protect present and future human health and safety or the environment as a result of the presence of hazardous materials in the subsurface below the Burdened Property. Each and all of the Restrictions shall run with the land, and pass with each and every portion of the Burdened Property, and shall apply to, inure to the benefit of, and bind the respective successors in interest thereof, for the benefit of the COFS and all Owners and Occupants. Each and all of the Restrictions are imposed upon the entire Burdened Property. Each and all of the Restrictions run with the land pursuant to section 1471 of the Civil Code. Each and all of the Restrictions are enforceable by the California Regional Water Quality Control Board for the San Francisco Bay Region (the "Board").
- 1.2 Concurrence of Owners and Lessees Presumed. All purchasers, lessees, or possessors of any portion of the Burdened Property shall be deemed by their purchase, leasing, or possession of such Burdened Property, to be in accord with the foregoing and to agree for and among themselves, their heirs, successors, and assignees, and the agents, employees, and lessees of such owners, heirs, successors, and assignees, that the Restrictions as herein established must be adhered to for the benefit of the COFS and the Owners and Occupants of the Burdened Property and that the interest of the Owners and Occupants of the Burdened Property shall be subject to the Restrictions contained herein.
- 1.3 Apportionment of Burden Among Multiple Owners. Where ownership of the Burdened Property is held by multiple persons, holding by several titles, the burdens imposed by this Covenant shall be apportioned between them proportionate to the value of the property held by each owner, if such value can be ascertained, and if not, then according to their respective interests in point of quantity. (Cal. Civ. Code, § 1467.)
- 1.4 Incorporation into Deeds and Leases. Covenantor desires and covenants that the Restrictions set out herein shall be incorporated in and attached to each and all deeds and leases of any portion of the Burdened Property. Recordation of this Covenant shall be deemed binding on all successors, assigns, and lessees, regardless of whether a copy of this Covenant and Agreement has been attached to or incorporated into any given deed or lease.
- 1.5 <u>Purpose</u>. It is the purpose of this instrument to convey to the COFS real property rights, which will run with the land, to facilitate the remediation of past environmental contamination and to protect human health and the environment by reducing the risk of exposure to residual hazardous materials.

ARTICLE II DEFINITIONS

- 2.1 <u>COFS</u>. "COFS" shall mean the City of Oakland Fire Services and shall include its successor agencies, if any.
- 2.2 <u>Board</u>. "Board" shall mean the California Regional Water Quality Control Board for the San Francisco Bay Region and shall include its successor agencies, if any.
- 2.3 <u>Improvements</u>. "Improvements" shall mean all buildings, roads, driveways, regradings, and paved parking areas, constructed or placed upon any portion of the Burdened Property.
- 2.4 Occupants. "Occupants" shall mean Owners and those persons entitled by ownership, leasehold, or other legal relationship to the exclusive right to use and/or occupy all or any portion of the Burdened Property.
- 2.5 Owner or Owners. "Owner" or "Owners" shall mean the Covenantor and/or its successors in interest, who hold title to all or any portion of the Burdened Property.

ARTICLE III

DEVELOPMENT, USE AND CONVEYANCE OF THE BURDENED PROPERTY

- 3.1 <u>Restrictions on Development and Use</u>. Covenantor promises to restrict the use of the Burdened Property as follows:
- a. Development of the Burdened Property shall be restricted to industrial, commercial or office space;
 - b. No residence for human habitation shall be permitted on the Burdened Property;
 - c. No hospitals shall be permitted on the Burdened Property;
- d. No schools for persons under 21 years of age shall be permitted on the Burdened Property;
- e. No day care centers for children or day care centers for Senior Citizens shall be permitted on the Burdened Property;
- f. No Owners or Occupants of the Burdened Property or any portion thereof shall conduct any excavation work on the Burdened Property, unless expressly permitted in writing by the COFS. Any contaminated soils brought to the surface by grading, excavation, trenching, or

backfilling shall be managed by Covenantor or his agent in accordance with all applicable provisions of local, state and federal law;

- g. All uses and development of the Burdened Property shall be consistent with any applicable Board Order or Risk Management Plan, each of which is hereby incorporated by reference including future amendments thereto. All uses and development shall preserve the integrity of any cap, any remedial measures taken or remedial equipment installed, and any groundwater monitoring system installed on the Burdened Property pursuant to the requirements of the COFS, unless otherwise expressly permitted in writing by the COFS. Any development of the Burdened Property will maintain a surface cap of the soil, exclusive of minor landscape areas, by buildings or paved surfaces.
- h. No Owners or Occupants of the Property or any portion thereof shall drill, bore, otherwise construct, or use a well for the purpose of extracting water for any use, including but not limited to, domestic, potable, or industrial uses, unless expressly permitted in writing by the Board.

3.1.1 Notifications/Access/Non Aggravation

- a. The Owner shall notify the COFS of each of the following: (1) The type, cause, location and date of any disturbance to any cap, any remedial measures taken or remedial equipment installed, and of the groundwater monitoring system installed on the Burdened Property pursuant to the requirements of the COFS, which could affect the ability of such cap or remedial measures, remedial equipment, or monitoring system to perform their respective functions and (2) the type and date of repair of such disturbance. Notification to the COFS shall be made by registered mail within ten (10) working days of both the discovery of such disturbance and the completion of repairs;
- b. The Covenantor agrees that the COFS, and/or any persons acting pursuant to COFS orders, shall have reasonable access to the Burdened Property for the purposes of inspection, surveillance, maintenance, or monitoring, as provided for in Division 7 of the Water Code.
- c. No Owner or Occupant of the Burdened Property shall act in any manner that will aggravate or contribute to the existing environmental conditions of the Burdened Property. All use and development of the Burdened Property shall preserve the integrity of any capped areas.
- 3.2 <u>Enforcement</u>, Failure of an Owner or Occupant to comply with any of the restrictions, as set forth in paragraph 3.1, shall be grounds for the COFS, by reason of this Covenant, to have the authority to require that the Owner modify or remove any Improvements constructed in violation of that paragraph. Violation of the Covenant shall be grounds for the COFS to file civil actions against the Owner as provided by law.
- 3.3 <u>Notice in Agreements</u>. After the date of recordation hereof, all Owners and Occupants shall execute a written instrument which shall accompany all purchase agreements or leases relating to the property. Any such instrument shall contain the following statement:

The land described herein contains hazardous materials in soils and in the ground water under the property, and is subject to a deed restriction dated as of <u>June 8</u>, 2000, and recorded an <u>Concurrently herewith 266</u>, in the Official Records of <u>Alameda</u> County, California, and restrictions on usage of the property described herein. This statement is not a declaration that a hazard exists.

ARTICLE IV VARIANCE AND TERMINATION

- 4.1 <u>Variance.</u> Any Owner or, with the Owner's consent, any Occupant of the Burdened Property or any portion thereof may apply to the COFS for a written variance from the provisions of this Covenant.
- 4.2 <u>Termination</u>. Any Owner or, with the Owner's consent, any Occupant of the Burdened Property or a portion thereof may apply to the COFS for a termination of the Restrictions as they apply to all or any portion of the Burdened Property which consent to termination shall not be unreasonably withheld.
- 4.3 <u>Term.</u> Unless terminated in accordance with paragraph 4.2 above, by law or otherwise, this Covenant shall continue in effect in perpetuity.

ARTICLE V MISCELLANEOUS

- 5.1 No Dedication Intended. Nothing set forth herein shall be construed to be a gift or dedication, or offer of a gift or dedication, of the Burdened Property or any portion thereof to the general public.
- 5.2 Notices. Whenever any person gives or serves any notice, demand, or other communication with respect to this Covenant, each such notice, demand, or other communication shall be in writing and shall be deemed effective (1) when delivered, if personally delivered to the person being served or official of a government agency being served, or (2) three (3) business days after deposit in the mail if mailed by United States mail, postage paid certified, return receipt requested:

If To: "Covenantor"
Nestlé USA, Inc.
Legal Department
800 North Brand Boulevard
Glendale, Ca. 91203

If To: "COFS"

City of Oakland Fire Services

Attention: Hazardous Materials Program Supervisor

1605 Martin Luther King Jr. Way

Oakland, California 94612

- 5.3 Partial Invalidity. If any portion of the Restrictions or terms set forth herein is determined to be invalid for any reason, the remaining portion shall remain in full force and effect as if such portion had not been included herein.
- 5.4 <u>Article Headings</u>. Headings at the beginning of each numbered article of this Covenant are solely for the convenience of the parties and are not a part of the Covenant.
- 5.5 Recordation. This instrument shall be executed by the Hazardous Materials Program Supervisor of the COFS. This instrument shall be recorded by the Covenantor in the County of Alamada within ten (10) days of the date of execution.
 - 5.6 References. All references to Code sections include successor provisions.
- 5.7 <u>Construction</u>. Any general rule of construction to the contrary notwithstanding, this instrument shall be liberally construed in favor of the Covenant to effect the purpose of this instrument and the policy and purpose of the Water Code. If any provision of this instrument is found to be ambiguous, an interpretation consistent with the purpose of this instrument that would render the provision valid shall be favored over any interpretation that would render it invalid.

IN WITNESS WHEREOF, the parties execute this Covenant as of the date set forth above.

Coveni	antor: NESTLE	USA, INC.
Ву:	Monder	Robert H. Sanders
Title:	V. P.	
Date:	6.8.00	

Agency:

City of Oakland Fire Services

By: LeRoy Griffin
Title: Hazardous Materials Program Supervisor

APPENDIX A

LEGAL DESCRIPTION DEED RESTRICTION AREA

ROBERT BEIN, WILLIAM FROST & ASSOCIATES 1981 N. Broadway, Suite 235 Walnut Creek, California 94596

LEGAL DESCRIPTION DEED RESTRICTION AREA

That certain parcel of land situated in the City of Oakland, County of Alameda, State of California described as follows:

Being a portion of Lots 4 through 23 and a portion Kirkham Street of the Scotchler Tract and Vicinity, Oakland, as shown on a map thereof filed in Book 7 of Maps at Page 21 on December 10, 1874 in the Office of the County Recorder of Alameda County more particularly described as follows:

BEGINNING at the intersection of said Kirkam Street and the northwest corner of lot 17, in block 584, as shown on the map of "Re-division of Blocks 584, 585, 601, 153 and 580-A, City of Oakland, County of Alameda, California", filed May 1, 1885, in Book 4 of Maps, at Page 25, in said office of the County Recorder;

Thence, along the northerly line of said Kirkham Street and said lots 13, 12, 11, 10, 9, 8, 7, 6 and 5, North 72°53'28" West 292.25 feet to the northwest corner of said lot 5, said point also being the northeasterly corner of that certain parcel of land described in the deed to the State of California, recorded May 12, 1955 in Volume 7658, of Official Records at Page 299, in said office of the County Recorder;

Thence, continuing along said northerly line of Kirkham Street, North 72°53'28" West 8.64 feet:

Thence, along said State of California parcel, along a non-tangent 1240 foot radius curve to the right, through a central angle of 2°59'04" to the easterly line of the parcel of land described in the deed to the State of California, recorded August 12, 1955 in Book 7749, of Official Records at Page 447, as Instrument Number AK-86901, in said office of the County Recorder;

Thence, along last said State of California parcel (7749 OR 447), along a non-tangent 1240 foot radius curve to the right from a tangent that bears South 10°54'36" West to the south line of said lot 22, said southerly line also being the north line of 15th Street, as shown on said map of the Scotchler Tract (7 M 21);

Thence, along said northerly line of 15th Street and the easterly prolongation of said north line. South 74°03'30" East 285.05 feet to the easterly line of said Kirkham Street;

Thence, along said easterly line, North 15°56'30" West 209.50 feet to the POINT OF BEGINNING.

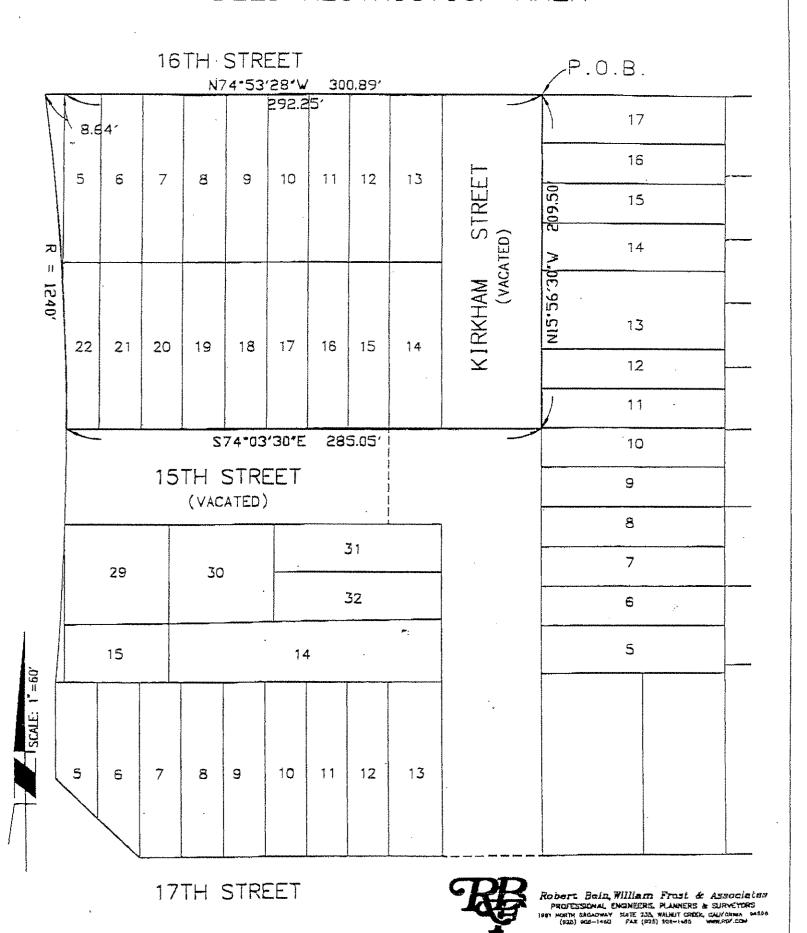
EXHIBIT attached and by this reference made a part hereof.

Patrick J. Tami, L.S. 5816

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April 19, 2000 (4:28PM)

DEED RESTRICTION AREA



County of Alameda		
County of France a		
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	pro	oved to me on the basis of satisfactory evidence
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	personally known to me
- -	Troved to me on the basis of satisfactory evidence
MARIA HAZEL PERRI	to be the person(s) whose name(s) is/are
Commission # 1113713	subscribed to the within instrument and
Notary Public — California E Los Angeles County	acknowledged to me that he/she/they executed the same in his/her/their authorized
My Comm. Expires Cat 13, 2000	capacity(ies), and that by his/her/their
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Appendix B

Construction Worker Risk/Hazard Calculation Summary

APPENDIX B CONSTRUCTION WORKER RISK/HAZARD CALCULATION SUMMARY

This appendix summarizes estimates of carcinogenic risk and noncarcinogenic hazards associated with exposure of construction workers to chemicals underlying the deed restricted portion of the former Nestle property. This portion of the former Nestle property is referred to as the "subject facility" in this document. The approach to estimating risks to construction workers followed the methodology outlined by the California Environmental Protection Agency Regional Water Quality Control Board - San Francisco Bay Region (RWQCB) guidelines for Application of Risk-Based Screening Levels to Sites with Impacted Soil and Groundwater (RWQCB 2000), wherein potential health risks to construction workers are assessed based on evaluation of direct exposure to chemicals of potential concern (COPCs) in soils. This methodology is similar to that adopted by the American Society for Testing and Materials ([ASTM] 1995 and 1998).

Details and assumptions behind RWQCB's approach to evaluation of risks to construction workers are documented in Appendix 1 of RWQCB (2000). A brief summary is provided below. Conservative exposure assumptions by RWQCB (2000) for evaluation of direct exposure of construction workers to COPCs are based on guidance presented in the USEPA Exposure Handbook (USEPA 1997), trench-worker risk-assessment guidance developed by the Massachusetts Department of Environmental Protection (MADEP 1994), and general directexposure assumptions included in the USEPA Region IX Preliminary Remediation Goals document (USEPA 1999), focusing on direct contact via ingestion, dermal, and inhalation routes of exposure. Key among these assumptions is the use of an exposure duration of 7 years, an exposure frequency of 20 days per year, a soil ingestion rate of 480 mg/day, and a particulate emission factor corresponding to a concentration of air-born dust of approximately 700 ug/m³. Based on these conservative assumptions, risks to construction workers are generally lower than those to commercial/industrial receptors, which in turn are lower than those to residential receptors. This pattern is primarily due to the assumed shorter exposure duration and frequency associated with the construction worker exposure, in comparison with the other two exposure scenarios. Exceptions to this pattern may occur for chemicals with high oral toxicity such as various heavy metals, none of which are considered COPCs at the "subject facility". Direct exposure to COPCs in onsite soils for commercial/industrial receptors was previously evaluated in the risk-based corrective action (RBCA) analysis for the "subject facility" portion of the former Nestle site (JCI 2000, as reported in ETIC 2001).

To estimate potential health risks to future construction workers at the Nestle facility, all chemicals detected in recent sampling of surface (<3 ft below ground surface [bgs]) and subsurface (>3 ft bgs), including saturated soils, were included as COPCs (see Table B.1). For each COPC, construction/trench worker risk-based screening levels (RBSLs) corresponding to a target carcinogenic risk level of 1 x 10⁻⁶ and a target hazard level of 1.0 were identified from the RWQCB guidance (see Table K-3 of Appendix 1 to RWQCB 2000). Based on these RBSLs, individual and cumulative carcinogenic risks and noncarcinogenic hazards for the COPCs were back-calculated using the site maximum COPC concentrations and the following formula:

CW Risk/Hazard = Cmax * Target R/H Level RBSL

Where:

CW Risk/Hazard = Carcinogenic risk or hazard to construction worker Cmax = Site maximum soil concentration Target R/H Level = Target risk (i.e., 1 x 10⁻⁶) or target hazard (i.e., 1.0) level RBSL = Construction worker RBSL (RWQCB 2000)

The results of this calculation for each COPC are summarized in Table B.1. As indicated in the table, the cumulative pathway risk associated with exposure of construction workers to COPCs in saturated and unsaturated soils approximates 4.76×10^{-6} , which is within the target risk range of 1×10^{-4} to 1×10^{-6} adopted by the USEPA. The risk associated with exposure to benzene corresponds to more than 99 percent of the total cumulative risk to construction workers.

As indicated in Table B.1, the cumulative pathway hazard associated with exposure of construction workers to the COPCs approximates 1.005, slightly exceeding the target hazard of 1.0 adopted by the USEPA. Approximately 63 percent of this hazard corresponds to exposure to TPH-g in soils.

The above estimates of carcinogenic risks and noncarcinogenic hazards should be used in support of developing a site-specific health and safety plan for future construction workers, as suggested by the risk management plan for the "subject facility" portion of the former Nestle site.

REFERENCES

ASTM (American Society for Testing and Materials). 1995. Standard Guide for Risk-Based Corrective Action Applied at Petroleum Release Sites, E 1739-95.

ASTM (American Society for Testing and Materials). 1998. Standard Provisional Guide for Risk-Based Corrective Action, PS 104-98.

ETIC (ETIC Engineering, Inc.). 2001. Comprehensive Site Characterization Report, Support for the Site as a Low-Risk Soil and Groundwater Case, Former Nestle USA, Inc. Facility, 1310 14th Street, Oakland, California. ETIC, Pleasant Hill, California. January.

JCI (Javaherian Consulting, Inc.). 2000. Technical Memorandum: Risk-Based Corrective Action Analysis, Nestle USA, Inc. Facility, 1310 14th Street, Oakland, California. JCI, San Francisco, California. 22 August.

MADEP (Massachusetts Department of Environmental Protection). 1994. Interim Final Petroleum Report: Development of Health-Based Alternative to the Total Petroleum Hydrocarbon Parameter, August.

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USEPA (U.S. Environmental Protection Agency). 1997. Health Effects Assessment Summary Tables (HEAST).

USEPA (U.S. Environmental Protection Agency). 1999. Region 9 Preliminary Remediation Goals Chemical Toxicity Database, on-line @ www.epa.gov/region09/waste/sfund/prg/.