Report of Additional Site Assessment UST 1, 2, 3 Site Santa Rita Correction Facility Dublin, California

Prepared for:

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This report of additional investigation has been prepared by Environmental Science and Engineering, Inc. (ESE) for the exclusive use of the Alameda County General Services Agency as it pertains to their site located at the UST 1,2,3 Site, Santa Rita Correctional Facility, Dublin, California. This report was prepared with that degree of care and skill ordinarily exercised by other geologists and engineers practicing in this field. No other warranty, either expressed or implied, is made as to professional advice in this report.

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1.0 Introduction

This report presents the results of site assessment activities performed by Environmental Science & Engineering, Inc. (ESE) for the Alameda County General Services Agency (GSA) at the UST 1,2,3 Area (site) during the month of January 1996 (Figure 1 - Location Map). ESE submitted a workplan dated November 16, 1995 to the GSA and the Alameda County Health Care Services Agency (HCSA) describing the tasks to be performed during this site assessment (ESE, 1994c).

ESE's primary objective was to assess petroleum hydrocarbons in the area of the three former underground storage tanks (USTs) at the site (Figure 2 -Site Map) and to properly abandon four existing ground water monitoring wells. These objectives were accomplished with the following work scope:

- Using Geoprobe technology, probe discrete intervals and sample soils for analytical and lithologic purposes;
- Using Geoprobe technology, probe to the occurrence of ground water and obtain a grab water sample; and
- Using a hollow-stem auger drilling rig, destroy four monitoring wells by over-drilling and grouting the holes to grade.

Presented in the following sections are a site history, a summary of the regional geology and hydrology, field methodologies for soil sampling and ground water sampling, and the reported analytical results for soil and ground water samples collected during this site assessment. This report also discusses the findings obtained from this investigation, presents conclusions, and provides recommendations for future site activities.

2.0 Site History

In March, 1988, Environmental Technology (ET) directed the removal of three USTs at the subject site under permit from the HCSA and the Dougherty Regional Fire Authority. ET removed one 3,000-gallon capacity UST (UST 1) used for the storage of diesel fuel and two 5,000-gallon capacity USTs (UST 2 and UST 3) used for the storage of Bunker C fuel oil. The fuels were used to operate a series of boilers formerly located at the site. Each UST was of single-wall carbon steel construction.

During the removal of the USTs, a representative from the HCSA witnessed the collection of eight soil samples from the base of the excavation. All samples were analyzed for total petroleum hydrocarbons as diesel fuel (TPH-D) and gasoline (TPH-G) using EPA Method 8015 (modified per CA LUFT) and total oil and grease (TOG) using Standard Method for the Examination of Water and Waste Water (SMWW) Method 503E. Four samples were reported to contain detectable concentrations of TPH-D ranging from 25 to 15,500 parts per million (ppm) and two samples were reported to contain TPH-G concentrations of 50 ppm and 195 ppm, respectively. All eight soil samples were reported to contain detectable concentrations of TOG ranging from 6 to 1,097 ppm.

A preliminary site assessment was performed by Gregg & Associates during March 1988 to determine the areal extent of soil impacted with petroleum hydrocarbons. One soil sample was collected at a depth of 15 feet from each of the four borings (1C, 3D, 3E, and 3F) drilled during the preliminary site assessment and analyzed for TPH-D. No detectable concentrations of TPH-D were reported to occur in the four samples. Detectable concentrations of TOG were reported for each sample and ranged from 22 to 42 ppm. Based on these findings, Gregg & Associates supervised the overexcavation of soil impacted with petroleum hydrocarbons. All findings were documented in an Underground Tank Removal and Site Remediation Report prepared by Gregg & Associates and submitted to the HCSA during May, 1988.

On November 3, 1993, ESE measured and mapped the stockpiled soil at the subject site. ESE estimated the total volume of the stockpiled soil at the site to be approximately 400 cubic yards. On November 24, 1993, ESE submitted a workplan to the HCSA for sampling the stockpiled soil. Subsequently, ESE collected soil samples from the stockpile on November 30, 1993 at a frequency of one sample for every 50 cubic yards and analyzed each for TPH-D and benzene, toluene, ethylbenzene, and total xylenes (BTEX) using EPA Method 8015 (modified per CA LUFT) and EPA Method 8020, respectively. Of the eight soil samples analyzed, one sample (SP-3-2.5') was reported to contain TPH-D at a concentration of 130 ppm. Results of the stockpile sampling were presented to the HCSA in a letter report dated December 7, 1993.

On May 11, 1994, ESE supervised the loading, hauling, and disposal of the 50 cubic yards of stockpiled soil reported to contain detectable concentrations of TPH-D. The impacted soil was hauled to the BFI-Vasco Road landfill for disposal. The remaining 350 cubic yards of stockpiled soil were spread at the site on the ground surface.

On June 24, 1994, ESE submitted a workplan to the GSA and HCSA describing the tasks to be performed to determine if petroleum hydrocarbons occur in the soil adjacent to the former USTs 1, 2, and 3. Subsurface investigation field activities were performed at the site on September 6, 1994. A Site Assessment Report was prepared by ESE and submitted to the GSA and HCSA on December 21, 1994. A comprehensive description including site history, regional geology, and regional hydrology was presented in this report. The Site Assessment Report also included analytical results for ground water samples collected from the four ground water wells installed. The results for the samples collected indicated no detectable concentrations of TPH-D, TOG, and BTEX. The report recommended that three additional quarters of ground water monitoring be performed at the site prior to requesting site closure from the HCSA.

Subsequently, three ground water monitoring events were performed at the site during December 1994, February 1995, and May 1995. No detectable concentrations of TPH-D or BTEX were reported to occur in any of the ground water samples collected during the monitoring events.

The location of the monitoring wells is approximately 300 feet east of the former USTs. GSA, HCSA, and ESE met at the site on November 8, 1995 to discuss the need for supplemental information on subsurface conditions closer to the former UST location. Both GSA and HCSA agreed that some additional subsurface investigation was appropriate.

3.0 Regional Geology and Regional Hydrology

3.1 Regional Geology

The site is located within the Coast Ranges geomorphic province (Norris and Webb, 1976) at the northern boundary of the Livermore Valley depression, located midway between the southern part of San Francisco Bay and the San Joaquin Valley. The Livermore Valley is approximately 13 miles long in an east-west direction and approximately four miles wide and is completely surrounded by hills of the Diablo Range.

The Livermore Valley fill materials are comprised of alluvial sediments of Quaternary age (DWR, 1974). Alluvium of Pleistocene to Holocene age has been deposited in the gently sloping central area of the Livermore Valley and adjacent to active streams in the ravines and canyons tributary to Livermore Valley. The alluvium consists of unconsolidated deposits of interbedded clay, silt, fine sand, and lenses of clayey gravel. These sediments are up to 200 feet in thickness and overlie sedimentary rocks of the Livermore Formation and the Tassajara Formation. Fine-grained alluvial fan deposits occur along the northern side of the Livermore Valley. These deposits consist of stratified beds of clay, silt, and sand, and were formed by deposition from streams draining upland areas composed of sandstone and shale of the Tassajara Formation.

The Livermore Valley is bisected by six major faults or fault groups and at least five other faults of a more local nature (DWR, 1974). The major faults are the Carnegie, Tesla, Mocho, Livermore, Pleasanton, and Calaveras faults. The minor faults include the Parks, Verona, and several unnamed faults. The site is located on a down-dropped block of land bounded by the Mocho Fault to the north, the Parks Fault to the south, and the Pleasanton Fault to the east.

3.2 Regional Hydrology

The water-bearing sediment series in the Livermore Valley consist of multi-layered systems having an unconfined upper aquifer over a sequence of leaky or semi-confined aquifers (State of California Department of Water Resources, 1974). The Livermore Valley Ground Water Basin has been divided into approximately 12 hydrologic sub-basins on the basis of fault traces and hydrologic discontinuities. The northern boundary of the Amador sub-basin is a permeability barrier formed by the interfingering of alluvial deposits and the southern boundary of the sub-basin is formed partly by the contact of the water-bearing Livermore Formation with nonwater-bearing rocks and partly by the drainage divide between Livermore Valley and Sunol Valley.

Regional ground water flow maps indicate that water in the uppermost aquifer beneath the site flow toward the southwest (Alameda County Flood Control and Water Conservation District,

1990; 1991). Ground water in the sub-basin has been analyzed by the State of California Department of Water Resources (1974) and is classified as sodium carbonate water of irrigation Class II quality.

The northern portion of the site is crossed by the east-southeast flowing Arroyo Las Positas and the southern portion of the site is crossed by the east-southeast flowing Arroyo Mocho. Both streams are considered to be major drainages for the Livermore Valley and are located in modified earth channels which converge just west of the site.

4.0 Field Investigation

Prior to beginning fieldwork, ESE obtained all necessary permits for drilling soil borings and installing ground water monitoring wells at the site. In addition, ESE reviewed the site-specific Health and Safety Plan (HASP) prepared for this investigation with the onsite personnel, subcontractors, and qualified visitors. ESE performed the fieldwork in accordance with Tri-Regional Water Quality Control Board guidelines (RWQCB, 1990) and other applicable State regulations and standards.

4.1 Soil and Ground Water Collection

ESE supervised the probing and sampling of three locations (HP-1, HP-2, and HP-3) using a truck-mounted Direct Penetration Technology (DPT) Geoprobe unit. Soil probes HP-1, HP-2, and HP-3 were completed on January 9, 1996 to a depth of approximately 35 feet below ground surface (bgs). Ground water was encountered at approximately 35 feet bgs in all probings. The locations of the probes in the area of the former USTs are presented in Figure 2 -Site Map. Soil samples were collected at approximately 35 feet below ground surface (bgs) in each of the probe bores. Probing activities were performed by EnProbe of Oroville, California.

On January 23, 1996, ESE supervised EnProbe in the re-probe of the site to collect soil samples not collected in the initial soil probing operations. Soil samples were collected at 10 and 20 feet bgs in the same area as the original soil probes HP-1, HP-2, and HP-3.

4.1.1 Soil Probing and Soil Sample Collection

On January 9, 1996 ESE collected one soil sample from each probing at the vadose zone-ground water interface (sample nos. HP-1-35, HP-2-35, and HP-3-35). On January 23, 1996, ESE collected two soil samples from each soil probing at 10 and 20 feet bgs. The DPT unit utilized a 2.25-inch outside diameter, stainless steel tube that was pushed into the subsurface. A four-foot continuous soil sampling device was used to collect relatively undisturbed soil samples. The stainless steel sample tube was then extracted and the polyacetate liner containing the soil was removed. When collection of an analytical sample was necessary another 2.25-inch outside diameter, stainless steel tube was used. The sampler release spring was disengaged and the stainless steel tube was pushed an additional two feet, collecting a relatively undisturbed soil sample. The stainless steel sample tube was then extracted and the brass liner containing the soil was removed. The ends of the brass liner were capped with teflon-lined plastic caps. The soil samples collected for chemical analysis were placed in a cooler with ice, and transported under

chain-of-custody to McCampbell Analytical (McCampbell), a State-certified analytical laboratory of Pacheco, California.

A portion of each soil sample was sealed in a clean individual Ziploc™ bag and set in direct sunlight to enhance the volatilization of volatile organic compounds (VOCs) present in the soil. After approximately ten minutes, each sample was screened for VOCs using a photoionization detector (PID).

Each soil boring was logged by an ESE field geologist in accordance with the Unified Soils Classification System (USCS). Additionally, the ESE field geologist noted soil color, relative density, moisture content, biologic content, and odor, if present. Geologic boring logs are presented as Appendix A.

4.1.2 Ground Water Sample Collection

On January 9, 1996, ground water samples were collected from HP-1 through HP-3 using a two-inch diameter stainless steel slotted tube pushed into the saturated subsurface by the DPT unit. After approximately ten minutes ground water infiltrated the tube and a sample was retrieved using a 3/8-inch stainless steel bailer. The samples were collected in appropriate laboratory-supplied glassware, labeled, and placed on ice under proper chain-of-custody documentation for transport to McCampbell, a State-certified analytical laboratory.

4.2 Analytical Methodologies

4.2.1 Soil Sample Analysis

All soil samples were analyzed for TPH-D using EPA Method 8015 (modified per CA LUFT) and benzene, toluene, ethylbenzene, and total xylenes (BTEX) using EPA Method 8020. TPH-D was detected then one sample with the highest concentration would be analyzed for polynuclear aromatics (PNAs) using EPA Method 8270.

4.2.2 Ground Water Sample Analysis

All ground water samples were analyzed for TPH-D using EPA Method 8015 (modified per CA LUFT) and benzene, toluene, ethylbenzene, and total xylenes (BTEX) using EPA Method 8020. TPH-D was detected then one sample with the highest concentration would be analyzed for polynuclear aromatics (PNAs) using EPA Method 8270.

4.3 Monitoring Well Destruction

On January 15, 1996, ESE supervised the destruction of four monitoring wells by Exploration Geoservices, Inc. (EGI) of San Jose, California. EGI used a mobile B-61 drilling rig with 10-inch hollow stem augers to overdrill the four-inch polyvinyl chloride (PVC) monitoring well casings. Upon drilling out the concrete and bentonite seals, the PVC casings were removed from their respective boreholes. Then a ten-sack cement slurry was emplaced by tremie from the base of the borehole to the ground surface. The PVC casing was disposed offsite by EGI.

5.0 Results

5.1 Soil and Ground Water Samples

5.1.1 Soil

Sediments of the unsaturated zone in the three soil probings are comprised of an interbedded sequence of silty clay, sandy clay, and sand (Appendix A). Brown stiff silty clays and sandy clays are predominant in the soil probes to a depth of approximately eight feet bgs. Fine, light brown sandy silts occur between approximately eight and 16 feet bgs. Light brown clays are present from approximately 16 feet to the base of the probe at 35 feet bgs. Below a depth of approximately 31 feet bgs the silty clay and clay sediments become wet.

The results of field screening drill cuttings with a PID indicated no detectable concentrations of volatile organic vapors. Soil samples collected at 10 feet bgs, 20 feet bgs, and at 35 feet bgs (the vadose zone-ground water interface) in all soil probings (HP-1, HP-2, and HP-3) were reported not to contain any detectable concentrations of petroleum hydrocarbon constituents. Soil sample analytical results are summarized in Table 1, and the analytical reports with the chain-of-custody documentation are presented in Appendix B.

5.1.2 Ground Water

TPH-D was detected in all three of the ground water samples submitted (HP-1, HP-2, and HP-3), at concentrations of 220 μ g/L, 220 μ g/L, and 61 μ g/L, respectively. Benzene and ethylbenzene were not detected at or above the method detection limits in any of the ground water samples submitted. Toluene was detected in all three of the ground water samples submitted (HP-1, HP-2, and HP-3), at concentrations of 3.3 μ g/L, 1.2 μ g/L, and 1.6 μ g/L, respectively. Total xylenes were detected in all three of the ground water samples submitted (HP-1, HP-2, and HP-3), at concentrations of 1.4 μ g/L, 0.58 μ g/L, and 1.1 μ g/L, respectively. Ground water sample HP-1 was analyzed for PNAs and was not found at a concentration at or above the method detection limit. The analytical results are summarized in Table 2 and the analytical report with the chain-of-custody documentation is presented in Appendix B.

6.0 Conclusions and Recommendations

6.1 Conclusions

Field screening results using a PID indicated no detectable concentrations of volatile organic compounds in any of the probe soil samples from HP-1, HP-2, and HP-3. In addition, the analytical results for the four soil samples collected at the vadose zone-saturated zone interface indicated no detectable concentrations of TPH-D or BTEX. Analytical results for ground water samples collected from the probes indicated no detectable concentrations of benzene, ethylbenzene, and PNAs, and only low concentrations of TPH-D, toluene, and total xylenes. The concentrations of toluene and total xylenes detected were well below the federal maximum contaminant levels (MCLs) of 150 μ g/L and 1,750 μ g/L, respectively. ESE concludes that based on the data collected, the shallow nature of the water-bearing zone and no known municipal production wells in the area of the site, the UST 1,2,3 site should be considered to be low risk.

6.2 Recommendations

Based on the results and conclusions of this site investigation at the UST 1,2,3 site, ESE recommends that the GSA should submit this report to the HCSA and request case closure.

7.0 References

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 Regional Ground Water Flow Maps of the Livermore Valley.
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- Environmental Science & Engineering, Inc. (ESE), 1994a. Workplan for Site Investigation, Alameda County General Services Agency (GSA), UST 1, 2, 3 Site; June 24, 1994.
- Environmental Science & Engineering, Inc. (ESE), 1994b. Report of Stockpiled Soil Spreading and Disposal, Alameda County General Services Agency (GSA), UST 1, 2, 3 Site; June 20, 1994.
- Gregg & Associates, Inc., 1988. Underground Tank Removal and Site Remediation Report; May, 1988.
- Norris, R.M., and Webb, R.W., 1976. Geology of California; John Wiley & Sons, Inc., New York. 365pp.
- State of California Department of Water Resources (DWR), 1974. Evaluation of Ground Water Resources: Livermore and Sunol Valleys; Bull. 118-2, pp.153.
- State of California Regional Water Quality Control Board (RWQCB), 1990. Tri-Regional Board Staff Recommendations for Preliminary Evaluation and Investigation of Underground Tank Sites; August 10, 1990.

TABLES

TABLE 1
SUMMARY OF SOIL SAMPLE ANALYTICAL RESULTS

Alameda County General Services Agency UST 1,2,3 Site, Santa Rita Correctional Facility Dublin, California

Sample	Sample	Depth	TPH-D	Benzene	Toluene	Ethylbenzene	Total Xylenes
HP-1-10	1/23/96	10	ND (10)	ND (0.005)	ND (0.005)	ND (0.005)	ND(0.005)
HP-1-20	1/23/96	20	ND (10)	ND (0.005)	ND (0.005)	ND (0.005)	ND (0.005)
HP-1-35'	1/9/96	35	ND (10)	ND (0.005)	ND (0.005)	ND (0.005)	ND (0.005)
HP-2-10	1/23/96	10	ND (10)	ND (0.005)	ND (0.005)	ND (0.005)	ND (0.005)
HP-2-20	1/23/96	20	ND (10)	ND (0.005)	ND (0.005)	ND (0.005)	ND (0.005)
HP-2-35'	1/9/96	35	ND (10)	ND (0.005)	ND (0.005)	ND (0.005)	ND (0.005)
HP-3-10	1/23/96	10	ND (10)	ND (0.005)	ND (0.005)	ND (0.005)	ND (0.005)
HP-3-20	1/23/96	20	ND (10)	ND (0.005)	ND (0.005)	ND (0.005)	ND (0.005)
HP-3-35'	1/9/96	35	ND (10)	ND (0.005)	ND (0.005)	ND (0.005)	ND (0.005)

Notes:

mg/Kg = milligrams per Kilogram;

TPH-D = Total Petroleum Hydrocarbons as Diesel, analyzed using EPA Method 8015 (modified per CA LUFT);

BTEX = Benzene, Toluene, Ethylbenzene, and Total Xylenes analyzed using EPA Method 8020;

ND (x.xx) = indicates not detected at method detection limit given as x;

Analytical Reports are presented in Appendix B of this report.

TABLE 2

SUMMARY OF GROUND WATER SAMPLE ANALYTICAL RESULTS

Alameda County General Services Agency UST 1,2,3 Site, Santa Rita Correctional Facility Dublin, California

Sample No.	Sample Dare	TPH-D (ag/L)	Benzene (µg/1.)	Toluëne (µg/L)	Ethylbenzene (µg/L)	Total Xylenes (μg/L)	PNA's (μg/L)
HP-1	1/9/96	220	ND (0.5)	3.3	ND (0.5)	1.4	ND
HP-2	1/9/96	220	ND (0.5)	1.2	ND (0.5)	0.58	
HP-3	1/9/96	61	ND (0.5)	1.6	ND (0.5)	1.1	
Trip	1/9/96		ND (0.5)	ND (0.5)	ND (0.5)	ND (0.5)	

Notes:

mg/L = milligrams per Liter;

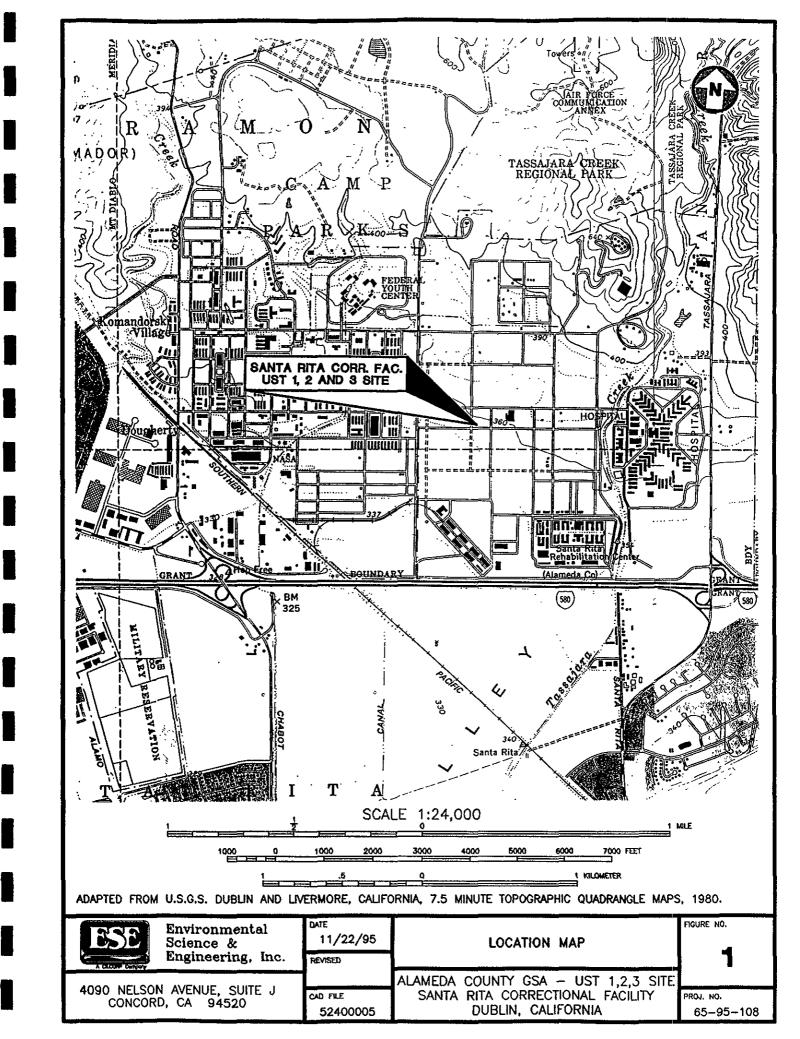
= Total Petroleum Hydrocarbons as Diesel, analyzed using EPA Method 8015 (modified per CA LUFT); TPH-D

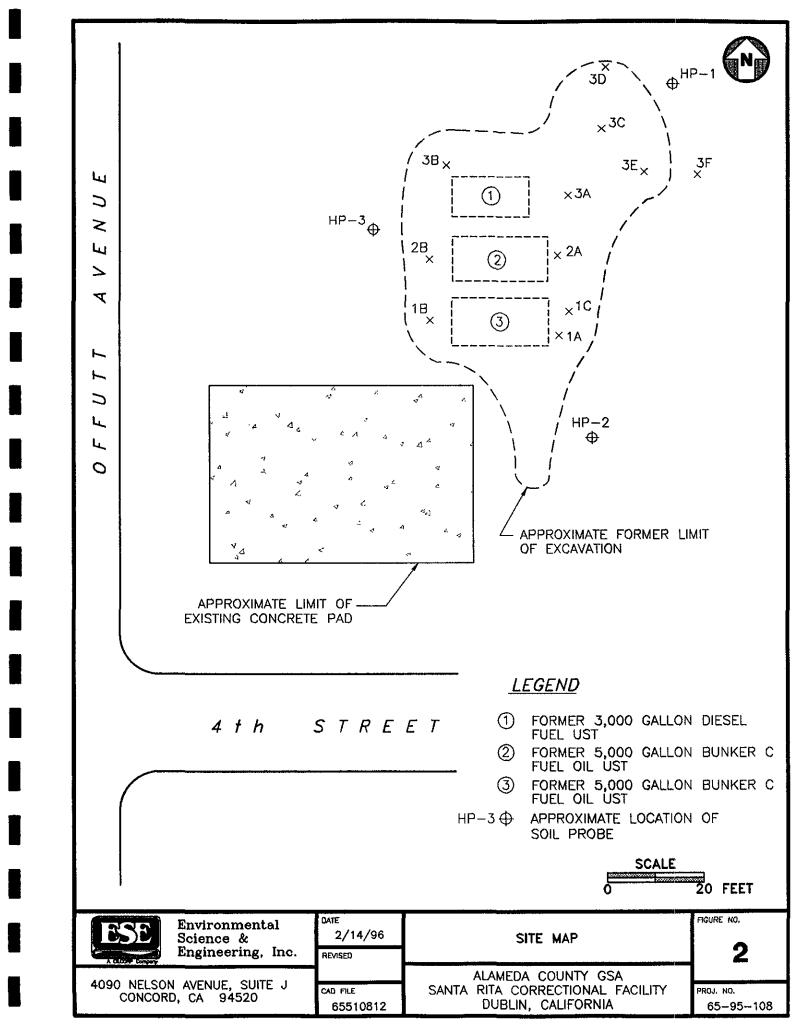
= Benzene, Toluene, Ethylbenzene, and Total Xylenes analyzed using EPA Method 8020; BTEX

ND (x.xx) = indicates not detected at method detection limit given as x;

Analytical Reports are presented in Appendix B of this report.

FIGURES





APPENDIX A SOIL BORING LOGS

	Environmental Science & Engineering, Inc.					L COMPL	ETI	OG AND ON SUMMARY	HP - 1
Со	ELL COMPLETION mptetion Depth: No Well installed Size/Type From sing:	То			ocation: U	16: Alameda Count ST 1, 2 & 3 anta Rita Correction ublin, California		Project No: 65-95-108	Page 1 of 1
Sci Filt Sei	reen: er:	37.01	Feet	H H	Method: Ge Iole Diame tef. Elevati	ter: 25 inches		Depth: 370 Feet	Dates: Start: 1-9-96 Finish; 1-9-96
Depth (ft)	Lithologic Description	oso	Sampl		aphic Log		Vapor	Remarks Water, drilling/completion, summ	
ది		<u> </u>	Blows	Liti	hology	Well Installation	·/>	water, or ming-completion, somm	кагу, заттріе туре
°	FILL MATERIAL	-						-	
_	-	-						-	
5 <u>-</u>		-			*****				
)	SILTY CLAY, dark brown, atiff, damp, 10-20% silts, no odor.	CL					2.1	-	
=		-						-	
10	<u>-</u>	_					2.8		
-	SANDY CLAY, light brown, stiff, damp, 10-20% fine grained sand,	- ■ CL)RE					_	
-	no odor.	-	OUS CC					-	
16-	-	-	CONTINUOUS CORE				42	- -	
-	• 	-	٥	Ì				-	
=		-							
20	_	<u>-</u>					2.6	_	
-	SAND, light brown, dense, damp, fine to medium grained, no odor.	SW						_	
=	- -							_ _	
25 —	j	-					2.4	<u>-</u>	
-		-						-	
	-	_						<u> </u>	
30	SILTY CLAY, light brown, stiff, damp, 10-20% sifts, no odor	CL					28	-	
<u>-</u>		-						- -	
35 -									

I

	Environmental Science & Engineering, Inc.				1	L COMPL	ET	OG AND ION SUMMARY	HP-2
Co	LL COMPLETION npletion Depth: No Well Installed Size/Type From	То			Location: U S	ne: Alameda Count ST 1, 2 & 3 anta Rita Correction ublin, California		Project No: 65-95-108	Page 1 of 1
Scr Filt Sec		35.0	Feet		Method: Ge	ter: 2.5 Inches ons:		Depth: 35.0 Feet	Dates: Start: 1-9-96 Finish: 1-9-96
Depth (ft)	Lithologic Description	nsc	Sam, Blov		Graphic Log	Well Installation	Vapor	Remarks Water, drilling/completion, sum	mary, sample type
	FILL MATERIAL SANDY CLAY, brown, stiff, dry, 20-30% fine grained sand, no odor.	CL					1.0		
10-	SANDY CLAY, brown, stiff, damp, 10-20% line grained sand, no odor	CL	ados sironina	CONTINUOUS CORE			22	- - - -	
15	CLAYEY SAND, light brown, dense, damp, 20-30% clay, fine grained, no odor.	SM	LINCO				2.9	- - - -	
25 —	CLAY, light brown, very stiff, damp, no odor.	CL					2.5	-	
25	SANDY CLAY, light brown, stiff, damp, 10-20% very fine sand, no odor.	CL						- -	
30	SILTY CLAY, light brown, stiff, damp, 10-20% silts, no odor	CL					3.5	- - - -	
35 =									

	Environmental Science & Engineering, Inc.			 WEL			OG AND ION SUMMARY	HP-3
Co	ELL COMPLETION mpletion Depth: No Well Installed SIze/Type From	То		 Location: L	me: Alameda Count IST 1, 2 & 3 anta Rita Correction Jublin, California		Project No: 65-95-108	Page 1 of 1
Sc Fill Se	sing: reent: er: er: al: Cement Slumy Grout 0.0 Feet	35 0	Feet	Method: G Hole Diame Ref. Elevat	eter: 2.5 inches		Depth: 35.0 Feet	Dates; Start: 1-9-96 Finish: 1-9-96
Depth (ft)	Lithologic Description	nsc	Samp	Graphic Log	Well Installation	Vapor	Remarks Water, drilling/completion, sum	mary, sample type
5	FILL MATERIAL SILTY CLAY, dark brown, stiff, damp, 10-20% sifts, no odor. SANDY CLAY, brown, stiff, dry, 20-30% fine grained sand, no odor. SANDY SILT, light brown, stiff, dry, 20-30% very fine sand, no odor.	CL.	Blow CONTINION SCORE			1.2		
20	SANDY CLAY, light brown, very stiff, damp, 20-30% medium sand, no odor. CLAY, light brown, stiff, damp, no odor.	Cl.				2.9		
_								

APPENDIX B ANALYTICAL REPORTS WITH CHAIN-OF -CUSTODY DOCUMENTS

01/19/96

Dear Eric:

Enclosed are:

- 1), the results of 7 samples from your Alameda Co. GSA-UST 1,2,3 Site project,
- 2). a QC report for the above samples
- 3). a copy of the chain of custody, and
- 4). a bill for analytical services.

If you have any questions please contact me. McCampbell Analytical Laboratories strives for excellence in quality, service and cost. Thank you for your business and I look forward to working with you again.

Yours truly,

Edward Hamilton

Environment	al Science & Eng.		•	meda Co.	GSA-UST	Date Sampled: 01/09/96					
4090 Nelson	Ave., Suite J	1,2,3 Site				Date Rece	ived: 01/10/	/96			
Concord, CA	94520	Client Co	ontact: Eric C	Farcia		Date Extracted: 01/10/96					
		Client P.	O:# SMSA-0	C-021	Date Analy	zed: 01/10/	/96				
EPA methods 50	•	(C6-C12) Volatile Hydrocarbons as Gasoline*, with BTEX* 020 or 602; California RWQCB (SF Bay Region) method GCFID (5030)									
Lab ID	Client ID	Matrix	TPH(g) ⁺	Benzene	Toluene	Ethylben- zene	Xylenes	% Rec. Surrogate			
60360	HP1-35'	S	204	ND	ND	ND	ND	108			
60361	HP2-35'	s		ND	ND	ND	ND	105			
60362	HP3-35'	S	***	ND	ND	ND	ND	106			
60363	HP1	w		ND	3.3	ND	1.4	103			
60364	HP2	w		ND	1.2	ND	0.58	108			
60365	HP3	w		ND	1.6	ND	1.1	99			
60366	Trip Blank	w		ND	ND	ND	ND	001			
Reporting	Limit unless other- ND means not de-	W	50 ug/L	0.5	0.5	0.5	0.5				
tected above	the reporting limit	S	1.0 mg/kg	0.005	0.005	0.005	0.005				

^{*} water and vapor samples are reported in ug/L, soil samples in mg/kg, and all TCLP extracts in mg/L

[#] cluttered chromatogram; sample peak coelutes with surrogate peak

⁺ The following descriptions of the TPH chromatogram are cursory in nature and McCampbell Analytical is not responsible for their interpretation: a) unmodified or weakly modified gasoline is significant; b) heavier gasoline range compounds are significant (aged gasoline?); c) lighter gasoline range compounds (the most mobile fraction) are significant; d) gasoline range compounds having broad chromatographic peaks are significant; biologically altered gasoline?; e) TPH pattern that does not appear to be derived from gasoline (?); f) one to a few isolated peaks present; g) strongly aged gasoline or diesel range compounds are significant; h) lighter than water immiscible sheen is present; i) liquid sample that contains greater than ~ 5 vol. % sediment; j) no recognizable pattern.

Environment	al Science & Eng.		oject ID: Alameda Co. GSA-UST	Date Sampled: 0	1/09/96
4090 Nelson	Ave., Suite J	1,2,3 Site		Date Received: 0	1/10/96
Concord, CA	94520	Client Co	ntact: Eric Garcia	Date Extracted: (01/12/96
		Client P.C	D: # SMSA-C-021	Date Analyzed: 0)1/12/96
EPA methods m	Diesel Ra nodified 8015, and 3550 or	nge (C10- 3510; Califo	C23) Extractable Hydrocarbons as rnia RWQCB (SF Bay Region) method Go	Diesel * CFID(3550) or GCFII	O(3510)
Lab ID	Client ID	Matrix	TPH(d) ⁺		% Recovery Surrogate
60360	HP1-35'	s	ND		101
60361	HP2-35'	S	ND		100
60362	HP3-35'	S	ND		104
60363	HP1	W	220,b,i		106
60364	HP2	W	220,b,i		106
60365	НР3	W	61,b,i		105
Reporting	Limit unless other; ND means not de-	W	50 ug/L		
tected above	e the reporting limit	s	10 mg/kg		

^{*} water samples are reported in ug/L, soil samples in mg/kg, and all TCLP and STLC extracts in mg/L

[&]quot; cluttered chromatogram resulting in coeluted surrogate and sample peaks, or; surrogate peak is on elevated baseline, or; surrogate has been diminished by dilution of original extract.

⁺ The following descriptions of the TPH chromatogram are cursory in nature and McCampbell Analytical is not responsible for their interpretation: a) unmodified or weakly modified diesel is significant; b) diesel range compounds are significant; no recognizable pattern; c) aged diesel? is significant); d) gasoline range compounds are significant; e) medium boiling point pattern that does not match diesel (?); f) one to a few isolated peaks present; g) oil range compounds are significant; h) lighter than water immiscible sheen is present; i) liquid sample that contains greater than ~ 5 vol. % sediment.

Date: 01/10/96

Matrix: Soil

	Concent	ration	(mg/kg)		% Reco	very	
Analyte	Sample (#56725)	MS	MSD	Amount Spiked	MS	MSD	RPD
TPH (gas) Benzene	0.000	1.778	1.872	2.03	88	92	5.2
Toluene	0.000	0.200	0.208 0.208	0.2	100	104	3.9
Ethylbenzene	0.000	0.202	0.208	0.2	101 101	104 104	2.9 2.9
Xylenes	0.000	0.602	0.618	0.6	100	103	2.6
TPH (diesel)	0	319	316	300	106	105	0.9
TRPH (oil & grease)	 N/A	N/A	N/A	N/A	N/A	N/A	N/A

% Rec. = (MS - Sample) / amount spiked x 100

Date: 01/10/96

Matrix: Water

	Concent	ration	(ug/L)	1	* Reco	very	
Analyte	Sample (#60334)	MS	MSD	Amount Spiked	 MS	MSD	RPD
TPH (gas)	0.0	95.0	94.6	100	95	95	0.4
Benzene	ļ 0	10.10	10.10	1.0	101.0	101.0	0.0
Toluene	0	10.60	10.70	10	106.0	107.0	0.9
Ethyl Benzene) 0	10.70	10.70	10	107.0	107.0	0.0
Xylenes	0.	33.20	33.20	30	110.7	110.7	0.0
TPH (diesel)	N/A	N/A	N/A	N/A	N/A	N/A	N/A
TRPH (oil & grease)	N/A	N/A	N/A	N/A	N/A	N/A	N/A

* Rec. = (MS - Sample) / amount spiked x 100

Date: 01/11/96-01/12/96 Matrix: Soil

	Concent	ration	(mg/kg)	1	% Reco	very	
Analyte	Sample (#56725)	MS	MSD	Amount Spiked	MS	MSD	RPD
TPH (gas)	0.000	1.906	1.770	2.03	94	87	7.4
Benzene Toluene	0.000	0.186	0.186	0.2	93	93	0.0
	0.000	0.192	0.190	0.2	96	95	1.0
Ethylbenzene	0.000	0.192	0.190	0.2	96	95	1.0
Xylenes	0.000	0.562	0.562	0.6	94	94	0.0
TPH (diesel)	0	307	310	300	102	103	1.1
TRPH (oil & grease)	0.0	19.9	22.3	20.8	96	107	11.4

% Rec. = (MS - Sample) / amount spiked \times 100

Date: 01/12/96

Matrix: Water

Concent	ration	(ug/L)]	% Reco	very	
Sample (#60334)	MS	MSD	Amount Spiked	MS	MSD	RPD
0.0	90.3	89.8	100	90	90	0.6
!			10	101.0	103.0	2.0
1				104.0	106.0	1.9
!			10	106.0	108.0	1.9
0	32.50	33.10	30	108.3	110.3	1.8
0	147	147	150	98	98	0.0
 	24900	25100	23700	105	106	0.8
	Sample (#60334) 0.0 0 0 0 0 0 0 0 0	Sample (#60334) MS 0.0 90.3 0 10.10 0 10.60 0 32.50 0 147	(#60334) MS MSD 0.0 90.3 89.8 0 10.10 10.30 0 10.40 10.60 0 10.60 10.80 0 32.50 33.10	Sample	Sample (#60334) MS	Sample (#60334) MS

% Rec. = (MS - Sample) / amount spiked x 100

CHROMALAB, INC.

Environmental Services (SDB)

January 19, 1996

Submission #: 9601132

MCCAMPBELL ANALYTICAL, INC.

Atten: Ed Hamilton

Project: E/A.C.

Project#: 5608

Received: January 16, 1996

re: One sample for Polynuclear Aromatic Hydrocarbons (PAH's) analysis.

Method: EPA 3510/625

SampleID: HP-1

Sample #: 116386 Matrix: WATER Extracted: January 16, 1996

Analyzed: January 18, 1996 Run: 10135-M Sampled: January 9, 1996

	RESULT	REPORTING LIMIT	BLANK RESULT	BLANK SPIKE RESULT
Analyte	(ug/L)	(ug/L)	(ug/L)	(%)
2-METHYLNAPHTHALENE	N.D.	2.0 2.0	N.D.	
NAPHTHALENE	N.D.	2.0	Ŋ.D.	
2-CHLORONAPHTHALENE	N.D.	2.0	Ŋ.D.	
ACENAPHTHYLENE	N.D.	2.0	Ŋ.D.	
ACENAPHTHENE	Ŋ.D.	2.0	Ŋ.D.	49
FLUORENE	N.D.	5.0 2.0 2.0	Ŋ.D.	
PHENANTHRENE	N.D.	2.0	N.D.	
ANTHRACENE	Ŋ.D.	2.0	N.D.	~ -
FLUORANTHENE	Ŋ.D.	2.0	Ŋ.D.	==
PYRENE	N.D.	2.0	N.D.	51
BENZO (A) ANTHRACENE	Ŋ.D.	2.0	N.D.	·
CHRYSENE	Ŋ.D.	2.0 2.0 2.0	N.D.	
BENZO (B) FLUORANTHENE	Ŋ.D.	2.0	N.D.	
BENZO (K) FLUORANTHENE	N.D.	2.0	Ŋ.D.	
BENZO (A) PYRENE	N.D.	2.0	N.D.	
INDENO (1, 2, 3-CD) PYRENE	N.D.	2.0	N.D.	
DIBENZO (A. H) ANTHRACENE	N.D.	2.0	N.D.	
BENZO (GHI) PERYLENE	N.D.	2.0	N.D.	

Michael Verona Chemist

Semivolatiles Supervisor

38 110 388 McCAMPBELL ANALYTICAL CHAIN OF CUSTODY RECORD 110 2nd AVENUE, # D7 RUSH 24 HOUR 48 HOUR TURN ARGUND TIME: (510) 798-1820 PACHECO, CA 94553 FAX (510) 798-1822 REPORT TO Ed HAMILTON BILL TO MAT ANALYSIS REQUEST COMPANY: Mc CAMpbell ANALYtiCAL Total Petroleum DI & Grease (5520 EEF/5520 B&P) SUBK #: 9601132 REP: 60 CLIENT: MCCAM Total Petroleun Hydrocarbons (418.1) DUE: 01/23/96 TELE ABOUE FAX In REF #:25986 PROJECT NAME: Z/A.C. PROJECT NUMBER 5608 EPA - Priority Pollutant Metals PROJECT LOCATION PLEASANCON LEAD (7240/7421/239,2/6010) SAMPLER SIGNATURE EPA 608/8080 - PCBs Unly COMMENTS BTEX & TPH as Gasoline EPA 624/8240/8260 TYPE CONTAINERS HETHOD SAMPLING MATRIX # CONTAINERS PRESERVED CAM - 17 Hetals DRGANIC LEAD RC1 EPA 602/8020 EPA 608/8080 EPA 625/8270 SAMPLE DIHERMICH EPA 601/8010 LOCATION ID SL.UDGE OTHER VATER DATE TIME SOft 축 HP-1 1-9-95 60363 40-2 60364 HP 3 60365 DATE RECEIVED BY TIME REMARKS: LAST DAY OF Hold PIME #1630 TIME RECEIVED BY 1-16-96 RECEIVED BY LABORATORY Mimie Pak

DATE 12/06/45 PAGE	; / OF /				CH	AIN	OF	CU	STO	DY R	EC	ORD				5608AESEXNA	
PROJECT NAME ALAMONA CO. G		Π	ANA:	LYS	ES '	го	BE	PER	FOR	MED		мат	RIX			14 14	ronmental nce &
ADDRESS_ <u>UST 1.2.3 51</u>	TE							[N C			neering, Inc.
Winth RITA CORRECT	TIONAL FACILITY	6252	18		26					.		M A		NON TAR		A CR.CORP Company	Phone (510) 685-4053
PROJECT NO. 65-95-108		Salveled	9C15m	8020	1-4	<u> </u>						A T R I		EA	Sui	(e)	
SAMPLED BY COG	1 1	1	_		3							χ		N.			Fax (510) 685-5323
LAB NAME McCampbell Analy		100	TPA-0	页	PWA		}							O E F S	(REMARKS	S ZE. ETC.)
SAMPLE # DATE TIME/ HPI-35 1.9.94 12-20	LOCATION	+	1	8	9	-	<u> </u>	 -				MAT			<u> </u>		60360
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2.		4.	Plu C	J						1-10-	96	3pm		EPOI	RT TO:	SPECIAL SHIPM REQUIREMENTS	MENT
3. ICET° 🚽	PRESERVATIVA	WAS	103 6	MEI	LSO	HER		· · · ·			+	 -	4	ħ		REQUIREMENTS	
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INSTRUCTIONS TO LABORA	TORY (hand	llir	ng,	ana	lys	es,	, st	ora	ıge,	et	c.)	:	,			CHAIN OF CUST	ODY SEALS
INVOICE TO ALAMEDA CO. GSA -	KOV MEITAG	٠.														REC'D GOOD CO	NOTN/COLD
NORMAL T.A.T.						<u>.</u>										CONFORMS TO F	ECORD (6)

02/02/96

Dear Eric:

Enclosed are:

- 1). the results of 6 samples from your # 6595108; Alameda Co. GSA, UST 1,2,3 project,
- 2). a QC report for the above samples
- 3). a copy of the chain of custody, and
- 4), a bill for analytical services.

If you have any questions please contact me. McCampbell Analytical Laboratories strives for excellence in quality, service and cost. Thank you for your business and I look forward to working with you again.

Yours truly,

Edward Hamilton

Ca thet

Environmental Science & Eng.		Client Project ID: # 6595108; Alameda Co. Date Sampled: 01/23/96 GSA, UST 1,2,3									
4090 Nelson	Ave., Suite J	GSA, US	11,2,3			Date Received: 01/23/96					
Concord, CA	. 94520	Client Co	ontact: Eric C	- Farcia	Date Extracted: 01/23/96						
		Client P.	O:# SMSA-0	C-021		Date Analy	zed: 01/23	/96			
EPA methods 5	Gasoline Range 030, modified 8015, and 8										
Lab ID	Client ID	Matrix	TPH(g) ⁺	Benzene	zene Toluene Ethylben- Vylanes		% Rec. Surrogate				
60663	HP-1-10	S		ND	ND	ND	ND	104			
60664	HP-1-20	S		ND	ND	ND	ND	105			
60665	HP-2-10	S	•••	ND	ND	ND	ND	104			
60666	HP-2-20	s		ND	ND	ND	ND	110			
60667	HP-3-10	s		ND	ND	ND	ND	104			
60668	HP-3-20	S		ND	ND	ND	ND	104			
								<u> </u>			
											
Reporting	Limit unless other-	W	50 ug/L	0.5	0.5	0.5	0.5	<u> </u>			
wise stated;	ND means not de- e the reporting limit	S	1.0 mg/kg	0.005	0.005	0.005	0.005	1			

^{*} water and vapor samples are reported in ug/L, soil samples in mg/kg, and all TCLP extracts in mg/L

[#] cluttered chromatogram; sample peak coelutes with surrogate peak

⁺ The following descriptions of the TPH chromatogram are cursory in nature and McCampbell Analytical is not responsible for their interpretation: a) unmodified or weakly modified gasoline is significant; b) heavier gasoline range compounds are significant(aged gasoline?); c) lighter gasoline range compounds (the most mobile fraction) are significant; d) gasoline range compounds having broad chromatographic peaks are significant; biologically altered gasoline?; e) TPH pattern that does not appear to be derived from gasoline (?); f) one to a few isolated peaks present; g) strongly aged gasoline or diesel range compounds are significant; h) lighter than water immiscible sheen is present; i) liquid sample that contains greater than ~ 5 vol. % sediment; j) no recognizable pattern.

Environmental Science & Eng.			ject ID: # 6595108; Alameda Co.	Date Sampled: 01/23/96			
4090 Nelson Ave	., Suite J	GSA, US	1 1,2,3	Date Received: 01/23/96 Date Extracted: 01/23/96 Date Analyzed: 01/23/96			
Concord, CA 945	520	Client Co	ntact: Eric Garcia				
		Client P.C):# SMSA-C-021				
EPA methods modifi			C23) Extractable Hydrocarbons as mia RWQCB (SF Bay Region) method Go		(3510)		
Lab ID	Client ID	Matrix	TPH(d) ⁺		% Recovery Surrogate		
60663	HP-1-10	S	ND		105		
60664	HP-1-20	S	ND		104		
60665	HP-2-10	s	ND		104		
60666	HP-2-20	S	ND		105		
60667	HP-3-10	s	ND		102		
60668	HP-3-20	S	ND		102		

Reporting Lim wise stated; ND tected above the	means not de-	W	50 ug/L				
tocieu above tile	roporting mint	S	10 mg/kg				

^{*} water samples are reported in ug/L, soil samples in mg/kg, and all TCLP and STLC extracts in mg/L

[#] cluttered chromatogram resulting in coeluted surrogate and sample peaks, or; surrogate peak is on elevated baseline, or; surrogate has been diminished by dilution of original extract.

⁺ The following descriptions of the TPH chromatogram are cursory in nature and McCampbell Analytical is not responsible for their interpretation: a) unmodified or weakly modified diesel is significant; b) diesel range compounds are significant; no recognizable pattern; c) aged diesel? is significant); d) gasoline range compounds are significant; e) medium boiling point pattern that does not match diesel (?); f) one to a few isolated peaks present; g) oil range compounds are significant; h) lighter than water immiscible sheen is present; i) liquid sample that contains greater than ~ 5 vol. % sediment.

Date: 01/23/96

Matrix: Soil

31	Concent	ration	(mg/kg)		% Reco		
Analyte	Sample (#56725)	MS	MSD	Amount Spiked	MS	MSD	RPD
TPH (gas) Benzene Toluene Ethylbenzene Xylenes	0.000 0.000 0.000 0.000	2.213 0.212 0.220 0.222 0.676	2.156 0.234 0.240 0.236 0.718	2.03 0.2 0.2 0.2 0.2	109 106 110 111	106 117 120 118 120	2.6 9.9 8.7 6.1 6.0
TPH (diesel)	0	327	316	300	109	105	3.4
TRPH (oil & grease)	N/A	N/A	N/A	N/A	N/A	N/A	N/A

* Rec. - (MS - Sample) / amount spiked x 100