

TEXACO REFINING AND MARKETING INC. 100 CUTTING BOULEVARD RICHMOND CA 94804

August 10, 1989

Mr. Rafat Shahid Alameda County Environmental Health Department Hazardous Materials Division 80 Swan Way, Room 200 Oakland, CA 94621

Dear Mr. Shahid:

Enclosed is a copy of our environmental assessment report dated June 22, 1989 for the former Texaco service station located at 2225 Telegraph Avenue, Oakland, California.

We know of no particular hydrocarbon leaks or spills at the site. No petroleum hydrocarbons were found in vadose zone soils at or near the site. Detectable quantities of benzene, toluene, ethylbenzene, and xylenes (BTEX), however, were found in water from all on-site wells. There is BTEX in water from one of three nearby off-site monitoring wells. The complete lateral extent of BTEX in ground water is not presently known because of restricted access into the bordering city streets.

Please call me at (415) 236-1770 if you have any questions.

Very truly yours,

R.R. ZIELINSKI Field Environmental

Supervisor

RRZ:kn

Enclosure

cc: Ms. Leslie Ferguson San Francisco Bay Regional Water Quality Control Board 1111 Jackson Street, Room 6040 Oakland, CA 94607

RR

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ENVIRONMENTAL ASSESSMENT FORMER TEXACO STATION NO. 62488000195 2225 TELEGRAPH AVENUE OAKLAND, CALIFORNIA April 6, 1989

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June 22, 1989 Env. Assesment 2 copies: Texaco Refining and Marketing, Inc. 1-2 100 Cutting Boulevard Richmond, California 94804 Attention: Mr. R.R. Zelinski 1 copy: Corporate Record Copy 3 Job File 1 copy: 4

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QUALITY CONTROL REVIEWER

Randolph Stone

Associate Hydrogeologist



A Report Prepared for

Texaco Refining and Marketing, Inc. 100 Cutting Boulevard Richmond, California 94804

ENVIRONMENTAL ASSESSMENT FORMER TEXACO STATION NO. 62488000195 2225 TELEGRAPH AVENUE OAKLAND, CALIFORNIA

HLA Job No. 02251,080.03

by

Gregory Fasiano Project Geologist

Steppen J. Osborne Civil Engineer 29555 PER SALES OF CALLED

Harding Lawson Associates 1355 Willow Way, Suite 109 Concord, California 94520 415/687-9660

June 22, 1989

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DISTRIBUTION

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I INTRODUCTION

On behalf of Texaco Refining and Marketing, Inc. Harding Lawson Associates (HLA) has evaluated the extent of a suspected impact to subsurface soils and shallow ground water at the former Texaco service station (No. 62488000195) located at 2225 Telegraph Avenue in Oakland, California (Plate 1). The investigation findings and description of the environment assessment program are detailed in this report.

II PREVIOUS INVESTIGATION

In May 1988, Texaco Refining and Marketing Inc. retained HLA to conduct a Sensitive Receptor Survey (SRS) for the Texaco station at 2225 Telegraph Avenue in Oakland, California. The purpose of the survey was to acquire and provide the following site-specific information:

- Proximity of wells for:
 - public water supply
 - private water supply
 - observation or monitoring
- Proximity of subways, basements and schools
- Proximity of surface-water bodies
- Types of local water supply
- Local aquifer classification
- Site and area maps

The information collected during the SRS is presented on a fact sheet included in Appendix A.

In June 1988, following the SRS, Texaco Refining and Marketing Inc. requested that HLA proceed with a limited subsurface investigation to evaluate whether petroleum hydrocarbons had affected the shallow soil or ground water. The subsurface investigation included the following tasks:

- 1. Drill, develop, and sample four 2-inch-diameter, shallow ground-water monitoring wells (MW-6A, MW-6B, MW-6C, and MW-6D)
- 2. Survey wells and gauge water levels
- 3. Determine the direction of ground-water flow

4. Analyze one ground-water sample from each monitoring well for benzene, toluene, ethylbenzene, and xylenes (BTEX)

The subsurface investigation results (documented in a report issued to Texaco on July 20, 1988) suggested that the ground water contained minor concentrations of petroleum hydrocarbon in two wells, MW-6C and MW-6D. Water from MW-6C and MW-6D contained benzene concentrations of 7400 and 220 parts per billion (ppb) respectively. Water from MW-6C also contained 2300 ppb total xylenes. Water from MW-6A and MW-6B contained no detectable concentrations of BTEX. The soil boring logs, well completion details and a summary of the chemical test results are included in Appendix A.

III LOCATION AND TOPOGRAPHY

The former Texaco (now Exxon) service station is located approximately 3.0 miles east of San Francisco Bay near the main business district of Oakland, California (Plate 1). The surrounding area consists of commercial/retail businesses, including a Chevron service station immediately across Telegraph Avenue, and a Beacon service station northeast of the site (Plate 2). Adjacent south of the site is the 1st Baptist Church of Oakland. There is an apartment building immediately west of the site which is currently occupied. The aerial photograph, presented in Appendix B, shows the immediate site vicinity.

Surface elevation at the site is approximately 20 feet above mean sea level. The land surface slopes gently to the southeast towards Lake Merritt, and the Oakland/Alameda Inner Harbor, to an area of old tidal flats that have been recently filled. This area has been extensively developed, and surface water runoff is mainly controlled by the municipal storm sewer system.

A. Regional Geology

The site lies within the East Bay Plain which is an alluvial plain that covers approximately 114 square miles in Western Alameda County. Most of the sediments that make up the unconsolidated deposits beneath the East Bay Plain were derived from the Diablo Range. Local shallow subsurface geology consists of the Alameda formation overlain by the Temescal Formation. Both are alluvial deposits laid down in early to middle Pleistocene time. Surface geology in the site vicinity is mapped as being within the Temescal Formation. The Temescal Formation is an alluvial fan deposit comprising interfingering lenses of clayey gravel, sandy silty clay, and sand-clay-silt mixtures.

Approximately 1000 feet west of the site is an outcrop of the Merritt Sand. The Merritt Sand is a beach or near-shore Pleistocene deposit of slightly clayey, silty sand.

IV HYDROGEOLOGY

The East Bay Plain has been divided into seven ground water subareas on the basis of areal differences (ie, faults and geologic conditions). The site lies within the Oakland upland and Alluvial Plain subarea as defined by the California Department of Water Resources (DWR). Most ground water in the East Bay Plain is used for irrigation or industrial purposes. The majority of domestic water is supplied by the East Bay Municipal Utility District (EBMUD). Little ground water is pumped for domestic purposes. The Alameda and Temescal Formations along with the Merritt Sand, with an aggregate thickness of more than 1100 feet, constitute the ground water reservoir. The regional ground-water flow direction is to the west-southwest towards the San Francisco Bay.

Ground-water recharge occurs as infiltration of rain, seepage of streams, and subsurface inflow from adjacent areas. Recharge to the east at higher elevations results in slightly confined conditions throughout the East Bay Plain.

The older alluvium (Alameda Formation) is the major ground-water reservoir in the Oakland Upland subarea. Wells that utilize water found within this zone typically range from 100 to 500 feet in depth.

V FIELD INVESTIGATION

HLA used several tools to investigate the presence of organic chemicals in the soil and ground water at, and adjacent to, the site. This section discusses the uses and applications of these tools to the site.

A. Soil-Gas Survey

A soil-gas survey involves sampling and analyzing the soil gas from the pore spaces of the unsaturated soils (vadose zone) above the water table. This reconnaissance tool helps measure the distribution of organic chemicals in soil and ground water. Because many petroleum hydrocarbons exhibit significant vapor pressures, their introduction into subsurface soil results in vapor-phase permeation and transport. If they reach the water table and travel with ground water, vapors can emanate into the overlying soil. Thus, measuring the concentrations of organic compounds in the soil gas can give some indication of their presence in soil or ground water.

Because of complex interactions between organic compounds and subsurface materials, it is seldom possible to quantitatively estimate concentrations of such compounds in soil or ground water from soil-gas data alone. These interactions are a function of soil particle size and mineralogy, natural soil organic content, soil moisture, temperature, lithology, and heterogeneity. However, a soil-gas survey can be an excellent relative indicator or screening tool.

On September 19, 1988, under the supervision of HLA, a soil-gas survey was conducted by Tracer Research Company (TRC) to assess the near-surface distribution of selected organic compounds. TRC's standard sample collection and analyses methods, described in Appendix C, were used. Each soil-gas sample was analyzed in the field

using gas chromatography for the following compounds: total petroleum hydrocarbons (TPH), benzene, toluene, ethylbenzene, and xylenes (BTEX). These compounds are common constituents of gasoline, and they exhibit high vapor pressure and therefore can be quantified by soil-gas measurements. Soil-gas probe locations are shown on Plate 3.

B. Soil-Sample Collection

HLA explored subsurface conditions on and off site by drilling and sampling nine soil borings during October an November, 1988. Five of the borings were completed as monitoring wells (MW-6E through MW-6I). Boring locations are shown on Plate 3, and logs are presented on Plates 4 through 12. Off site exploration was not conducted to the north and east of the site because of restricted access into Telegraph and West Grand Avenues imposed by the City of Oakland and the existing underground Bay Area Rapid Transit (BART) tunnel (see Plate 2).

The borings were advanced using truck-mounted, 8-inch (borings), and 12-inch (wells) diameter hollow-stem auger drilling equipment and sampled using a 2.5-inch-diameter (I.D.) Sprague and Henwood (S&H) split-barrel sampler lined with three, 6-inch long, brass tubes. Drilling was performed under the direction of an HLA field geologist, who logged the borings in accordance with the Unified Soil Classification System presented on Plate 13. Soil samples were screened in the field with either a Photovac TIP-I photoionization detector (PID) or a Johnson Gas Detector (Gastech), Model 1314. Vapor readings, which indicate relative concentrations of volatile organic compounds in soil, are presented on the logs.

All drill cuttings were placed in 55-gallon drums for subsequent disposal.

Sampling equipment was washed with a phosphate-free detergent solution and rinsed

with clean water between sampling intervals. All drilling equipment was cleaned using a high-pressure, hot-water wash (steam-cleaned) before and after each boring.

C. Monitoring Well Construction

The monitoring wells were constructed with steam-cleaned 4-inch-diameter, Schedule 80, flush-threaded PVC casing and screen (see well construction details, Plates 14 through 18). The wells were constructed under permit from the Alameda County Flood Control and Water Conservation District. The annular space between the screen and the borehole wall was filled with No. 3 Monterey sand to approximately 2 feet above the top of the screen. A bentonite seal was placed above the sand pack, and the remainder of the annulus was filled with cement/bentonite grout to just below the ground surface. The top of each well was placed slightly below the ground surface. The wells were equipped with locking, water-tight caps (OPW 634-TTM) to minimize intrusion of surface water. Over each well, a locking, water-tight traffic box (EMCO Wheaton A-721) was installed, which extends slightly above the surrounding grade.

D. Water Quality Sampling

On October 19, 1988, the recently installed Monitoring Wells MW-6E and MW-6F were developed, sampled, and surveyed by an HLA technician. All other existing onsite wells were also sampled at this time. On December 7, 1988, Monitoring Wells MW-6G through MW-6I were developed, sampled, and surveyed by an HLA technician. Before and after development, a clear lucite bailer was lowered into each well to check for free product. Each well was developed by purging and bailing with a PVC bailer; at least six well volumes were removed from each well. After development, we purged three additional well volumes, while monitoring temperature, pH, and conductivity, and

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then sampled the ground water. Purged water was placed in 55-gallon drums for subsequent disposal as requested by Texaco Refining and Marketing.

Ground-water samples were collected from each well with a clean, stainless steel bailer. A representative sample was decanted into laboratory-prepared, 40-milliliter volatile organic analysis (VOA) vials. The vials were immediately sealed, labeled, and placed in a cooler with blue ice until delivery for chemical testing to ChemWest Analytical Laboratories, Inc., in Sacramento, California. All sampling equipment was washed with a phosphate-free detergent solution and rinsed in clean water and distilled water between wells.

Appropriate quality assurance and quality control (QA/QC) measures were employed during the field investigation. HLA maintains an internal QA/QC program that includes provisions for avoiding cross-contamination during site investigation and procedures for decontamination, sample handling, preservation, and chain-of-custody.

VI SUBSURFACE CONDITIONS

A. Geologic Profile

All soils encountered within the borings were described in accordance with the USCS (Plate 13). Borings B-1 through B-3 and Monitoring Wells MW-6A through MW-6F were located on site within paved areas. Borings B-4 and wells MW-6G through MW-6I were located off-site in the adjacent sidewalks. As shown on Plates 4 through 12, subsurface materials generally consist of stiff, silty clay underlain by a dense silty sand layer of variable thickness. The top of the sand occurs at between 10 and 15 feet deep. In some borings, the sand layer is in turn underlain by more silty clay or silt. Ground water was initially encountered in the borings at approximately 13 feet below grade.

Relatively strong gasoline odors were observed in Borings B-1 and B-3 and MW-6H in soils very near the water table. In all other borings only trace amounts of gasoline vapors were found in vadose zone soils. Two soil samples from each boring, and one soil sample from each boring where a well was installed, were submitted for chemical testing. The laboratory results are discussed in Section VII of this report.

The BART right-of-way is located beneath the intersection of Telegraph Avenue and West Grand Avenue. This portion of the BART system is underground. A small portion of this right-of-way encompasses the extreme eastern corner of the site.

B. Ground-Water Flow Patterns

The tops of well casings were surveyed to a temporary datum located at the western end of the dispenser island nearest West Grand Avenue with an assumed elevation of 100.0 feet (HLA datum, see Plate 3). Well monitoring and survey data are

presented in Table 1. The estimated direction of ground-water flow is to the southwest, with a gradient of 0.004 feet per foot. As shown on the groundwater surface map, Plate 19, a slight variation in the ground-water flow direction is observed near the northern portion of the site. At this location, groundwater appears to flow to the west.

Table 1. Well Monitoring and Survey Data

	Top of		Ground-Water	
	Casing	Depth to	Surface	
Well	Elevation*	Ground Water**	Elevation+	
No.	(feet)	(feet)	(feet)	Comments
MW-6A	98.99	13.40	85.59	no odor
MW-6B	98.81	12.94	85.87	no odor
MW-6C	99.89	14.10	85.79	hydrocarbon odor and sheen
MW-6D	98.78	13.44	85.34	hydrocarbon oder and sheen
MW-6E	98.99	13.70	85.29	no odor
MW-6F	99.91	14.48	85.43	no odor
MW-6G	99.16	12.22	86.94	no odor
MW-6H	97.93	12.36	85.57	hydrocarbon odor
MW-6I	97.60	12.83	84.77	no odor

Notes:

- * Elevation relative to HLA temporary benchmark located at the western end of the dispenser island nearest West Grand Avenue with an arbitrary elevation of 100.0 feet (see Plate 3).
- ** Depth to ground water on December 15, 1988.
- + Ground-water surface elevation = top of casing elevation depth to water.

VII DISCUSSION OF CHEMICAL TEST RESULTS

A. Chemical Test Methods

Soil-gas, and/or ground-water samples obtained from soil-gas probes, were chemically analyzed in the on-site mobile laboratory for their concentration of BTEX and TPH. Ambient air samples were also obtained at the start and finish of work and were tested for the same compounds.

To evaluate the presence of petroleum products in the vadose zone soils and the ground water, we scheduled the following chemical analyses:

Soil -

TPH as gasoline, using DHS Method - LUFT Field Manual, procedures, and BTEX, using EPA Test Method 8020.

Ground Water -

BTEX, using EPA Test Method 602.

B. Distribution of Chemicals in Soil-gas

Detectable concentrations of petroleum products in soil-gas samples were limited to probe locations SG-3 and SG-4 (Plate 3). At these locations relative high concentrations of xylenes and TPH were detected in soil-gas samples obtained from depths of 12 and 13 feet at probe locations SG-3 and SG-4, respectively. Both SG-3 and SG-4 were located along the edge of Telegraph Avenue which possesses underground sewer and storm drain lines. The chromatograph resulting from these samples did not match that of gasoline. The on-site chemist informed us that it more closely resembled the pattern that a paint or varnish would produce. An example of a typical chromatograph produced by gasoline, and the chromatograph produced from samples obtained from SG-3 and SG-4 is presented in Appendix C.

Concentrations of petroleum products were less than the detection limit at probe locations SG-2, WS-5 and SG-6. Because of tight clays encountered at probe locations SG-1 and SG-7, we were not able to obtain soil-gas samples. A summary of the analytical results of the soil-gas survey is presented in Table 2.

Table 2. Analytical Results of Soil-Gas Survey
Conducted on September 19, 1988

Concentrations in micrograms per liter (ug/L)

						Total
	Depth		Ethyl-			Petroleum
<u>Sample</u>	(ft)	<u>Benzene</u>	<u>Benzene</u>	<u>Toluene</u>	Xy lenes	Hydrocarbons
Air	N/A	<0.7	<0.8	<0.8	<0.8	<0.7
SG-01						
SG-02	5.0	<0.7	<0.8	<0.8	<0.8	<0.7
SG-03	12.0	10	4	<0.8	2,800	6,100
SG-04	13.0	<0.7	<0.8	<0.8	140	780
WS-05*	12.0	<75	<76	<77	<77	<75
SG-06	13.0	<0.7	<0.8	<0.8	<0.8	<0.7
SG-07						
Air	N/A	<0.7	<0.8	<0.8	<0.8	<0.7
	Not able to obtain sample					

N/A -not applicable

Air -ambient air sample

WS-05 was a sample of ground water

C. Soil Sample Test Results

Soil samples were obtained for chemical testing from on and off-site borings at the sample depths with the highest PID vapor readings. Laboratory results of chemical analyses on soil are presented in Table 3. Laboratory reports are presented in Appendix D.

Table 3. Laboratory Results of Soil Analyses Concentrations in milligrams per kilogram (mg/kg)

Sample Number	Depth (ft)	<u>Benzene</u> ¹	Ethyl <u>Benzene²</u>	<u>Toluene³</u>	Xylenes ³	TPH as Gasoline ⁴
B-1	8.0	0.05	ND	ND	ND	ND
B-1	13.0	ND(5)	10(10)	16(10)	41(10)	2,000(1,000)
B-2	7.0	ND	ND	ND	ND	ND
B-2	13.5	ND	ND	ND	ND	ND
B-3	7.0	0.06	ND	ND	ND	ND
B-3	13.5	₄₀ (25)	84(50)	39 0 (50)	₃₇₀ (50)	11,000(5,000)
B-4	13.5	ND	ND	ND	ND	ND
MW-6E	13.0	ND	ND	ND	ND	ND
MW-6F	13.0	ND	ND	ND	NĎ	ND
MW-6G	13.5	ND	ND	ND	ND	5.2
MW-6H	13.5	11(0.5)	3.8(2)	3.2(1)	19(1)	$1,000^{(495)}$
MW-6I	13.5	ND	ND	ND	ND	ND

ND = Not detected.

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Detection limit 0.05 mg/kg except as noted in parentheses.

Detection limit 0.2 mg/kg except as noted in parentheses.

³ Detection limit 0.1 mg/kg except as notes in parentheses.

⁴ Detection limit 10 mg/kg except as noted in parentheses.

Test results indicate the absence of BTEX or TPH in the vadose zone soils at the locations sampled. Relatively high concentrations of BTEX and TPH are found in soils very near the water table. The soil sample from boring B-3, from a depth of 13.5 feet, had the highest concentrations of BTEX (11,000 parts per million [ppm] TPH)*.

Soil from the same depth (approximately 13 feet) in B-1 and MW-6H also contained relatively high concentrations of TPH at 2,000 and 1,000 ppm respectively. Soil obtained from all other boring locations had either nondetectable concentrations, or concentrations very near the detection limit for BTEX and TPH.

D. Water Quality Results

Monitoring wells MW-6A through MW-6F were sampled on October 19, 1988 and wells MW-6G through MW-6I were sampled on December 7, 1988 using the procedures described in Section V. Results of chemical analyses on ground water samples are summarized in Table 4. Laboratory reports are presented in Appendix E.

^{*}Indicated on Table 3 as milligrams per kilogram

Table 4. Laboratory Results of Ground-Water Analyses
Concentrations in micrograms per liter (ug/L)

EPA TEST METHOD 602

Well <u>Number</u>	Date <u>Sampled</u>	<u>Benzene</u>	Ethyl- <u>Bensene</u>	<u>Toluene</u>	Xylenes
MW-6A	10/20/88	$0.6^{(0.5)}$	ND(2)	ND(1)	ND(1)
MW-6B	10/20/88	4.1(0.5)	ND(2)	2.5(1)	ND(1)
MW-6C	10/20/88	9,500 ⁽⁵⁰⁾	170(2)	65(100)	850(1)
MW-6D	10/20/88	710(5)	22(20)	74(10)	110(10)
MW-6E	10/20/88	1.1(0.5)	$_{ m ND}(2)$	ND(1)	3.4(1)
MW-6F	10/25/88	$ND^{(0.5)}$	ND(2)	ND(1)	2.4(1)
MW-6G	12/07/88	ND ^(0.5)	$_{ m ND}^{}(2)$	ND ⁽¹⁾	ND(1)
MW-6H	12/07/88	1200(25)	110(20)	320(10)	220(10)
MW-6I	12/07/88	$_{ m ND}^{(0.5)}$	ND(2)	ND(1)	ND(1)

ND = Not detected.

Detection limits given in parentheses.

Detectable concentrations of BTEX were found in ground water samples from all on-site wells. Detectable concentrations of BTEX were also found in off-site well MW-6H. No BTEX was detected in off-site wells MW-6G and MW-6I.

VIII AQUIFER TESTING

The hydraulic conductivity of shallow saturated earth materials was estimated from the results of tests using three monitoring wells located generally downgradient of the underground fuel tanks on-site. Three single-well slug tests were performed using monitoring wells MW-6H, MW-6D, and MW-6E. At the time of the slug tests, the equilibrium water levels in the wells were 11.88, 12.59, and 12.58 feet below the top of casing in each well, respectively. The water levels were compared to the stratigraphic log of the wells to classify the most permeable statum adjacent to the screen in the saturated zone as hydraulically confined or unconfined.

Various techniques were used to rapidly remove from or inject into the wells a volume (slug) of water. Depending on the circumstances, slugs were removed by pumping them out of the wells with either a centrifugal suction pump or a submersible turbine pump. Slug injection was accomplished by rapidly releasing a volume of clean drinking water into the well. Table 5 summarizes the conditions of the slug tests at this site.

Table 5. Slug Test Conditions

Well Number	Test <u>Type</u>	Pump Type	Approximate Slug Volume (gallons)	Initial Water Level (feet)	Classification of Stratum
MW-6D	Injection		2	12.59	Confined
MW-6E	Withdrawal	Submersible	3	12.58	Confined
MW-6H	Withdrawal	Suction	1.5	11.88	Unconfined

A pressure transducer, placed near the bottom of the wells was used to measure the water level recoveries following the slug injections or withdrawals. The output of the transducer was interpreted and recorded by a data logger for subsequent analysis. The methods of slug-test analysis are discussed in Appendix F.

The slug-test recovery hydrographs are shown in Plates 21, 22, and 23 for tests in MW-6D, MW-6E, and MW-6H, respectively. Table 6 lists the hydraulic conductivity estimates derived from the tests.

Table 6. Slug Test Results

Well <u>Number</u>	Lithology of Most Permeable Stratum	Thickness of Stratum <u>(feet)</u>	Estimated Hydraulic Conductivity of Stratum (feet/day)
MW-6D	sand	2	5.9
MW-6E	sand, fine-grained	2.5	1.2
MW-6H	sand, medium-grained	6	4.8

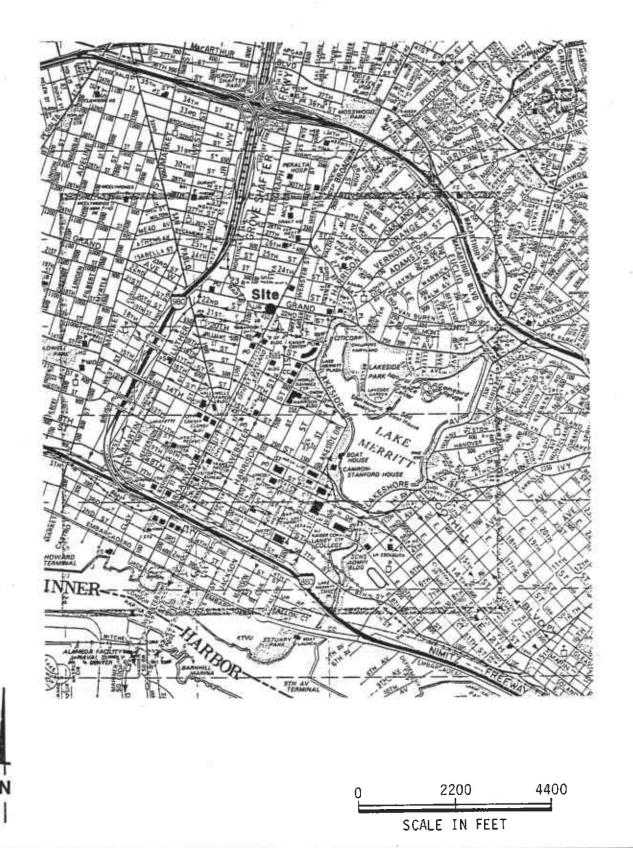
IX SUMMARY OF FINDINGS

Conclusions and observations are summarized as follows:

- 1. The shallow subsurface soils consist of 12 to 14 feet of silty and sandy clay underlain by 2 to 8 feet of sand. Clay was again encountered beneath the sand unit where the sand was relatively thin. The water table surface is approximately 12 to 13 feet below the ground surface.
- 2. The ground water flow direction is to the southwest with a gradient of 0.004 feet per foot. A slight variation in the ground-water flow pattern is observed near the northern portion of the site. At this location, groundwater appears to flow to the west.
- 3. The hydraulic conductivity of shallow fine-to medium-grained saturated sand beneath the site is estimated to range from 1.2 to 5.9 feet per day.
- 4. BTEX and TPH (as gasoline) have been detected exclusively in soils at 12 to 13 feet below the ground surface. This interval is within the zone of fluctuation of the groundwater table.
- 5. Elevated concentration of TPH and xylenes detected in SG-3 and SG-4 can most likely be attributed to the close proximity of these probe locations to the sanitary sewer and storm drain lines beneath Telegraph Avenue. The resulting chromatograph from vapor samples collected from these two probes more closely resembles the pattern a paint product or varnish would produce, neither of which would have commonly been stored or used on site.
- 6. As shown on Plate 20, shallow groundwater beneath the site contains detectable quantities of BTEX. BTEX has also been detected in the ground water off-site, in the down gradient direction. The lateral extent of BTEX in the ground water is not known at this time because of restricted access into Telegraph and West Grand Avenues imposed by the city of Oakland. Subsurface exploration north of the site, into West Grand Avenue, was also restricted because of the existing BART tunnel.
- 7. The highest concentrations of BTEX in the ground water from onsite wells is found in MW-6C and MW-6D, which are very near the underground fuel tanks or directly downgradient of them. The underground tank complex most likely represented an onsite source. The relatively high concentrations of BTEX found in MW-6H, and at lower concentrations in MW-6B, may represent an unidentified off-site source, MW-6B is upgradient of the tanks and MW-6H is crossgradient.

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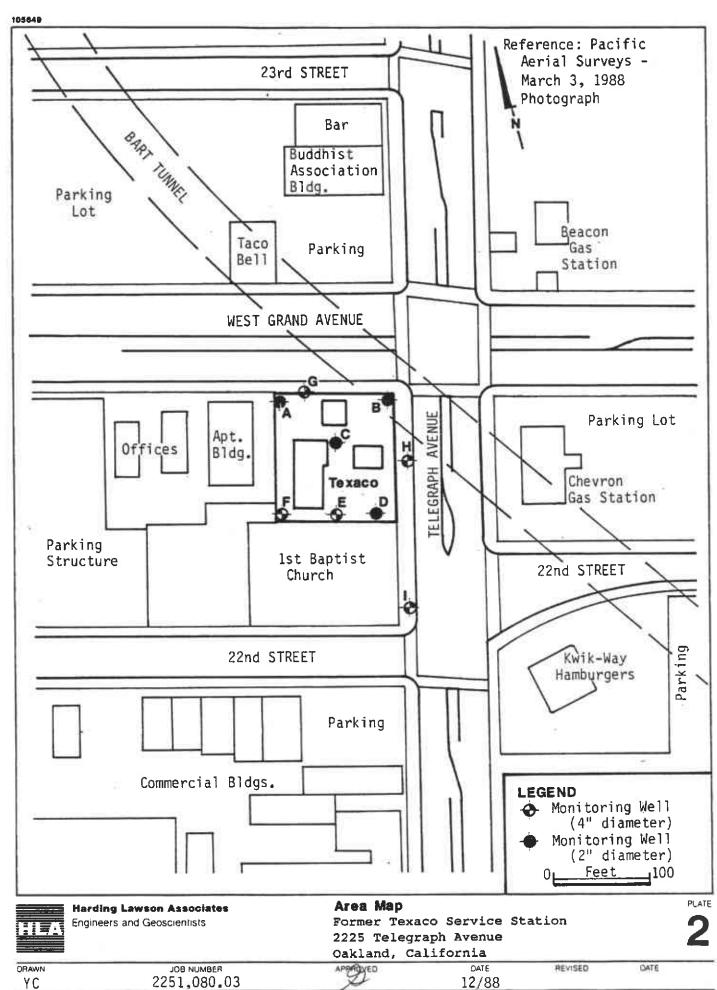
Engineers and Geoscientists

Vicinity Map

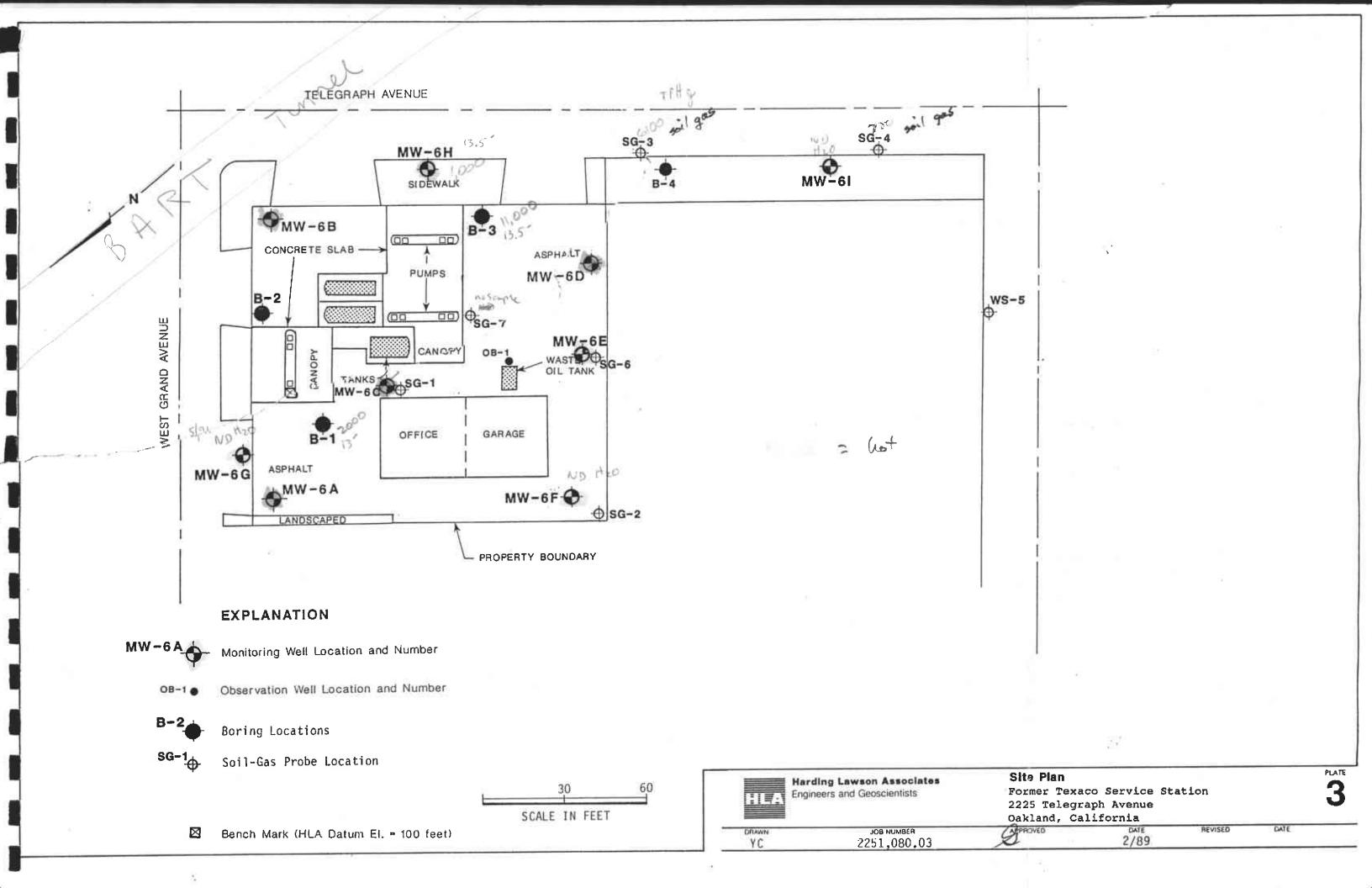
Former Texaco Service Station 2225 Telegraph Avenue Oakland, California PLATE

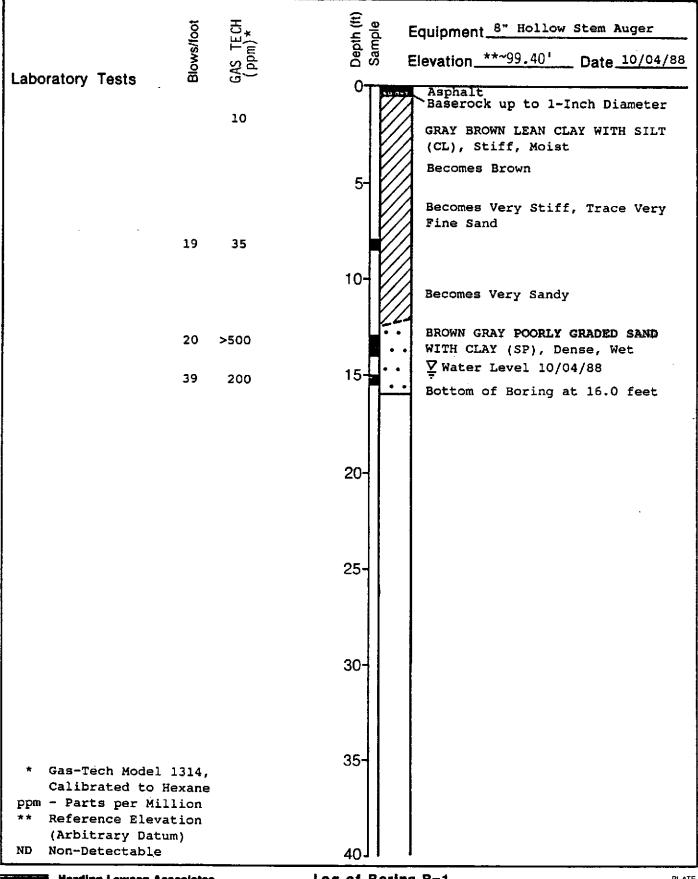
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Log of Boring B-1

Former Texaco Service Station 2225 Telegraph Avenue

4

Oakland, California

DRAWN JOB NUMBER APPROVED DATE REVISED DATE
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Labor	atory Tests	GAS TECH (ppm)*	Equipment 8" Hollow Stem Auger But Equipment 8" Hollow Stem Auger But Elevation ** 798.80' Date 10/04/88 Asphalt Baserock
	16	35	GREENISH GRAY LEAN CLAY WITH SILT (CL), Stiff, Moist Fine Sand Lens Becomes Mottled Greenish Gray
			GREENISH GRAY POORLY GRADED SAND WITH SILT (SP), Dense, Moist, Medium-Grained
	. 41	30	BROWN LEAN CLAY WITH SAND (CL), Very Stiff, Wet Absence of Sand
	10	20	Bottom of Boring at 19.5 feet
		·	25-
			30-
* ppm **	Gas-Tech Model 1314, Calibrated to Hexane - Parts per Million Reference Elevation (Arbitrary Datum)		35-
ND	Non-Detectable		40]]
	Harding Lawson Associates		Log of Boring B-2 PLATE

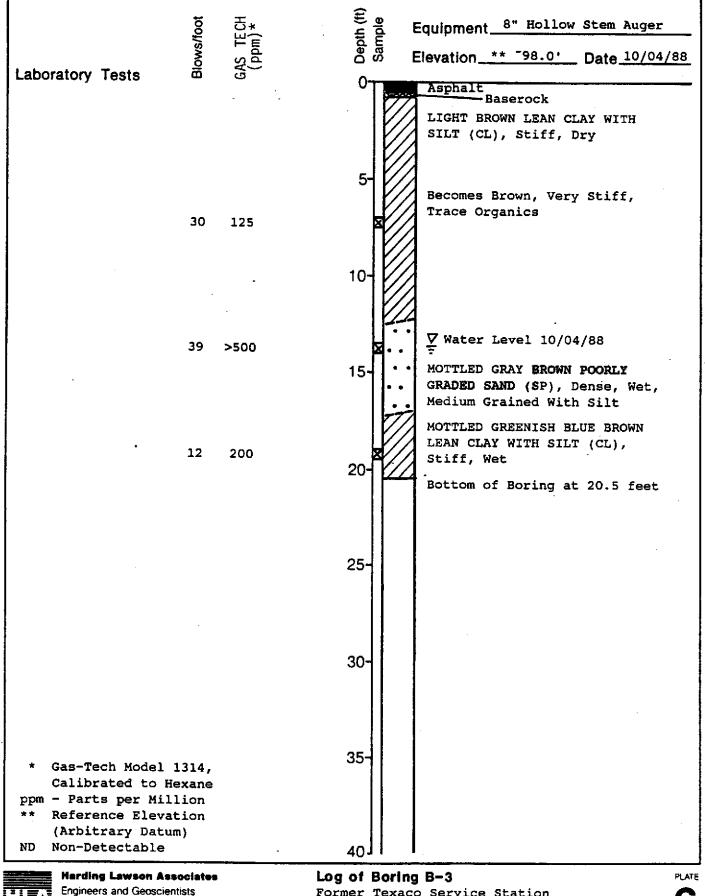


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Log of Boring B-2
Former Texaco Service Station

2225 Telegraph Avenue Oakland, California

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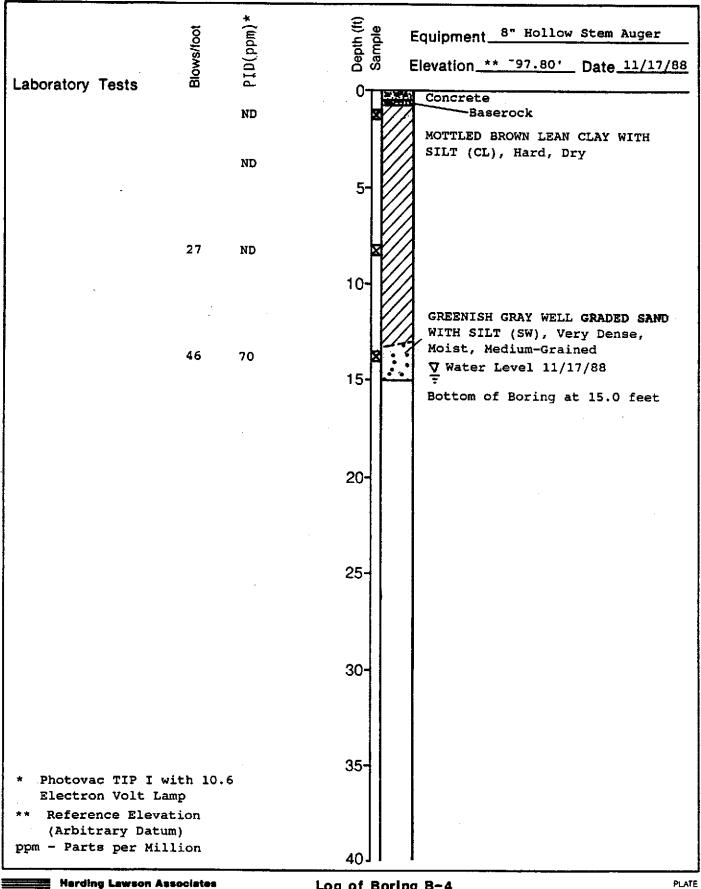
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YC

Former Texaco Service Station 2225 Telegraph Avenue

Oakland, California

REVISED DATE 12/88



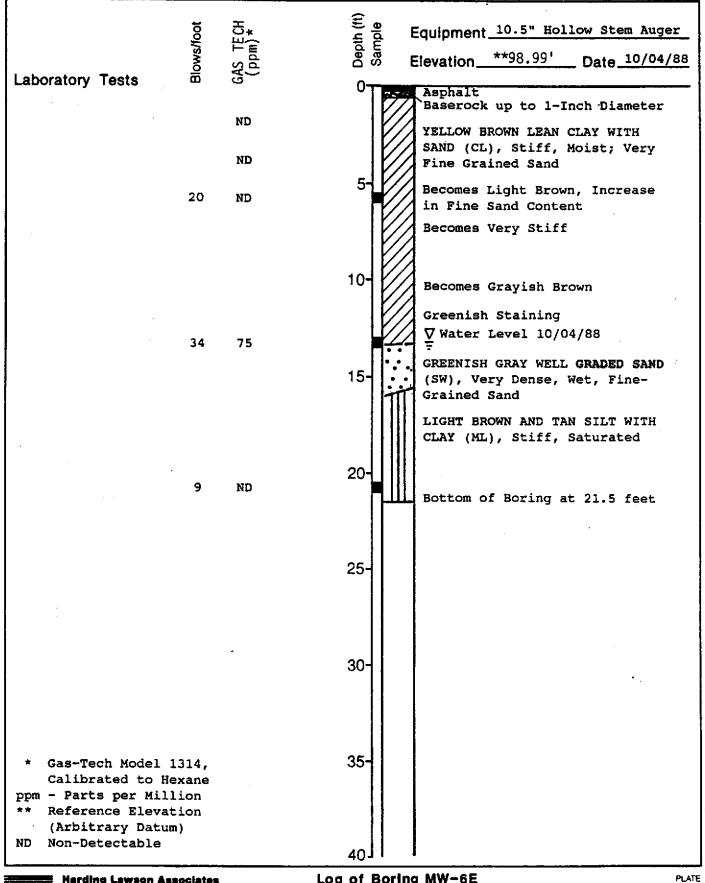


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Log of Boring B-4

Former Texaco Service Station 2225 Telegraph Avenue

Oakland, California JOB NUMBER 2251,080.03 DRAWN APPROVED DATE REVISED YC 12/88



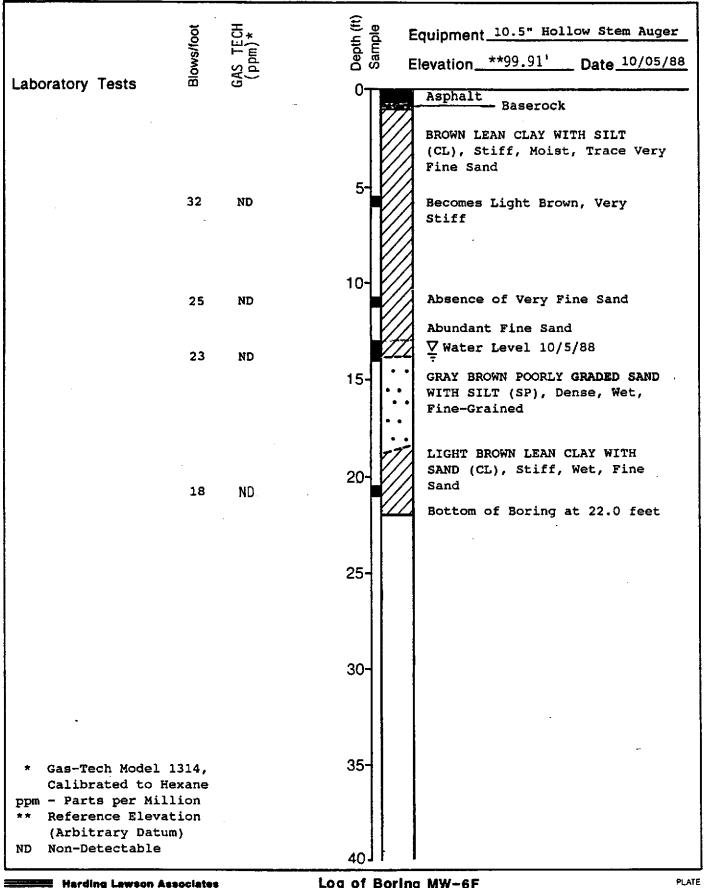


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Log of Boring MW-6E

Former Texaco Service Station 2225 Telegraph Avenue Oakland, California

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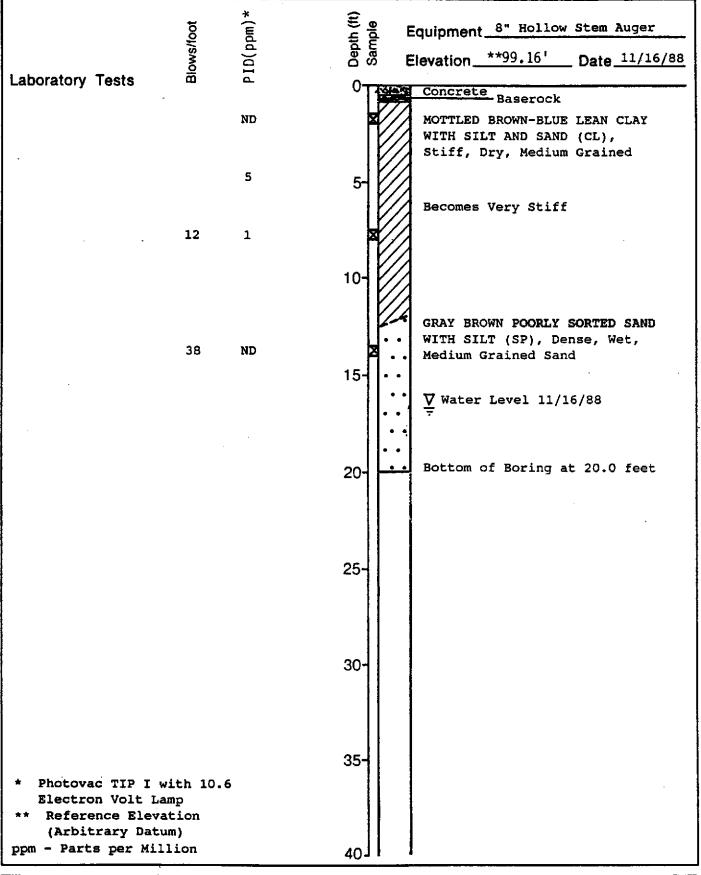


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Log of Boring MW-6F Former Texaco Service Station 2225 Telegraph Avenue Oakland, California

9

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Log of Boring MW-6G

Former Texaco Service Station 2225 Telegraph Avenue

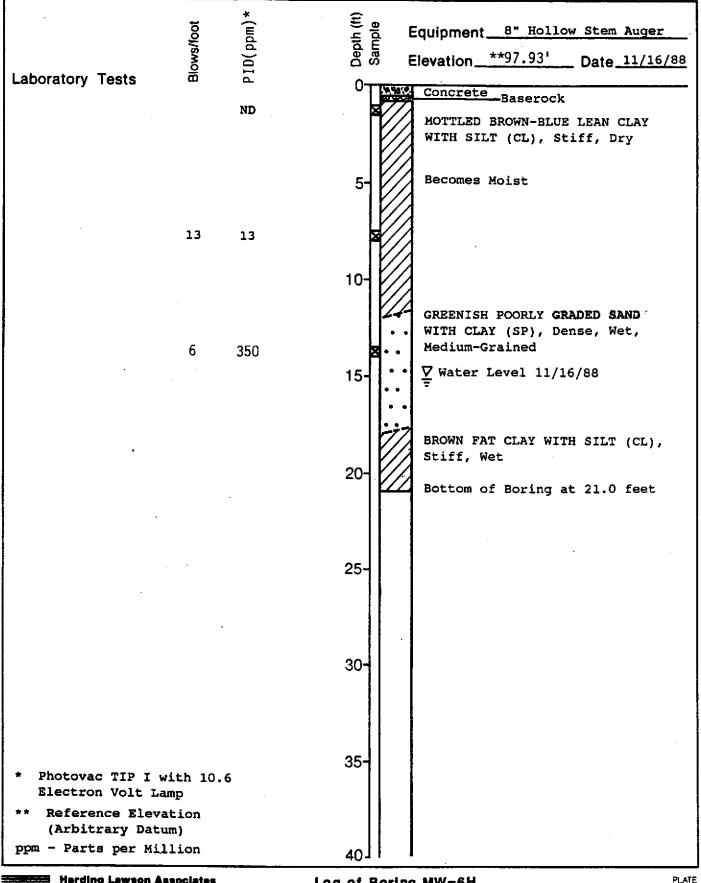
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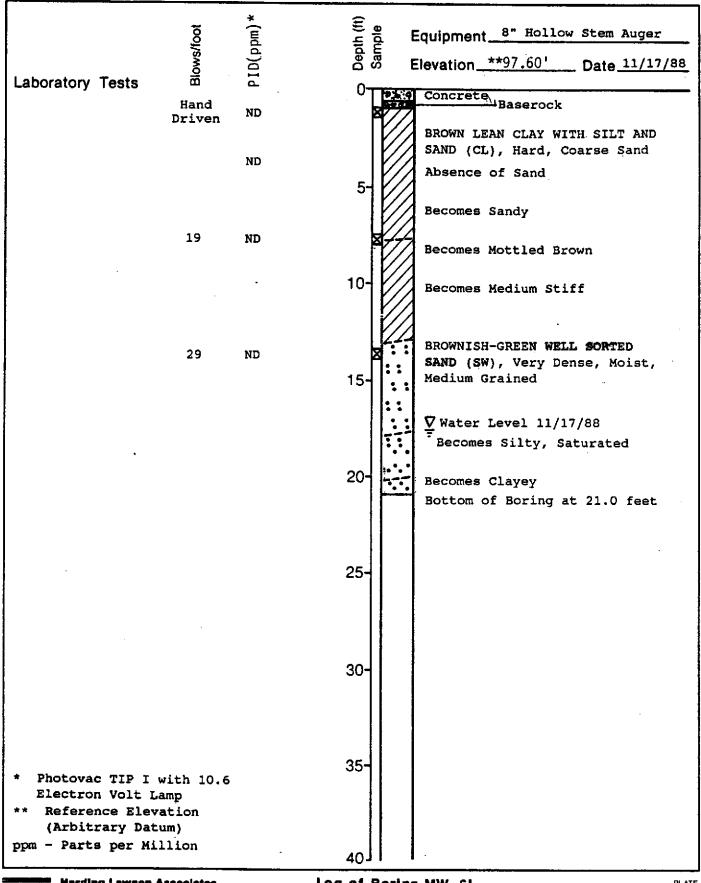
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Log of Boring MW-6H

Former Texaco Service Station 2225 Telegraph Avenue

Oakland, California

DRAWN JOB NUMBER DATE REVISED DATE Y C 2251,080.03 12/88





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Log of Boring MW-61

Former Texaco Service Station 2225 Telegraph Avenue 12

Oakland, California

DRAWN JOB NUMBER APPROVED DATE
YC 2251,080.03 12/88

	MAJOR DIV	ISIONS		TYPICAL NAMES
SOILS OARSER Ve		CLEAN GRAVELS WITH	gw	WELL GRADED GRAVELS WITH OR WITHOUT SAND, LITTLE OR NO FINES
	GRAVELS MORE THAN HALF COARSE FRACTION IS LARGER THAN NO. 4 SIEVE SIZE	LITTLE OR NO FINES	GP	POORLY GRADED GRAVELS WITH OR WITHOUT SAND, LITTLE OR NO FINES
		GRAVELS WITH OVER 12% FINES	GM	SILTY GRAVELS, SILTY GRAVELS WITH SAND
LF IS C			GC	CLAYEY GRAVELS, CLAYEY GRAVELS WITH SAND
COARSE—GRAINED SOII MORE THAN HALF IS COARSEI THAN NO. 200 SIEVE		CLEAN SANDS WITH	sw	WELL GRADED SANDS WITH OR WITHOUT GRAVEL, LITTLE OR NO FINES
	SANDS MORE THAN HALF COARSE FRACTION IS SMALLER THAN NO. 4 SIEVE SIZE	LITTLE OR NO FINES	SP	POORLY GRADED SANDS WITH OR WITHOUT GRAVEL, LITTLE OR NO FINES
		SANDS WITH OVER 12% FINES	ѕм	SILTY SANDS WITH OR WITHOUT GRAVEL
			sc	CLAYEY SANDS WITH OR WITHOUT GRAVEL
FINE—GRAINED SOILS MORE THAN HALF IS FINER THAN NO. 200 SIEVE	SILTS AND CLAYS LIQUID LIMIT 50% OR LESS			INORGANIC SILTS AND VERY FINE SANDS, ROCK FLOUR, SILTS WITH SANDS AND GRAVELS
				INORGANIC CLAYS OF LOW TO MEDIUM PLASTICITY, CLAYS WITH SANDS AND GRAVELS, LEAN CLAYS
				ORGANIC SILTS OR CLAYS OF LOW PLASTICITY
	SILTS AND CLAYS LIQUID LIMIT GREATER THAN 50%			INORGANIC SILTS, MICACEOUS OR DIATOMACIOUS, FINE SANDY OR SILTY SOILS, ELASTIC SILTS
				INORGANIC CLAYS OF HIGH PLASTICITY, FAT CLAYS
				ORGANIC SILTS OR CLAYS OF MEDIUM TO HIGH PLASTICITY
HIGHLY ORGANIC SOILS			Pt	PEAT AND OTHER HIGHLY ORGANIC SOILS

UNIFIED SOIL CLASSIFICATION - ASTM D2487-85

Perm Permeability		Shear Strength (psf)		Confining Pressure		g Pressure		
Consol	_	Consolidation	TxUU	3200	(2600)	_	Unconsolidated Undrained Triaxial Shear	
LL	_	Liquid Limit (%)	(FM) or (S)				(field moisture or saturated)	
PI	_	Plastic Index (%)	TxCU	3200	(2600)		Consolidated Undrained Triaxial Shear	
G _s	_	Specific Gravity	(P)				(with or without pore pressure measurement	
-		· ·	TxCD	3200	(2600)	_	Consolidated Drained Triaxial Shear	
MA	_	Particle Size Analysis	SSCU	3200	(2600)	_	Simple Shear Consolidated Undrained	
	_	"Undisturbed" Sample	(P)				(with or without pore pressure measureme	
\boxtimes	_	Bulk or Classification Sample	SSCD	3200	(2600)	_	Simple Shear Consolidated Drained	
		•	DSCD	2700	(2000)		Consolidated Orained Direct Shear	
			ŲC	470		_	Unconfined Compression	
			LVS	700		_	Laboratory Vane Shear	

KEY TO TEST DATA



Harding Lawson Associates

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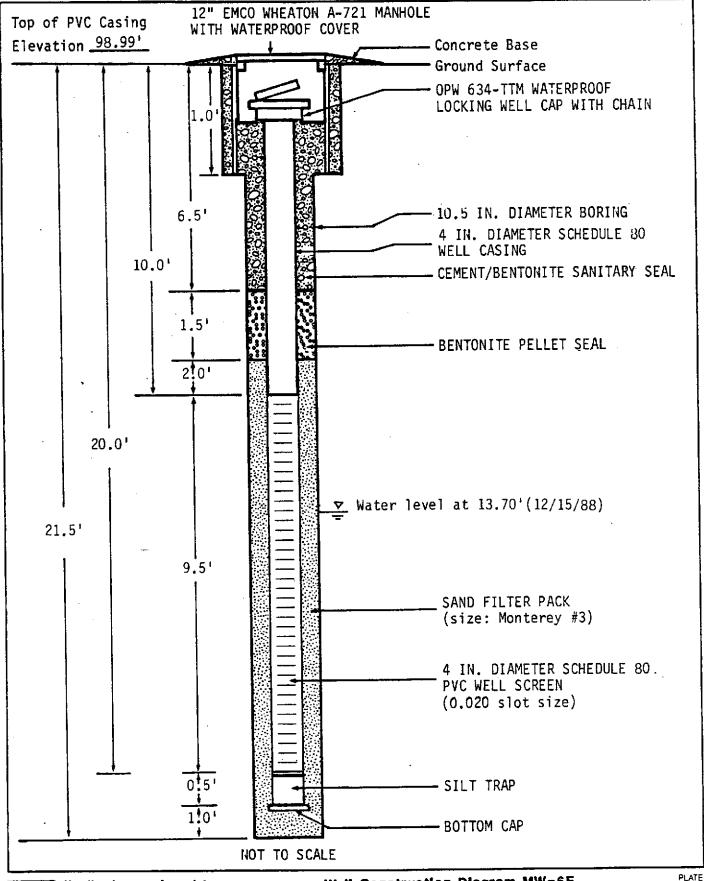
Soil Classification and Test Data Key Former Texaco Service Station 2225 Telegraph Avenue Oakland, California

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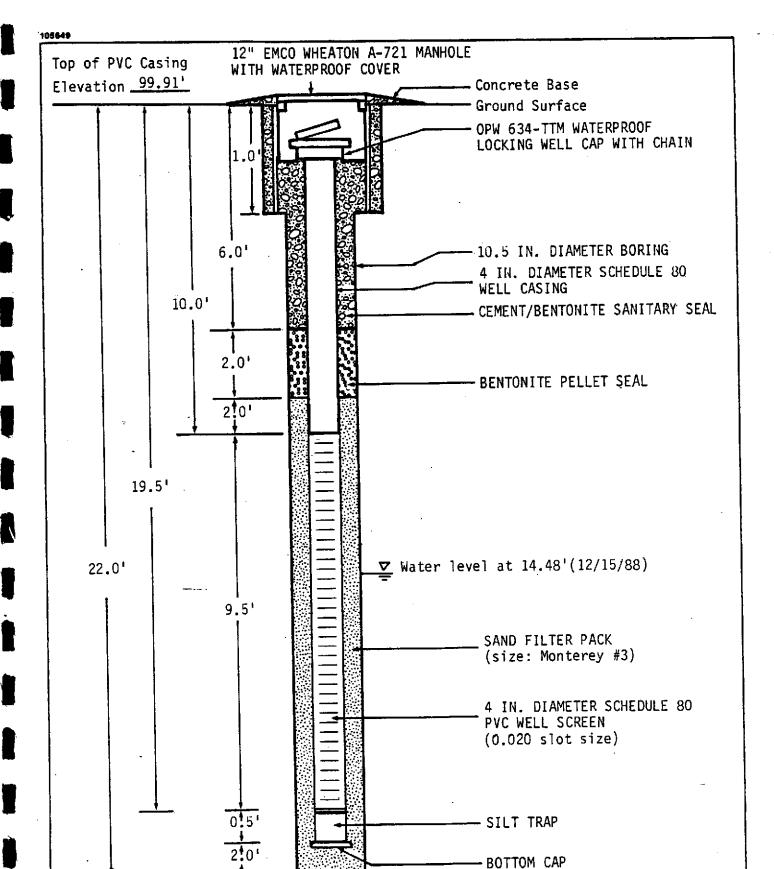
Harding Lawson Associates Engineers and Geoscientists

Well Construction Diagram MW-6E Former Texaco Service Station

2225 Telegraph Avenue

Oakland, California

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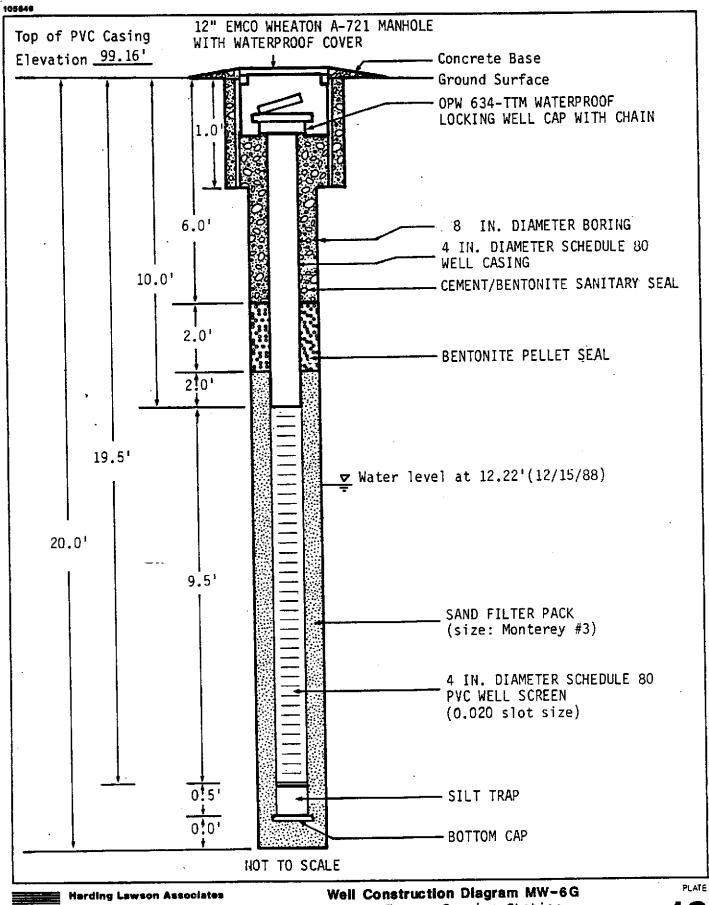
Harding Lawson Associates Engineers and Geoscientists Well Construction Diagram MW-6F

Former Texaco Service Station 2225 Telegraph Avenue Oakland, California 15

PLATE

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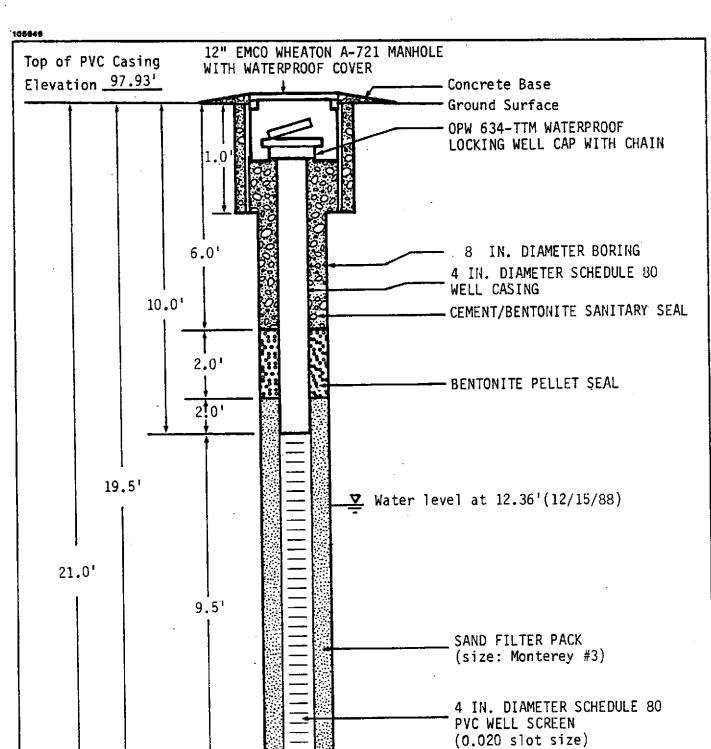
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Engineers and Geoscientists

Former Texaco Service Station 2225 Telegraph Avenue Oakland, California

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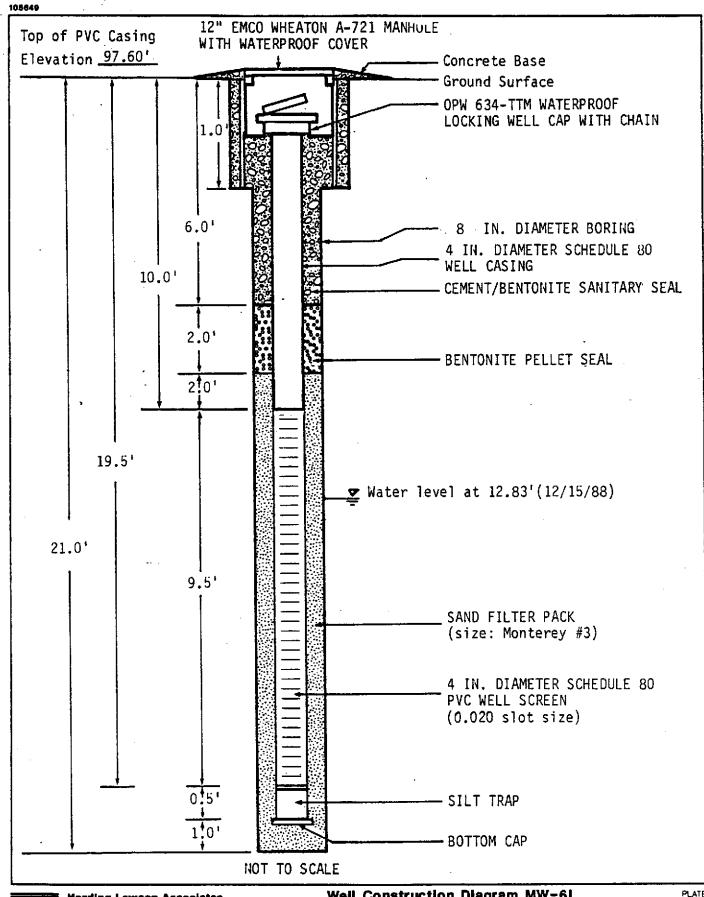
SILT TRAP -BOTTOM CAP NOT TO SCALE

Harding Lawson Associates Engineers and Geoscientists

Well Construction Diagram MW-6H Former Texaco Service Station 2225 Telegraph Avenue

Oakland, California

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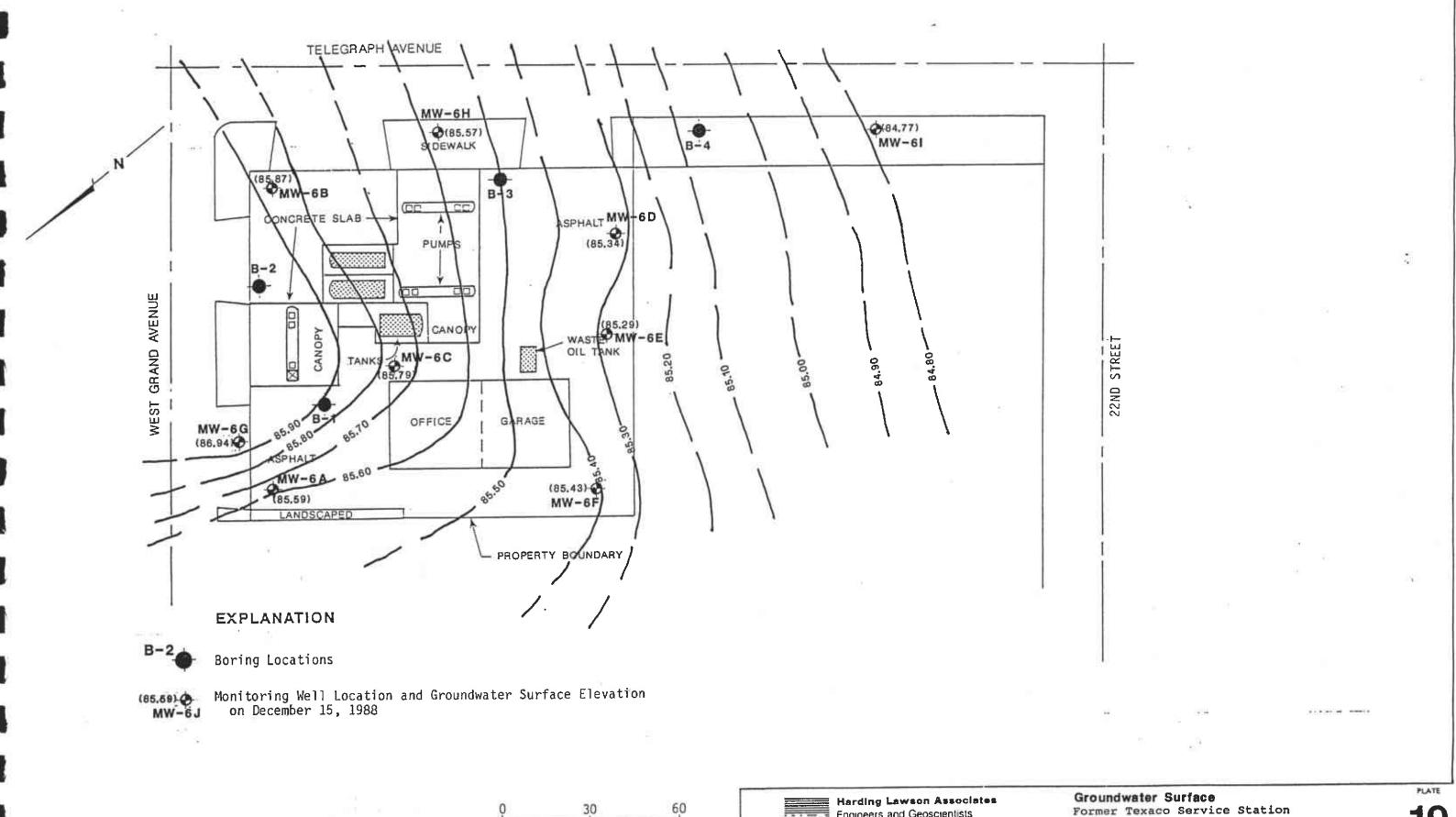
Harding Lawson Associates
Engineers and Geoscientists

Well Construction Diagram MW-61 Former Texaco Service Station

2225 Telegraph Avenue Oakland, California

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Bench Mark (HLA Datum El. = 100 feet)

SCALE IN FEET

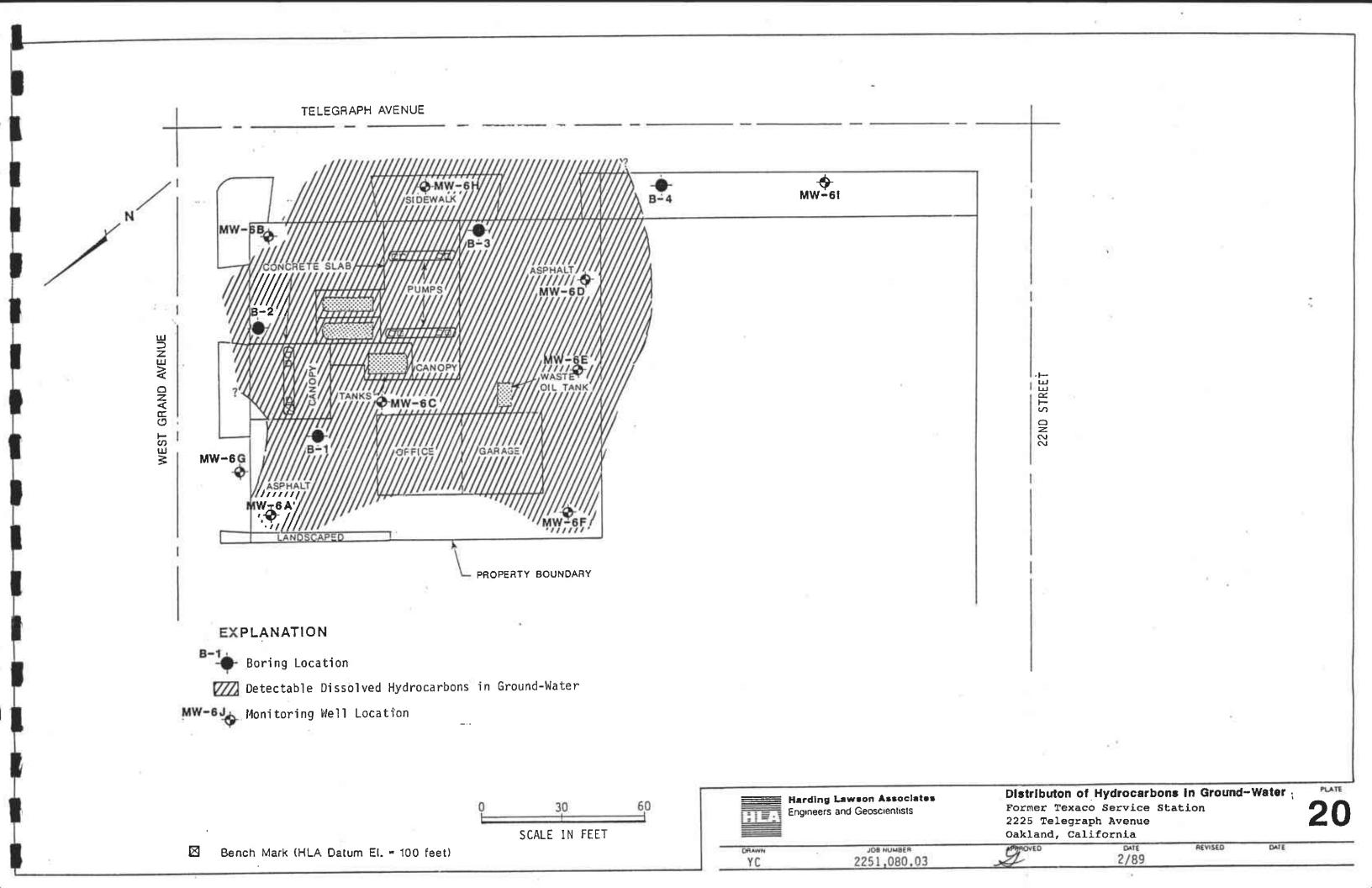
Engineers and Geoscientists

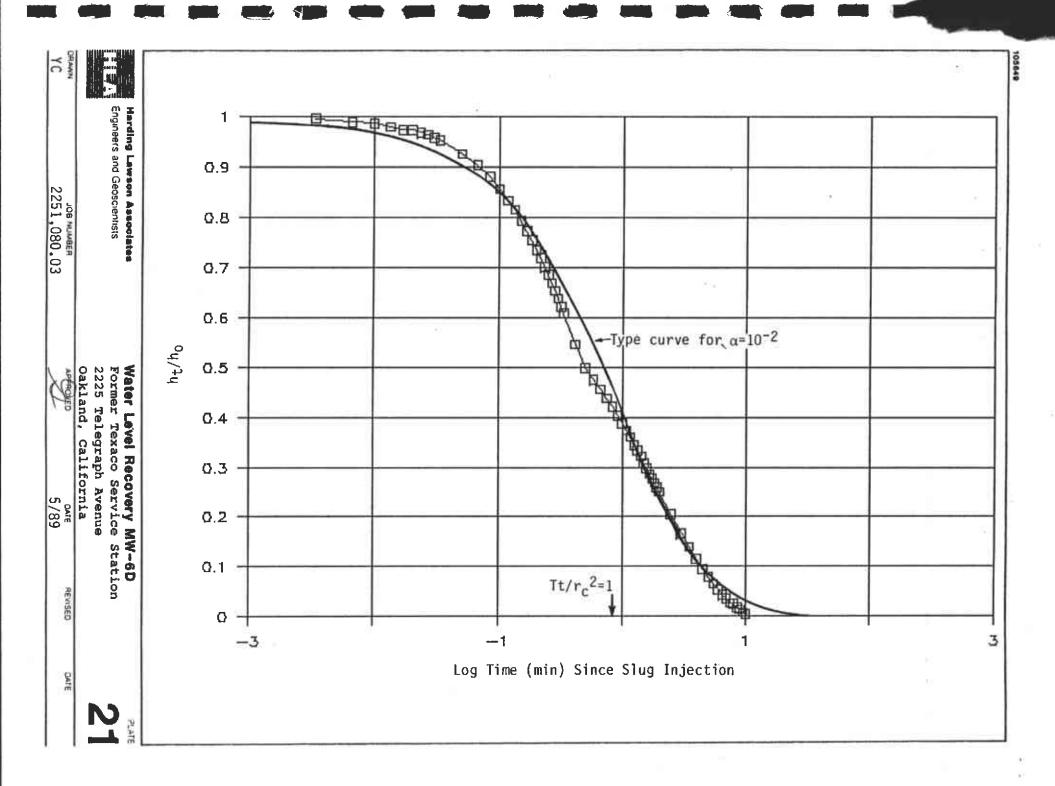
2225 Telegraph Avenue Oakland, California

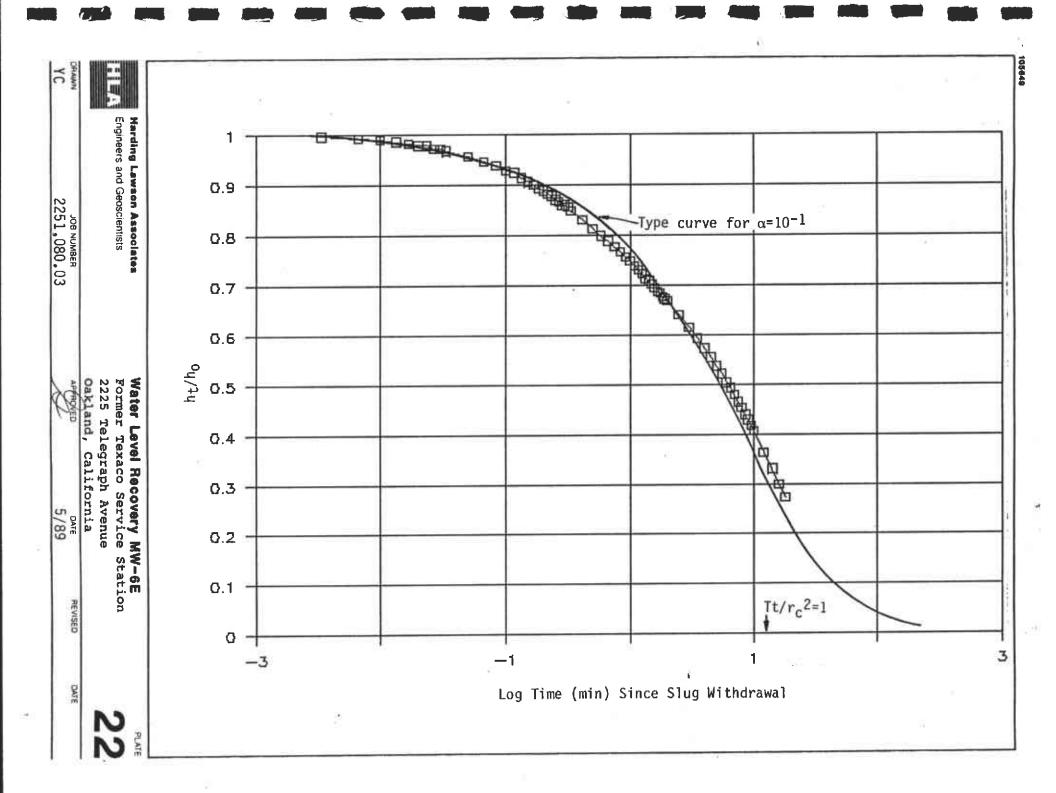
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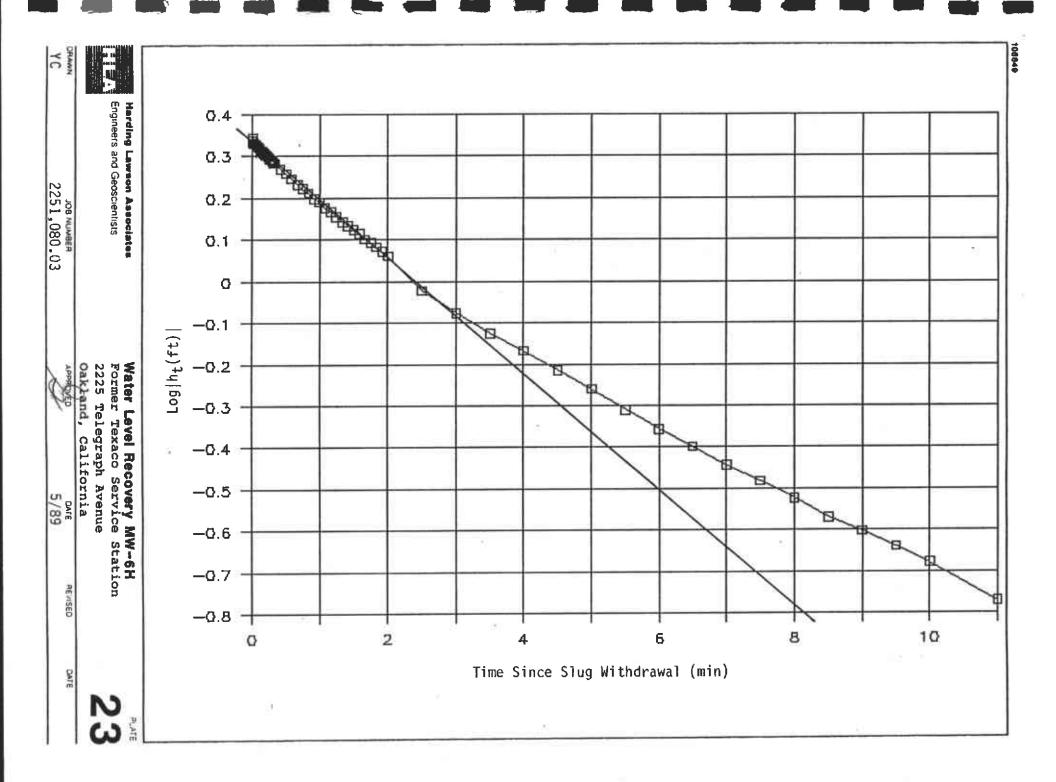
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APPENDIX A

SENSITIVE RECEPTOR SURVEY FACT SHEET, RESULTS OF GROUND-WATER ANALYSES (JUNE, 1988), BORING LOGS & WELL CONSTRUCTION DETAILS

SENSITIVE RECEPTORS - SITE INVESTIGATION AND RISK ASSESSMENT

	Location #: 62488000195 Address: 2225 Telegraph City/State: Oakland CA County: Alameda
I I	Provide answers to the following questions to the extent reasonably known:
	A. Is there a public water supply well within 2500'? (Y/N) No If Yes, distance (FT)
	B. Is there a private water supply well within 1000'? (Y/N) No If Yes, distance (FT)
	C. Is there a subway within 1000'? If Yes, distance (FT) 600
	D. Is there a basement within 500'? If Yes, distance (Y/N) Yes (FT) 25
	E. Is there a school within 1000'? If Yes, distance (Y/N) Yes (Y/N) 800
ĺ	F. Is there a surface body of water within 500'? (i.e., lake, river, ocean) If Yes, distance (FT)
1	Describe type of local water supply: Public Suppliers' Name Suppliers' Source Bokelumne Aqueduc Distance to Site Private
II	Aquifer Classification, if available:
] 	Class I - Special Ground Waters - Irreplaceable Drinking Water Source - Ecologically Vital Class II - Current and Potential Drinking Water Sources
	Class III - Not Potential Source of Drinking Water
J	Describe observation wells, if any: Number $\frac{1}{\text{Free Product (Y/N)}}$
	Provide a site diagram or a local/topographic (USGS) map of the area.
ī	Report should consist of this fact sheet, the site or area map, and a cover letter.
II	Signature of Preparer: Date: Date:

Table 2. Results of Ground-Water Analyses* (concentrations in micrograms per liter $[\mu g/l]$)

Well No.	Benzene	Ethyl- benzene	Toluene	Xylenes	
MW-6A	ND (0.5)	ND (2)	ND (1)	ND (I)	
MW-6B	ND (0.5)	ND (2)	ND (1)	5.0	
MW-6C	7400	170	7.1	2300	
MW-6D	220	ND (20)	27	ND (10)	
DWAL	0.7	680	100	620	

ND = Nondetectable.

Detection limits are given in parentheses. * Samples obtained June, 1988

Laboratory Tests	** Blows/foot	PID * Reading (ppm)	Depth (ft) Sample	Equipment 8-inch Hollow Stem Auger Elevation 99.4 feet*** Date 6/15/88
	ш	1	0	asphaltic concrete pavement aggregate baserock GRAY SILTY CLAY WITH SAND (CH) stiff, moist
*	21	1	5	GRAY SILTY SAND TRACE CLAY (SM) dense, moist
	14	<1	10-	BROWN SILTY CLAY (CL) stif, moist
	20/5"	<1		BROWN SAND (SP) very dense, saturated
				GRAYISH BROWN CLAYEY SAND (SC) medium dense, saturated BLUISH GRAY SILTY CLAY (CH)
	11	<1	20	very stiff, moist
				bottom of boring 21.5 feet boring cleaned out to 20 feet
			25-	stabilized water level at 13.5 feet on 7/11/88
*PID = photo ionization detector, HNU PI 101 ppm = parts per million			30-	
**S&H Sampler bl to SPT blow co		converted	- 11	
***Reference Elevation (arbitrary datum)		35-		
			40	
Harding Lawson	Associates		Log of E	Boring MW-6A



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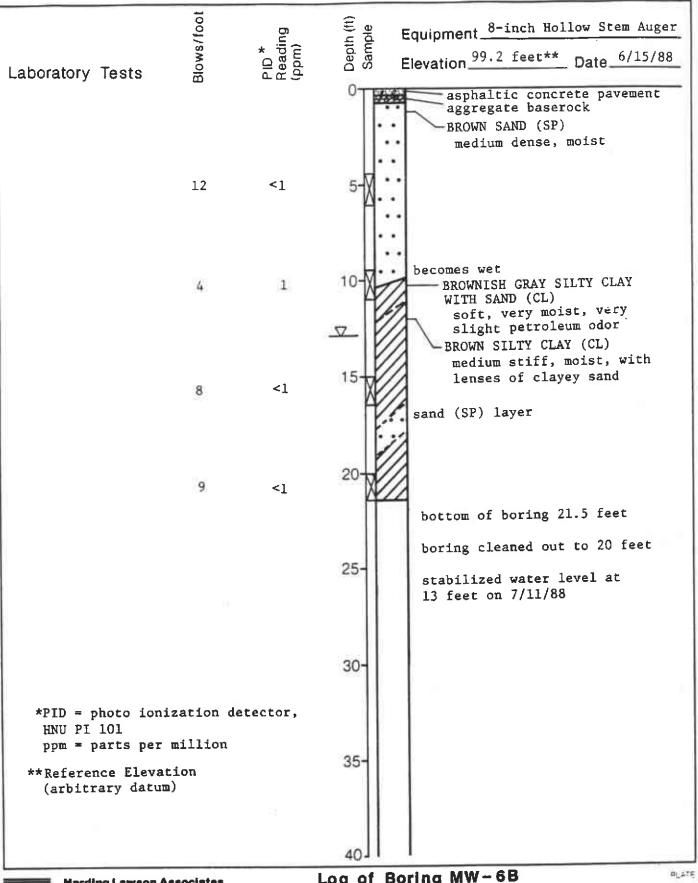
Texaco Station - 62488000195 2225 Telegraph Avenue

JOB NUMBER 2251,080.03 DRAWN RS

Oakland, California

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APPROVED DATE 2/89 DATE





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Log of Boring MW-6B

Texaco Station - 62488000195 2225 Telegraph Avenue

Oakland, California

APPROVED DRAWN JOB NUMBER RS 2251,080.03 2/89

aboratory Tests	Blows/foot	PID * Reading (ppm)	Depth (ft) Sample	Equipment 8-inch Hollow Stem Auger Elevation 100.2 feet** Date 6/15/88
		2 <1	°T	asphaltic concrete pavement aggregate baserock GRAYISH BROWN SILTY CLAY WITH SAND (CL)
	35	1	5-7	very moist, with slight petroleum odor stained gray between 2 and 2-1/2 feet, with petroleum odor
	28	25	10-	becomes brown GRAY MOTTLED BROWN SANDY CLAY (CL) very stiff, moist, very slight petroleum odor BROWN SILTY CLAY TRACE SAND (CL)
	48	60	15-7	very stiff, moist, very slight petroleum odor GRAY CLAYEY SAND (SC) dense, GRAY SILTY SAND (SM) dense, moist
	30	10	20-	GRAYISH BROWN SILTY CLAY (CL) hard, moist bottom of boring 21.5 feet boring cleaned out to 20 feet
			25-	stabilized water level at 14.5 feet on 7/11/88
			30-	
*PID = photo ion: HNU PI 101 ppm = parts per		letector,		
**Reference Eleva (arbitrary datu			35-	
			40	



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Oakland, California

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Laboratory Tests	Blows/foot	PID * Reading (ppm)	Equipment 6-inch Flight Auger By September 199 feet** Date 7/6/88
	22	<1	asphaltic concrete pavement aggregate baserock -YELLOW BROWN CLAY (CL)
	14	5	GRAYISH BROWN CLAY TRACE SAND (CL) very stiff, moist strong petroleum odors between 12.5 and 15.5 feet SAND (SP)
	<u></u>	70	dense, saturated GRAYISH BROWN SILTY CLAY (CL) WITH SAND very stiff, moist no petroleum odors
	12	3	bottom of boring 20 feet
			stabilized water level at 13.5 feet on 7/11/88
*PID = photo ion HNU PI 101	nization (letector,	30-
**Reference Elev (arbitrary dat			35-
(90)			Log of Boring MW-6D

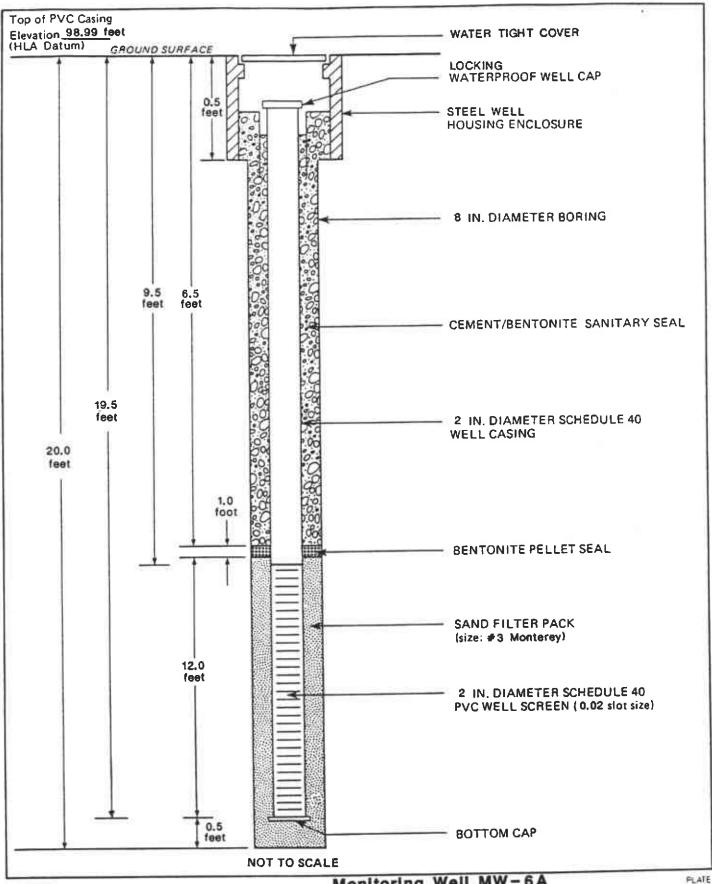


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Texaco Station - 62488000195

2225 Telegraph Avenue

Oakland, California DATE: REVISED JOB NUMBER 2251,080.03 RS RS



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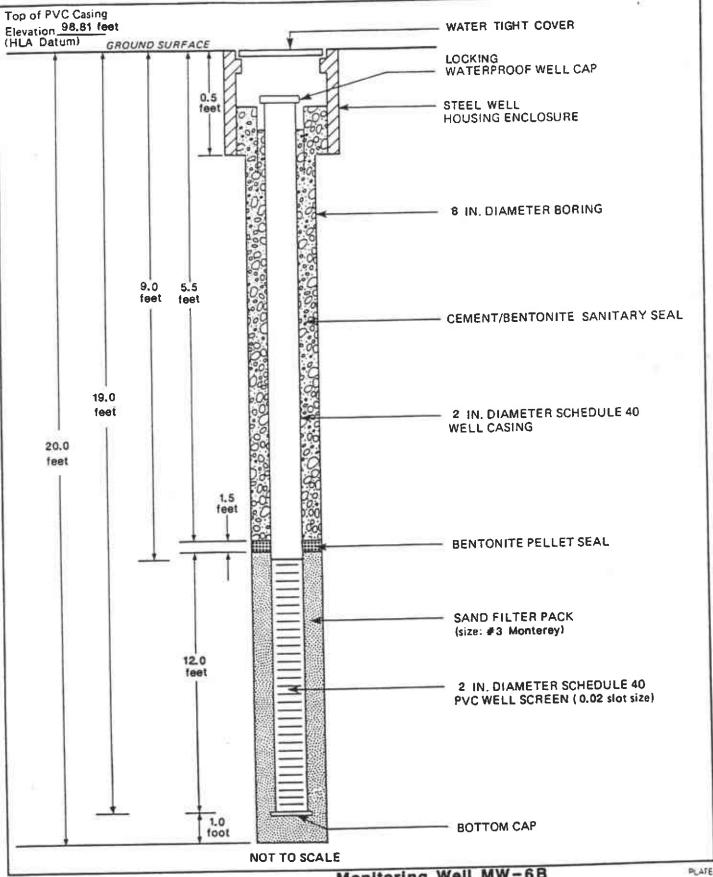
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Engineers, Geologists & Geophysicists

Monitoring Well MW-6A Completion Detail

Texaco Station - 62488000195 2225 Telegraph Avenue Oakland, California

DATE REVISED DATE APPROVED 20 2/89



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Monitoring Well MW-6B Completion Detail

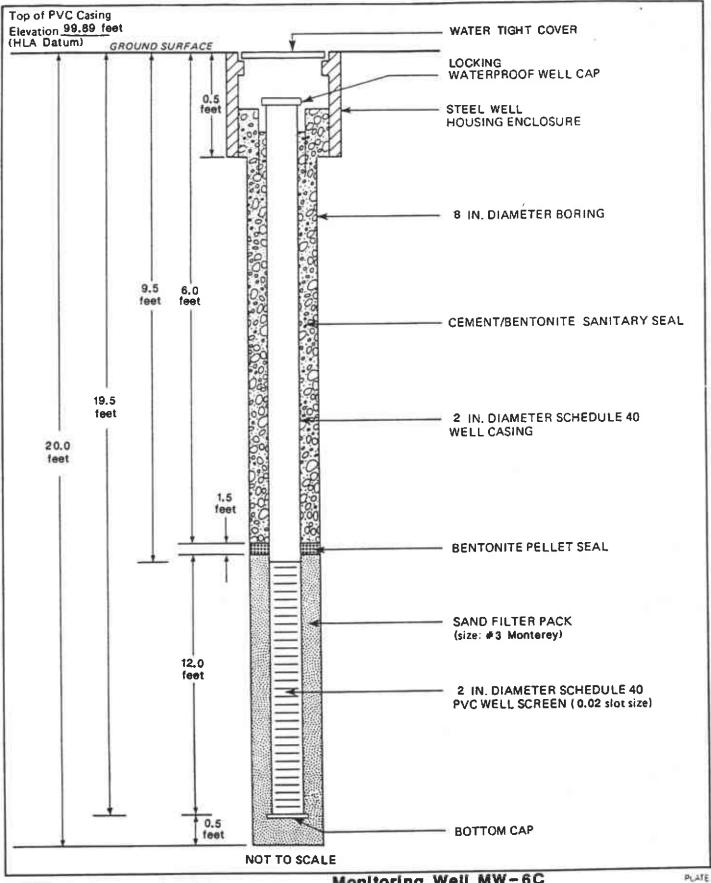
Texaco Station - 62488000195 2225 Telegraph Avenue Oakland, California

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DATE 2/89 REVISED

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FORM GW3





Harding Lawson Associates Engineers, Geologists

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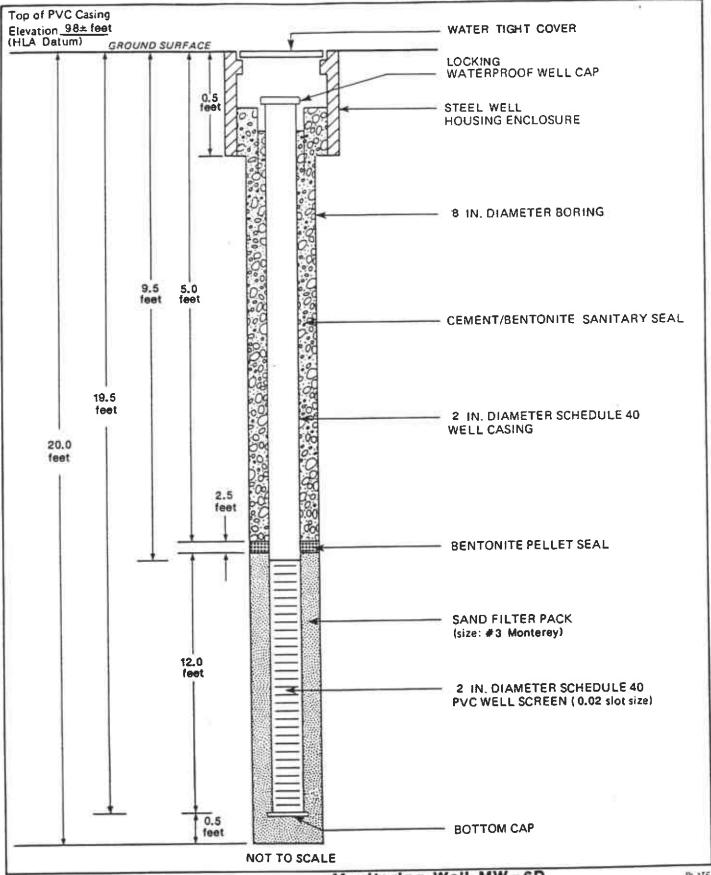
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Monitoring Weil MW-6C Completion Detail Texaco Station - 62488000195 2225 Telegraph Avenue Oakland, California

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Engineers, Geologists & Geophysicists

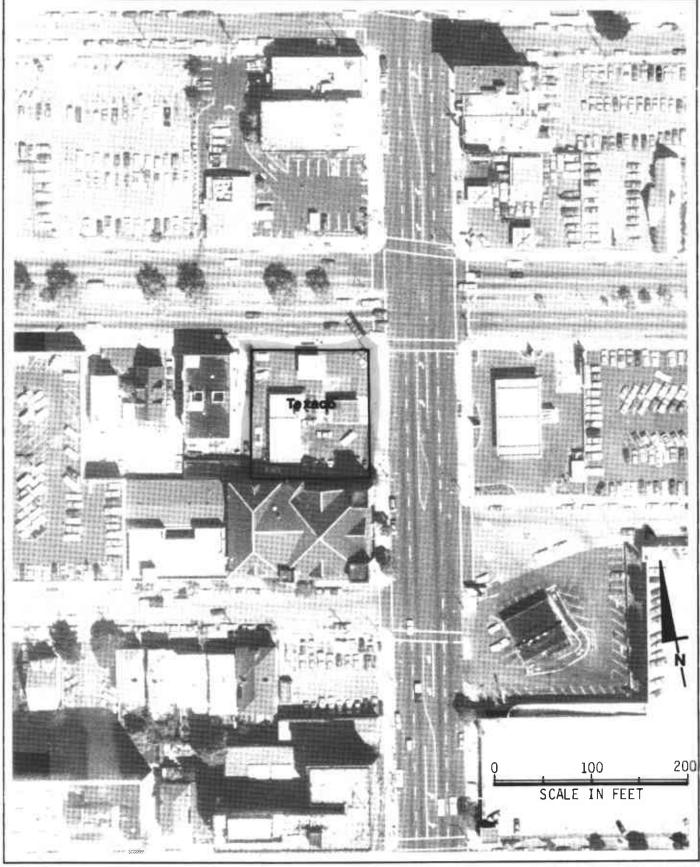
Monitoring Well MW-6D Completion Detail

Texaco Station - 62488000195 2225 Telegraph Avenue Oakland, California

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FORM GW3

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Harding Lawson Associates Engineers and Geoscientists

Aorial Photograph
Former Texaco Service Station
2225 Telegraph Avenue
Oakland, California

6/89

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JOB NUMBER 2251,080.03

APPENDIX C

TRACER RESEARCH REPORT - SOIL-GAS INVESTIGATION

SHALLOW SOIL GAS/GROUNDWATER
INVESTIGATION
AT THE
TEXACO SITES
NORTHERN, CALIFORNIA

SEPTEMBER/OCTOBER 1988

PREPARED FOR:

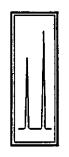
Harding Lawson Associates 1355 Willow Way, Suite 109 Concord, California 94520 SUBMITTED BY:

Tracer Research Corporation



TABLE OF CONTENTS

INTRODUCTION	1
SHALLOW SOIL GAS INVESTIGATION-METHODOLOGY	2
EQUIPMENT AND SAMPLING PROCEDURES	3
ANALYTICAL PROCEDURES	4
QUALITY ASSURANCE/QUALITY CONTROL PROCEDURES	5
APPENDIX A	
CONDENSED DATA	7
APPENDIX B	
CUDOMATOCOAME	0



INTRODUCTION

Several shallow soil gas/groundwater investigations were performed by Tracer Research Corporation (TRC) at the following Texaco gas station sites in the Northern California:

2225 Telegraph Ave., Oakland, CA
2200 E. Twelth St., Oakland, CA
500 Grand Ave., Oakland, CA
495 El Camino Real, San Carlos, CA
800 El Camino Real, San Bruno, CA
196 Main St., Half Moon Bay, CA
595 Munras St., Monterey, CA
1550 Fremont St., Seaside, CA
334 San Antonio Rd., Mountain View, CA
975 Stierlin Rd., Mountain View, CA
5153 Redwood Hwy., Petaluma, CA

The investigation was conducted between September 19 and October 19, 1988 under contract to Harding Lawson Associates. The purpose of the investigation was to determine the possible presence of volatile organic compounds (VOCs) in the subsurface and groundwater as part of an environmental site assement.

For this survey, a total of 126 soil gas samples and 11 groundwater samples were collected and analyzed in the field. The samples were analyzed for the following compounds:

benzene
toluene
ethyl benzene
xylenes
total hydrocarbons (THC)

The compounds in this suite were chosen because of their extensive use at the sites and their suspected presence in the subsurface.



SHALLOW SOIL GAS INVESTIGATION - METHODOLOGY

Soil gas contaminant investigation refers to a method developed by TRC for investigating underground contamination from volatile organic chemicals (VOCs) such as industrial solvents, cleaning fluids and petroleum products by looking for their vapors in the shallow soil gas. The method involves pumping a small amount of soil gas out of the ground through a hollow probe driven into the ground and analyzing the gas for the presence of volatile contaminants. The presence of VOCs in shallow soil gas indicates the observed compounds may either be in the vadose zone near the probe or in groundwater below the probe. The soil gas technology is most effective in mapping low molecular weight petroleum hydrocarbons halogenated solvent chemicals and possessing high vapor pressures and low aqueous solubilities. These compounds readily partition out of the groundwater and into the soil gas as a result of their high gas/liquid partitioning coefficients. Once in the soil gas, VOCs diffuse vertically and horizontally through the soil to the ground surface where they The contamination acts as a dissipate into the atmosphere. source and the above ground atmosphere acts as a sink, and typically a concentration gradient develops between the two. concentration gradient in soil gas between the source and ground surface may be locally distorted by hydrologic and geologic anomalies (e.g. clays, perched water); however, soil gas mapping because distribution of the effective generally remains contamination is usually broader in areal extent than the local geologic barriers and is defined using a large data base. The presence of geologic obstructions on a small scale tends to create anomalies in the soil gas-groundwater correlation, but generally does not obscure the broader areal picture of the contaminant distribution.



EQUIPMENT

Research Corporation utilized a one ton Ford Tracer equipped with one gas field which was van analytical SP4270 computing Physics Spectra chromatograph and two In addition, the van has two built-in gasoline integrators. powered generators which provide the electrical power (110 volts AC) to operate all of the gas chromatographic instruments and field equipment. A specialized hydraulic mechanism consisting of two cylinders and a set of jaws was used to drive and withdraw the sampling probes. A hydraulic hammer was used to assist in driving probes past cobbles and through unusually hard soil.

SAMPLING PROCEDURES

Sampling probes consist of 7-foot lengths of 3/4 inch diameter hollow steel pipe which are fitted with detachable drive points. Soil gas samples were collected after driving the steel probe to a depth between 2 and 15 feet into the ground. The above-ground end of the sampling probes were fitted with a steel reducer and a length of polyethylene tubing leading to a vacuum pump. To adequately purge the volume of air within the probe, 5 to 10 liters of gas were evacuated with a vacuum pump. During the soil gas evacuation, samples were collected in a glass syringe by inserting a syringe needle through a silicone rubber segment in the evacuation line and down into the steel probe. Ten milliliters of gas were collected for immediate analysis in the TRC analytical field van. Soil gas was subsampled (duplicate injections) in volumes ranging from 1 µL to 2 mL, depending on the VOC concentration at any particular location.

Groundwater samples were collected by driving a hollow steel probe to a depth between 5 and 12 into the ground or by direct sampling of an observation or monitoring well. A length of polyethylene tubing was inserted to the bottom of the probe. The

tubing was attached to a peristaltic pump. Approximately 40 mL of water was collected in bottles with teflon lined septum caps so as to exclude air. Any sediment collected with the groundwater was allowed to settle. The water was subsampled (duplicate injections) in volumes ranging from 1 µL to 10 µL.

ANALYTICAL PROCEDURES

A Varian 3300 gas chromatograph equipped with a flame ionization detector (FID) was used for the soil gas analyses. The FID was used to analyze for benzene, toluene, ethyl benzene, xylenes and total hydrocarbons. Xylenes are reported as the total of the three xylene isomers and total hydrocarbons are approximately C4-C9 aliphatic, alicyclic and aromatic compounds. Nitrogen was used as the carrier gas.

Detection limits for the compounds of interest are a function of the injection volume as well as the detector sensitivity for individual compounds. Thus, the detection limit varies with the sample size. Generally, the larger the injection size the greater the sensitivity. However, peaks for compounds interest must be kept within the linear range of the analytical equipment. If any compound has a high concentration, it is necessary to use small injections, and in some cases to dilute the sample to keep it within linear range. This may cause decreased detection limits for other compounds in the analyses. For example, during this investigation, a few of the soil gas samples had high concentrations of benzene; toluene and xylenes. To bring the peak for these compounds within linear range, it was This had the effect of necessary to make small injections. decreasing the detection limits for ethyl benzene in these samples.

The detection limits range down to 0.08 kg/L for compounds such as benzene and toluene depending on the conditions of the



measurement, in particular, the sample size. If any component being analyzed is not detected, the detection limit for that compound in that analysis is given as a "less than" value (e.g. $(0.08~\mu g/L)$. Detection limits obtained from GC analyses are calculated from the current response factor, the sample size, and the estimated minimum peak size (area) that would have been visible under the conditions of the measurement.

QUALITY ASSURANCE/QUALITY CONTROL PROCEDURES

Tracer Research Corporation's normal quality assurance procedures were followed in order to prevent any cross-contamination of soil gas samples.

- . Steel probes are used only once during the day and then washed with high pressure soap and hot water spray or steam-cleaned to eliminate the possibility of cross-contamination. Enough probes are carried on each van to avoid the need to reuse any during the day.
- Probe adaptors (steel reducer and tubing) are used once during the course of the day and cleaned at the end of each working day by baking in the GC oven. The tubing is replaced periodically as needed during the job to insure cleanliness and good fit.
- Silicone tubing (connecting the adaptor to the vacuum pump) is replaced as needed to insure proper sealing around the syringe needle. This tubing does not directly contact soil gas samples.
- . Glass syringes are usually used for only one sample per day and are washed and baked out at night. If they must be used twice, they are purged with carrier gas (nitrogen) and baked out between probe samplings.
- Septa through which soil gas samples are injected into the chromatograph are replaced on a daily basis to prevent possible gas leaks from the chromatographic column.
- Analytical instruments are calibrated each day by the use of chemical standards prepared in water by serial dilution from commercially available pure chemicals. Calibration checks are also run after approximately every five soil gas sampling locations.



- 2 cc subsampling syringes are checked for contamination prior to sampling each day by injecting nitrogen carrier gas into the gas chromatograph.
- Prior to sampling each day, system blanks are run to check the sampling apparatus (probe, adaptor, 10 cc syringe) for contamination by drawing ambient air from above ground through the system and comparing the analysis to a concurrently sampled air analysis.
- All sampling and 2 cc subsampling syringes are decontaminated each day and no such equipment is reused before being decontaminated. Microliter size subsampling syringes are reused only after a nitrogen carrier gas blank is run to insure it is not contaminated by the previous sample.
- . Soil gas pumping is monitored by a vacuum gauge to insure that an adequate gas flow from the vadose zone is maintained. A negative pressure (vacuum) of 2 in. Hg less than the maximum capacity of the pump (evacuation rate >0.02 cfm) usually indicates that a reliable gas sample cannot be obtained because the soil has a very low air permeability.

Tracer Research Corporation

APPENDIX A: CONDENSED DATA

HARDING LAWSON ASSOCIATES/2225 TELEGRAPH AVENUE/OAKLAND, CALIFORNIA

Sample	Depth Date	Benzene (ug/1)	Toluene (ug/l)	Ethyl Benzene (ug/l)	Xylenes (ug/l)	Total Hydroc. (ug/l)	_
Air S6-02 . S6-03	09/19 5' 09/19 12' 09/19	<0.7 <0.7 10	<0.8 <0.8 4	<0.8 <0.8	<0.8 <0.8 2,800	<0.7 <0.7 6,100	
S6-04 WS-05 S6-06	13' 09/19 12' 09/19 13' 09/19	<0.7 <75 <0.7	<0.8 <76 <0.8	<0.0 <77 <0.8	140 <77 <0.8	780 <75 <0.7	
Air	09/19	<0.7	<0.8	<0.8	<0.8	<0.7	

Tracer Research Corporation

Notations: I interference with adjacent peaks NA not analyzed Analyzed by K. Tolman
Checked by R. Sheldrake
Proofed by X. Mixlandu



Tracer Research Corporation

APPENDIX B: CHROMATOGRAMS

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TOTAL	100.		5222974			

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9209550

TOTAL

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APPENDIX D

SOIL SAMPLE CHEMICAL TEST DATA



October 24, 1988

Harding Lawson Associates 1355 Willow Way, Suite 109 Concord, CA 94520

Attention: Mr. Randy Stone

Subject: Report of Data - Case Number 2382

Dear Mr. Stone:

The technical staff at CHEMWEST is pleased to provide our report for the analyses you requested: Total Petroleum Hydrocarbons, Purgeable (gasoline) -DHS Method, LUFT Field Manual; and BTEX - EPA Method 602.

Eight soil samples for Project Texaco #6; Telegraph Oakland, Project Number 02251,080.03 were received October 6, 1988 in good condition. Results of the analyses, along with the analytical methodology and appropriate reporting limits, are presented on the following pages.

Thank you for choosing CHEMWEST Laboratories. Should you have questions concerning this data report or the analytical methods employed, please do not hesitate to contact Toni Weeks, our Technical Service Representative, or your project manager. We hope that you will consider CHEMWEST Laboratories for your future analytical support and service requirements.

Sincerely,

Jil B. Henes, Ph.D.

Vice President of Technical Services

and

Kirk Poćan

Project Manager

KP:bw

cc: Joel Bird, President File

Phone (916) 923-0840

FAX (916) 923-1938

ANALYTICAL METHODOLOGY

BTEX (Benzene, Toluene, Ethyl Benzene, and Xylenes) by Purge & Trap and GC-PID

WATER - Method 602 or 8020

A 5 ml sample volume, or 5 ml of a suitable dilution, is purged on a suitable purge and trap system with helium. The purged sample is analyzed on a Gas Chromatograph equipped with a Photoionization Detector (PID). A packed column is used to separate the compounds.

SOIL - Method 8020

A 10 gram, or other appropriate aliquot of soil, is weighed into a clean VOA vial. Soils received in brass core tubes are sampled by discarding 2-5 centimeters of soil from each end of the tubes (this is done to reduce the possibility of analyzing a portion of soil that has been exposed to sampling technique contamination). Equal aliquots of soil are then removed from each end of the tube and combined in the VOA vial. Soil in jars or bags is aliquoted using a similar technique, which discards exposed sample surfaces. A 10 ml, or other appropriate volume of methanol, is added to the soil and the soil is shaken with the solvent. 100 ul of the extract, or a reduced aliquot or volume of a suitable dilution, is injected into 5 ml of laboratory blank water and analyzed by the same technique used for water samples.

ANALYTICAL METHODOLOGY

Total Petroleum Hydrocarbons by Purge & Trap and GC-FID

WATER - DHS Method - Luft Field Manual

A 5 ml sample volume, or 5 ml of a suitable dilution, is purged on a suitable purge and trap system with helium. The purged sample is analyzed on a Gas Chromatograph equipped with a Flame Ionization Detector (FID). A packed column is used to separate the compounds.

SOIL - DHS Method - Luft Field Manual

A 10 gram, or other appropriate aliquot of soil, is weighed into a clean VOA vial. Soils received in brass core tubes are sampled by discarding 2-5 centimeters of soil from each end of the tubes (this is done to reduce the possibility of analyzing a portion of soil that has been exposed to sampling technique contamination). Equal aliquots of soil are then removed from each end of the tube and combined in the VOA vial. Soil in jars or bags is aliquoted using a similar technique, which discards exposed sample surfaces. A 10 ml, or other appropriate volume of methanol, is added to the soil and the soil is shaken with the solvent. 100 ul of the extract, or a reduced aliquot or volume of a suitable dilution, is injected into 5 ml of laboratory blank water and analyzed by the same technique used for water samples.

Client I.D. : MW-6E enso Date Analyzed: 10/09/88 Date Extracted: 10/09/88

CHEMWEST I.D.: 2382-1

Matrix : Soil

Compound	Amount Detected (mg/Kg)	RL (mg/Kg)
Benzene	BRL	- 0.05
Toluene	BRL	0.1
Ethyl Benzene	BRL	0.2
Total-Xylenes (1)	BRL	0.1
Total Petroleum Hydrocarbon (Purgeable)	n BRL	10

Surrogate	% Recovery	Acceptance Window
ortho-Chlorotoluene	72%	50-150%

BRL: Below Reporting Limit.

RL: Reporting Limit.

Client I.D. : Bl 13.0 Date Analyzed : 10/16/88 Date Extracted: 10/09/88 CHEMWEST I.D.: 2382-2

Matrix : Soil

	Amount etected (mg/Kg)	RL (mg/Kg)
Benzene	BRL	5
Toluene	16	10
Ethyl Benzene	19	10
Total-Xylenes (1)	41	10
Total Petroleum Hydrocarbon (Purgeable)	2000	1000

Surrogate	% Recovery	Acceptance Window
ortho-Chlorotoluene	85%	50-150%

BRL: Below Reporting Limit.

RL: Reporting Limit.

Client I.D. : B1 8.0 Date Analyzed: 10/14/88 Date Extracted: 10/09/88 CHEMWEST I.D.: 2382-3 Matrix : Soil

Compound	Amount Detected (mg/Kg)	RL (mg/Kg)
Benzene	0.05	0.05
Toluene	BRL	0.1
Ethyl Benzene	BRL	0.2
Total-Xylenes (1)	BRL	0.1
Total Petroleum Hydrocarbon (Purgeable)	n BRL	10

Surrogate	% Recovery	Acceptance Window
ortho-Chlorotoluene	65%	50-150%

BRL: Below Reporting Limit.

RL: Reporting Limit.

Client I.D. : B2 7.0-7.5 Date Analyzed: 10/09/88 Date Extracted: 10/09/88

CHEMWEST I.D.: 2382-4 Matrix : Soil

	Amount etected (mg/Kg)	RL (mg/Kg)
Benzene	BRL	0.05
Toluene	BRL	0.1
Ethyl Benzene	BRL	0.2
Total-Xylenes (1)	BRL	0.1
Total Petroleum Hydrocarbon (Purgeable)	BRL	10

Surrogate	% Recovery	Acceptance Window
ortho-Chlorotoluene	67%	50-150%

BRL: Below Reporting Limit.

RL: Reporting Limit.

Client I.D. : B2 13.5-14.0

CHEMWEST I.D.: 2382-5

Date Analyzed: 10/14/88

Matrix : Soil

Date Extracted: 10/09/88

Compound	Amount Detected (mg/Kg)	RL (mg/Kg)
Benzene	BRL	0.05
Toluene	BRL	0.1
Ethyl Benzene	BRL	0.2
Total-Xylenes (1)	BRL	0.1
Total Petroleum Hydrocarbon (Purgeable)	n BRL	10

Surrogate	Recovery	Acceptance Window
ortho-Chlorotoluene	66%	50-150%

BRL: Below Reporting Limit.

RL: Reporting Limit.

Client I.D. : B3 7.0-7.5 Date Analyzed : 10/09/88 CHEMWEST I.D.: 2382-6

Date Extracted: 10/09/88

Compound	Amount Detected (mg/Kg)	RL (mg/Kg)
Benzene	0.06	0.05
Toluene	BRL	0.1
Ethyl Benzene	BRL	0.2
Total-Xylenes (1)	BRL	0.1
Total Petroleum Hydrocarbon (Purgeable)	BRL	10

Surrogate	% Recovery	Acceptance Window
ortho-Chlorotoluene	66%	50-150%

BRL: Below Reporting Limit.

RL: Reporting Limit.

Client I.D. : B3 13.5-14.0

CHEMWEST I.D.: 2382-7

Date Analyzed: 10/14/88

Matrix : Soil

Date Extracted: 10/09/88

	Amount Detected (mg/Kg)	RL (mg/Kg)
Benzene	40	25
Toluene	390	50
Ethyl Benzene	84	50
Total-Xylenes (1)	370	50
Total Petroleum Hydrocarbon (Purgeable)	11000	5000

Surrogate	% Recovery	Acceptance Window
ortho-Chlorotoluene	77%	50-150%

BRL: Below Reporting Limit.

RL: Reporting Limit.

Client I.D. : MW 6F Date Analyzed: 10/09/88 Date Extracted: 10/09/88 CHEMWEST I.D.: 2382-8

Matrix : Soil

Compound	Amount Detected (mg/Kg)	RL (mg/Kg)
Benzene	BRL	0.05
Toluene	BRL	0.1
Ethyl Benzene	BRL	0.2
Total-Xylenes (1)	BRL	0.1
Total Petroleum Hydrocarbon (Purgeable)	n BRL	10

Surrogate	% Recovery	Acceptance Window
ortho-Chlorotoluene	73%	50-150%

BRL: Below Reporting Limit. RL: Reporting Limit.

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Sacramento, California 95834	Compl. Date
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November 30, 1988

Harding Lawson 1355 Willow Way, Suite 109 Concord, CA 94520

Attention: Mr. Randy Stone

Subject: Report of Data - Case Number 2708

Dear Mr. Stone:

The technical staff at CHEMWEST is pleased to provide our report for the analyses you requested: Total Petroleum Hydrocarbons, Purgeable (gasoline) - DHS Method, LUFT Field Manual; and BTEX - EPA Method 602.

Four soil samples for Project Texaco Assessment #6, Project Number 02251,080.03 were received November 18, 1988 in good condition. Results of the analyses, along with the analytical methodology and appropriate reporting limits, are presented on the following pages.

Thank you for choosing CHEMWEST Laboratories. Should you have questions concerning this data report or the analytical methods employed, please do not hesitate to contact Toni Weeks, our Technical Service Representative, or your project manager. We hope that you will consider CHEMWEST Laboratories for your future analytical support and service requirements.

Sincerely,

Jill B. Henes, Ph.D.

Vice President of Technical Services

and

Kirk Pocan

Project Manager

KP:bw

cc: Joel Bird, President File

ANALYTICAL METHODOLOGY

BTEX (Benzene, Toluene, Ethyl Benzene, and Xylenes) by Purge & Trap and GC-PID

WATER - Method 602 or 8020

A 5 ml sample volume, or 5 ml of a suitable dilution, is purged on a suitable purge and trap system with helium. The purged sample is analyzed on a Gas Chromatograph equipped with a Photoionization Detector (PID). A packed column is used to separate the compounds.

SOIL - Method 8020

A 10 gram, or other appropriate aliquot of soil, is weighed into a clean VOA vial. Soils received in brass core tubes are sampled by discarding 2-5 centimeters of soil from each end of the tubes (this is done to reduce the possibility of analyzing a portion of soil that has been exposed to sampling technique contamination). Equal aliquots of soil are then removed from each end of the tube and combined in the VOA vial. Soil in jars or bags is aliquoted using a similar technique, which discards exposed sample surfaces. A 10 ml, or other appropriate volume of methanol, is added to the soil and the soil is shaken with the solvent. 100 ul of the extract, or a reduced aliquot or volume of a suitable dilution, is injected into 5 ml of laboratory blank water and analyzed by the same technique used for water samples.

ANALYTICAL METHODOLOGY

Total Petroleum Hydrocarbons by Purge & Trap and GC-FID

WATER - DHS Method - Luft Field Manual

A 5 ml sample volume, or 5 ml of a suitable dilution, is purged on a suitable purge and trap system with helium. The purged sample is analyzed on a Gas Chromatograph equipped with a Flame Ionization Detector (FID). A packed column is used to separate the compounds.

SOIL - DHS Method - Luft Field Manual

A 10 gram, or other appropriate aliquot of soil, is weighed into a clean VOA vial. Soils received in brass core tubes are sampled by discarding 2-5 centimeters of soil from each end of the tubes (this is done to reduce the possibility of analyzing a portion of soil that has been exposed to sampling technique contamination). Equal aliquots of soil are then removed from each end of the tube and combined in the VOA vial. Soil in jars or bags is aliquoted using a similar technique, which discards exposed sample surfaces. A 10 ml, or other appropriate volume of methanol, is added to the soil and the soil is shaken with the solvent. 100 ul of the extract, or a reduced aliquot or volume of a suitable dilution, is injected into 5 ml of laboratory blank water and analyzed by the same technique used for water samples.

Client I.D.: MW-G

CHEMWEST I.D.: 2708-1

Date Analyzed: 11/23/88 Date Extracted: 11/21/88 Matrix : Soil

Compound	Amount Detected (mg/Kg)	RL (mg/Kg)
Benzene	BRL	0.05
Toluene	BRL	0.1
Ethyl Benzene	BRL	0.2
Total-Xylenes (1)	BRL	0.1
Total Petroleum Hydrocarbon (Purgeable)	5.2	10

Surrogate	% Recovery	Acceptance Window
ortho-Chlorotoluene	93%	50-150%

BRL: Below Reporting Limit.

RL: Reporting Limit.

Client I.D.: MW-H

CHEMWEST I.D.: 2708-2

Matrix : Soil

Date Analyzed: 11/23/88 Date Extracted: 11/21/88

Compound	Amount Detected (mg/Kg)	RL (mg/Kg)
Benzene	11	0.5
Toluene	3.2	1
Ethyl Benzene	8.8	2
Total-Xylenes (1)	19	1
Total Petroleum Hydrocarbon (Purgeable)	1000	495
		

Surrogate	% Recovery	Acceptance Window
ortho-Chlorotoluene	147%	50-150%

BRL: Below Reporting Limit.

RL: Reporting Limit.

Client I.D.: MW-I

CHEMWEST I.D.: 2708-3

Date Analyzed: 11/23/88

Matrix : Soil

Date Extracted: 11/21/88

Compound	Amount Detected (mg/Kg)	RL (mg/Kg)
Benzene	BRL	0.05
Toluene	BRL	0.1
Ethyl Benzene	BRL	0.2
Total-Xylenes (1)	BRL	0.1
Total Petroleum Hydrocarbon (Purgeable)	BRL	10

Surrogate	% Recovery	Acceptance Window
ortho-Chlorotoluene	70%	50-150%

BRL: Below Reporting Limit.

RL: Reporting Limit.

Client I.D.: B-4

CHEMWEST I.D.: 2708-4

Date Analyzed: 11/23/88

Matrix : Soil

Date Extracted: 11/21/88

Compound	Amount Detected (mg/Kg)	RL (mg/Kg)
Benzene	BRL	0.05
Toluene	BRL	0.1
Ethyl Benzene	BRL	0.2
Total-Xylenes (1)	BRL	0.1
Total Petroleum Hydrocarbo (Purgeable)	n BRL	10

Surrogate	% Recovery	Acceptance Window
ortho-Chlorotoluene	80%	50-150%

BRL: Below Reporting Limit.

RL: Reporting Limit.

CHEM WEST ANALYTICAL LABORATORIES, INC. 600 West North Marker Bivd. Secremento. California 98834 (916) 923-0840 RAX (916) 923-1938 CLIENT. Harding Sauran (1994-1995) Concard, CA 941520 Concard, CA 941520 ANALYSIS. Jaw and samples her'd ingless Change for TPH ENTRY GC-FID & BTEX. Seven day T/AH Sample ID 13.5'-140' "Illustration of Contact of		0700
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Concard, CA 94620 Concard, CA 94620 PONO Consert Standy Starne Consert Standy Starne Phone 9(4/5) 1687-9660 ANALYSIS: Four sail samples rec'd under Chain of contry in 6" Mital case tuber to be analyzed fast TPH EXTRY GC-FID & BTEX. Samen day 1/H Somele In Depth Date Analysis matrix Contrinses 2708-1 MW-G 13.5: 140' ulmissistared, BTEX/INSOI 6 Contrins -2 MW-H 13.5: 140' ulmissistared, BTEX/INSOI 6 Contrins -4 B -4 13.5: 140' ulmissistared, BTEX/INSOI 6 Contrins AMENDED Rex Randy Stone of HIA on 11/21/88 @ 1620 Inter Change the analyses to BTEX TEH only on all 4 samples. The ulfills of the samples of BTEX TEH only on all 4 samples.	Unit of the second	Project Name: TEXACO ASSESSMENT THE
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Harding Lawson association 1355 Willow Way, Suite 109 Concord, California 94520 415/687-9660 Telecopy: 415/687-9673

Lab:	Chism	WEST
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				Samplers: DA	wid R. Huser	ANALYSIS REQUESTED
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APPENDIX E

WATER SAMPLE CHEMICAL TEST DATA



November 10, 1988

Harding Lawson Associates 1355 Willow Way, Suite 109 Concord, CA 94520

Attention: Mr. Randy Stone

Subject: Report of Data - Case Number 2535

Dear Mr. Stone:

The technical staff at CHEMWEST is pleased to provide our report for the analysis you requested: BTEX - EPA Method 602.

One water sample for Project Texaco #6; Telegraph, Project Number 02251,080.03 was received October 26, 1988 in good condition. Results of the analysis, along with the analytical methodology and appropriate reporting limits, are presented on the following pages.

Thank you for choosing CHEMWEST Laboratories. Should you have questions concerning this data report or the analytical methods employed, please do not hesitate to contact Toni Weeks, our Technical Service Representative, or your project manager. We hope that you will consider CHEMWEST Laboratories for your future analytical support and service requirements.

Sincerely,

J:113Henes

Jill B. Henes, Ph.D. Vice President of Technical Services

and

Kirk Pocan

Project Manager

KP:bw

cc: Joel Bird, President File

ANALYTICAL METHODOLOGY

BTEX (Benzene, Toluene, Ethyl Benzene, and Xylenes) by Purge & Trap and GC-PID

WATER - Method 602 or 8020

A 5 ml sample volume, or 5 ml of a suitable dilution, is purged on a suitable purge and trap system with helium. The purged sample is analyzed on a Gas Chromatograph equipped with a Photoionization Detector (PID). A packed column is used to separate the compounds.

SOIL - Method 8020

A 10 gram, or other appropriate aliquot of soil, is weighed into a clean VOA vial. Soils received in brass core tubes are sampled by discarding 2-5 centimeters of soil from each end of the tubes (this is done to reduce the possibility of analyzing a portion of soil that has been exposed to sampling technique contamination). Equal aliquots of soil are then removed from each end of the tube and combined in the VOA vial. Soil in jars or bags is aliquoted using a similar technique, which discards exposed sample surfaces. A 10 ml, or other appropriate volume of methanol, is added to the soil and the soil is shaken with the solvent. 100 ul of the extract, or a reduced aliquot or volume of a suitable dilution, is injected into 5 ml of laboratory blank water and analyzed by the same technique used for water samples.

CHEMWEST ANALYTICAL LABORATORIES BENZENE, TOLUENE, ETHYL BENZENE, XYLENES

Client I.D.: MW-6F-1&2 Date Analyzed: 11/01/88

CHEMWEST I.D.: 2535-1 Matrix : Water

Compound	Amount Detected (ug/L)	RL (ug/L)
Benzene	BRL	0.5
Toluene	BRL	1
Ethyl Benzene	BRL	2
Total-Xylenes (1)	2.4	1

Surrogate	% Recovery	Acceptance Window
ortho-Chlorotoluene	57%	50-150%

BRL: Below Reporting Limit.

RL: Reporting Limit.

CHEM WEST ANALYTICAL LABORATORIES,INC 600 West North Market Blvd. Sacramento, California 95834 (916) 923-0840 FAX (916) 923-1938	Order No. 2535 Date Rec'd 10/20/88@190 Compl. Date Section VIII Docum
1355 Willaw Long Switch 109 Project Concord, CA 94520 P.O. NO. Contact. Phone 1	Name: 2251,080.03 No. Texaco #6, Telegration 415) 1087-96100
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November 11, 1988

Harding Lawson Associates 1355 Willow Way, Suite 109 Concord, CA 94520

Attention: Mr. Randy Stone

Subject: Report of Data - Case Number 2498

Dear Mr. Stone:

The technical staff at CHEMWEST is pleased to provide our report for the analysis you requested: BTEX - EPA Method 602.

Five water samples for Project Texaco #6; 2225 Telegraph, Project Number 2251,080.03 were received October 21, 1988 in good condition. Results of the analysis, along with the analytical methodology and appropriate reporting limits, are presented on the following pages.

Thank you for choosing CHEMWEST Laboratories. Should you have questions concerning this data report or the analytical methods employed, please do not hesitate to contact Toni Weeks, our Technical Service Representative, or your project manager. We hope that you will consider CHEMWEST Laboratories for your future analytical support and service requirements.

Sincerely,

Jill B. Henes, Ph.D.

1:11 Bleves

Vice President of Technical Services

and

Kirk Pocan

Project Manager

KP:bw

cc: Joel Bird, President File

ANALYTICAL METHODOLOGY

BTEX (Benzene, Toluene, Ethyl Benzene, and Xylenes) by Purge & Trap and GC-PID

WATER - Method 602 or 8020

A 5 ml sample volume, or 5 ml of a suitable dilution, is purged on a suitable purge and trap system with helium. The purged sample is analyzed on a Gas Chromatograph equipped with a Photoionization Detector (PID). A packed column is used to separate the compounds.

SOIL - Method 8020

A 10 gram, or other appropriate aliquot of soil, is weighed into a clean VOA vial. Soils received in brass core tubes are sampled by discarding 2-5 centimeters of soil from each end of the tubes (this is done to reduce the possibility of analyzing a portion of soil that has been exposed to sampling technique contamination). Equal aliquots of soil are then removed from each end of the tube and combined in the VOA vial. Soil in jars or bags is aliquoted using a similar technique, which discards exposed sample surfaces. A 10 ml, or other appropriate volume of methanol, is added to the soil and the soil is shaken with the solvent. 100 ul of the extract, or a reduced aliquot or volume of a suitable dilution, is injected into 5 ml of laboratory blank water and analyzed by the same technique used for water samples.

Client I.D.: MW-6A
Date Analyzed: 10/25/88

CHEMWEST I.D.: 2498-1
Matrix : Water

Compound	Amount Detected (ug/L)	RL (ug/L)
Benzene	0.6	0.5
Toluene	BRL	1
Ethyl Benzene	BRL	2
Total-Xylenes (1)	BRL	1

Surrogate	% Recovery	Acceptance Window	
ortho-Chlorotoluene	66%	50-150%	
			

BRL: Below Reporting Limit.

RL: Reporting Limit.

Client I.D.: MW-6B Date Analyzed: 10/25/88 CHEMWEST I.D.: 2498-2

Matrix

: Water

Compound	Amount Detected (ug/L)	RL (ug/L)	
Benzene	4.1	0.5	
Toluene	BRL	1	
Ethyl Benzene	2.5	2	
Total-Xylenes (1)	BRL	1	

Surrogate	% Recovery	Acceptance Window	
ortho-Chlorotoluene	58%	50-150%	

BRL: Below Reporting Limit.

RL: Reporting Limit.

Client I.D.: MW-6C Date Analyzed: 10/27/88 CHEMWEST I.D.: 2498-3

Matrix

: Water

Compound	Amount Detected (ug/L)	RL (ug/L)
Benzene	9500	. 50
Toluene	170	100
Ethyl Benzene	65	2
Total-Xylenes (1)	850	1

Surrogate	% Recovery	Acceptance Window	
ortho-Chlorotoluene	94%	50-150%	

BRL: Below Reporting Limit.

RL: Reporting Limit.

Client I.D.: MW-6D Date Analyzed: 10/25/88 CHEMWEST I.D.: 2498-4
Matrix : Water

Compound	Amount Detected (ug/L)	RL (ug/L)
Benzene	710	5
Toluene	22	10
Ethyl Benzene	74	20
Total-Xylenes (1)	110	10

Surrogate	% Recovery	Acceptance Window	
ortho-Chlorotoluene	69%	50-150%	

BRL: Below Reporting Limit.

RL: Reporting Limit.

Client I.D.: MW-6E Date Analyzed: 10/25/88 CHEMWEST I.D.: 2498-5

Matrix : Water

Compound	Amount Detected (ug/L)	RL (ug/L)	
Benzene	1.1	0.5	
Toluene	BRL	1	
Ethyl Benzene	BRL	2	
Total-Xylenes (1)	3.4	1	

Surrogate	% Recovery	Acceptance Window	
ortho-Chlorotoluene	74%	50-150%	

BRL: Below Reporting Limit.

RL: Reporting Limit.

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METHOD OF SHIPMENT

DEC 29 1988



December 23, 1988

Harding Lawson Associates 1355 Willow Way, Suite 109 Concord, CA 94520

Attention: Mr. Randy Stone

Subject: Report of Data - Case Number 2849

Dear Mr. Stone:

The technical staff at CHEMWEST is pleased to provide our report for the analysis you requested: BTEX - EPA Method 602.

Three water samples for Project Texaco Assessment #6 Project # 02251,080.03 were received December 9, 1988 in good condition. Results of the analysis, along with the analytical methodology and appropriate reporting limits, are presented on the following page(s).

Thank you for choosing CHEMWEST Laboratories. Should you have questions concerning this data report or the analytical methods employed, please do not hesitate to contact Toni Weeks, our Technical Service Representative or your project manager. We hope that you will consider CHEMWEST Laboratories for your future analytical support and service requirements.

Sincerely,

Jill B. Henes, Ph.D.

Vice President of Technical Services

and

Kirk Pocan

Project Manager

KP:pjg

cc: Joel Bird, President

File

ANALYTICAL METHODOLOGY

BTEX (Benzene, Toluene, Ethyl Benzene, and Xylenes) by Purge & Trap and GC-PID

WATER - Method 602 or 8020

A 5 ml sample volume, or 5 ml of a suitable dilution, is purged on a suitable purge and trap system with helium. The purged sample is analyzed on a Gas Chromatograph equipped with a Photoionization Detector (PID). A packed column is used to separate the compounds.

SOIL - Method 8020

A 10 gram, or other appropriate aliquot of soil, is weighed into a clean VOA vial. Soils received in brass core tubes are sampled by discarding 2-5 centimeters of soil from each end of the tubes (this is done to reduce the possibility of analyzing a portion of soil that has been exposed to sampling technique contamination). Equal aliquots of soil are then removed from each end of the tube and combined in the VOA vial. Soil in jars or bags is aliquoted using a similar technique, which discards exposed sample surfaces. A 10 ml, or other appropriate volume of methanol, is added to the soil and the soil is shaken with the solvent. 100 ul of the extract, or a reduced aliquot or volume of a suitable dilution, is injected into 5 ml of laboratory blank water and analyzed by the same technique used for water samples.

Client I.D.: MW6G1/MW6G2 Date(s) Analyzed: 12/14/88 CHEMWEST I.D.: 2849-1 Matrix : Water

Compound	Amount Detected (ug/L)	RL (ug/L)
Benzene	BRL	0.5
Toluene	BRL	1
Ethyl Benzene	BRL	2
Total-Xylenes (1)	BRL	1

urrogate	% Recovery	Acceptance Window
ortho-Chlorotoluene	100%	50-150%

BRL: Below Reporting Limit.

RL: Reporting Limit.

Client I.D.: MW6H1/MW6H2 Date(s) Analyzed: 12/20/88

CHEMWEST I.D.: 2849-2 Matrix : Water

Compound	Amount Detected (ug/L)	RL (ug/L)
Benzene	1200	25
Toluene	. 110	10
Ethyl Benzene	320	20
Total-Xylenes (1)	220	10

Surrogate	Recovery	Acceptance Window
ortho-Chlorotoluene	95%	50-150%

BRL: Below Reporting Limit.

RL: Reporting Limit.

APPENDIX F
METHODS OF SLUG TEST ANALYSIS

APPENDIX F

METHODS OF SLUG TEST ANALYSIS

The slug tests of the confined strata were analyzed according to the method of Cooper et al. (1967). The water level recovery data were matched to a recovery-type curve, and the value of time on the data graph where $Tt/r_c^2 = 1$ on the type curve was noted. Here T is transmissivity (L^2/t) , t is time after slug withdrawal or injection, and r_c is the equivalent interior radius of the well casing (L).

Transmissivity was then estimated from

$$T = \frac{1.0 \text{ r}_{c}^{2}}{t}$$

Where t is the value of time on the data graph where $Tt/r_c^2 = 1$ on the type curve. The hydraulic conductivity, K, of the stratum tested was estimated from

$$K = \frac{T}{b}$$

where b is the thickness of the stratum.

The slug tests of the unconfined strata were analyzed according to the method of Bouwer and Rice (1976). The hydraulic conductivity is estimated directly from the relationship

$$K = \frac{r_c^2 \ln (R_e/r_w)}{2L} \frac{1}{t} \ln \frac{h_0}{h_1}$$

where R_e is the radial distance over which the head change h is dissipated in the flow system, r_w is the radial distance between the undisturbed stratum (aquifer) and the well center, L is the height of the portion of well through which water enters, h_0 is the initial vertical distance between the water level in the well and the equilibrium water table (hydraulic head) in the tested stratum immediately after slug withdrawal or injection, and h_t is that vertical distance at some time after slug withdrawal or injection.

The term $1/t \ln h_0$ is evaluated from the early straight-line portion of a graph of h_t

the logarithm of h_t as a function of time. For the case where D, the vertical distance from the water table to the bottom of the permeable stratum, is equal to H, the vertical distance from the water table to the bottom of the interval through which water enters the well, the term $\ln R_e/r_w$ is evaluated from the empirical relation

$$\ln R_e/r_w = \left(\frac{1.1}{\ln (H/r_w)} + \frac{C}{L/r_w}\right)^{-1}$$

Here C is an empirical coefficient that depends on L/r_w . Bouwer and Rice (1976) provide a graphical representation of the dependency of C on values of L/r_w .

Client I.D.: MW6I1/MW6I2 Date(s) Analyzed: 12/14/88 CHEMWEST I.D.: 2849-3
Matrix : Water

Compound	Amount Detected (ug/L)	RL (ug/L)
Benzene	BRL	0.5
Toluene	BRL	1
Ethyl Benzene	BRL	2
Total-Xylenes (1)	BRL	1

Surrogate	% Recovery	Acceptance Window
ortho-Chlorotoluene	99%	50-150%
		

BRL: Below Reporting Limit.

RL: Reporting Limit.

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METHOD OF SHIPMENT

CHEM WEST ANALYTICAL LABORATORIES, INC. 600 West North Market Blvd. Sacramento, California 95834 (916) 923-0840 FAX (916) 923-1938 CLIENT: Harding Lawson Association 1355 Willow Way, Suite 109 Contord, California 94520 ANALYSIS: Shull water Damples are relid under the VOA vials (b) to be analyzed for BTEX. (7-	Order No. 2849 Date Rec'd. 12/9/88 Compl. Date Section KIRK FULLY Project Name: 12XALO ASSESSMENT #6 Project No. 02251,080.03 P.O. NO. Contact Randy Stone Phone (415) 687-9660 Day T/A
SAMPLE ID DATE TIME	ANALYSIS MATEIX CONTAINER
2949-1 MWGG1/MWGG2 12/7/88 1100	BTEX Water 2-40 ml vcA
-2 MW6H1/MW6H2 1230	
-3 MW6II/MW6I2 + 1505	
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