

Texaco Refining and Marketing Inc 108 Cutting Poulevard Richmond CA 91804

April 21, 1993

ENV-STUDIES, SURVEYS & REPORTS 930 Springtown Blvd., Livermore, CA

Ms. Eva Chu Alameda County Department of Environmental Health 80 Swan Way, Room 200 Oakland, CA 94621

Dear Ms. Chu:

Enclosed are the following reports for the former Texaco Service Station located at the above referenced site:

- 1) <u>Quarterly Groundwater Monitoring Letter Report</u>, dated February 4, 1993, covering the third quarter 1992
- 2) <u>Quarterly Groundwater Monitoring Letter Report</u>, dated March 31, 1993, covering the fourth quarter 1992
- 3) <u>Extraction Well Installation and Feasibility Testing</u> <u>Report</u>, dated January 5, 1993

If you have any questions, I may be reached at (510) 236-3611.

Sincerely, Texaco Environmental Services Kall Detter

Karel Detterman, R.G. Project Coordinator

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Attachments

cc: HRPearson-RRZielinski

Regional Water Quality Control Board 2101 Webster Street, Suite 500 Oakland, CA 94612 Mr. Bob Vasquez The Southland Corporation 5820 Stoneridge Mall Road, Suite 310 Pleasanton, CA 94588-3201

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## EXTRACTION WELL INSTALLATION AND FEASIBILITY TESTING REPORT

at

Former Texaco Service Station 930 Springtown Boulevard Livermore, California

prepared for

Ms. Karel Detterman Environmental Geologist Texaco Environmental Services 108 Cutting Boulevard Richmond, CA 94808

prepared by

Weiss Associates 5500 Shellmound Street Emeryville, California 95608

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January 5, 1993



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## EXTRACTION WELL INSTALLATION AND FEASIBILITY TESTING REPORT

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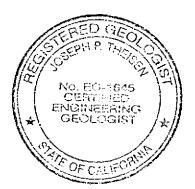
Former Texaco Service Station 930 Springtown Boulevard Livermore, California

prepared for

Texaco Environmental Services 108 Cutting Boulevard Richmond, CA 94808

Bob Riddell, P.E. #C049629 Project Engineer

Weiss Associates work for the Former Texaco Service Station, 930 Springtown Boulevard, Livermore, California, was conducted under my supervision. To the best of my knowledge, the data contained herein are true and accurate and satisfy the scope of work prescribed by the client for this project. The data, findings, recommendations, specifications or professional opinions were prepared in accordance with generally accepted professional engineering and geologic practice. We make no other warranty, either expressed or implied.



Joseph P. Theisen, C.E.G. #1645 Senior Hydrogeologist

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#### **SUMMARY**

In October and November 1992, Weiss Associates' installed three wells and conducted feasibility testing at the Former Texaco Service Station located at 930 Springtown Boulevard, Livermore, California. The objective of the well installation and feasibility testing was to assess the effectiveness of ground water extraction, vapor extraction and air sparging technologies at this site and to obtain data necessary to properly design and operate a future remediation system.

During the well installation phase, WA drilled two soil borings and installed one ground water extraction well, one vapor extraction well and one air sparge well. Sediments encountered during drilling consisted of sand, gravelly sand and silty sand interbedded with silty and clayey units. The highest hydrocarbon concentration detected in soil was 1,200 ppm TPH-G in boring B-1 at 14.4 ft depth. The highest hydrocarbon concentrations were detected just below the water table, which was at about 13 ft below grade.

Aquifer test results indicate that ground water extraction from well EW-1 may capture most of the hydrocarbon-bearing ground water beneath the site. Also, because EW-1 may be screened in a coarse-grained channel deposit, extracting ground water from this zone should mitigate offsite migration of hydrocarbons. If ground water extraction from well EW-1 is implemented, we recommend continued monitoring of ground water levels and contaminant concentrations in onsite and offsite monitoring wells to assess whether ground water extraction from EW-1 sufficiently removes and/or contains the dissolved contaminants beneath the site.

Vapor extraction test results indicate that soil vapor extraction (SVE) from existing wells should effectively remove hydrocarbons from the subsurface, but that the effectiveness of SVE varies in site wells. For example, SVE from well MW-5 achieved a hydrocarbon removal rate of about 127 pounds per day (ppd) while SVE from wells MW-B and VE-1 achieved about 12 and 0.3 ppd, respectively. A comparison of these recent test results with the previous testing indicates that hydrocarbon concentrations in vapor extracted during the recent tests were higher from both well MW-5 and MW-B, with the concentrations from MW-B significantly exceeding the previous test results. Although the effective radii of influence are considered low to moderate, SVE from site wells would remove hydrocarbons most readily from the high permeability materials and would encourage diffusion from the low permeability materials and ground water into the high permeability materials within the SVE system's zone of influence. Additional wells may be required to effect the entire subsurface area of concern. Also, oxygen circulation caused by SVE should enhance natural biodegradation of subsurface hydrocarbons in ground water and soil.

Air sparging test results indicate that air sparging with vapor extraction effectively removes hydrocarbons from the subsurface at this site. During this air sparging testing the hydrocarbon concentrations in extracted vapor increased about ten-fold and then decreased when air sparging ceased. The effective radius of influence for air sparging from air sparge well SP-1 is about 8 to 15 ft based on subsurface pressure/vacuum and water level measurements. Air sparging effectiveness based on other monitored parameters was essentially inconclusive since these parameters usually require testing for one week or longer to be effective indicators. Test results indicate air sparging would be effective at this site, although



it would be more effective if site soils were more permeable and if higher vapor extraction flow rates were achievable.

Because each technology is relatively effective at this site, the chosen technology or combination of technologies depends on remediation goals and schedule and extent of hydraulic control desired. The technology comparison in Section 7 presents information for selecting the desired remediation approach for this site.





#### 1. INTRODUCTION

This report presents the results of Weiss Associates' extraction well installation and feasibility testing at the former Texaco Service Station located at 930 Springtown Boulevard, Livermore, California (Figure 1). The primary objective of the well installation and feasibility testing was to assess the effectiveness of ground water extraction, vapor extraction and air sparging technologies at this site and to obtain data necessary to properly design and operate a future remediation system.

#### 1.1 SCOPE OF WORK

Weiss Associates' scope of work for this project was to:

- Prepare a Site Safety Plan,
- Install one ground water extraction well, one vapor extraction well and one air sparge well,
- Develop ground water extraction well EW-1 and analyze the water samples for TPH-G and BETX,
- Conduct an aquifer test for 24 hours from ground water extraction well EW-1 and analyze an effluent ground water sample,
- Conduct a vapor extraction test from vapor extraction well VE-1, ground water extraction well EW-1, and existing monitoring wells MW-A, MW-B and MW-5,
- Conduct an air sparging test from the air sparge and vapor extraction well,
- Analyze selected soil samples for total petroleum hydrocarbons as gasoline (TPH-G) and benzene, ethylbenzene, toluene and xylenes (BETX),
- Provide suitable of f-gas control for the vapor extraction and air sparging testing,

- Provide temporary surface protection to the installed wells and restore the site,
- Interpret the data and make recommendations for future site remediation, and
- Prepare a report suitable for regulatory submittal which presents the results of the well installation and feasibility testing.

## 1.2 SITE HISTORY AND PREVIOUS INVESTIGATIONS

The former Texaco service station is located adjacent to Highway 580 in a mixed residential and commercial neighborhood in northeast Alameda County in Livermore. The station retailed regular leaded, regular unleaded and premium unleaded gasoline from three underground storage tanks via one pump island. The underground storage tanks were removed on June 26, 1985. The site is currently owned by Southland Corporation and operated as a 7-11 convenience store.

Several subsurface investigations have been primarily conducted at the site. A total of twelve soil borings were drilled at the site. Ground water monitoring wells were installed in ten of the soil borings. All soil and water sample analytic results from these investigations are summarized in the Soil and Ground Water Remediation Work Plan prepared by Groundwater Technology Incorporated (GTI, 1991). The boring logs, well construction details and geologic cross-section from this work plan are presented in Appendix B. Analytic results suggest that hydrocarbons are present in soil near the underground storage tanks and in saturated soil near monitoring wells MW-A, MW-B, and MW-5, and soil boring SB-1. Analytic results for ground water suggest that elevated hydrocarbons in ground water occur mainly beneath the planter area between wells MW-5 and MW-B.



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#### 2. WELL INSTALLATION

On October 19 and 20, 1992, WA installed ground water extraction well EW-1, soil vapor extraction well VE-1 and air sparge well SP-1. The objective of the well installation was to allow testing of the effectiveness of ground water extraction, vapor extraction and air sparging technologies at this site and to obtain information for the future design of a remediation system. The wells were installed near the locations of the highest detected historical hydrocarbon concentrations in ground water and soil and were located at strategic distances from each other and existing wells for influence monitoring during the feasibility testing. The ground water extraction well was screened to recover hydrocarbons from the upper 15 to 20 ft of saturated soils. The air sparging well was screened above a low permeability zone that may have prevented the upward migration of injected air and caused spreading of hydrocarbons laterally. The vapor extraction well was screened immediately above the sparging well to recover hydrocarbon-bearing vapors displaced during air sparging. Since the air sparging and vapor extraction wells were completed in the same borehole, a hydrated bentonite seal was placed between the screened zones to prevent short circuiting of air between the screened zones in the borehole annulus. This section describes well installation activities and site restoration.

#### 2.1 SITE SETTING

Geographic Location:	The site is located in Livermore, California, about one- quarter mile northeast of Arroyo Seco.
Topography:	The site is about 520 ft above mean sea level and situated near the base of a small hill. Local topography slopes eastward.
Surroundings:	Mixed commercial and residential development.
Site Geology:	The shallow sediments in the site vicinity are primarily alluvial sands and silty sands interbedded with clayey and silty units.

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### 2.2 DRILLING AND SOIL SAMPLING

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Drilling Dates:	October 19 and 20, 1992.
Drilling Geologist:	Eric Anderson, Weiss Associates Senior Staff Geologist.
Drilling Method:	Six-inch diameter hollow-stem augers for sampling the borings and twelve-inch diameter hollow-stem augers for well installation. Drilling and sampling procedures are described in Appendix A.
Number of Borings:	2 (one boring B-1 for well EW-1, and boring B-2 for both VE-1 and SP-1) (Figure 1).
Boring Depths:	19 to 33.5 ft.
Soil Sampling Method:	Steam-cleaned, split-barrel drive samplers lined with steam-cleaned brass or stainless steel tubes (Appendix A).
Soil Analyses:	TPH-G and BETX by EPA Methods 5030 using gas chromatography (GC) with flame ionization detection (FID) and by EPA Method 8020 using GC with photo ionization detection (PID).
Sediments Encountered:	Sand and silty sands interbedded with clayey silt and silty clay units. The boring logs and well construction details are presented in Appendix B.
Waste Disposal:	Soil cuttings were stored on plastic sheeting and covered by additional sheeting, then hauled to Zanker Road Landfill in San Jose, California after chemical characterization. Steam clean rinsate, purge water, and well development water were stored in 55-gallon drums, then pumped through granular activated carbon (GAC) with the aquifer test effluent and discharged to the sanitary sewer according to discharge permit requirements.

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#### 2.3 WELL CONSTRUCTION, DEVELOPMENT AND GROUND WATER SAMPLING

Number of Wells:	3 (Figure 1).
Well Materials:	Ground water extraction well (EW-1): 6-inch diameter Schedule 40 PVC well casing with 0.020-inch slotted screen; Monterey #3 sand.
	Air sparge well (SP-1): 1-inch diameter Schedule 80 PVC well casing with 0.010-inch slotted screen; Monterey #3 sand.
	Vapor extraction well (VE-1): 2-inch diameter Schedule 40 PVC well casing with 0.020-inch slotted screen; Monterey #3 sand.
Screened Interval:	Ground water extraction well: Approximately 8 to 33 ft depth (Appendix B).
	Air sparge well: Approximately 16 to 18 ft depth (Appendix B).
	Vapor extraction well: Approximately 7 to 12 ft depth (Appendix B).
Well Development Method:	The ground water extraction well was developed by surge block agitation and air-lift evacuation. Well development and sampling procedures are described in Appendix A. The vapor extraction and air sparge wells do not require development.
EW-1 Ground Water Evacuation Rate:	8.5 gallons per minute during well development.
Ground Water Sampling Method:	Steam-cleaned Teflon bailer. Sample collected on October 27, 1992 before aquifer and air sparge testing.
Analyses for Ground Water:	TPH-G and BETX by EPA Methods 5030, GC/FID, and 2by EPA Method 602, GC/PID.
Ground Water Depth:	Approximately 13 below grade.
Ground Water Flow Direction:	Generally northward with a gradient of 0.005 ft/ft.

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## 2.4 ANALYTIC RESULTS FOR SOIL

Based on the results of this investigation, the highest hydrocarbon concentration detected in soil was 1,200 ppm TPH-G in boring B-1 at 14.5 ft depth (Table 1, Appendix C). The highest hydrocarbon concentrations for the previous and this investigation were generally detected downgradient of the former underground fuel storage tanks near the water table, which was about 13 ft below grade on the date of the drilling for our investigation. No hydrocarbons were detected in low permeability soil samples at 9.7 ft and 18.5 ft, but TPH-G (at 3.0 ppm) was detected in a high permeability soil unit at 24.7 ft depth, but was not detected in the same soil unit at 29.7 ft depth.

#### 2.5 SITE RESTORATION

Site restoration consisted of disposing of all waste soil, and relandscaping by planting new ivy to match the existing ivy and replacing the wooden planter edging.



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### 3. AQUIFER TESTING

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WA conducted aquifer testing of well EW-1 on November 17 and 18, 1992. The test objectives were to determine the hydraulic characteristics of the shallow water-bearing zone beneath the site and to determine the optimal number and placement of ground water extraction wells for future site remediation. The aquifer test results indicate that ground water extraction from well EW-1 should capture most of the hydrocarbon-bearing ground water beneath the site.

### 3.1 PERMITTING

Aquifer test effluent was treated and discharged to City of Livermore, Water Reclamation Plant in accordance with a ground water discharge permit dated November 3, 1992 and presented in Appendix D.

### 3.2 AQUIFER TEST ANALYSIS AND GROUND WATER FLOW MODELING

A 24-hour drawdown test was conducted on November 17 and 18, 1992. Ground water was extracted from well EW-1 at an average flow rate of about 7.85 gallons per minute (gpm). Water level changes were recorded in test well EW-1, and monitoring wells MW-A, MW-B, MW-1, MW-2, MW-3, MW-5, and MW-7. The initial flow rate of 8 gpm was reduced to 7.5 gpm for the last 5 hours of the test to prevent dewatering the test well.

The pump test data was analyzed using the Cooper-Jacob (Cooper, 1946) semi-log method. The hydraulic responses in the monitoring wells are shown in Figure 2. Estimates of the hydraulic parameters from each well, and the distances to the extraction well are provided in Table 2. As shown in Table 2, the average transmissivity of the aquifer is estimated to be 3,400 gallons per day per foot (gpd/ft). Although most of the monitoring wells are screened over a length of 20 feet, boring logs indicate that the more permeable, sandy gravel zone is about 15 ft thick. Using this thickness, an average hydraulic conductivity value of 225 gpd/ft<sup>2</sup> (0.021 ft/min), and a specific storage of 0.001  $ft^{-1}$  are estimated for this aquifer. Monitoring well MW-7 did not show any response during the extraction test, therefore, no hydraulic analysis was conducted for this well. The drawdown in the extraction well, EW-1, was significantly larger (15.5 ft) than that observed in the monitoring wells, which are very close to the test well. The excessive drawdown in the pumping well is probably due to poor well efficiency, and large well losses across the screen and within the borehole. For these reasons, the hydraulic parameters estimated for well EW-1 were not used in estimating average aquifer values.

To estimate the long term effects of pumping well EW-1, ground water extraction from the well was simulated using the analytical element models, EQUIPLOT and CAPTURE (McEdwards, 1986). EQUIPLOT calculates water level changes in a gridded area due to pumping using the Theis (1935) transient drawdown equation and by superimposing the effects of extraction at every grid cell. CAPTURE also uses the same relation but reports resulting flow paths due to pumping. Both programs require hydraulic conductivity (ft/min), regional gradient (ft/ft), direction of regional flow, specific storage (ft<sup>-1</sup>), location and pumping rates (gpm) of extraction wells as input parameters.

Estimates of the hydraulic parameters from the hydraulic test analysis described above were used for model input (i.e. a hydraulic conductivity of 0.021 ft/min, a specific storage value of 0.001 ft<sup>-1</sup>, and an aquifer thickness of 15 ft). The regional ground water gradient was reported as 0.05 ft/ft (GTI, 1991). This steep gradient is considered to be the result of consistently higher ground water elevations in monitoring well MW-7, which is upgradient from the other wells. Because monitoring well MW-7 did not respond to the extraction test, this well is believed to be completed in a lower hydraulic conductivity zone that is not in hydraulic communication with the wells on the northern portion of the site. This interpretation is supported by the lithologies at the site. The boring log of MW-7 in Appendix B indicates that sediments screened by this well are clay and clayey gravel, whereas the remaining site wells screen more permeable sediments. If the water level for MW-7 is disregarded, a much smaller gradient of 0.005 ft/ft is obtained from the remaining water level elevation data. The direction of regional flow is generally to the north.

The model was calibrated using EQUIPLOT and simulating the 24-hour extraction test with an average flow rate of 7.85 gpm. We compared the observed and simulated drawdowns to check the accuracy of the selected model parameters. As previously discussed, estimates of

the drawdown within well EW-1 were disregarded, because of the apparent extensive well losses. Agreement between the observed and predicted drawdowns indicate that the parameters determined from the hydraulic test analyses are representative of the simulated aquifer.

Using the hydraulic parameter values from the calibrated model (listed in Table 3), a long term pumping and capture area simulation was conducted using CAPTURE. The modelling results show that a long term pumping flow rate of 5 gpm may be sustainable without dewatering extraction well EW-1. We assumed a long-term, sustainable flow rate of 5 gpm because long-term pumping rates are usually less than pumping rates obtained during shortterm testing. As shown in Figure 3, the capture area predicted for this flow rate covers the entire region east of Lassen Road and south of Springtown Boulevard. In addition, the downgradient part of the capture zone nearly extends to monitoring well MW-5.

If a steeper regional gradient of 0.05 ft/ft is used with the same hydraulic parameters used to create Figure 3, a much narrower capture zone is obtained. The capture area encompasses monitoring wells MW-A, MW-B, MW-1, MW-2, and MW-3, but does not extend to monitoring wells MW-5 or MW-7. Because a gradient of 0.05 ft/ft is calculated with emphasis on a single data point (from MW-7), it is not considered to be realistic for the entire aquifer.

However, during the aquifer test, the largest drawdowns were observed in MW-1 and MW-3, suggesting these wells may be in better hydraulic communication with the extraction well, EW-1, as compared to the other monitoring wells. Therefore, ground water extraction from EW-1 may preferentially withdraw ground water from a possible channel deposit, which would result in a smaller capture than that shown in Figure 3. However, the third largest drawdown was observed in MW-5, suggesting this well may also be screeened near a potential channel deposit.

#### 4. VAPOR EXTRACTION TESTING

WA conducted vapor extraction testing on October 27, 1992 to provide information for possible SVE remediation system design and to select cost-effective vapor treatment. Although, the test results indicate that vapor extraction from existing wells should effectively remove hydrocarbons from the subsurface, the effectiveness of vapor extraction varies in site wells. In addition to evaluating vapor extraction effectiveness from new wells VE-1 and EW-1, our test also provided data to re-evaluate vapor extraction effectiveness in wells MW-A, MW-B and MW-5. These wells were tested previously on July 24, 1991 (GTI, 1991).

#### 4.1 TEST CRITERIA AND PROCEDURES

WA conducts vapor extraction testing to obtain data on vapor extraction flow rates, vacuum requirements, hydrocarbon concentrations in extracted vapor, hydrocarbon removal rates and extent of vacuum influence. The extraction locations and durations, test equipment, and data collection and interpretation methods for our October 27 test are described below.

#### 4.1.1 Extraction Locations and Durations

WA extracted soil vapor from vapor extraction well VE-1, ground water extraction well EW-1, and existing monitoring wells MW-A, MW-B and MW-5. We monitored vacuum influence in these and other site wells (Figure 1). These wells are located adjacent to the underground storage tanks near the highest detected hydrocarbon concentrations in soil.

We extracted soil vapor for about 15 to 30 minutes from each well, except VE-1. We performed a 2.5 hour step test on VE-1 that involved initially determining the maximum achievable extraction flow rate from the well, and then reducing the flow rate to about half this initial flow. We then resumed extraction at the full flowrate. We tested from well VE-1 longer than the other wells to help stabilize hydrocarbon concentrations in extracted vapor before air sparge testing on the following day.

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#### 4.1.2 Equipment

The test extraction equipment included a positive-displacement vacuum pump powered by a 3 horsepower explosion-proof motor, rated to produce a vacuum of 160 inches of water at various flow rates. For off-gas control, we routed extracted vapor through the test system's moisture collection drum and two 200 lb GAC adsorption vessels connected in series in accordance with vapor extraction testing requirements of the Bay Area Air Quality Management District.

#### 4.1.3 Data Collection and Interpretation

Data collection included recording the test system's operation parameters, collecting influent and effluent vapor samples, and measuring the induced vacuum and/or pressure in site wells. The influent vacuum gauge indicated the vacuum applied to each test well. We estimated vapor extraction flow rates based on applied vacuum, differential pressure gauge readings and performance curves supplied by the vacuum equipment manufacturer.

To record vacuum influence, we sealed site wellheads and measured the induced vacuum in each location with a differential pressure gauge which indicates the difference between well pressure and atmospheric pressure. Before and after each extraction event, we measured background vacuum in all sealed wells to distinguish between pressure differences induced by atmospheric change versus pressure differences induced by vapor extraction.

To normalize vacuum influence data from each well, we subtracted the initial measured background vacuum from the final vacuum influence measurement. In some cases the background vacuum influence did not fully stabilize due to residual vacuum, atmospheric fluctuations, subsurface conditions and brief intervals between extraction tests, so we noted this on the vacuum influence data table.

WA measured hydrocarbon concentrations in extracted vapor and after each carbon adsorption vessel with a portable organic vapor analyzer/flame ionization detector (OVA/FID). We also submitted one-liter vapor samples to a state-certified analytical laboratory for analysis for TPH-G by modified EPA Method 5030 and for BTEX by modified EPA Method 8020. We collected samples for laboratory analysis near the end of each extraction event. We collected one sample at the end of both step test intervals for extraction from vent point VE-1.

Hydrocarbon removal rates were calculated using data on stabilized hydrocarbon concentrations in extracted vapor and vapor flow rates for each extraction event. Actual hydrocarbon removal rates for a dedicated SVE system may vary depending on the applied vacuum, extraction flow rates, and hydrocarbon concentration fluctuation during SVE project duration.

#### 4.2 TEST RESULTS

The vacuum applied by the test equipment to each well ranged from 73 to 141 inches of water and induced vapor extraction flow rates ranging from about 1 to 20 standard cubic feet per minute (scfm). Hydrocarbon concentrations in extracted vapor ranged from 190 parts per million by volume (ppmv) to 20,450 ppmv TPH-G. Hydrocarbon removal rates ranged from 0.3 pound per day (ppd) to 127 ppd TPH-G.

During extraction from each well, the applied vacuum induced water upwelling into the equipment's water collection drum. Accordingly, we decreased the applied vacuum until vapor extraction flow rates maximized.

The hydrocarbon concentrations in extracted vapor were highest from MW-5 (at 20,450 ppmv TPH-G) and second highest in MW-B (at 18,450 ppmv TPH-G). However, vapor extraction is more effective from MW-5 than from MW-B because the vapor extraction flow rate was significantly higher from MW-5 (at 19.4 scfm) than from MW-B (at 2 scfm).

Table 5 summarizes test data and estimates hydrocarbon removal rates for extraction from each well. Table 6 presents test data and vacuum influence data for extraction from each well. Figure 4 presents the isobarometric contours for extraction from well VE-1. The analytical report and chain of custody forms are presented in Appendix C. We converted the hydrocarbon concentrations reported as milligrams per cubic liter of air to ppmv by dividing by the compound's molecular weight and multiplying by 24.45 which is the volume one grammole of perfect gas occupies at the standard temperature of 25 degrees centigrade and the standard pressure of 760 millimeters of mercury.

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#### 5. AIR SPARGING TESTING

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WA conducted an air sparging test on October 28, 1992. Test results indicate that air sparging with vapor extraction effectively removes hydrocarbons from the subsurface at this site. During this air sparging testing the hydrocarbon concentrations in extracted vapor increased about ten-fold and then decreased when air sparging ceased. The effective radius of influence for air sparging from air sparge well SP-1 is about 8 to 15 ft based on subsurface pressure/vacuum and water level measurements. Air sparging effectiveness based on other monitored parameters was essentially inconclusive since these parameters usually require testing for one week or longer to be effective indicators. This longer term testing of air sparging effectiveness is often performed during a trial operation period of an installed system.

#### 5.1 AIR SPARGING BACKGROUND

Air sparging involves injecting air below the water table to strip volatile organic compounds (VOCs) from the ground water. The VOC-laden vapors are then captured by a vapor extraction system. Air injection and vapor extraction is performed using either appropriately screened vertical wells, horizontal wells or trenches. The effective radius of influence from a vertical air sparging well is typically about 5 to 20 ft in coarse materials. Air sparging also oxygenates ground water, thereby possibly encouraging in-situ biodegradation of VOCs in ground water.

The effectiveness of air sparging is sensitive to the lithology and stratigraphy of the saturated and unsaturated zones, and to the effectiveness of vapor extraction. In highly stratified soils, injected air may travel laterally substantially before reaching the vadose zone. Generally, the lateral migration of air within the saturated zone spreads the dissolved contaminants in ground water laterally also. Vapor extraction should be designed to capture vapors emanating from the saturated zone.

Since, at the former Texaco site, an apparent confining layer from about 18 to 24 ft bgs separates the shallower, more contaminated, clayey sand and gravel unit from the deeper more permeable sand unit, hydrocarbon-bearing ground water would be remediated more efficiently from an air sparge well screened only in the shallower unit. Also, air sparging into the sandy unit below the apparent confining layer could produce air pockets which would laterally displace ground water and allow diffusion of vapor-phase hydrocarbons into previously uncontaminated ground water. Therefore, we installed the air sparge well only in the shallower clayey sand and gravel unit to reduce the risk of spreading hydrocarbons laterally below the confining layer. We screened the air sparge well from 16 to 17 ft below grade surface (bgs) based on an evaluation of the local lithology during well installation. This screen interval is about 3 to 4 ft below the static water level and is above the clayey silt sediments found at about 18 to 22 ft bgs in the bore hole. Air sparging from this well allowed more effective monitoring and evaluating of air sparging than installing and air sparging from a deeper well screened beneath the clayey silty sediments at 18 to 22 ft bgs.

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For these reasons, we conducted a short-term air sparging test from a shallow air sparge well nested with a vapor extraction well to minimize the possible lateral spreading of hydrocarbons in the saturated zone.

#### 5.2 TEST CRITERIA

WA performed the following tasks to evaluate air sparging effectiveness:

- Monitored the hydrocarbon concentration in extracted soil vapor,
- Monitored the subsurface pressure/vacuum regime,
- Monitored the water table to determine whether the air sparging caused rising water levels,
- Measured the dissolved oxygen concentrations in ground water, and
- Analyzed ground water samples from the adjacent well EW-1 for TPH-G and BTEX before and after the test.

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#### 5.3 PROCEDURES

WA extracted soil vapor from the vapor extraction well for about 2.5 hours to determine the background hydrocarbon concentrations in extracted soil vapor before air sparging. We then simultaneously injected air (sparging) and extracted vapors for about one hour while monitoring the extracted vapors for increasing hydrocarbon concentrations. We then ceased air sparging and monitored the extracted soil vapor for about one-half hour.

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To help ensure full capture of the injected air, we first determined the vapor extraction rate using the test equipment, and then air sparged at a lower flow rate. This ensured that the volume of air injected was less that the total volume removed.

#### 5.4 RESULTS

The vapor extraction flow rate from well VE-1 was about 3 scfm, therefore, we injected air at 1 scfm. The air pressure required to inject 1 scfm of air was 10 pounds per square inch (psi) initially but decreased to 5.5 psi within 10 minutes. During air sparging at about 1.1 scfm, the vapor extraction flow rate increased from 3.3 scfm to 3.7 scfm for the approximate constant applied vacuum of 95 inches of water. After air sparging the vapor extraction flow rate decreased to 3.6 scfm.

The hydrocarbon concentrations in extracted vapor increased about six-fold when air sparging started and then decreased when air sparging ceased. Figure 5 shows the TPH-G concentrations in extracted vapor before, during and after air sparging. After 2 hours of vapor extraction and before air sparging began, the hydrocarbon concentrations in extracted vapor were about 600 ppmv TPH-G. After one hour of air sparging, the hydrocarbon concentrations in extracted vapor increased to about 6,100 ppmv TPH-G. About 25 minutes after air sparging ceased, the hydrocarbon concentrations in extracted vapor decreased to about 3,500 ppmv TPH-G.

The subsurface pressure/vacuum changed most in nearby well EW-1, located about 8 ft from VE-1/SP-1, and significantly less in well MW-A, located about 20 ft away. The pressure/vacuum readings from EW-1 indicated a vacuum of 0.06 inches of water before air sparging, positive air pressure of 0.24 inches of water during air sparging, and a vacuum of 0.11 inches of water after air sparging. The pressure/vacuum readings from MW-A indicated a vacuum of 0.01 inches of water before air sparging and neither pressure nor a vacuum during air sparging. Air pressure measurements from wells EW-1 and MW-A are presented in Tables 7 and 8, respectively.

Similarly, the water levels responded most in nearby well EW-1 located about 8 ft from VE-1/SP-1 and significantly less in well MW-A located about 20 ft away. The water level in EW-1 rose 0.03 ft during vapor extraction before air sparging and rose further to 0.18 ft above the initial water level during the first 13 minutes of air sparging before lowering to 0.06 ft above the initial water level by the end of the air sparging. When air sparging ended, the water level dropped to 0.05 ft below the initial water table elevation. About one-half hour after vapor extraction and air sparging ended the water level was 0.11 ft above the initial water level elevation. The water level in well MW-A was essentially unchanged until the air sparging began, when the water level rose 0.07 ft above the initial water table elevation. After vapor extraction and air sparging the water level in MW-4 returned to the initial water table elevation. Water level measurements from wells EW-1 and MW-A are presented in Tables 7 and 8, respectively.

Dissolved oxygen concentrations in ground water fluctuated during feasibility testing as presented in Table 9. These concentration fluctuations are described in the conclusion section 6.3.

Ground water samples from well EW-1 contained 11 ppm TPH-G before the air sparging/vapor extraction test and 13 ppm TPH-G after the test.

#### 6. CONCLUSIONS

This section discusses the conclusions of the aquifer, vapor extraction and air sparge testing.

#### 6.1 AQUIFER TESTING

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The modeling based on the aquifer test results suggests that pumping from extraction well EW-1 may be a sufficient remedial design alternative, because the capture zone should influence the regions of high contamination and much of the contaminated ground water may be extracted. Modeling results also indicate that EW-1 may be capable of a long-term, sustainable flow rate of 5 gpm. The predicted capture area for this flow rate covers the entire region east of Lassen Road and south of Springtown Boulevard. In addition, the downgradient part of the capture zone extends beneath Springtown Boulevard and nearly extends to monitoring well MW-5. Long-term pumping from EW-1 may capture ground water from well MW-5.

However, the extraction well may be screened within a channel deposit, hence the capture area may be smaller than that shown in Figure 3. A coarser-grained channel deposit would provide a preferred migration pathway for contaminants beneath the site. Extracting ground water from this zone would therefore mitigate further offsite migration of the contaminants.

In conclusion, we recommend utilizing EW-1 as an extraction well to remove contaminated ground water beneath the site. We also recommend continued monitoring of ground water levels and contaminant concentrations in onsite and offsite monitoring wells to assess whether ground water extraction from EW-1 sufficiently removes and/or contains the dissolved contaminants beneath the site.

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#### 6.2 VAPOR EXTRACTION TESTING

SVE test results indicate that SVE from existing wells should effectively remove hydrocarbons from the subsurface, but that the effectiveness of SVE varies in site wells. For example, SVE from well MW-5 achieved a hydrocarbon removal rate of about 127 pounds per day (ppd) while SVE from wells MW-B and VE-1 achieved about 12 and 0.3 ppd, respectively. Also, because the influence area from most wells is considered low to moderate, additional wells may be required to effect the subsurface area of concern. A comparison of these recent test results with the previous testing indicates that hydrocarbon concentrations in extracted vapor the recent tests were higher from both well MW-5 and MW-B, with the concentrations from MW-B significantly exceeding the previous test results.

The varying effectiveness and irregular vacuum influence measurements suggest that the coarser grained interbedded materials in the site subsurface may be acting as subsurface conduits for vapor flow. For example, the vacuum of 73 inches of water applied to MW-5 induced about 0.06 inches of water in well MW-B and MW-3, while it induced only 0.01 inches of water in MW-A, located closer to MW-5 than either MW-B or MW-3. Vacuum influence readings during extraction from well VE-1 were more predictable. The vacuum of 141 inches of water applied to VE-1 induced about 3.5 inches of water in well EW-1 and 0.1 inches of water in well MW-A. The effective radii of influence for extraction from MW-5 and VE-1, typically defined as one percent of the applied vacuum, is about 30 and 12 ft, respectively.

Although the effective radii of influence seem relatively low, SVE from site wells would remove hydrocarbons most readily from the high permeability materials and would encourage diffusion from the low permeability materials and ground water into the high permeability materials within the SVE system's zone of influence. Also, oxygen circulation caused by SVE should enhance natural biodegradation of subsurface hydrocarbons in ground water and soil.

SVE effectiveness could be enhanced by installing additional wells or a trench. However, trench installation would be very disruptive and vapor extraction from a trench would predominantly remove hydrocarbons from the higher permeability materials similar to extraction from the site wells. SVE effectiveness would also be enhanced if it were performed in conjunction with ground water extraction as described in Technology Comparison Section 7 below.

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Selection of the most cost-effective vapor treatment method for SVE is based on the estimated vapor flow rates and hydrocarbon concentrations in extracted soil vapor over the anticipated project duration. Although it is not possible to predict long term hydrocarbon concentrations based on this one day test, hydrocarbon concentrations are expected to remain high initially and decrease with time. Since hydrocarbon mass removal rates experienced during the test peaked at 127 ppd TPH-G, we recommend using an available Texaco-owned catalytic or thermal/catalytic oxidizer until the hydrocarbon mass removal rates decrease to about 5 to 10 pounds per day. This may require about 30 to 60 days at this site. For a hydrocarbon mass removal rate of about 10 pounds per day, carbon adsorption is typically the most cost-effective vapor treatment method. However, it may be more cost-effective to use the oxidizer for the duration of the project than to design, permit and install a carbon adsorption system. Conversely, if Texaco does not have a catalytic or thermal/catalytic oxidizer available, it may be more cost-effective to permit, install and operate a carbon adsorption system for the project duration.

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#### 6.3 AIR SPARGE TESTING

Test results indicate that air sparging with vapor extraction effectively removes hydrocarbons from the subsurface at this site. During this air sparging testing the hydrocarbon concentrations in extracted vapor increased about ten-fold and then decreased when air sparging ceased. The effective radius of influence for air sparging from air sparge well SP-1 is about 8 to 15 ft based on subsurface pressure/vacuum and water level measurements. Air sparging effectiveness based on other monitored parameters was essentially inconclusive since these parameters usually require testing for one week or longer to be effective indicators.

Test results indicate air sparging would be effective at this site, although it would be more effective if site soils were more permeable and if higher vapor extraction flow rates were achievable.

The ten-fold increase in TPH-G concentrations in extracted vapor indicates that sparging increased hydrocarbon removal from the saturated zone. This suggests that hydrocarbons are removed by stripping them from ground water or by the creation of new vapor flow pathways through the saturated and unsaturated zones. The air sparging well may be essentially serving as an air inlet well within the saturated zone where hydrocarbons are adsorbed to soil. With this air sparging well, more air is drawn from within the saturated and lower unsaturated zone, and less air is drawn from the unsaturated zone and the ground surface.

Based on the subsurface pressure/vacuum and water level measurements, the effective radius for air sparging radius from SP-1 is between 8 to 15 ft. We estimate this range because the subsurface pressure/vacuum and water levels changed most in nearby well EW-1 located about 8 ft from VE-1/SP-1 and significantly less in well MW-A located about 20 ft away. In both monitored wells, the water level rose during air sparging despite the increased pressure within the wells. This suggests that the air sparging caused enough of a water table rise in this area to overcome the increased pressure in the wells caused by air sparging.

The subsurface pressure/vacuum and water levels may also suggest that air sparging caused more water table rise initially, but then as the injected air developed preferential flow paths it displaced ground water around the well and depressed water in the subsurface due to the increased subsurface pressure. This suggests that longer term testing at different flow rates could test the effectiveness of air sparging at different flow rates. This also suggests that air sparging effectiveness is enhanced by cycling operation on and off to create different flow paths thereby effecting more soil and ground water. Although this may also suggest that deeper air sparging wells would be more effective for air sparging, air sparging from deeper wells at this site may penetrate the apparent confining layer at 18 to 22 ft bgs and cause lateral spreading of hydrocarbons in ground water.

The effective radius for air sparging could be assessed further during a longer test or during system operation by evaluating the dissolved oxygen and hydrocarbon concentrations and hydrocarbon-degrading microbial populations in ground water. Increasing dissolved oxygen concentrations and hydrocarbon-degrading microbial populations, as well as decreasing hydrocarbon concentrations in monitored wells would indicate air sparging was effective in those wells.

If an air sparging system is installed at this site, we would recommend cycling the system on and off to vary the vapor flow paths within the saturated zone, to minimize the possible lateral spreading of hydrocarbons in the saturated zone, and to minimize energy costs. We would also recommend installing one or two additional air sparging and vapor extraction wells between MW-A and MW-5, and installing one air sparge well near MW-5.

#### 7. TECHNOLOGY COMPARISON

Test results indicate that each of the evaluated technologies may be effective at this site. However, the remediation effectiveness of each technology will vary, depending on the specific advantages and limitations of the individual techniques. To select the optimal remedial approach for the site, WA compared each technology and combinations of technologies. The results of this review are presented below.

#### 7.1 GROUND WATER EXTRACTION

Ground water extraction (GWE) is a proven and reliable remedial technology to hydraulically contain and remediate hydrocarbon-bearing ground water. GWE would remove hydrocarbons from the saturated zone, but would not remediate hydrocarbons in the unsaturated zone. Therefore, using this technique alone may allow hydrocarbons in unsaturated soil to continue to impact ground water quality. The effectiveness of this method is dependent upon the extraction well locations relative to the mass of hydrocarbon-bearing ground water and the subsurface hydrogeology.

The aquifer test results indicate that ground water extraction from EW-1 should hydraulically contain the hydrocarbon-bearing ground water at this site. If hydrocarbons have migrated off-site further than estimated, then GWE from EW-1 alone may not draw these offsite hydrocarbons back to the site. GWE typically requires a minimum of 5 years to achieve remediation objectives.

#### 7.2 SOIL VAPOR EXTRACTION

SVE is a proven and reliable remedial technology for removing hydrocarbons from the unsaturated zone. SVE from site wells would remove hydrocarbons most readily from the high permeability materials and would encourage hydrocarbon diffusion from the low permeability materials into the high permeability materials within the SVE system's zone of influence. SVE



should also improve ground water quality at the site by inducing benzene and hydrocarbon diffusion from ground water and possibly by supplying oxygen to naturally occurring micoorganisms that degrade organic materials, therefore, encouraging in situ biodegradation of hydrocarbons. Although SVE may improve ground water quality, it would not hydraulically contain hydrocarbon-bearing ground water at the site. Unfortunately, the SVE test results indicated limited SVE effectiveness except from well MW-5. SVE would be most effective with a series of extraction and air inlet trenches although this may not be very cost-effective and would certainly entail significant site disruption. SVE typically requires about 6 months to 2 years to achieve unsaturated zone remediation objectives.

#### 7.3 COMBINED GROUND WATER AND SOIL VAPOR EXTRACTION

Combined ground water and soil vapor extraction is a proven technology for removing hydrocarbons from the saturated and unsaturated zones and achieving remediation objectives faster than ground water extraction alone. The vacuum applied by SVE would enhance ground water extraction flow rates thereby enhancing hydraulic control. Also, GWE would depress the water table, limiting the upwelling effects caused by SVE and exposing previously saturated soil to the air stream induced by SVE. This is especially useful since ground water fluctuations create a zone of high residual hydrocarbon-saturated soil both above and below the average water table elevation. Combined ground water and soil vapor extraction also offers the same advantages as the individual techniques discussed above.

#### 7.4 AIR SPARGING WITH SOIL VAPOR EXTRACTION

Air sparging with soil vapor extraction is an innovative and relatively new technology for removing hydrocarbons from ground water and saturated soil. Test results indicate that air sparging would remove hydrocarbons from the saturated zone by stripping hydrocarbons from ground water or by creating vapor flow through the saturated zone. Air sparging should also oxygenate ground water, thereby, possibly encouraging in-situ biodegradation of VOCs in ground water.

A trial period of air sparging could be performed in conjunction with SVE to more fully evaluate air sparging effectiveness for this site. If air sparging with SVE effectively remediates the saturated zone then ground water extraction would be unnecessary. However, air sparging would not hydraulically contain the hydrocarbon-bearing ground water.

Test results indicate that the effective radius for air sparging is about 8 to 15 ft. Therefore, to implement this technology at this site, we would recommend installing one air sparge well near MW-5 and one or two air sparge and vapor extraction wells between MW-A and MW-5. We would recommend cycling the system on and off to vary the vapor flow paths within the saturated zone, to minimize the possible lateral spreading of hydrocarbons in the saturated zone, and to minimize energy costs. We would also recommend installing one or two additional air sparge and vapor extraction wells between MW-A and MW-5, and installing one air sparge well near MW-5.

Unfortunately, the effectiveness of air sparging is sensitive to the lithology and stratigraphy of the saturated and unsaturated zones, and to the effectiveness of vapor extraction. In the highly stratified soil at this site, injected air may travel laterally substantially before reaching the vadose zone, and spread the dissolved contaminant plume. Vapor extraction should be designed to capture vapors emanating from the saturated zone.

#### 7.5 COMBINED GROUND WATER AND SOIL VAPOR EXTRACTION AND AIR SPARGING

Implementing all the technologies simultaneously should prove very effective and should achieve the remediation objectives quickly. However, this combination approach would probably be more expensive than the other approaches. This combined approach would have the advantages of each of the individual techniques discussed above.

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#### 8. REFERENCES

## Cooper, H.,H., and C. E. Jacob, 1946, A generalized graphical method for evaluating formation constants and summarizing well field history, Am. Geophys. Union Trans. 27, pp. 526-534.

- McEdwards, D., 1986, Equiplot: A program for generating a grid of ground water elevations for extraction wells in uniform ground water flow, Data Services, Petaluma, CA.
- Groundwater Technology Inc., 1991, Work Plan for Soil and Ground Water Remediation at Former Texaco Service Station, 930 Springtown Boulevard, Livermore, California, 9 pp. and appendices.

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Boring/	Sample	<b>B</b> - 4 -	A.v 1		TPH-G	В	E	T	x
Sample ID	Depth Date (ft)Sampled		Analytic Sat/ Method Unsat		<		mg/kg (ppm)		·····>
8-1	9.7	10/19/92	5030/8020	Unsat	<1.0	<0.005	<0.005	<0.005	<0.005
(EV-1)	14.5	10/19/92	5030/8020	Sat	1,200	6.6	15	21	50
-	24.7	10/19/92	5030/8020	Sat	3	0.017	0.050	0.051	0.21
	29.5	10/19/92	5030/8020	Sat	<1.0	<0.005	<0.005	<0.005	<0.005
B-2	14.5	10/20/92	5030/8020	Sat	1,000	7.t	13	22	56
(VE-1/	16.7	10/20/92	5030/8020	Sat	990	2.9	14	15	53
SP-1)	18.5	10/20/92	5030/8020	Sat	<1.0	0.007	<0.005	0.029	<0.005

Abbreviations:

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TPH-G = Tatal petroleum hydrocarbons as gasoline by Modified EPA Method 8015

B = Benzene by EPA Method 8020

E = Ethylbenzenerby EPA Method 8020

T = Toluene by EPA Method 8020 X = Xylenes by EPA Method 8020

ppm = parts per million

ppin - parts per mittron

#### Analytical Laboratory:

Mobile Chem Labs Inc., Martinez, California

#### Analytic Methods

5030 = Purge and trap by EPA Method 5030 with flame ionization detector (FID) for TPN-G

8020 = EPA Method 8020 for BETX

Well	Transmissivity (gpd/ft)	Hydraulic Conductivity (gpd/ft2)	Specific Storage (ft <sup>-1</sup> )	Distance to Pumping wel (ft)	
			· · · · · · · · · · · · · · · · · · ·		
EW-1*	340*	23*	-	-	
MW-A	4,700	310	0.05	12	
MW-B	5,600	370	0.004	50	
MW-1	910	60	0.0002	78	
MW-2	5,950	400	0.002	88	
MW-3	920	61	0.0008	45	
MW-5	2,200	150	0.0002	75	
MW-7*	-	-	-	130	
Average	3,400	225	0.001	-	

## Table 2. Hydraulic Parameters Estimated Using Cooper-Jacob (1946).

\*Not used in averaging.

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### Table 3. Hydraulic Parameters Used in EQUIPLOT and CAPTURE Models.

#### Parameter

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Value used in the simulations

K - Hydraulic conductivity

b - Aquifer thickness

i - Local gradient

Q - Extraction rate

0.021 ft/min 15.0 ft 0.005 ft/ft to North 5.0 gpm at steady state

Well	Date Sampled	Analytic Method	Analytic Laboratory	ТРК-G <	8 part:	E s per billion (	τ μg/l)	X
EW-1 Before Air Sparge Test	10/27/92	602/5030	MC	11,000	410	540	2,000	2,100
EW-1 After Air Sparge Test	10/27/92	602/5030	MC	13,000	840	580	2,400	1,900
EW-1 (Pump Test Effluent After 24 Hours Before Treatment)	11/18/92	602/5030	MC	4,300	140	96	340	560
EW-1 (Pump Test Effluent After 24 Kours After Treatment)	11/18/92	602/5030	MC	<50	<0.5	<0.5	<0.5	<0.5
bbreviations: PH-G = Total perroleum hydroca 8015	<u>Analytical Laboratory:</u> MC = Mobile Chem Labs, Inc., Martinez, California							
B = Benzene by EPA Method 602 E = Ethylbenzene by EPA Method 602 F = Toluene by EPA Method 602 K = Xylenes by EPA Method 602 Kn = Not detected at detection limits of n ppb				<u>Analytical Methods:</u> 602 = EPA Method 602 for BETX 5030 = EPA Method 5030 for TPK-G				

### Table 4. Analytic Results for Ground Water · Former Texaco Service Station, 930 Springtown Boulevard, Livermore, California

Table 5. Soil Vapor Extraction Test Data, Former Texaco Service Station, 930 Springtown Boulevard, Livermore, California

WELL	EXPOSED SCREEN DEPTH (a)	VACUUM APPLIED	FLOW	ELAPSED TIME	CONCENT	(DROCARBON RATIONS(b) (ppmv)	(lbs	MOVAL RATE(c) /day)
ID	(ft-ft)	(inches water)	(scfm)	(min)	TPH-G	BENZENE	TPH-G	BENZENE
EW-1	8-12.0	101	5.6	0-18	425	4	0.8	0.007
MW-A	4-12.7	92	1.1	0-24	900	9	0.3	0.003
MW-B	4-11.6	84	2	0-14	18,450	350	11.8	0.204
MW-5	5-11.9	73	19.4	0-22	20,450	110	127.3	0.621
VE-1	7-12.0	79 141	2.2 5.2	0-34 34-138	1,300 190	18 3	0.9 0.3	0.012 0.005

Notes:

(a) = Depth interval below grade surface between top of well screen and ground water (nearest 0.1 feet).

(b) = Analysis for Total Petroleum Hydrocarbons as Gasoline (TPH-G) and Benzene by Modified EPA Method 5030 and 8020. The concentrations reported as milligrams per cubic meter are converted to ppmv by dividing by the compound molecular weight and multiplying by 24.45 which is the volume one gram-mole of perfect gas occupies at the standard temperature and pressure of 25 degrees centigrade and 760 millimeters of mercury.

(c) = Mass removal rate based on Bay Area Air Quality Management District's Manual of Procedures for Soil Vapor Extraction dated July 17, 1991. Rate = concentration(ppmv) x flowrate(scfm) x (1 lb-mole/386ft3) x molecular weight (86 lb/lb-mole for TPH-G as Hexane, 78 for Benzene) x 1440 min/day.

scfm = Standard cubic feet per minute.

ppmv = Parts per million by volume.

Flow Rate	(#) cuum ("H <sub>2</sub> O)		VE 1 7 2. 0-	9 2	VE 2 14 5. 34-	2	EW 1 10 5. 0-1	1 6
Probe/ Well ID	Exposed Screen Depth <sup>8</sup> (ft-ft)	Est. Perm. in Exposed Screen	Distance <sup>C</sup> from Test Point (ft)	Vacuum <sup>d</sup> Influence ("स <sub>2</sub> 0)	Distance <sup>C</sup> from Test Point (ft)	Vacuum <sup>d</sup> Influence ("H <sub>2</sub> 0)	Distance <sup>C</sup> from Test Point (ft)	Vacuum <sup>d</sup> Influence ("K <sub>2</sub> O)
EW-1	8-12.0	L,M	8	1.0	8	3.5		
M₩-A	4-12.7	L	20	0.02	20	0.10	13	0.5
MH-B	4-11.6	L	42	0.005	42	0.005	51	0.005
MW-2	5-11.8	L,M	78	0.005	78	0.005	87	
MW-3	5-12.8	L,M	39	0.005	39	0.005	45	0
MW-5	5~11.9	L	85	0.005	85	0.0075	76	0.005
MW-6	5-15.8	L,H	139		139		130	•••
ve-1	7-12.0	L,M					8	Р

Table 6. SVE Vacuum Influence Data, Former Texaco Service Station, 930 Springtown Boulevard, Livermore, California

Notes:

a = Depth interval below grade surface between top of well screen and ground water (nearest 0.1 feet).

b = Estimated permeability of materials in unsaturated zone around well screen expressed as low, moderate or high; L = Low; M = Moderate; H = Kigh

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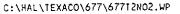
c = Distances between well(s) were taken from site map.

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d = Vacuum influence is reported at final recorded vacuum measurement minus initial background vacuum measurement.

P = Pressure measured or vacuum decreased at monitoring point during test.

--- = Not measured



Test Location (ID) Step Test (#) Applied Vacuum ("H <sub>2</sub> D) Flow Rate (scfm) Step Test Duration (min)			MW-A 1 92 1.1 0-24		MW-B 1 84 2 0-14		MW-5 1 73 19.4 0-22	
Probe/ Well 1D	Exposed Screen Depth <sup>a</sup> (ft-ft)	Est. Perm. in Exposed Screen <sup>D</sup>	Distance <sup>C</sup> from Test Point (ft)	Vacuum <sup>d</sup> Influence ("H <sub>2</sub> O)	Distance <sup>C</sup> from Test Point (ft)	Vacuum <sup>d</sup> Influence ("H <sub>2</sub> O)	Distance <sup>C</sup> from Test Point (ft)	Vacuum <sup>d</sup> Influence ("H <sub>2</sub> 0)
EW-1	8-12.0	L,M	13	0.025	51	0.02	76	p
MW-A	4-12.7	L			60	Ρ	67	0.01
MW-B	4-11.6	L	60	e			126	0.06
MW-2	5-11.8	L,M	96	0	39	0	164	
MW-3	5-12.8	L,M	50	٩	30	0.01	115	0.055
MW-5	5-11.9	L	. 67	0.005	126	Ô	•	•••
MW-6	5-15.8	L,H	120		182	•••	60	
VE-1	7-12.0	L,M	20	0.005	42	0	85	0.005

Table 6. SVE Vacuum Influence Data, Former Texaco Service Station, 930 Springtown Boulevard, Livermore, California (continued)

Notes:

a = Depth interval below grade surface between top of well screen and ground water (nearest 0.1 feet).

b = Estimated permeability of materials in unsaturated zone around well screen expressed as low, moderate or high; L = Low; M = Moderate; H = High c = Distances between well(s) were taken from site map.

d = Vacuum influence is reported at final recorded vacuum measurement minus initial background vacuum measurement.

e = Vacuum influence measurements during test were less than the initial background reading.

P = Pressure measured or vacuum decreased at monitoring point during test.

--- = Not measured

Clock Time	Test Time (minutes)	Pressure ("H2O)	Depth to Water (ft)	Change From Initial H20 Level (ft)	Comments
01:15 PM	0	0.00	12.21	0	Soil Vapor Extraction Start
02:43 PM	88	-0.06	12.18	0.03	
03:29 PM	134	-0.06			
03:39 PM	144	·			Air Sparging Start
03:43 PM	148	0.07	12.08	0.13	
03:44 PM	149	0.105	12.03	0.18	
03:52 PM	157	0.13	12.04	0.17	
04:00 PM	165	0.145	12.09	0.12	
04:15 PM	180	0.175	12.12	0.09	
04:30 PM	195	0.23	12.14	0.07	
04:45 PM	210	0.24	12.15	0.06	
)4:47 PM	212				Air Sparging Off
04:57 PM	222	-0.11	12.26	-0.05	
)5:14 PM	239				Soil Vapor Extraction Off
)5:45 PM	270	0.00	12.10	0.11	

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Table 7. Air Pressure and Water Level Measurements in Well EW-1 During Air Sparging Test

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Clock Time	Test Time (minutes)	Pressure ("H2O)	Depth to Water (ft)	Change From Initial H20 Level (ft)	Comments
12:30 PM	0	-0.005	14.00	0	
01:00 PM	30	-0.01	14.03	-0.03	
01:15 PM	45				Soil Vapor Extraction Start
02:45 PM	135	-0.02	14.02	-0.02	
03:30 PM	180	-0.01	14.02	-0.02	
03:39 PM	189	·			Air Sparging Start
03:47 PM	197	0.00	13.93	0.07	
04:52 PM	262				Air Sparging Off
05:14 PM	284				Soil Vapor Extraction Off
05:20 PM	290	0.00	14.04	-0.04	

Table 8. Air Pressure and Water Level Measurements in MW-A During Air Sparging Test

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Well	Before SVE Test 10/27/92 11:00 AM	Before SVI Sparge Tes 10/28/92 08:45 AM	st	After SVE/ Sparge Test 10/28/92 05:26 PM	Before Pump Test 11/09/92 01:25 PM
EW-1	5.5	3.7	a	5.2	4.2
		4.9	ь		7.4
MW-A	2.9	1.3		2.3	3.3
MW-B	3.9	2.8		2.4	3.7

Notes:

a = Prior to ground water sampling b = After ground water samplingSVE = Soil Vapor Extraction

#### C:\KAO\TEXT3.WQP



Table 9. Dissolved Oxygen Measurements During Air Sparge Testing

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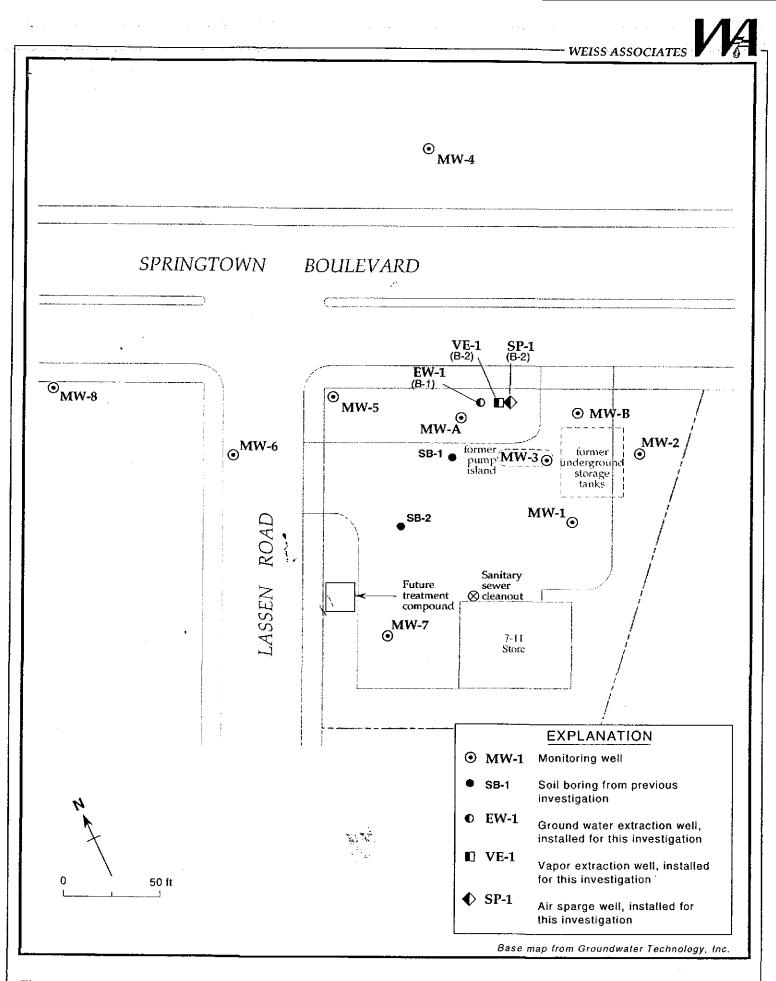
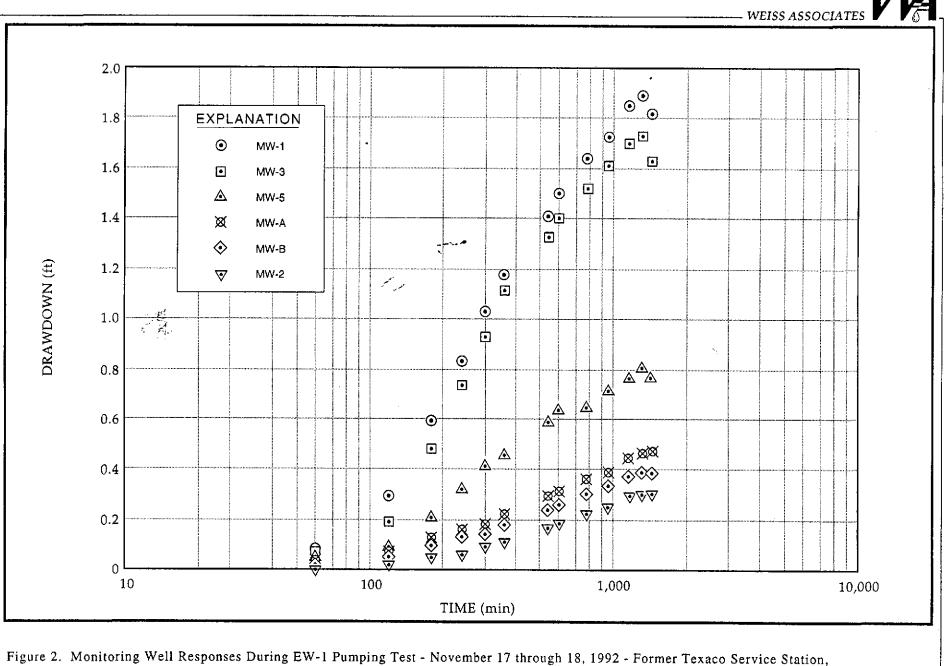


Figure 1. Site Plan and Well Location Map - Texaco Refining and Marketing Inc., 930 Springtown Boulevard, Livermore, Califonia



930 Springtown Boulevard, Livermore, California

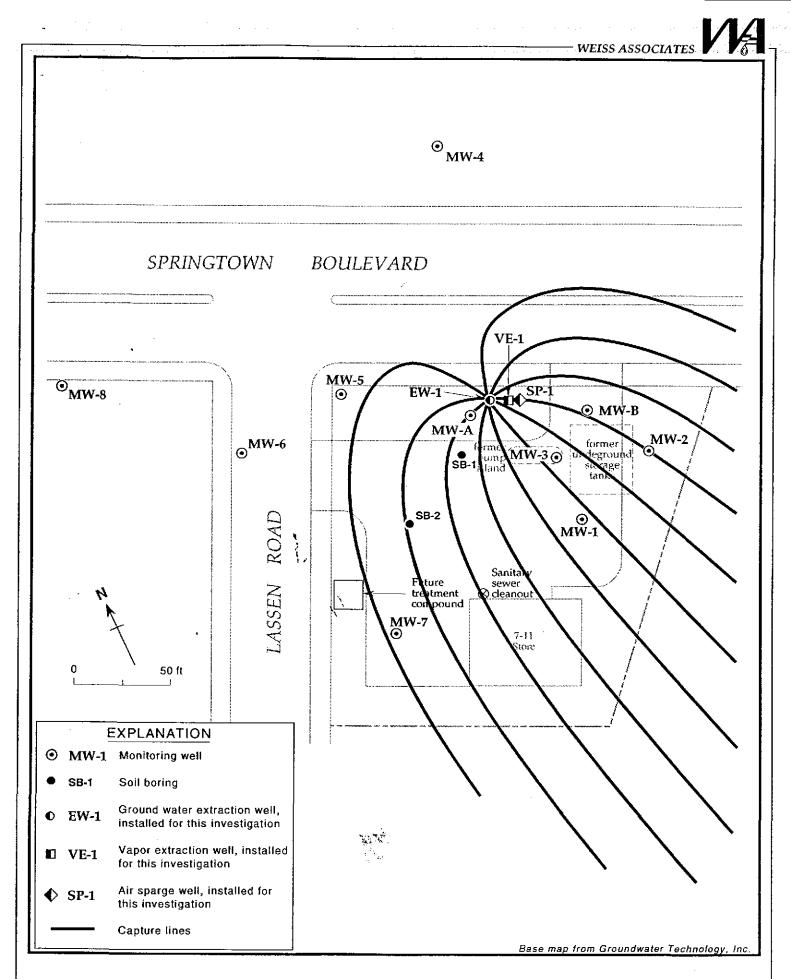


Figure 3. Hydraulic Capture Area for Long Term Pumping of Well EW-1 at 5.0 gpm flow rate -Former Texaco Service Station, 930 Springtown Boulevard, Livermore, Califonia

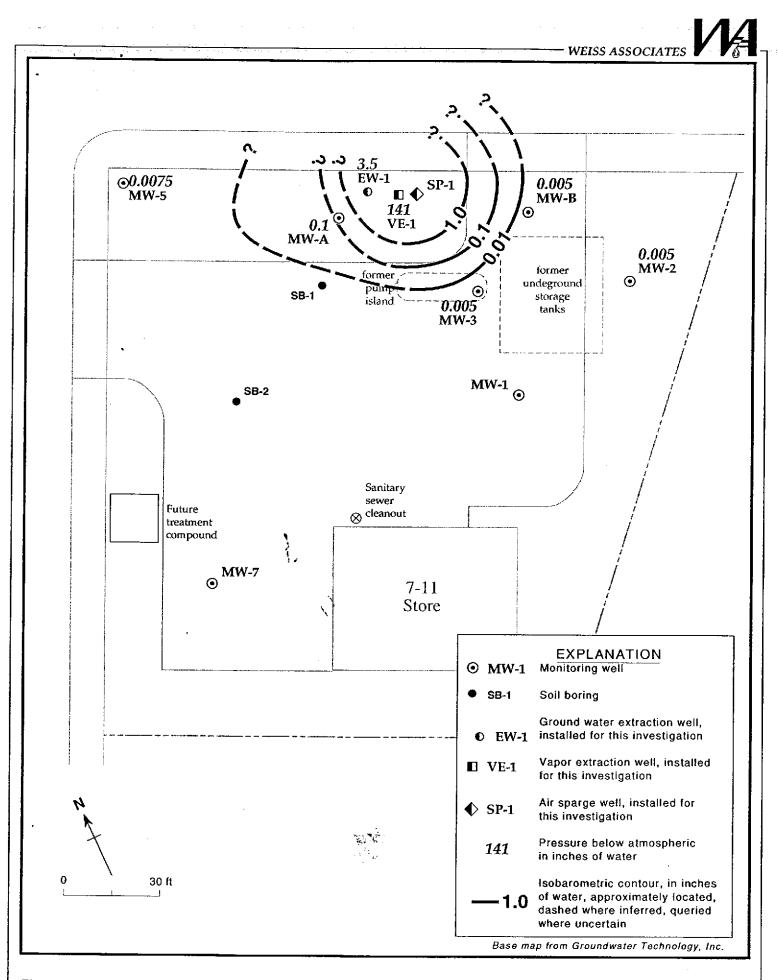
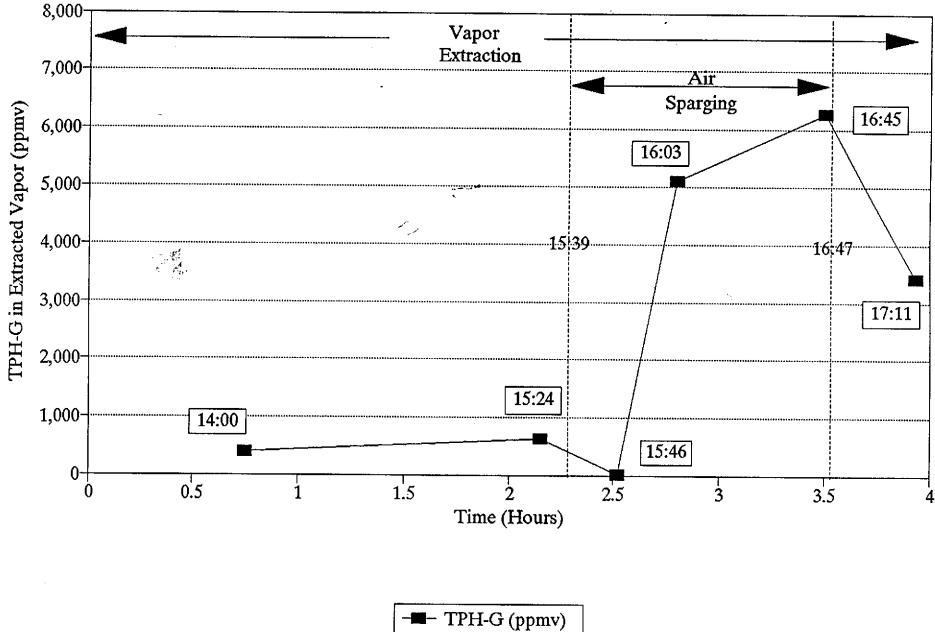
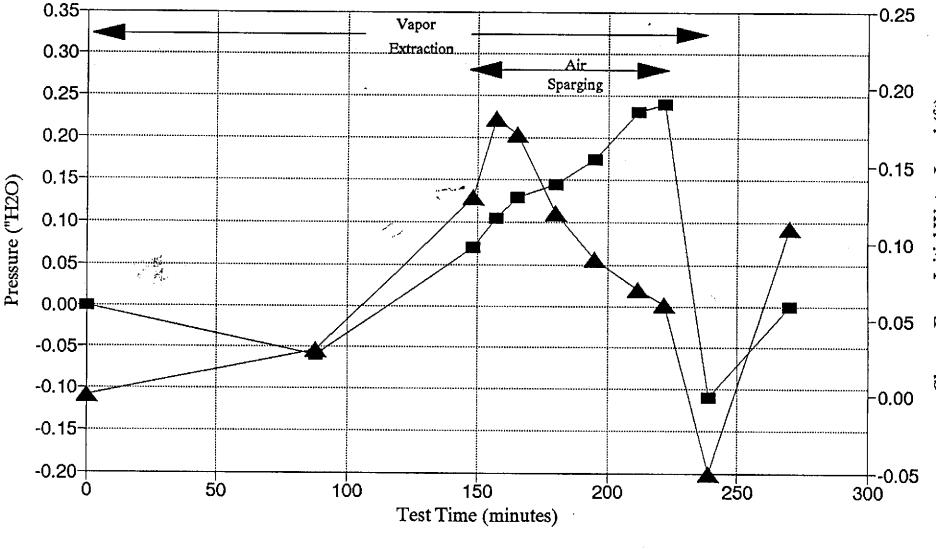


Figure 4. Isobarometric Contour for Vapor Extraction from Well VE-1 - Texaco Refining and Marketing Inc., 930 Springtown Boulevard, Livermore, Califonia

# Figure 5. TPH-G Concentrations in Extracted Vapor During Air Sparging



### Figure 6. Air Pressure and Water Levels in Well EW-1 During Air Sparging



- Pressure ("H2O) - H20 Change (ft)

Change From Initial Water Level (ft)

· · · Weiss Associates

#### APPENDIX A

#### STANDARD FIELD PROCEDURES

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#### STANDARD FIELD PROCEDURES

Weiss Associates (WA) has developed standard procedures for drilling and sampling soil borings and installing, developing and sampling ground water monitoring wells. These procedures comply with Federal, State and local regulatory guidelines. Specific procedures are summarized below.

#### SOIL BORING AND SAMPLING

#### Objectives/Supervision

 Prove the second se second sec

Soil sampling objectives include characterizing subsurface lithology, assessing whether the soils exhibit obvious hydrocarbon or other compound vapor or staining and collecting samples for analysis at a State-certified laboratory. All borings are logged using the Unified Soil Classification System by a trained geologist working under the supervision of a California Registered Geologist (RG) or a Certified Engineering Geologist (CEG).

#### Soil Boring and Sampling

Soil borings are typically drilled using hollow-stem augers. To collect soil samples, split-barrel samplers lined with steam-cleaned brass or stainless steel tubes are driven through the hollow auger stem into undisturbed sediments at the bottom of the borehole using a 140 pound hammer dropped 30 inches. Soil samples can also be collected without using hollow-stem augers by progressively driving split-barrel soil samplers to depths of up to 20 ft.

Soil samples are collected at least every five ft to characterize the subsurface sediments and for possible chemical analysis. Near the water table and at lithologic changes, the sampling interval may be less than five ft.

Drilling and sampling equipment is steam-cleaned prior to drilling and between borings to prevent cross-contamination. Sampling equipment is washed between samples with trisodium phosphate or an equivalent EPA-approved detergent.

 $\sum_{i=1}^{n} \sum_{j=1}^{n}$ 

#### Sample Analysis

After noting the lithology at each end of the sampling tubes, the tube chosen for analysis is immediately trimmed of excess soil and capped with teflon tape and plastic end caps. The sample is labelled, stored at or below 4°C, and transported under chain-ofcustody to a State-certified analytic laboratory.  $(\mathbf{x}, \mathbf{x}) = (\mathbf{x}, \mathbf{y}) + (\mathbf{x$ 

#### Screening

One of the remaining tubes is partially emptied leaving about one-third of the soil in the tube. The tube is capped with plastic end caps and set aside to allow hydrocarbons to volatilize from the soil. Alternatively, soil from the tube is placed in a scaled plastic bag and set in the light. After ten to fifteen minutes, a portable photoionization detector (PID) measures volatile hydrocarbon vapor concentrations in the tube headspace, extracting the vapor through a slit in the cap or hole in the plastic bag. PID measurements are used along with the stratigraphy and ground water depth to select soil samples for analysis.

#### Grouting

If the borings are not completed as wells, the borings are filled to the ground surface with cement grout poured or pumped through a tremie pipe. If wells are completed in the borings, the well installation, development and sampling procedures summarized below are followed.

#### MONITORING WELL INSTALLATION, DEVELOPMENT AND SAMPLING

#### Well Construction and Surveying

Wells are installed to monitor ground water quality and determine the ground water elevation, flow direction and gradient. Well depths and screen lengths are based on ground water depth, occurrence of hydrocarbons or other compounds in the borchole, stratigraphy and state and local regulatory guidelines. Well screens typically extend 15 ft below and 5 ft above the static water level at the time of drilling. However, the well screen will generally not extend into or through a clay layer that is at least three to five ft thick.

Well casing and screen are flush-threaded, Schedule 40 PVC. Screen slot size varies according to the sediments screened, but slots are generally 0.010 or 0.020 inches wide. A rinsed and graded sand occupies the annular space between the boring and the well screen to about one to two ft above the well screen. A two ft thick hydrated bentonite seal separates the sand from the overlying sanitary surface seal composed of cement with 3-5% bentonite.

Well-heads are secured by locking well-caps inside traffic-rated vaults finished flush with the ground surface. A stovepipe may be installed between the well-head and the vault cap for additional security. The well top-of-casing elevation is surveyed with respect to mean sea level and the well is surveyed for horizontal location with respect to an onsite or nearby offsite landmark.

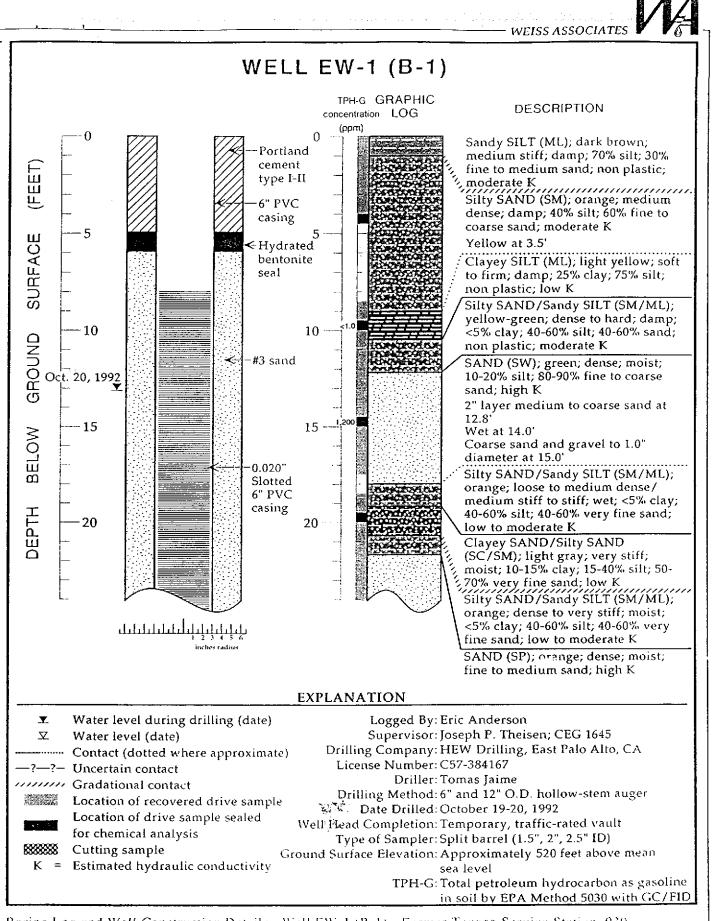
#### Well Development

After at least 24 hours, the wells are developed using a combination of ground water surging and extraction. Surging agitates the ground water and dislodges fine sediments from the sand pack. After about ten minutes of surging, ground water is extracted from the well using bailing, pumping and/or reverse air-lifting through an eductor pipe to remove the sediments from the well. Surging and extraction continue until at least ten well-casing volumes of ground water are extracted and the sediment volume in the ground water is negligible. All equipment is steam-cleaned prior to use and air used for air-lifting is filtered to prevent oil entrained in the compressed air from entering the well. Wells that are developed using air-lift evacuation are not sampled until at least 24 hours after they are developed.

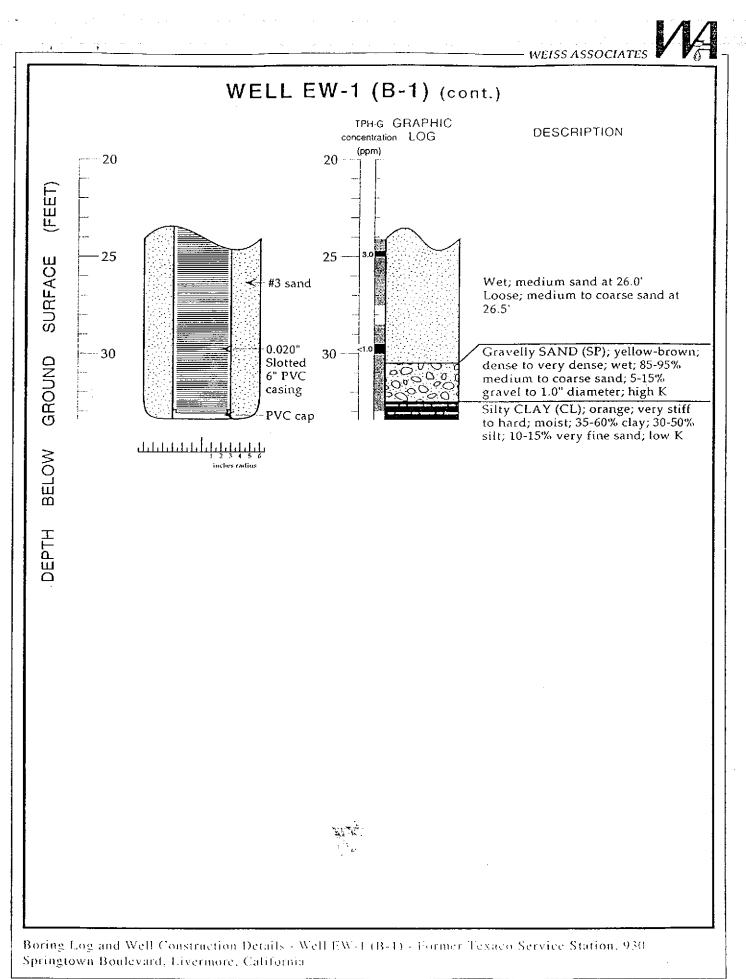
Weiss Associates

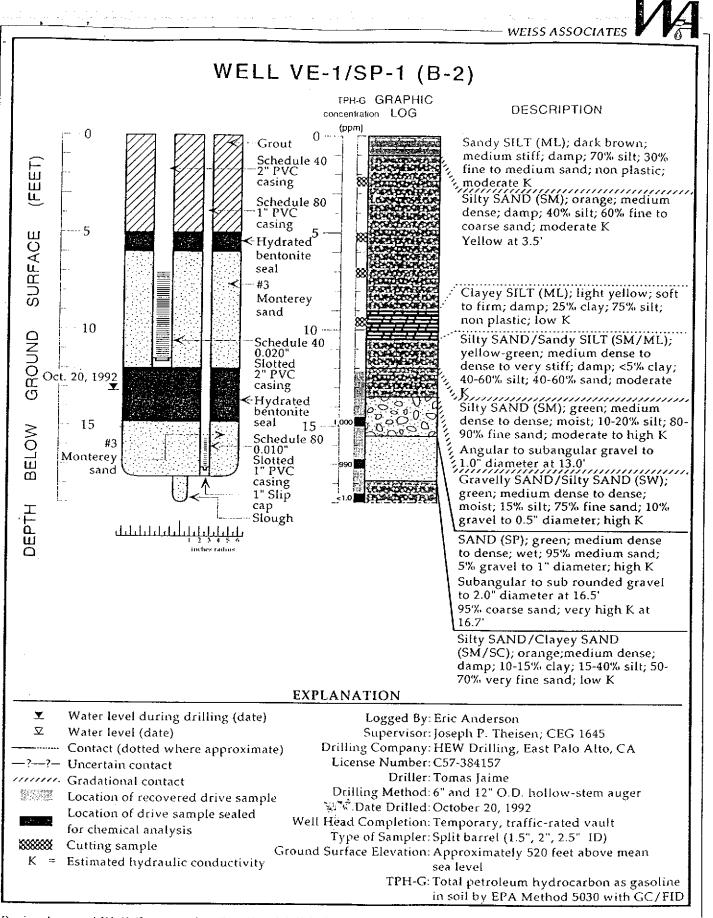
#### Ground Water Sampling

Depending on local regulatory guidelines, three to four well-casing volumes of ground water are purged prior to sampling. Purging continues until ground water pH, conductivity, and temperature have stabilized. Ground water samples are collected using bailers or pumps and are decanted into the appropriate containers supplied by the analytic laboratory. Samples are labelled, placed in protective foam sleeves, stored at or below 4°C, and transported under chain-of-custody to the laboratory. Laboratory-supplied trip blanks accompany the samples and are analyzed to check for cross-contamination. An equipment blank may be analyzed if non-dedicated sampling equipment is used.

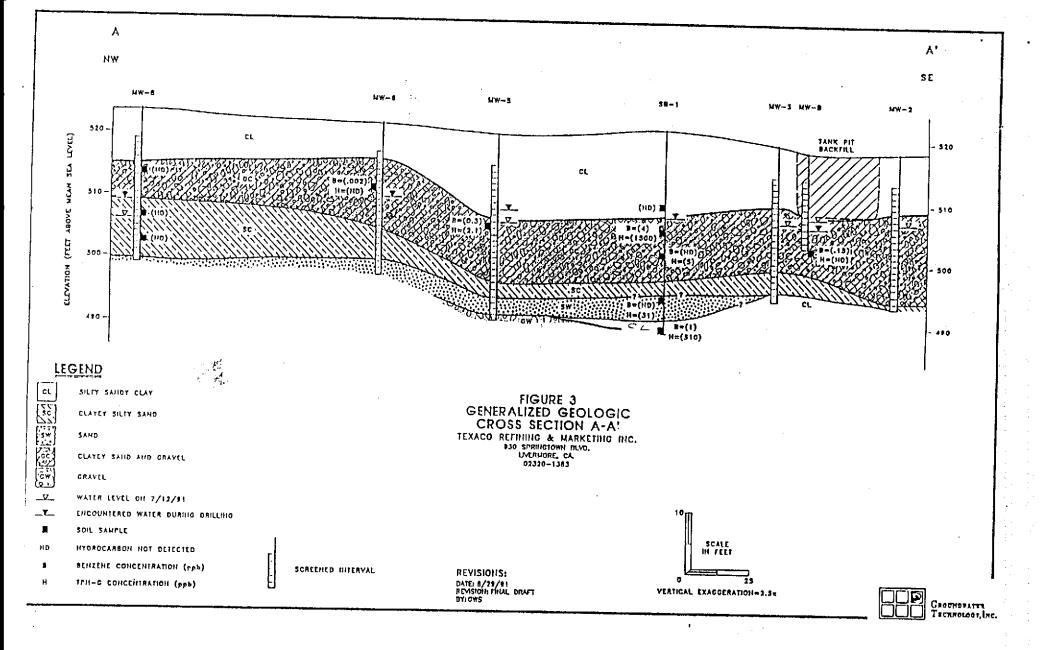


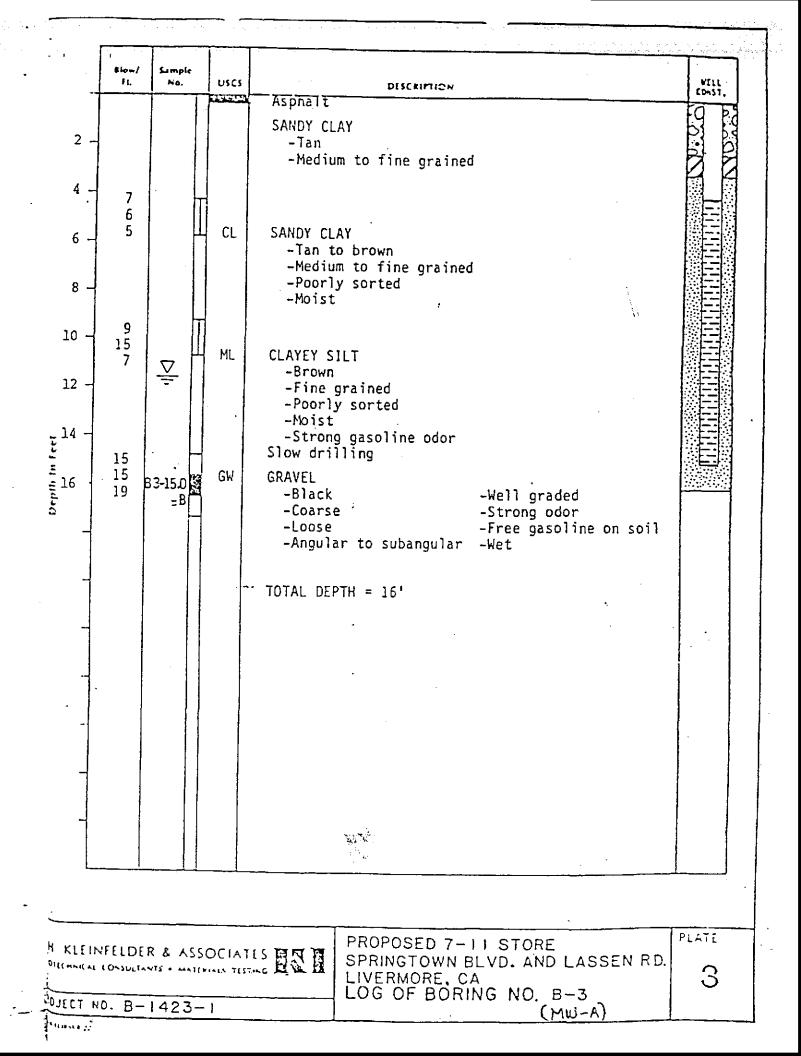
Boring Log and Well Construction Details - Well EW-1 (B-1) - Former Texaco Service Station, 930 Springtown Boulevard, Livermore, California





Boring Log and Well Construction Details - Well VE-1/SP-1 (B-2) - Former Texaco Service Station, 930 Sprintown Boulevard, Livermore, California





Blow/ Sample VILL CGAST, USCS Ŧı, No. DESCRIPTION 3.55 Asphalt SANDY CLAY 2 --Brown to tan -Poorly sorted -Medium to fine grained 4 --Subangular 6 5 CL SANDY CLAY 7 6 --Brown -Poorly sorted -Medium to fine grained 8. -Moist -Strong odor No free gasoline. 8 10 -6 ML CLAYEY SILT 15 -Dark brown Ā 12 --Fine grained -Poorly sorted -Strong odor 14 -Slow drilling 15 Depth In-Feet 22 B4-15 13 GW ; GRAVEL 16 ~B -Black -Wet -Coarse -Free gasoline -Subangular -Strong odor -Loose TOTAL DEPTH = 16' PLATE PROPOSED 7-11 STORE H KLEINFELDER & ASSOCIATES SPRINGTOWN BLVD. AND LASSEN RD 4 LIVERMORE, CA LOG OF BORING NO. 8-4 ОJECT NO. B-1423-1 (MW-B) ---

	GROU TECH Division of Di		WATI .OGY	ER in the second second second	n en segne de la sectar de la sectaria de la secta En la sectaria de la s
				Well Number 1	Drilling Log
Project Texa	co/Livermor	e	Owner	Southland Corp.	Sketch Map
Location Spr.	ingtown&Las	sen	Project	Number 20-4051	
Date Drilled	6-20-85	Total Dep	oth of Hole	25 ft. Diameter 7.5 in.	
				24-hrs <u>11.68</u>	
Screen: Dia.	4-inch	Lengih	<u>20-fe</u>	<u>et</u>	
Casing: Dia	4-inch	Length	<u>5-fe</u>	et Type PVC	· · ·
Drilling Compar	<sub>ny</sub> S <u>ierra Pac</u>	<u>ific</u>	Drilling	Method H.S. Auger	Notes
Driller Lynn	Pera		Log by	Cori Condon	
Depth (Faet)	seitor Notes Salar Cant	Sample Number	Graphic Log	(Color, Texlu	oil Classification re. Structures)
-1- -2- -6- -7.5- -10- -12- -12- -15- 20- 25- -15- -25- -15-	11-12-24	#1 #2		Brown-green silty fine s and gravels, moist, no o	no odor. th subangular white and, stiff.damp, no odor. and with rounded cobbles dor. ine sand, moist, no odor. silt, less cobbles and no odor. wet, no odor. wet, no odor, clay. feet blank
· -11			┝╾╴╺┥		

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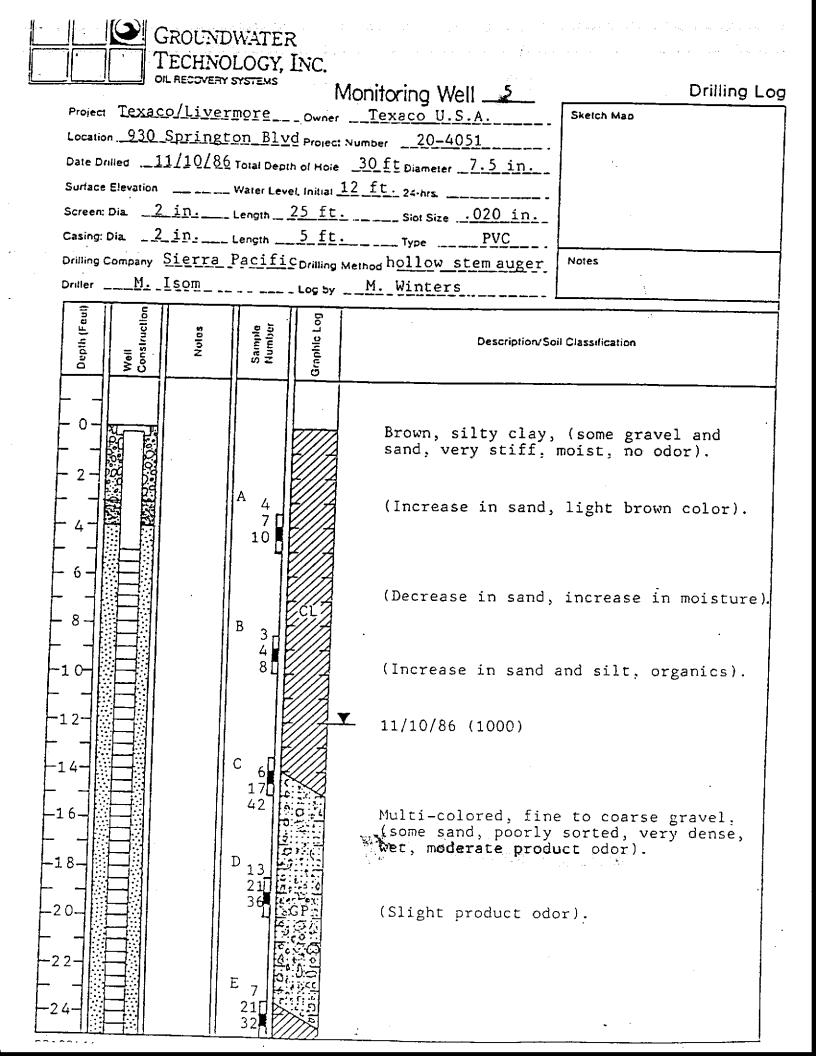
GROUNDWATER TECHNOLOGY Division of Oil Recovery Systems, Inc.

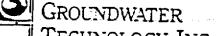
				Well Number	Drilling Log
Project Texac	o/Livermon	ce	_ Owner	Southland Corp.	Sketch Map
				Number 20-4051	х
				24 ft. Diameter 7.5 in.	
-				24-hrs 10.30	
Screen: Dia.	4-inch	Length	20-feet		
Casing: Dia.	4-inch	Length	4-feet	Type PVC	· · · · · · · · · · · · · · · · · · ·
Drilling Company	<u>Sierra Pa</u>	<u>cific</u>	_ Drilling	Method H.S. Auger	Noles
Driller Lynn	<u>Pera</u>	· · · · · · · · · · · · · · · · · · ·	_ Log by .	<u>Cori Condon</u>	
Depth (Feet) Well Construction	ž Ž Blav Cart	Sample Number	Graphic Log	Description/So (Color, T <del>e</del> xtur	Il Classification e, Structurea)
- 1 -				Asphalt and fill.	
				Red-brown clayey sand, oc no odor.	casional gravel, damp,
-9.5- - 10- 	21-33-35	#3		Gray sand and gravel, wet	, no odor.
- 15-	9-25-25	#4		Gray sand and gravel, gravery slight gas odor.	ding to cobbles, wet,
- 20-  - 25-      -	14–56+	Lost Sample		Gray sand and gravel, wet with sandy clay. Drilled 25 feet Cased 20 feet slotted, 4 f Aquarium sand to 3 feet Cement seal to surface Finished with steel manhol	
		- -			· · · ·

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		TECH	_			Drilling Lo
Project	Tevaco	liver	<b>r</b> 0	Churan	Southland Corp	Sketch Map
					Number	
					24_ft_ Diameter7_5_in	
					24-hrs59	
					<u>et</u> Slot Size <u>020. in</u>	
					<u>et</u> Type <u>PVC</u>	
					Method <u>H.S. Auger</u>	Notes
					Cori_Condon	
		11	11			
Depth (Feel)	Well Construction	Notes	Sample Number	ic Log	Description/Sc	bil Classification
Oept	Vell	l 2 Blow Canty	Nun	Graphic	(Color, Textu	re, Structures)
	-0		Ĩ┝		Asphalt and fill.	
- 1-					-	
					Light brown sandy clay w. damp, no odor.	ith occasional gravel,
					Light brown sandy clay wi	th ecceptional evenual
					moist, gasoline odor.	ten occasional glavel,
- 10-						
		13-27-37	# 5	╎┝╴╶┥	Gray sand and gravel, wet	t, slight gasoline odor.
				╟╴╶┥		
- 15-		6-9-19	#6	╠╴╶╡		
				╠╴╶┥	with sandy clay.	t, slight gas odor, contact
┝						
- 20-		5-7-12	#7			
					Mottled sandy clay, moist	, slight gasoline odor.
- 25-		8-22-25	#8			
				<u></u>	Gray sand, wet, no odor.	
-26.5				+ +		
-					Drilled 25 feet	
				F 1	-Cased 20 feet slotted	l, 4 feet blank
				F 1	Aquarium sand to 3 fe Cement seal to surface	
					Finished with steel m	



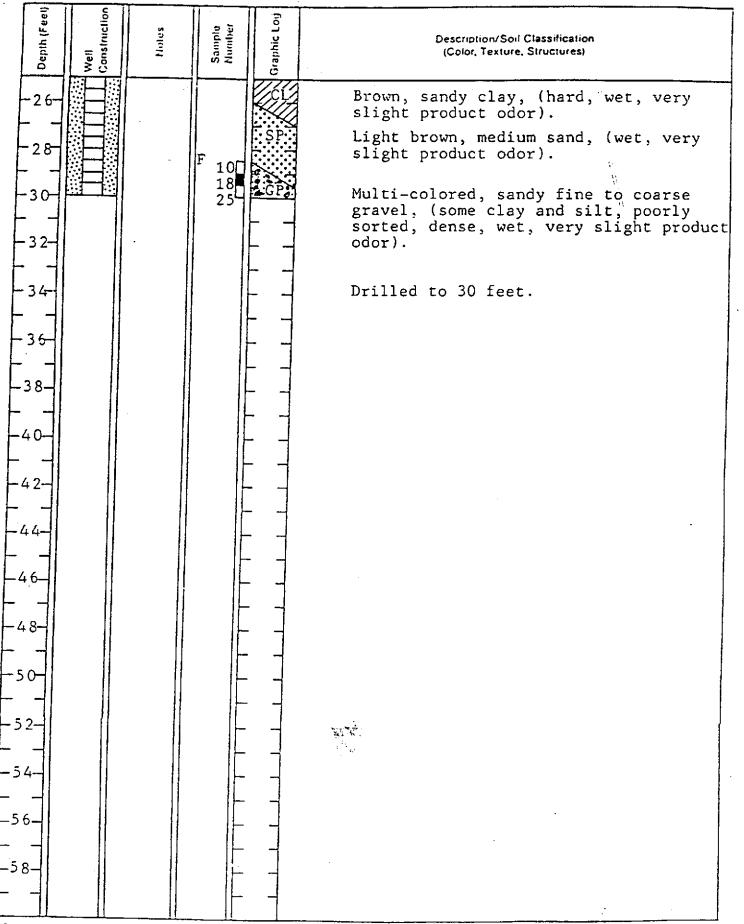


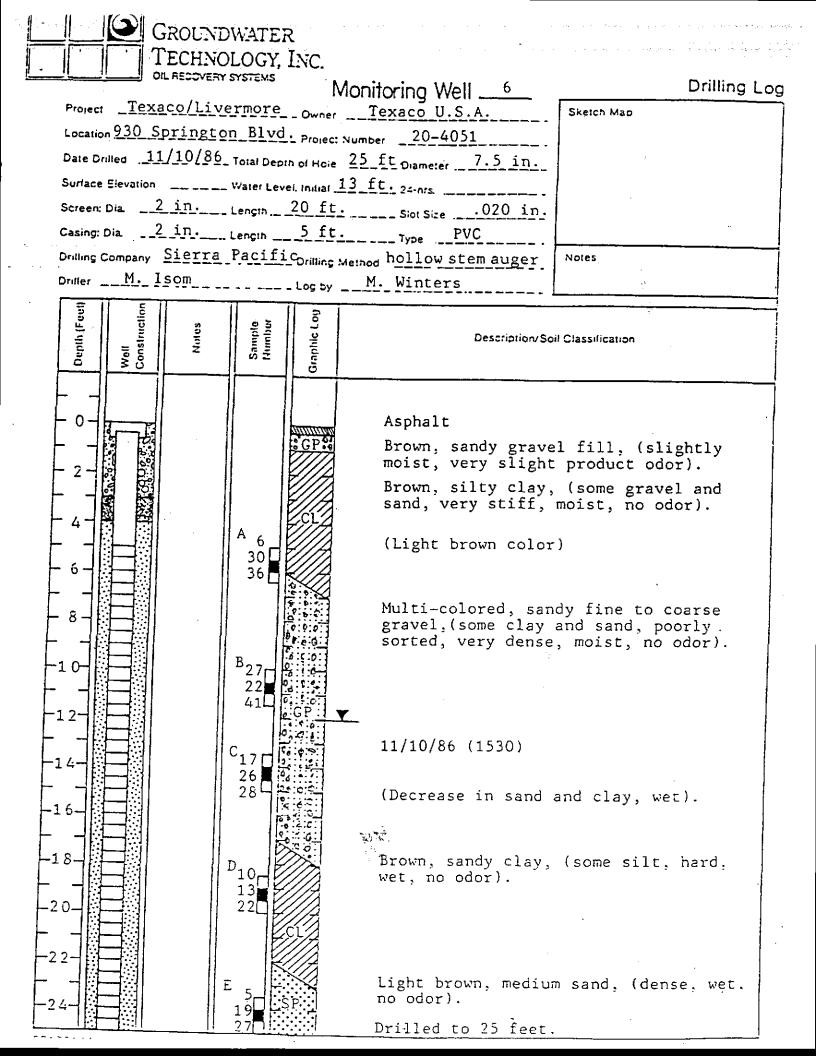
OIL RECOVERY SYSTEMS

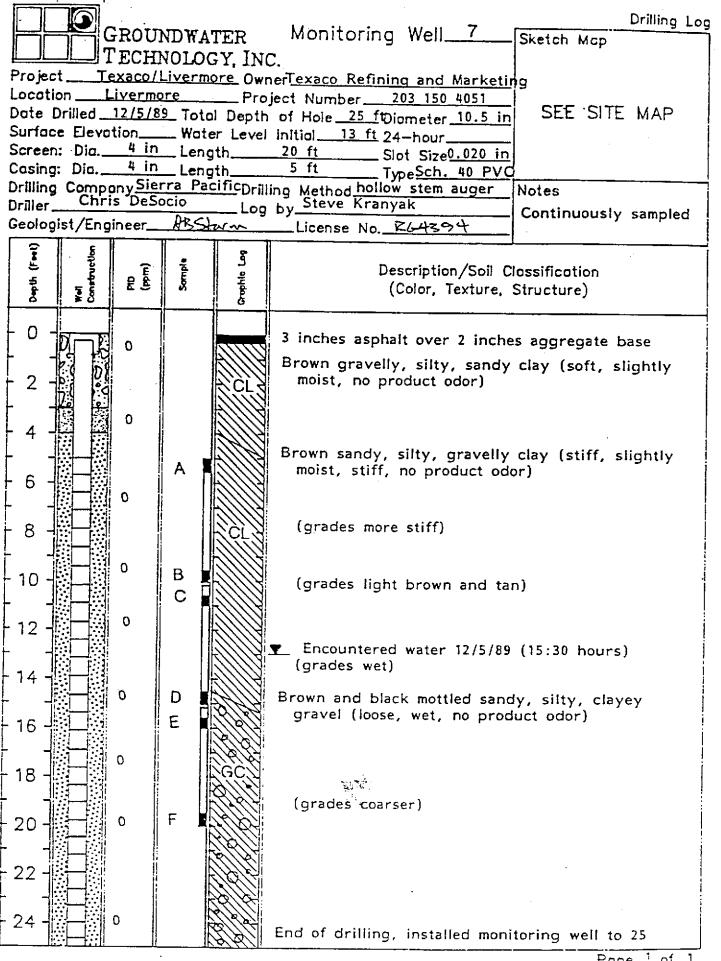
TECHNOLOGY, INC. Monitoring Well <u>5</u>

Drilling Log

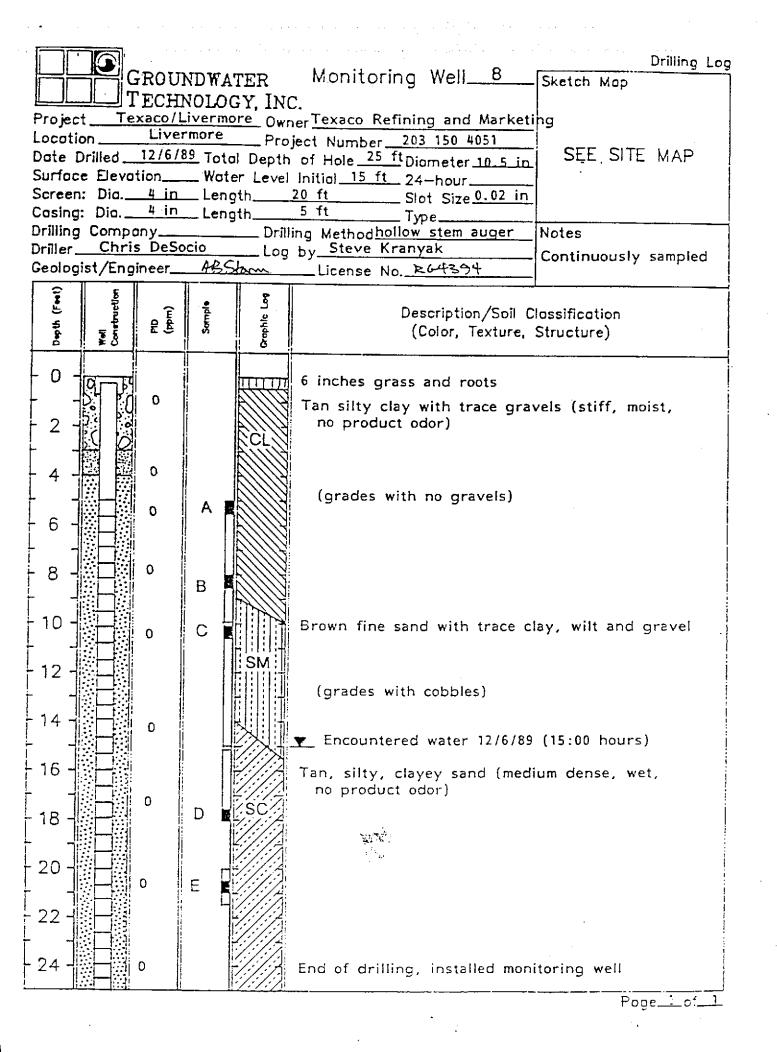
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Weiss Associates



### APPENDIX C

### ANALYTIC REPORTS FOR SOIL, GROUND WATER AND SOIL VAPOR

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Weiss Associates

### MOBILE CHEM LABS INC.

5021 Blum Road, Suite 3 • Martinez, CA 94553 Phone (415) 372-3700 • Fax (415) 372-6955

#### T69-677-01\1342\012201

5500 Shellmound St.		
Emeryville, CA 94611	Date Sampled:	
Attn: Eric Anderson	Date Received:	10-22-92
Project Manager	Date Analyzed:	10-27-92

Sample Number 102281 Sample Description -----Project # T69-677-01 Texaco - Livermore 930 Springstown Blvd. B-1-9.7' SOIL

ANALYSIS

	Detection Limit	Sample Results
	ppm	ppm
Total Petroleum Hydrocarbons as Gasoline	1.0	<1.0
Benzene	0.005	<0.005
Toluene	0.005	<0.005
Xylenes	0.005	<0.005
Ethylbenzene	0.005	<0.005

QA/QC: Sample blank is none detected

Note: Analysis was performed using EPA methods 5030 and TPH LUFT with method 8020 used for BTX distinction. (ppm) = (mg/kg)

.

MOBILE CHEM LABS



5021 Blum Road, Suite 3 • Martinez, CA 94553 Phone (415) 372-3700 • Fax (415) 372-6955

Weiss Associates 5500 Shellmound St. Emeryville, CA 94611 Attn: Eric Anderson Project Manager

т69-677-01	1342	\012201
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Date Sampled: 10-19-92 Date Received: 10-22-92 Date Analyzed: 10-27-92

Sample Number 102282

ANALYSIS

	Detection Limit	Sample Results	
	ppm	ppm	
Total Petroleum Hydrocarbons as Gasoline	1.0	1,200	
Benzene	0.005	6.6	
Toluene	0.005	21	
Xylenes	0.005	50	
Ethylbenzene	0.005	15	

QA/QC: Sample blank is none detected

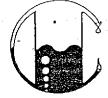
Note:

: Analysis was performed using EPA methods 5030 and TPH LUFT with method 8020 used for BTX distinction. (ppm) = (mg/kg)

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MOBILE CHEM LABS

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5021 Blum Road, Suite 3 • Martinez, CA 94553 Phone (415) 372-3700 • Fax (415) 372-6955

Weiss Associates 5500 Shellmound St. Emeryville, CA 94611 Attn: Eric Anderson Project Manager T69-677-01\1342\012201

and a second second

Date Sampled:	10-19-92
Date Received:	10-22-92
Date Analyzed:	10-27-92

Sample Number

102283

Sample Description ------Project # T69-677-01 Texaco - Livermore 930 Springstown Blvd. B-1-24.7' SOIL

ANALYSIS

		·
	Detection Limit	Sample Results
	ppm	ppm
Total Petroleum Hydrocarbons as Gasoline	1.0	3.0
Benzene	0.005	0.017
Toluene	0.005	0.051
Xylenes	0.005	0.21
Ethylbenzene	0.005	0.050

QA/QC: Sample blank is none detected

Note: Analysis was performed using EPA methods 5030 and TPH LUFT with method 8020 used for BTX distinction. (ppm) = (mg/kg)

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MOBILE CHEM LABS

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5021 Blum Road, Suite 3 • Martinez, CA 94553 Phone (415) 372-3700 • Fax (415) 372-6955

Weiss Associates 5500 Shellmound St. Emeryville, CA 94611 Attn: Eric Anderson Project Manager T69-677-01\1342\012201

Date Sampled: 10-19-92 Date Received: 10-22-92 Date Analyzed: 10-27-92

Sample Number

102284

Sample Description ------Project # T69-677-01 Texaco - Livermore 930 Springstown Blvd. B-1-29.5' SOIL

ANALYSIS

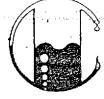
	Detection Limit	Sample Results	
	ppm	ррт	
Total Petroleum Hydrocarbons as Gasoline	1.0	<1.0	
Benzene	0.005	<0.005	
Toluene	0.005	<0.005	
Xylenes	0.005	<0.005	
Ethylbenzene	0.005	<0.005	

QA/QC: Sample blank is none detected

Note: Analysis was performed using EPA methods 5030 and TPH LUFT with method 8020 used for BTX distinction. (ppm) = (mg/kg)

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5021 Blum Road, Suite 3 • Martinez, CA 94553 Phone (415) 372-3700 • Fax (415) 372-6955

Weiss Associates 5500 Shellmound St. Emeryville, CA 94611 Attn: Eric Anderson Project Manager

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Date	Sampled:	10-19-92
Date	Received:	10-22-92
Date .	Analyzed:	10-27-92

Sample Number

102285

Sample Description
Project # T69-677-01
Texaco - Livermore
930 Springstown Blvd.
B-2-14.5' SOIL

ANALYSIS

	Detection Limit	Sample Results	
	ррш	. ppm	
Total Petroleum Hydrocarbons as Gasoline	1.0	1,000	
Benzene	0.005	7.1	
Toluene	0.005	· 22	
Xylenes	0.005	56	
Ethylbenzene	0.005	13	

QA/QC: Sample blank is none detected

Note: Analysis was performed using EPA methods 5030 and TPH LUFT with method 8020 used for BTX distinction. (ppm) = (mg/kg)

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MOBILE CHEM LABS



5021 Blum Road, Suite 3 • Martinez, CA 94553 Phone (415) 372-3700 • Fax (415) 372-6955

Weiss Associates 5500 Shellmound St. Emeryville, CA 94611 Attn: Eric Anderson Project Manager т69-677-01\1342\012201

Date Sampled:	10-19-92
Date Received:	10-22-92
Date Analyzed:	10-27-92

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Sample Number

102286

Sample Description Project # T69-677-01 Texaco - Livermore 930 Springstown Blvd. B-2-16.7' SOIL

ANALYSIS

	Detection Limit	Sample Results		
	ppm	ppm		
Total Petroleum Hydrocarbons as Gasoline	1.0	<b>990</b>		
Benzene	0.005	2.9		
Toluene	0.005	15		
Xylenes	0.005	53		
Ethylbenzene	0.005	14		

QA/QC: Sample blank is none detected

Note: Analysis was performed using EPA methods 5030 and TPH LUFT with method 8020 used for BTX distinction. (ppm) = (mg/kg)

MOBILE CHEM LABS

Ronald G. Evans

5021 Blum Road, Suite 3 • Martinez, CA 94553 Phone (415) 372-3700 • Fax (415) 372-6955

Weiss Associates 5500 Shellmound St. Emeryville, CA 94611 Attn: Eric Anderson Project Manager т69-677-01\1342\012201

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Date Sampled: 10-19-92 Date Received: 10-22-92 Date Analyzed: 10-27-92

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Sample Number

102287

Sample Description Project # T69-677-01 Texaco - Livermore 930 Springstown Blvd. B-2-18.5' SOIL

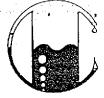
ANALYSIS

	Detection	Sample	
	Limit	Results	
	ppm	ppm	
Total Petroleum Hydrocarbons as Gasoline	1.0	<1.0	
Benzene	0.005	0.007	
Toluene	0.005	0.029	
Xylenes	0.005	<0.005	
Ethylbenzene	0.005	<0.005	

QA/QC: Sample blank is none detected

Note: Analysis was performed using EPA methods 5030 and TPH LUFT with method 8020 used for BTX distinction. (ppm) = (mg/kg)

MOBILE CHEM LABS



5021 Blum Road, Suite 3 • Martinez, CA 94553 Phone (415) 372-3700 • Fax (415) 372-6955

Weiss Associates 5500 Shellmound St. Emeryville, CA 94611 Attn: Eric Anderson Project Manager

#### т69-677-01\1342\012201

Date Sampled:	10-19-92
Date Received:	10-22-92
Date Analyzed:	10-27-92

Sample Number

102288

Sample Description

Project # T69-677-01 Texaco - Livermore 930 Springstown Blvd. SP1 SOIL

#### ANALYSIS

	Detection	Sample
	Limit - <b></b>	Results
	ppm	ppm
Total Petroleum Hydrocarbons as Gasoline	1.0	66

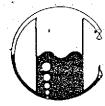
QA/QC: Sample blank is none detected

Note: Analysis was performed using EPA methods 5030 and TPH LUFT with method 8020 used for BTX distinction. (ppm) = (mg/kg)

.

MOBILE CHEM LABS

Ronald G. Evans



5021 Blum Road, Suite 3 • Martinez, CA 94553 Phone (415) 372-3700 • Fax (415) 372-6955

Weiss Associates 5500 Shellmound St. Emeryville, CA 94611 Attn: Eric Anderson Project Manager T69-677-01\1342\012201

Date Sampled: 10-19-92 Date Received: 10-22-92 Date Analyzed: 10-27-92

Sample Number 102289 ANALYSIS

Detection Limit	Sample Results
ppm	ppm
1.0	96

Total Petroleum Hydrocarbons as Gasoline

QA/QC: Sample blank is none detected

Note: Analysis was performed using EPA methods 5030 and TPH LUFT with method 8020 used for BTX distinction. (ppm) = (mg/kg)

MOBILE CHEM LABS



5021 Blum Road, Suite 3 • Martinez, CA 94553 Phone (415) 372-3700 • Fax (415) 372-6955

Weiss Associates 5500 Shellmound St. Emeryville, CA 94611 Attn: Bob Riddell Project Manager T69-677-02\1342\012272

Date Sampled:	11-17-92
Date Received:	11-19-92
Date Analyzed:	11-23-92

Sample Number 112301 Sample Description Project # T69-677-02 Texaco - Livermore 930 Springtown Influent WATER

ANALYSIS

	Detection Limit	Sample Results		
	ppb	 dqq.		
Total Petroleum Hydrocarbons as <b>Gaso</b> line	50	4,300		
Benzene	0.5	140		
Toluene	0.5	340		
Xylenes	0.5	560		
Ethylbenzene	0.5	96		

QA/QC: Sample blank is none detected Duplicate Deviation is 6.8%

Note: Analysis was performed using EPA methods 5030 and TPH LUFT with method 602 used for BTX distinction. (ppb) =  $(\mu g/L)$ 

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MOBILE CHEM LABS

CURING-



5021 Blum Road, Suite 3 • Martinez, CA 94553 Phone (415) 372-3700 • Fax (415) 372-6955

Weiss Associates 5500 Shellmound St. Emeryville, CA 94611 Attn: Bob Riddell Project Manager **Т69-677-02\1342\012272** 

Date Sampled: 11-17-92 Date Received: 11-19-92 Date Analyzed: 11-23-92

Sample Number

112302

Sample Description Project # T69-677-02 Texaco - Livermore 930 Springtown Effluent WATER

ANALYSIS

	Detection Limit	Sample Results
	ppb	ppb
Total Petroleum Hydrocarbons as Gasoline	50	<50
Benzene	0.5	<0.5
Toluene	0.5	<0.5
Xylenes	0.5	<0.5
Ethylbenzene	0.5	<0.5

QA/QC: Sample blank is none detected

Note: Analysis was performed using EPA methods 5030 and TPH LUFT with method 602 used for BTX distinction. (ppb) = ( $\mu$ g/L)

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MOBILE CHEM LABS

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WEISS ASSOCIATES 5500 Shellmound Street, Emeryville, CA 94608 Phone: 4155475420 Fax: 4155475043 CHAIN-OF-CUSTODY RECORD AND ANALYTIC INSTRUC		orm to:		<ul> <li>PLEASE INCLUDE QA/QC DATA IF I CHECKED.</li> <li>Specify analytic method and detection limit in report.</li> <li>Notify us if there are any anon peaks in GC or other scans.</li> <li><u>ANY</u> QUESTIONS/CLARIFICATIONS: <u>US.</u></li> </ul>	omalous
Sampled by: CHARLES	Vol <sup>2</sup> Fil <sup>3</sup> Ref <sup>4</sup> Preservative (specify)	CHEM. LAB Analyze for TPH. G/BETX	Analytic Method XOIS / 803 b ,	Turn <sup>5</sup> COMMENTS	
1 Released by (Signature), Date 1/1/2-92 Affiliation 2/10-10-10-11-17-92 Received DM(Signature), Date 2/10-10-10-10-11-17-92 Received DM(Signature), Date 2/10-10-10-10-10-10-10-10-10-10-10-10-10-1	3 Detterment II 19/9 Released by (Sygnature), Date 3 MUSS Affiliation 4 Law Law Me Shipping Carrier, Method, Date 4 Mobile Cham Labs Inc. Affiliation	2 5 Released by (Sign 5 Affiliation 6 Received by Lab P 6 Affiliation, Tele	ersonnel, Date	x Seal intact?	

1 Sample Type Codes: W = Water, S = Soil, Describe Other; Container Type Codes: V = VOA/Teflon Septa, P = Plastic, C or B - Clear/Brown Glass, Describe Other; Cap Codes: PT = Plastic, Teflon Lined 2 = Volume per container; 3 = Filtered (Y/N); 4 = Refrigerated (Y/N) Page <u>1</u> of <u>1</u>

5 Turnaround [N = Normal, W = 1 Week, R = 24 Hour, HOLD (write out)]

ADDITIONAL COMMENTS, CONDITIONS, PROBLEMS:

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• Weiss Associates 03/08/91



5021 Blum Road, Suite 3 • Martinez, CA 94553 Phone (415) 372-3700 • Fax (415) 372-6955

т69-677-02\1223\012220

Weiss Associates
5500 Shellmound St.
Emeryville, CA 94611
Attn: Bob Riddell
Project Manager

🕂 🕅 Date Sampled:	10-27-92
Date Received:	
Date Analyzed:	11-11-92

Sample Number 102420 Sample Description Project # T69-677-02 Texaco - Livermore 930 Springtown 102200184 WATER

#### ANALYSIS

	Detection Limit  ppb	Sample Results ppb
Total Petroleum Hydrocarbons as Gasoline	50	11,000
Benzene	0.5	410-
Toluene	0.5	2,000
Xyleneş	0.5	2,100
Ethylbenzene	0.5	540

QA/QC: Sample blank is none detected Duplicate Deviation is 12.1%

Note: Analysis was performed using EPA methods 5030 and TPH LUFT with method 602 used for BTX distinction. (ppb) =  $(\mu g/L)$ 

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MOBILE CHEM LABS

Ronald G. Evans Lab Director



5021 Blum Road, Suite 3 • Martinez, CA 94553 Phone (415) 372-3700 • Fax (415) 372-6955

т69-677-02\1223\012220

Weiss Associates 5500 Shellmound St. Emeryville, CA 94611 Attn: Bob Riddell Project Manager

Date Sampled: 10-27-92 Date Received: 10-29-92 Date Analyzed: 11-11-92

Sample Number ------102421 (1,1,1,1) , we are a single state of the second state  $\Psi_{1}$  , then  $\Psi_{2}$ 

ANALYSIS

	Detection Limit	Sample Results
	ppb	bby
Total Petroleum Hydrocarbons as Gasoline	50	13,000
Benzene	0.5	840
Toluene	0.5	2,400
Xylenes	0.5	1,900
Ethylbenzene	0.5	580

QA/QC: Sample blank is none detected

Note: Analysis was performed using EPA methods 5030 and TPH LUFT with method 602 used for BTX distinction. (ppb) =  $(\mu g/L)$ 

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5021 Blum Road, Suite 3 • Martinez, CA 94553 Phone (415) 372-3700 • Fax (415) 372-6955

T69-677-02\1342\012219

Weiss Associates 5500 Shellmound Street Emeryville, CA 94608 Attn: Eric Anderson Project Manager

Date Sampled: Date Received:	10-27-92
Date Received:	10-27-92
Date Reported:	10-29-92

Sample Number 102398 المتحدية والمناجر المراجع والمروحية ومنجوا كالمراجع والمروح

ANALYSIS

a. 1.

	Detection Limit	Sample Results
	mg/m <sup>3</sup>	mg/m <sup>3</sup>
Total Petroleum Hydrocarbons as Gasoline	2.0	<b>`8</b> ∳ <b>∂∂90</b>
Benzene	0.05	30
Toluene	0.05	7.8
Xylenes	0.05	34
Ethylbenzene	0.05	21

QA/QC: Sample blank is none detected

Note: Analysis was performed using EPA methods 5030 and TPH LUFT with method 8020 used for BTX distinction.

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INDRA



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т69-677-02\1342\012219

Weiss Associates 5500 Shellmound Street Emeryville, CA 94608 Attn: Eric Anderson Project Manager

Date Sampled: 10-27-92 Date Received: 10-27-92 Date Reported: 10-29-92

Sample Description Project #T69-677-02 Texaco - Livermore 930 Springtown MM-D AIR

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#### ANALYSIS

	Detection Limit	Sample Results
	 mg/m <sup>3</sup>	mg/m <sup>3</sup>
Total Petroleum Hydrocarbons as Gasoline	2.0	65 <b>, 990</b>
Benzene	0.05	1,100
Toluene	0.05	260
Xylenes	0.05	210
Ethylbenzene	0.05	150

QA/QC: Sample blank is none detected

Note: Analysis was performed using EPA methods 5030 and TPH LUFT with method 8020 used for BTX distinction.

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Ronald G. Evans Lab Director



5021 Blum Road, Suite 3 • Martinez, CA 94553 Phone (415) 372-3700 • Fax (415) 372-6955

т69-677-02\1342\012219

Weiss Associates 5500 Shellmound Street Emeryville, CA 94608 Attn: Eric Anderson Project Manager

Date Sampled: Date Received:	10-27-92
Date Received:	10-27-92
Date Reported:	10-29-92

Sample Number 102400 Sample Description Project #T69-677-02 Texaco - Livermore 930 Springtown MW-5

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ANALYSIS

	Detection Limit	Sample Results
	mg/m <sup>3</sup>	mg/m³
Total Petroleum Hydrocarbons as Gasoline	2.0	72,000
Benzene	0.05	360
Toluene	0.05	14
Xylenes	0.05	56
Ethylbenzene	0.05	5.3

QA/QC: Sample blank is none detected Duplicate Deviation is 6.5%

Note: Analysis was performed using EPA methods 5030 and TPH LUFT with method 8020 used for BTX distinction.

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т69-677-02\1342\012219

Weiss Associates	
5500 Shellmound Street	
Emeryville, CA 94608	
Attn: Eric Anderson	
Project Manager	

Date Sampled:	10-27-92
Date Received:	10-27-92
Date Reported:	10-29-92

Sample Number

102401

Sample Description Project #T69-677-02 Texaco - Livermore 930 Springtown VE-1 Start AIR

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#### ANALYSIS

	Detection Limit	Sample Results
	 mg/m <sup>3</sup>	mg/m <sup>3</sup>
Total Petroleum Hydrocarbons as Gasoline	2.0	4,600
Benzene	0.05	56
Toluene	0.05	67
Xylenes	0.05	60
Ethylbenzene	0.05	22

QA/QC: Sample blank is none detected

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Note: Analysis was performed using EPA methods 5030 and TPH LUFT with method 8020 used for BTX distinction.

MOBILE CHEM LABS

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Ronald G. Evans Lab Director



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т69-677-02\1342\012219

Weiss Associates 5500 Shellmound Street		
Emeryville, CA 94608 Attn: Eric Anderson Project Manager	Date Sampled: Date Received: Date Reported:	

Sample Number 102402

Sample Description
Project #T69-677-02
Texaco - Livermore
930 Springtown
VE-1 End AIR

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ANALYSIS

	Detection Limit	Sample Results	
	 mg/m <sup>3</sup>	 mg/m <sup>3</sup>	
Total Petroleum Hydrocarbons as Gasoline	2.0	680	
Benzene	0.05	8.4	
Toluene	0.05	6.4	
Xylenes	0.05	5.9	
Ethylbenzene	0.05	2.6	

QA/QC: Sample blank is none detected

Note: Analysis was performed using EPA methods 5030 and TPH LUFT with method 8020 used for BTX distinction.

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Stwan r

Ronald G. Evans Lab Director



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т69-677-02\1342\012219

Weiss Associates
5500 Shellmound Street
Emeryville, CA 94608
Attn: Eric Anderson
Project Manager

Date Sampled:	10-27-92
Date Sampled: Date Received:	10-27-92
Date Reported:	10-29-92

Sample Number

102403

Sample Description
Project #T69-677-02
Texaco - Livermore
930 Springtown
EW-1 AIR

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ANALYSIS

	Detection Limit	Sample Results	
	mg/m³	mg/m <sup>3</sup>	
Total Petroleum Hydrocarbons as Gasoline	2.0	1,500	
Benzene	0.05	12	
Toluene	0.05	10	
Xylenes	0.05	12	
Ethylbenzene	0.05	5.0	

QA/QC: Sample blank is none detected

Note: Analysis was performed using EPA methods 5030 and TPH LUFT with method 8020 used for BTX distinction.

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MOBILE CHEM LABS

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WEISS ASSOCIATES 5500 Shellmound Street, Emeryville, CA 94608 Phone: 415-547-5420 Fax: 415-547-5043 CHAIN-OF-CUSTODY RECORD AND ANALYTIC INSTRUCTION			<ul> <li>PLEASE INCLUDE QA/QC DATA IF BOX IS CHECKED.</li> <li>Specify analytic method and detection limit in report.</li> <li>Notify us if there are any anomalous peaks in GC or other scans.</li> <li>ANY QUESTIONS/CLARIFICATIONS: CALL US.</li> </ul>
	ol <sup>2</sup> Fil <sup>3</sup> Ref <sup>4</sup> Preservative	ANTIMET CA Analyze for Analytic	Turn <sup>5</sup> COMMENTS
Containers       Type       Date $2$ $MW-A$ $Arr       10-27-92/M 2 MW-B 11 11 2 MW-S 11 11 2 MW-S 11 11 2 MW-S 11 11 2 VE-1 START       11 11 2 VE-1 END       11 11 11 $	(specify) $(specify)$ $(spe$	Method           S, BETX         SOIS, 8020           (1         1/           11         1/           11         11           11         11           11         11           11         11           11         11           11         11           11         11           11         11           11         11           11         11	N=72-HBS Hold DrPE $(1)$ $($
$\frac{1}{10000000000000000000000000000000000$	= Volume per container; 3 = Filtered (Y	Released by (Signature), Date 5 Affiliation 6 Received by Lab Personnel, Date 6 Affiliation, Telephone = VOA/Teflon Septa, P = Plastic, C or	x Seal intact? B - Clear/Brown Glass, Describe Other; Page 1 of 1

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• Weiss Associates 03/08/91



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т69-677-02\1342\012220

Weiss Associates 5500 Shellmound Street Emeryville, CA 94608 Attn: Eric Anderson Project Manager

Date Sampled:	10-28-92
Date Received:	10-29-92
Date Reported:	10-30-92

Sample Number 102422 Sample Description Project #T69-677-02 Texaco - Livermore 930 Springtown VE-1 1400 AIR

ANALYSIS

	Detection Limit	Sample Results
	mg/m³	mg/m³
Total Petroleum Hydrocarbons as Gasoline	2.0	1,400
Benzene	0.05	11
Toluene	0.05	10
Xylenes	0.05	14
Ethylbenzene	0.05	4.4

QA/QC: Sample blank is none detected

Note: Analysis was performed using EPA methods 5030 and TPH LUFT with method 8020 used for BTX distinction.

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т69-677-02\1342\012220

Weiss Associates			
5500 Shellmound	Street		
Emeryville, CA	94608		
Attn: Eric Ande	rson		
Project Manager			

MA Date Sampled:	10-28-92
Date Received:	10-29-92
Date Reported:	10-30-92

Sample Number 102423

Sample Desc	ription
Project #T6	
Texaco - Li	vermore
930 Springt	lown
VE-1 1524	AIR

A Sugar St.

#### ANALYSIS

	Detection Limit	Sample Results	
	$mg/m^3$	mg/m³	
Total Petroleum Hydrocarbons as Gasoline	2.0	2,200	
Benzene	0.05	17	
Toluene	0.05	8.1	
Xylenes	0.05	13	
Ethylbenzene	0.05	4.3	

QA/QC: Sample blank is none detected

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Note: Analysis was performed using EPA methods 5030 and TPH LUFT with method 8020 used for BTX distinction.

MOBILE CHEM LABS

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Ronald G. Evans Lab Director



5021 Blum Road, Suite 3 • Martinez, CA 94553 Phone (415) 372-3700 • Fax (415) 372-6955

т69-677-02\1342\012220

Weiss Associates 5500 Shellmound Street Emeryville, CA 94608 Attn: Eric Anderson Project Manager

Date Sampled: 10-28-92 Date Received: 10-29-92 Date Reported: 10-30-92

Sample Description Project #T69-677-02 Texaco - Livermore 930 Springtown VE-1 1546 AIR

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ANALYSIS

	Detection Limit	Sample Results
	mg/m <sup>3</sup>	mg/m³
Total Petroleum Hydrocarbons as Gasoline	10.0	<10
Benzene	0.05	<0.05
Toluene	0.05	<0.05
Xylenes	0.05	<0.05
Ethylbenzene	0.05	<0.05

QA/QC: Sample blank is none detected

Note: Analysis was performed using EPA methods 5030 and TPH LUFT with method 8020 used for BTX distinction.

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5021 Blum Road, Suite 3 • Martinez, CA 94553 Phone (415) 372-3700 • Fax (415) 372-6955

т69-677-02\1342\012220

Weiss Associates 5500 Shellmound Street Emeryville, CA 94608 Attn: Eric Anderson Project Manager

Date Sampled: 10-27-92 Date Received: 10-29-92 Date Reported: 10-30-92

Sample Number 102425 Sample Description Project #T69-677-02 Texaco - Livermore 930 Springtown VE-1 1603 AIR

ANALYSIS

<b></b>	Detection Limit	Sample Results
	mg/m <sup>3</sup>	mg/m <sup>3</sup>
Total Petroleum Hydrocarbons as Gasoline	2.0	18,000
Benzene	0.05	63
Toluene	0.05	10
Xylenes	0.05	10
Ethylbenzene	0.05	4.6

QA/QC: Sample blank is none detected

Note: Analysis was performed using EPA methods 5030 and TPH LUFT with method 8020 used for BTX distinction.

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5021 Blum Road, Suite 3 • Martinez, CA 94553 Phone (415) 372-3700 • Fax (415) 372-6955

Weiss Associates 5500 Shellmound Street Emeryville, CA 94608 Attn: Eric Anderson Project Manager т69-677-02\1342\012220

Date Sampled: 10-27-92 Date Received: 10-29-92 Date Reported: 10-30-92

Sample Number 102426 Sample Description Project #T69-677-02 Texaco - Livermore 930 Springtown VE-1 1645 AIR

#### ANALYSIS

	Detection Limit	Sample Results
	 mg/m <sup>3</sup>	 mg/m <sup>3</sup>
Total Petroleum Hydrocarbons as Gasoline	2.0	22,000
Benzene	0.05	260
Toluene	0.05	130
Xylenes	0.05	60
Ethylbenzene	0.05	29

QA/QC: Sample blank is none detected

Note: Analysis was performed using EPA methods 5030 and TPH LUFT with method 8020 used for BTX distinction.

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5021 Blum Road, Suite 3 • Martinez, CA 94553 Phone (415) 372-3700 • Fax (415) 372-6955

т69-677-02\1342\012220

Weiss Associates	3
5500 Shellmound	Street
Emeryville, CA	94608
Attn: Eric Ander	rson
Project Ma	anager

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Date Sampled:	10-27-92
Date Received:	10-29-92
Date Reported:	10-30-92

Sample Number 102427 Sample Description Project #T69-677-02 Texaco - Livermore 930 Springtown VE-1 1711 AIR

#### ANALYSIS

	Detection Limit	Sample Results
	 mg/m <sup>3</sup>	mg/m <sup>3</sup>
Total Petroleum Hydrocarbons as Gasoline	2.0	12,000
Benzene	0.05	170
Toluene	0.05	120
Xylenes	0.05	54
Ethylbenzene	Û.05	29

QA/QC: Sample blank is none detected

Note: Analysis was performed using EPA methods 5030 and TPH LUFT with method 8020 used for BTX distinction.

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WEISS ASSOCIATES 5500 Shellmound Street, Emeryville, CA 94608 Phone: 4155475420 Fax: 4155475043 CHAIN-OF-CUSTODY RECORD AND ANALYTIC INSTRUCTION	Please send analytic results and a cop of the signed chain of custody form to <u>BOB</u> <u>RIDDELL</u> Project ID: <u>T69-677</u> S	ру о:	Personnel:	CHECKED. ) Specify and detection l ) Notify us i peaks in GC	UDE QA/QC DATA IF BOX IS lytic method and imit in report. f there are any anomalous or other scans. NS/CLARIFICATIONS: <u>CALL</u>
Sampled by: <u>D.C./P.R.M</u> . No. of Sample ID Container Sample Vo	Laboratory Name: <u>MOBIL CHER</u> 1 <sup>2</sup> Fil <sup>3</sup> Ref <sup>4</sup> Preservative	Analyze for	Analytic	Turn <sup>5</sup>	COMMENTS
Containers       Type '       Date $3$ $102 - EWI BH$ $W/V$ $10 - 28 \cdot 92$ $40$ $3$ $102 \cdot EWI - AFTER$ $11$ $10 - 28 \cdot 92$ $40$ $2$ $VE - I$ $1400$ $TEDLAR$ $IL$ $2$ $VE - I$ $1400$ $TEDLAR$ $IL$ $2$ $VE - I$ $1524$ $11$ $11$ $1$ $VE - I$ $1546$ $11$ $11$ $2$ $VE - I$ $1645$ $11$ $12$ $2$ $VE - I$ $1711$ $11$ $V$	$(specify)$ $= \underbrace{N} \underbrace{Y} \underbrace{HCL} \underbrace{T}$ $= \underbrace{N} \underbrace{NONE}$ $= \underbrace{V} \underbrace{V} \underbrace{V}$ $= \underbrace{V} \underbrace{V} \underbrace{V}$ $= \underbrace{V} \underbrace{V} \underbrace{V}$		Method <u>PA 80/5/8020</u> V		25.
Affiliation 2 Robert - Browey Received by (Signature), Date 0/29/92 2 UDISS ASSOC. 0200 Affiliation	hipping Carrier, Method, Date	Released by (Signatur AffiNiation Received by Lab Person Affiliation, Telephor	DINE 10-2 Donnel, Date	Seal intact?	
1 Sample Type Codes: W = Water, S = Soil, Desc Cap Codes: PT = Plastic, Teflon Lined 2 = 5 Turnaround [N = Normal, W = 1 Week, R = 24 Ho ADDITIONAL COMMENTS, CONDITIONS, PROBLEMS:	<pre>Volume per container; 3 = Filtered (Y</pre>	= VOA/Teflon Septa, P = '/N); 4 = Refrigerated (	= Plastic, C or B - (Y/N)	Clear/Brown Gl	ass, Describe.Other; Page <u>(</u> of <u>/</u>
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#### APPENDIX D

#### GROUND WATER DISCHARGE PERMIT

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November 3, 1992

ADMINISTRATION BUILLUW 1052 South Livermore Avenus Livermore, CA 94550 (415) 373-5100 FAX (415) 373-5110

Texaco Refining and Marketing 108 Cutting Blvd. Richmond Ca, 94804

Attention: Bob Riddell

Subject: Groundwater Discharge Permit

Dear Mr. Riddell,

The City has reviewed your groundwater discharge permit application and has found it to be complete. Enclosed you will find the following information:

- 1. 1992-93 Groundwater Discharge Permit
- 2. Discharge Permit Fee Statement
- 3. Permit Conditions and Prohibitions

As we discussed, the city will require that one sample is collected at the completion of the pumping tests. This sample shall be analyzed for Total Petroleum Hydrocarbons, and the results must be submitted to:

#### City of Livermore Water Resources Division 10 Rickenbacker Circle Livermore, CA 94550

Analytical results must include the volume of groundwater discharged to the sanitary sewer. For the purposes of this permit, a limit on Total Petroleum Hydrocarbons of 250 ug/L will be imposed. Based on the information in the permit application, this limit will be easily obtained with the proposed treatment technology.

This permit is being issued for a 24-hour pumping test only. If a treatment system and remediation plan are developed within the duration of this permit, the permit may be modified to cover discharges from this system. If you would like to pursue this option, please submit additional information on the treatment system when you are ready to begin remediation.

If you have any questions regarding the permit or any of the attached information, please contact me at (510) 373-5230.

Sincerely,

22

14.

Darren Greenwood Source Control Inspector



#### WATER RECLAMATION PLANT

1250 Kitty Hawk Road Livermore, California 94550

(510) 373-5230

AUTHORIZATION: The below named party is hereby authorized to discharge wastewater to the City of Livermore community sewer subject to compliance with the City of Livermore wastewater control ordinance and the conditions set forth in this permit.

CITY OF LIVERMORE

GROUNDWATER DISCHARGE PERMIT

PERMITTEE	Texaco Refining and Marketing			
ADDRESS	930 Springtown Blvd.			
	Livermore			
	California	zip94550		

- PERMIT CONDITIONS -

The above named shall report to the City of Livermore Water Reclamation Plant any change, (permanent or temporary) to the premise or operation that significantly change the quality or volume of the Groundwater discharge or deviate from the terms and conditions under which this permit is granted.

EFFECTIVE DATE:	November 3,	1992	EXPIRATION DATE:	November 2, 1993
DATED:	November 3.	1992	APPROVED BY:	Charlow of

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POST PERMIT IN PLAIN VIEW

#### WASTEWATER PERMIT CONDITIONS FOR GROUNDWATER DISCHARGER APPLICANTS

- 1. The City of Livermore Water Reclamation Plant issues the groundwater permit on a temporary and conditional basis only. The groundwater permit will not exceed one year in duration. The permit is conditional and may be revoked at any time by the WRP superintendent or his representative. The permit is non-transferrable.
- 2. Permittee shall abide by all applicable provisions of the City of Livermore Municipal Code or any applicable local, State, or Federal code or regulation. Any Violation of any provision of said codes or regulations will be just cause for revoking this permit.
- 3. Permittee shall not discharge to the sanitary system any materials or liquid wastes which may be harmful to the system or create a hazard or nuisance as defined in Section 13.32.060. Permittee may be required to bear the costs of any damage to the sanitary system attributable to the permittee.
- 4. The pH shall be no lower than 6.0 or higher than 9.0 at any time.
- 5. Any accidental discharge to the City of Livermore sanitary system must be reported immediately. Non-reporting of spill or slug incidents will be cause for administrative permit review.
- 6. All liquid or solid waste stored or hauled from the permittee's premise must meet all applicable local, State, and Federal rules and regulations. Certain RCRA regulations may apply to hazardous waste treated, stored, or generated on permittee's premise.
- 7. Any sludge generated by permittee is specifically prohibited from introduction into the City of Livermore sanitary system.
- 8. Conditional pollutant concentration limits for specific pollutants may be temporarily established by the Water Reclamation Plant Superintendent, and are subject to review and change without prior notification.
- Permittee shall not discharge wastewater containing in excess of:

0.06	mg/l	arsenic	0.20	mg/l	lead
0.14	mg/l	cadmium	0.01	mg/l	mercury
0.61	mg/l	nickel	0.20	mg/l	silver
0.62	mg/l	total chromium	3.00	mg/l	zinc
1.00	mg/l	copper	0.04	mg/l	cyanide
1.00	mg/l	total toxic or	Janics		

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From Section 13.32.100 of the Livermore Municipal Code

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