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Soil and Groundwater Characterization Study 187 North L Street Livermore, California

Prepared for

Don-Sul, Inc. 187 North L Street Livermore, CA 94550

June 12, 1991

Prepared by

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Mr. Tony Sullins
Don-Sul Inc.
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Subject: Soil and Groundwater Characterization Study

187 North L Street Livermore, California

Dear Mr. Sullins:

We are pleased to present the results of the Soil and Groundwater Characterization Study performed at the subject site. The purpose of this study is to explore the extent of petroleum contamination in soil and groundwater, and to evaluate the source of contamination. Two soil borings and five groundwater monitoring wells were installed during this study.

The following report presents the findings and conclusions of this study. Laboratory test results and supporting documentation are attached to the report. We are available to meet with you to discuss the results of this study and respond to questions.

Sincerely,

WOODWARD-CLYDE CONSULTANTS

Albert P. Ridley, CEG 926

Senior Associate

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Senior Staff Engineer

TABLE OF CONTENTS

Section		Page
	EXECUTIVE SUMMARY	1
	INTRODUCTION	3
	Scope of Work	4
	Task 1 - Alternate Source Evaluation	4
	Task 2 - Exploratory Borings and Groundwater Wells	5
	Task 3 - Groundwater Sampling	5
	Task 4 - Laboratory Analysis	5
	Task 5 - Aquifer Tests	6
	Task 6 - Analysis and Report	6
	Site Description	6
	Previous Studies	6
	FIELD INVESTIGATION	7
•	Soil Gas Study	7
	Borings and Monitoring Wells	8
	Soil Sampling	9
	Monitoring Well Development and Sampling	11
	In-Situ Hydrologic Tests	12
	LABORATORY TESTS RESULTS	12
-	RESULTS	13
	Extent of Soil and Groundwater Contamination	13
	Soil Contamination	13
	Groundwater Contamination	14
	Contaminant Sources	14
	EVALUATION OF MITIGATING MEASURES	17
	Soil Excavation	17
	Soil Vapor Extraction	18
	Bioremediation of Groundwater	18
	Groundwater Extraction and Treatment	19

		Т	ABLE OF CONTENTS (continued)
	EVII	DENCE OF SITE HISTORY	. 19
	DISC	USSION	20
	SUM	MARY OF CONCLUSIONS	21
	LIMI	TATIONS	23
	REF	ERENCES	24
List of Ta	<u>ıbles</u>		
		ratory Tests of Soil Samples ratory Tests of Water Samples	
List of Fi	gures		
Figure 3 Figure 4 Figure 5 Figure 6	Ground Ground Gross Cross Vertice Gross Gross	Plan Individual Plan Individual Plan Individual Plan Plan Plan Individual Plan Plan Plan Plan Plan Plan Plan Plan	
Appendice	<u>es</u>		
Appendi Appendi Appendi Appendi Appendi Appendi	ix B ix C ix D ix E	Tracer Research Report Boring and Well Logs Laboratory Analyses and Field Aqu Tank Integrity Test Historical Evidence of Mobil Service Declarations of Michael Comer and	ce Station

SOIL AND GROUNDWATER CHARACTERIZATION STUDY 187 NORTH L STREET LIVERMORE, CALIFORNIA

EXECUTIVE SUMMARY

The Arrow Rentals site, located at 187 North L Street in Livermore, California, was a former Mobil gas station from about 1951 to at least 1968. Arrow Rentals purchased the property in 1972. Three of the five underground fuel tanks were removed by Arrow Rentals in 1972, because they failed integrity tests. Two tanks that passed integrity tests remained in use until 1984, when they were also removed. No stains or gasoline odors were observed during removal of these two tanks. A new 1,000-gallon underground fuel tank was installed to meet the new California underground tank regulations. The new tank installation includes a vapor monitoring well adjacent to the tank. In 1985 it is reported (Declaration, Gary Pinks, Appendix F) that about 600 gallons of fuel was poured down the vapor well by a fuel delivery truck operator from Petcock Petroleum. Mr. Sullins reports that water was poured into the well from a garden hose some time after the spill.

Because the Livermore Redevelopment Agency is interested in purchasing this site as part of proposed redevelopment of this area, a study of possible site contamination was performed in 1988 and 1989 by Woodward-Clyde Consultants (WCC) on behalf of the Livermore Redevelopment Agency. Those studies showed detectable petroleum contamination in soil and groundwater at the Arrow Rentals site. The Alameda County Health Department has requested Arrow Rentals to perform additional site characterization to evaluate the extent of contamination.

Scope of Work

This study included drilling two exploratory borings (B-1A and B-F) and installing five groundwater monitoring wells. Prior to drilling, a soil gas study was performed by Tracer Research Corporation to explore the distribution of petroleum vapors in the soil. One well

(W-C) is located upgradient of the site, and two wells (W-D, W-E) are located downgradient of the site. One on-site well (W-B) is located downgradient of the former Mobil tanks, and a second on-site well (W-A) is located between the former Mobil tanks and the existing 1,000-gallon tank. Soil and groundwater samples were collected from the boring and wells, and were tested in an analytical laboratory for gasoline, diesel, and petroleum products. Water levels were measured in each of the wells to evaluate the groundwater gradient. These results were evaluated and presented in this report.

Results of Study

Gasoline was detected in soil samples from well W-A near the former Mobil tanks, and from Boring B-1A near the vapor monitoring well at the existing 1,000-gallon tank. Petroleum contamination was found in groundwater from well W-A (6,800 ppb benzene), and from well W-B (22,000 ppb benzene). No petroleum contamination was found in groundwater from wells W-C, W-D, and W-E. Floating gasoline was sampled from well W-1 located east of the former Mobil tanks, and downgradient of the vapor well.

Comparison of fuel fingerprints from the existing 1,000-gallon tank and from well W-1 shows that these materials do not match. Analysis of the product sample from well W-1 by Chevron Research and Technology Laboratory shows that on the basis of organic lead content, the floating product in W-1 is a gasoline produced in 1985. Analysis of organic lead content of gasoline in soil from Boring B-F, at a depth of 16 feet, shows that the gasoline was produced prior to 1985.

Evaluation of groundwater gradient and flow direction shows that the floating product is located in the downgradient direction of the reported 1985 release of Chevron Fuel into the vapor well.

Conclusions

Petroleum contamination in soil in the area of W-1 is most likely the result of leaks from a former pipeline extending from the five former Mobil station tanks to the former pump island closer to L Street. This is supported by detecting gasoline in soil at shallower depths (15 feet) and of higher concentration (1200 ppm) in W-1 east of the former Mobil tanks, and at lower concentrations (120 ppm) and deeper (20 feet) in B-1 and B-3 (25 feet) under the former Mobil tanks. That piping was not used by Arrow Rentals. In 1972 Arrow Rentals

installed a pump at the location of B-F and connected new pipes to the two remaining tanks. The nondetection of gasoline at a depth of 2 feet beneath this pump location in B-F, and only 16 ppm gasoline at a depth of 16 feet in B-F, indicates this pump and piping, operated between 1972 to 1984, is most likely not the source of gasoline in soil near W-1.

Floating product in W-1 is located on site, and has a lead content consistent with a gasoline produced in 1985, and spilled into the vapor well in 1985.

The location of floating product in W-1 is located about 40 feet downgradient of the vapor well and appears consistent with the reported spill in the vapor well in 1985.

The location of dissolved gasoline in groundwater extending a distance of more than 200 feet downgradient of well W-1 appears consistent with an older pipe leak during the period from 1951 to 1968.

The estimated 200- to 500-gallon volume of floating product appears consistent with the approximate volume of the spill in the vapor well in 1985.

About 85 percent of the soil contamination is estimated to be the result of leaks from pipes connected to the former Mobil tanks prior to 1968. About 15 percent of the soil contamination is estimated to be the 1985 Petcock Petroleum spill, into the vapor well.

About 97 to 99 percent of the petroleum groundwater contaminant is estimated to be the result of the Petcock Petroleum spill; about 1 to 3 percent is estimated to be from leaks from pipes connected to the former Mobil tanks prior to 1968.

While a number of options are available for remediation, the soil vapor extraction appears to be most feasible for soil contamination, and groundwater extraction and treatment appears most feasible for groundwater remediation.

INTRODUCTION

The Arrow Rental Site is located at 187 North L Street in Livermore, California. The site was a Mobil gas station for many years prior to 1968. Arrow Rentals purchased the property

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in 1972. In 1972 there were five underground fuel storage tanks remaining from the previous gas station operation. Three of the tanks were removed in 1972 because they failed leak detection tests. The two remaining tanks, a 6,000-gallon and a 4,000-gallon tank, remained in use until their removal in 1984. A pump was installed by Arrow Rentals near Boring B-F and connected with new piping to the two remaining tanks in 1972. A new 1,000-gallon underground tank was installed in 1984 to replace the removed tanks. The 1,000-gallon tank is currently in use for the Arrow Rentals operations. In 1985, it is reported that about 600 gallons of Chevron leaded regular gasoline was poured down a vapor monitoring well near the 1,000-gallon tank by the operator of the fuel delivery truck. Following that spill it is reported that water was poured into the vapor well with a garden hose, for about 24 to 30 hours (pers. comm., Tony Sullins).

In 1988 and 1989, the City of Livermore Redevelopment Agency retained Woodward-Clyde Consultants (WCC) to perform soil and groundwater exploration and testing to explore for potential contamination, based on the past site history (WCC 1988, 1989a, 1989b). Those studies detected petroleum contamination in soil and groundwater. The detection of petroleum contamination was reported to Arrow Rentals and to the Alameda County Department of Environmental Health. The Health Department subsequently requested Arrow Rentals to perform additional site characterization, to explore the extent of petroleum contamination, and form the basis for a Site Remediation Plan. Arrow Rentals retained the services of WCC to perform the requested work. The following report describes the results of the site characterization study, which was conducted in accordance with a work plan approved by Alameda County.

Scope of Work

The site characterization study consisted of site exploration, laboratory testing and analysis of the data and preparation of a report. This work, described below, was divided into a number of tasks.

<u>Task 1 - Alternate Source Evaluation</u>. Possible alternate sources of petroleum contamination were evaluated by testing for the presence of former piping, performing a precision integrity test on the 1,000-gallon tank and performing a soil vapor study of the site.

Task 2 - Exploratory Borings and Groundwater Wells. Prior to beginning field exploration a Health and Safety Plan (HSP) was prepared. The HSP considers potential health and safety risks and describes procedures and protective equipment needed for field personnel. Two exploratory boring and five groundwater monitoring wells were installed during this study (see Figure 1). Boring B-1A was located near the existing underground fuel tank, and Boring B-F was located at the former Arrow Rentals pump site. Wells W-A and W-B were drilled on-site, and wells W-C, W-D and W-E were drilled off-site. Well W-A is located in the down groundwater gradient direction from the existing underground fuel tank, and upgradient from W-1 and the former Mobil tanks. Well W-B is located near the north property line, downgradient from W-1 and the former Mobil tanks. Well W-C is located in North L Street, upgradient from the site. Well W-D is located in the rear of a residential property on Chestnut Street. Well W-E is located at the end of M Street Northwest of the site (Figure 1).

Task 3 - Groundwater Sampling. Each of the wells were developed by using a development rig to extract water from the wells to remove sediment. Water samples were collected from each well and transported to the laboratory on ice under chain-of-custody procedures. The water removed during development was stored in drums on site for proper disposal. Prior to sampling the stabilized groundwater level was measured and evidence of floating product was evaluated. The top of the well casings were surveyed by Bissel and Karn, Engineers, San Leandro, California. Measured groundwater depths were used to evaluate groundwater elevations and prepare a groundwater gradient map (Figure 2).

Task 4 - Laboratory Analysis. Selected soil samples from the borings and wells and water samples from wells were tested by Friedman and Bruye Laboratory, Seattle, Washington. A sample of fuel from the existing underground tank (Chevron Fuel) was tested in the Friedman and Bruye Laboratory. Tests included fuel fingerprinting, testing for total petroleum hydrocarbons as gasoline and as diesel (EPA 8015/8020). Some soil and water samples were tested for organic lead content, and selected metals. One soil sample from Boring B-F was tested by Coast to Coast Analytical Laboratories for tetraethyl lead and total petroleum as gasoline.

Water, floating product and fuel from the existing underground tank were submitted to Chevron Laboratories, Richmond, California. Tests were performed to evaluate if the floating product from the monitoring well contained characteristics of the 600 gallons of Chevron leaded fuel reportedly spilled by the delivery truck operator.

<u>Task 5 - Aquifer Tests</u>. Selected on-site wells were tested using slug test methods to evaluate the conductivity of the water bearing units under the site.

<u>Task 6 - Analysis and Report</u>. This report was prepared describing the results of the exploration and testing. The report describes the evaluation of the extent of soil and groundwater contamination, source(s) of petroleum contamination and possible mitigation methods.

Site Description

The 187 North L Street site is located on the west side of L Street, as shown in Figure 1. The site is about 100 feet wide, and 200 feet deep, and is entirely paved with asphaltic concrete. The former Mobil Station building is currently in use as an office, storage building by Arrow Rentals. The site is fenced and most of the rear (west) portion of the site is used to store rental equipment. A 1,000-gallon underground gasoline tank is located at the southeast corner of the site. Property to the south and west are undeveloped. The Western Pacific Railroad Right-of-Way borders the north side of the property. Residences are located north of the railroad property.

Previous Studies

Previous exploratory borings and wells were drilled by WCC at the site in 1988 and 1989 (WCC 1988, 1989a, 1989b) as part of the initial work for the Livermore Redevelopment Agency. Those borings and wells are shown on Figure 1 as B-1 through B-5, B-7, B-8 and W-1, W-2 and W-3. Gasoline contamination in the soil was detected at 170 parts per million (ppm) at a depth of 20 feet and 220 ppm at 25 feet in B-1. Gasoline was detected in soil boring B-2 at 3.5 ppm at a depth of 2 feet, 8.2 ppm at 5 feet and 1.7 ppm at 25 feet. Gasoline was not detected in soil in B-3 until a depth of 25 feet, where only 1.3 ppm was detected. The greatest concentration of gasoline in soil was detected in W-1, where gasoline was detected at 1200 ppm at 15 feet, 350 ppm at 20 feet, 490 ppm at 25 feet, 160 ppm at 30 feet, 370 ppm at 35 feet and 16,000 ppm at 40 feet. The highest concentration of gasoline at 40 feet is explained as an accumulation of gasoline floating on the groundwater, which was found at a depth of 43 feet.

No gasoline was detected in soil from depths of 2, 5, 10 and 15 feet in B-4 and 5 and 10 feet in B-7. Both B-4 and B-7 were drilled at opposite ends of the former pump island (Figure 1). No significant concentrations of gasoline (1.9 ppm at 5 feet, 1.7 ppm at 25 feet) were found in soil from Boring B-5, located near the existing 1,000-gallon fuel tank.

No gasoline contamination was detected in soil above the water table in Well W-3 and only 1.2 ppm gasoline was detected in Well W-2, with no detection to a depth of 50 feet. Groundwater from Well W-1 contained 210 ppm gasoline, 29 ppm benzene, 30 ppm toluene, 5.4 ppm ethylbenzene and 24 ppm xylenes. Groundwater from Well W-2 contained 0.36 ppm gasoline, 0.0067 ppm benzene, 0.0021 ppm toluene, 0.00047 ppm ethylbenzene and 0.0013 ppm xylenes. Water from Well W-3 contained 11 ppm gasoline, 0.290 ppm benzene, 0.120 ppm toluene, 0.150 ppm ethylbenzene and 0.140 ppm xylenes.

Measured groundwater depths ranged from 43.16 feet in Well W-1 and 44.24 feet in W-2, to 44.5 in W-3. The elevations of the groundwater surface indicated a northwest groundwater flow direction.

FIELD INVESTIGATION

Soil Gas Study

Tracer Research Corporation (TRC) of Tucson, Arizona was subcontracted to perform a soil gas investigation at the Arrow Rentals site on June 11 and 12, 1990. Samples were collected by driving a 3/4-inch-diameter stainless steel probe approximately eight to 10 feet into the subsurface. A truck-mounted hydraulic apparatus was used to push the probe into the soil. The probe end was protected from gathering soil during the driving procedure by inserting a pointed tip which is removed by lifting the probe after reaching the approximate depth of sampling. A fitting attached to the top of the steel probe is connected to a pump and approximately 2 to 5 liters of gas were evacuated from the steel probe. Samples were collected during the purging of the probe by inserting a syringe into the fitting at the top of the probe.

The samples were analyzed on the day sampled using a Varian 3300 gas chromatograph equipped with a flame ionization detector (FID). Samples were analyzed for benzene,

toluene, ethylbenzene, xylenes (BTEX) and total petroleum hydrocarbons. Quality assurance/quality control procedures are described in the report contained in Appendix A.

TRC reported no detection of BTEX above the stated detection limit in all of the samples. Total petroleum hydrocarbons were detected in all of the samples, however only up to 1 μ g/L. Figure 2 in the TRC Soil Gas Investigation report dated June 1990 shows an isoconcentration map of the total petroleum hydrocarbons (TPH) at the site.

Borings and Monitoring Wells

Between July 10 and 13, 1990, WCC installed five groundwater monitoring wells at and near the site (monitoring wells W-A, W-B, W-C, W-D, and W-E). Locations for these monitoring wells were based on consideration of the estimated groundwater flow direction from the data from previously built wells W-1, W-2, and W-3. The locations were selected to provide additional monitoring wells in the upgradient and downgradient directions of the known contamination. The wells were completed and screened into the upper, or first groundwater zone. The locations of these monitoring wells are shown on Figures 1 and 2. In addition, one boring was drilled adjacent to the vapor well of the 1,000-gallon underground regular gasoline storage tank in the southeast corner of the property.

The monitoring wells and the boring were advanced under WCC's field observation. Monitoring wells W-C and W-E were drilled using a truck-mounted drilling rig equipped with 8-inch outside-diameter, hollow-stem, continuous flight augers. Wells W-A and W-B were drilled using 12 inch diameter hollow stem augers, to allow installation the 4 inch diameter PVC well casing. These two wells have larger diameter well casing to allow for possible installation of pumps for groundwater extraction. For each well installation, a WCC engineer or hydrogeologist observed the drilling operations and prepared a log of the soils encountered in the monitoring well borings, which are presented in Appendix B. Upon completion of each boring, the monitoring wells were installed as described below. The wells were constructed of 2- or 4-inch-diameter flush threaded, schedule 40 polyvinyl chloride (PVC) casing. The screened intervals of the wells within the upper groundwater zone were placed between about 40 and 60 feet below the ground surface.

The screened portions of the well casings were constructed of slotted 2- or 4-inch-diameter PVC pipe, which contained standard factory-milled slots with a 0.010 inch aperture. The

remainder of each well casing was constructed of unslotted blank PVC well pipe. The filter pack, which consists of No. 2/12 Monterey Type, Lonestar sand, was placed in the annular space between the borehole wall and well casing from the base of the borehole to about one foot above the top of the slotted interval of the well casing. The filter pack was then sealed with approximately 1 foot of 3/8-inch-diameter bentonite pellets placed above the top of the sand pack. The remaining portion of the annulus was sealed with Portland cement grout. A water-tight, locking well cap was placed inside the casing and a traffic-rated Christie box was placed over each well. The well construction details are shown on the log of each monitoring well, in Appendix B.

Drilling equipment including the augers, soil samplers, drop hammer, plug, etc., where decontaminated by steam-cleaning prior to use at each boring. The soil sampler was also decontaminated between each sampling interval in a boring. All PVC well construction materials were also steam-cleaned prior to each installation. Steam cleaning operations were conducted so as to contain the resulting washwater. The washwater was subsequently transferred to drums which were sealed, labeled, and left on-site for proper disposal following completion of all drilling and sampling activities.

Soil Sampling

Soil samples were collected during the advancement of the monitoring well borings. Samples were generally collected at 5-foot intervals in each boring by advancing a 2-inch, outside-diameter, modified California sampler through the hollow stem of the augers. The sampler was driven 18 inches, using a 140 pound hammer with a 30-inch drop. The number of blows required to drive the sampler through each 6-inch portion of the 18-inch drive interval are shown on the boring logs in Appendix B.

The soil samples were retained in four 4-inch-long, 2-inch-diameter brass liners contained within the sampler. The ends of the tube samples were examined by WCC's engineer and the soil was visually classified using the Unified Soil Classification System. The soil descriptions are included in the boring logs. The depth to first encountered groundwater is also shown on the logs.

The soil samples in the lowermost brass liners were retained for potential chemical analyses, while the samples from the adjacent liner were generally used to perform a headspace

analysis in the field for volatile organic compounds. Chemical analyses was performed on soil samples when headspace analyses indicated a presence of volatiles in soils. The headspace test procedure involved emptying the contents of the brass liner into a Ziplock® plastic bag and sealing the bag. The soil was allowed to out-gas inside the sealed bag for approximately 5 minutes. The bag was then pierced with a probe and the air within the bag was tested for total organic vapor with an HNu photoionization detector.

Sampling equipment such as the brass liners and samplers were decontaminated between uses by washing in an Alconox solution followed by two tap water and one distilled water rinses.

The elevations of all new well casings were surveyed, relative to Mean Sea Level (MSL) datum, by means of a closed loop level traverse. The top of each well casing was surveyed to the nearest 0.01 foot. A notch was marked on the casing at the measuring point such that future well level measurements can be taken from the same point. In addition, all wells were surveyed for horizontal control. The surveying was performed by Bissel & Karn Engineers, a licensed land surveyor. All new groundwater monitoring wells were permitted and installed in accordance with Alameda County Flood Control and Water Conservation District regulations and guidelines.

Boring B-1A, located on the east side of the existing (1,000-gallon) underground tank, encountered sandy gravel grading to clayey silt at 25 feet. Field measurements of organic vapors from soil samples from the borings shows no significant detection until a depth of 20 feet, where 50 ppm organic vapors were detected with an HNU photoionization detector (PID). Readings increased to 225 ppm at a depth of 40 feet in B-1A. Groundwater was encountered at a depth of about 42 feet in B-1A.

Boring B-F was drilled at the location of the former gasoline pump, used by Arrow Rentals between 1972 and 1984. That boring encountered clayey gravel from beneath the concrete slab to a depth of 16 feet. The clay content increased near the bottom of the boring. No gasoline odors were observed in soil from Boring B-F to a depth of about 15 feet, where a slight gasoline odor was observed.

Field measurements of organic vapors detected 20 ppm at a depth of 20 feet in Well W-A, increasing to 110 ppm at 25 feet. A strong gasoline odor was observed from soil at a depth of 30 feet continuing to the top of the groundwater at about 44 feet.

Well W-B, located downgradient of the former Mobil underground tanks (Figure 1), did not encounter soil with a significant gasoline odor. The highest HNU reading from soil in W-B was only 30 ppm, at a depth of 40 feet. Groundwater was encountered at a depth of about 44 feet.

Well W-C is located upgradient of the site in the pavement of L Street. No gasoline odors were encountered from soil from W-C. The well was constructed using 2-inch diameter PVC well casing. A meter box cover was installed in the street over the well for protection and security.

Well W-D is located in the rear yard of the Armstrong property at 1951 Chestnut Street. No gasoline odors were detected in soil samples from W-D. A HNU reading of 150 ppm was recorded during drilling at a depth of 45 feet. Groundwater was encountered at a depth of about 42 feet. A 2-inch-diameter well casing was installed in W-D.

Well W-E is located in the pavement of M Street northwest of the site, as shown in Figure 1. No gasoline odors or significant HNU readings were detected during drilling of W-E. Groundwater was encountered at a depth of about 43 feet.

Monitoring Well Development and Groundwater Sampling

Following installation, Datum Exploration was subcontracted to develop the monitoring wells. The five groundwater monitoring wells were developed on July 26 and 27, 1990 by pumping or bailing groundwater and by applying a surge block to the water column. Pumping and bailing were performed with intermittent surging until the resultant groundwater was clear. However, if the well yield was very low, the well was considered developed when a total of at least 10 casing volumes of groundwater was removed and some improvement in well yield was observed. Copies of the well development logs are included in Appendix C.

Groundwater monitoring wells Nos. W-A, W-B, W-C, W-D and W-E were sampled by WCC personnel. The sampling procedure utilized included: 1) measurement of the depth to

groundwater; 2) removal of at least 5 wetted casing columns of groundwater from each well; and 3) allowing the groundwater level within the well to recover to at least 80 percent of its static level prior to obtaining samples. While purging the well, the groundwater from each well was evacuated with either a Teflon® bailer or a suction type pump. Water quality parameters including temperature, pH, and specific conductance were recorded throughout the well purging process. The physical parameters of the purged water, such as turbidity, color and odor, were noted.

Following the well purging activities, each well was allowed to recover to at least 80 percent of the original static groundwater level. Prior to sampling, the groundwater level and water quality parameters were rechecked and verified against the data obtained during the purging activities. Stabilized water quality parameters were used as a general check that the groundwater obtained for the samples was fresh formational water (i.e., had not stagnated in the near vicinity of the well). Monitoring well sampling data including well depth, depth to groundwater, groundwater elevation, conductivity, temperature, pH, and turbidity are shown in Appendix C. The depth to groundwater was established using a Solinst power sounder, with the depth measurements recorded to the nearest 0.01 foot. Groundwater samples were obtained with a clean Teflon® bailer and were carefully decanted into appropriate laboratory prepared sample containers. Samples analyzed for metals were filtered in the field and then decanted into the sample containers. The samples were then placed in laboratory-prepared containers and transported to the analytical laboratory on the day of sampling, under standard chain-of-custody procedures.

In-Situ Hydrologic Tests

Slug tests were performed on Wells W-A and W-B, Figure 1. These wells were selected for slug tests to measure the hydraulic conductivity of the water-bearing strata under the site. A pressure transducer was placed in the well and a selected volume of water was introduced into the well. The time for the water level to return to normal was measured. The results are shown in Appendix C and are discussed in the text of this report.

LABORATORY TESTS

Selected soil and water samples from borings and wells were tested in a commercial analytical laboratory (Friedman & Bruye, Inc.) for total petroleum hydrocarbons as gasoline,

and diesel (EPA 8015 modified) and for benzene, toluene, ethylbenzene, and xylenes (BTEX) (EPA 8020). Gas chromatograph methods were also used to perform fuel fingerprint tests on soil and water samples. Soil and water samples were also tested for organic lead content, cadmium, chromium, copper, nickel, vanadium, and zinc. Selected samples were also analyzed for the presence of phenol, naphthalene, 2-methyl naphthalene, and dye. A fuel fingerprint and analysis for BTEX and organic lead was performed on a sample of fuel from the existing 1,000-gallon tank filled with Chevron fuel. A sample of floating product from well W-1 (Figure 3), and water from wells W-A and W-B were analyzed in the Chevron Research and Technology Company Laboratory for characteristics which might identify the fuel as Chevron leaded gasoline from August of 1985. Chevron Research tested the samples using gas chromatography and a flame ionization detector to determine hydrocarbon composition and an electron capture detector to measure lead alkyl concentration. Detergent additive was analyzed by Sep-Pak extraction and infrared spectroscopy. The results of laboratory tests are included in Appendix C. One soil sample from Boring B-F was analyzed by Coast to Coast Analytical Laboratories for tetraethyl lead and total petroleum hydrocarbons as gasoline. Using a method developed by Dr. Brian Andresen, consultant, the lead content per gallon of gasoline in this soil was evaluated.

RESULTS

Extent of Soil and Groundwater Contamination

<u>Soil Contamination</u>. Laboratory analyses of soil samples from borings and wells indicate that gasoline contaminated soil is located on the site. No gasoline contamination was detected in soil samples from off-site wells W-C, W-D, and W-E, as shown in Table 1.

Soil samples from a depth of 15 feet in Well W-1 contained 1,200 parts per million (ppm) Total Petroleum Hydrocarbons (TPH) as gasoline. The detected concentrations of TPH as gasoline ranged from 160 to 370 ppm in soil from 20 to 35 feet in well W-1. At 40 feet in W-1 the soil contained 16,000 ppm TPH as gasoline. Laboratory analyses detected 2 ppm gasoline at 30 feet, and 1,000 ppm gasoline at 40 feet in Well W-A, located southeast of W-1 (Figure 5). A field vapor analyzer (HNU photo ionization detector) measured 50 ppm organic vapors at 20 feet in Boring B-1A. The HNU detected 130 ppm at 35 feet, and 225 ppm at 40 feet in Boring B-1A. Laboratory tests of soil from 10, 15, 20, 30, and 35 feet in B-1A, for total petroleum hydrocarbons as gasoline showed no detection (less than 5 ppm). Soil

from a depth of 40 feet in B-1A contained 350 ppm gasoline. At 45 feet in B-1A soil contained 54 ppm gasoline. No gasoline was detected in soil from a depth of 2 feet in Boring B-F. Analysis of soil from a depth of 16 feet in Boring B-F showed 10 parts per billion (ppb) tetraethyl lead and 16 ppm TPH gasoline. No gasoline was detected above the water table in soil samples from Well W-B located northwest of W-1, at the north property line of the site (Figures 3, 4 and 5). The estimated distribution of soil contamination under the site is shown in Cross-Sections A-A' and B-B' (Figures 5 and 7).

Groundwater Contamination. Analysis of groundwater from off site wells W-C (upgradient), W-D, and W-E (both downgradient) shows no significant detection of petroleum contamination. The detection of 1 part per billion (ppb) benzene, 2 ppb toluene, 1 ppb ethylbenzene, and 1 ppb xylenes in well W-D may be a result of contamination during transport of the water samples, since a travel blank shows detection of 7 ppb benzene, 37 ppb toluene, 14 ppb ethylbenzene, and 43 ppb xylenes. The travel blank may have been contaminated during transport to the laboratory, since a pure fuel sample was also placed in the ice chest with W-D.

The most significant gasoline contamination was found in well W-1, where 4 inches of gasoline (Figure 6) was found floating on the groundwater. After well development the thickness in the well increased to several feet, however, this may be a localized thickness in the well screen. TPH as gasoline was detected at 10 ppm in W-A, and 21 ppm in W-B. As shown in Table 2, benzene was detected at 22,000 ppb in water from W-B, and 6,800 in water from W-A. Toluene, ethylbenzene, and xylenes found in W-A and W-B up to about 7,900 ppb. No measurable floating product was detected in the downgradient well W-B, or the upgradient well W-A. Previous analyses of groundwater in wells W-2 and W-3 detected 6.7 ppb benzene in W-2 and 290 ppb benzene in W-3. The approximate limit of groundwater contamination is shown in Figure 3.

Contaminant Sources

The source of soil and groundwater contamination appears to be from two separate sources. One is a suspected leak between 1951 and at least 1968 from a pipe connected to the former Mobil tanks. A second source is the sudden spill of gasoline into the vapor well in 1985.

Exploration shows the location of floating product to be about 40 feet downgradient of the reported 1985 fuel spill. Floating product detected in well W-1 at about 4-inches thick, is not found in W-A or W-B. Fuel fingerprints of floating product from W-1 does not match the fingerprint of the Chevron fuel in the existing tank or soil from W-A at 40 feet (see Friedman and Bruya report Appendix C). The gas chromatograph peaks for water from W-A and W-B represent the soluble fraction of gasoline.

As shown in Figures 5 and 7, gasoline contaminated soil occurs from a depth of 15 feet downward to the top of groundwater in W-1. Cross-section B-B' shows that gasoline contamination is shallower (15 feet) and has a higher concentration (1200 ppm) in soil from W-1 located east of the former Mobil tanks, and along the approximate pipe route to the Mobil pump island. The gasoline contamination is deeper (20 feet) and less concentrated (120 ppm) in soil from B-1 under the former Mobil tanks. Since only 16 ppm gasoline was detected in soil from 16 feet in Boring B-F, the former Arrow Rentals pump does not appear to be the source of gasoline in soil. Evaluation of the amount of tetraethyl lead in soil from 16 feet in Boring B-F shows the gasoline contained 1.8 grams per gallon, which is ttypical of a pre-1985 gasoline.

Field measurements of organic vapors in soil samples from Boring B-1A adjacent to the vapor monitoring well at the tank storing Chevron fuel shows detection of 50 ppm at 20 feet, 25 ppm at 25 feet, 50 ppm at 30 feet, 130 ppm at 35 feet, and 225 ppm at 40 feet. However, laboratory tests of soil at those depths, for TPH as gasoline, shows no detection (less than 10 ppm), until a depth of 40 feet where 350 ppm gasoline is detected. The low concentration of gasoline detected in soil near the reported June, 1985, fuel spill into the vapor monitoring well may be explained as a result of the reported pouring of a large volume of tap water into the vapor well using a garden hose, following the spill.

The laboratory tests performed on floating product in W-1, and dissolved product in W-A and W-B provide the most conclusive evidence for the origin of the fuel in the groundwater. Chevron Research (Appendix C) reports that the lead content of the fuel in W-1 is 0.18 g/gal. Considering a 10 percent evaporation of the gasoline the fuel originally had a lead content of about 0.16 g/gal. Chevron reports that this lead content is consistent with gasolines produced between late 1985 and 1988. Chevron (Appendix C) reports that gasolines spilled before 1985 typically retain most of their lead, with lead levels of 0.5 to 3.0 g/gal. Tests

indicate that the gasoline from W-1 contains a polybutene-amine detergent. Chevron used a polybutene-amine-containing F310 detergent in its leaded gasolines until 1987.

Chevron Research reports that Mobil also used a polybutene-amine-containing detergent. Chevron Research report dated October 16, 1990 (Appendix C) also reports that there is nothing in the chromatogram of gasoline from W-1 to indicate that it could not have come from the Richmond refinery. In 1985 Chevron was adding 0.22 g/gal lead to its leaded regular gasolines. The November 1985 gasoline contained more than 95 percent of its lead as tetraethyl isomer. The W-1 gasoline is reported to contain 83 percent of its lead as tetraethyl isomer (TEL). This implies the gasoline from W-1 is not Chevron's. However, this may simply be a result of exposure to the environment or other factors as discussed by Dr. Brian Andresen in an April 30, 1991 letter, Appendix C.

The location of the floating product about 40 feet downgradient from the vapor well (Figure 3) is consistent with the anticipated location after five years. Using the conductivity data from slug tests ($k = 4 \times 10^{-4}$ cm/sec), and the northwest gradient (0.0071 ft/ft), and gasoline spill at the vapor monitoring well could have moved about 15 feet northwest since 1985. With a local and temporary increase in gradient, by pouring water down the vapor monitoring well (pers. comm., Tony Sullins), or from rainfall, the gasoline could have moved up to 40 feet northwest since 1985. Addition of water from a hose at an estimated 5 gallons per minute for about 30 hours could have added 9,000 gallons of water to the vapor monitoring well. That volume of water could easily have washed the gasoline from the area of the vapor monitoring well and moved it to its present location. The approximate volume of floating product is also consistent with the approximate 600 gallons of gasoline reported spilled in the vapor monitoring well. Using a measured thickness of about 4 inches, an estimated diameter of floating product of 25 feet, and a soil porosity of about 40 percent, about 200 to 500 gallons of gasoline could be floating on the groundwater at W-1. A tank integrity test performed on the 1,000-gallon tank (Appendix D) shows the tank is "tight" per National Fire Protection Association Standards.

The low concentration of lead in gasoline from W-1 (0.18 g/gal) is the strongest indicator that the W-1 gasoline is a gasoline produced around 1985. Gasolines produced before 1985 typically contained 0.5 to 3.0 g/gal lead. Therefore, the floating product clearly is not a pre-

1985 gasoline, and whether it is a Chevron product or not it is likely to be the gasoline delivered by Petcock Petroleum and spilled into the vapor monitoring well.

Based on the estimated groundwater flow rate of between 3 and 8 feet per year, the dissolved gasoline plume, extending over 200 feet downgradient from W-1, is not likely to be from the 1985 spill. This evaluation supports the conclusion that the source of that dissolved gasoline plume is from the earlier pipe leak during the period 1951 to 1968. A leak in 1960 from the vicinity of the Mobil pipes could have moved up to 250 feet downgradient by 1991. Poesthis mean contain in w-2 and w-3 are from pre 1985?

EVALUATION OF MITIGATION MEASURES

Soil and groundwater containing petroleum hydrocarbons can be remediated using techniques that have proven effective and have been used successfully at other sites in the San Francisco Bay Area. Soil remediation methods include excavation and removal of contaminated soil and transportation to an approved waste site, excavation and on-site bioremediation using natural soil bacteria and removal to a waste disposal site, in-situ bioremediation, or in-situ vapor extraction. Each of these methods have limitations as to their feasibility.

Soil Excavation

Soil contaminated with gasoline extends to a depth of 43 feet in W-1 and B-1A where the groundwater table is encountered. The soil contamination is assumed to extend laterally from W-1 in about a 25-foot radius. The volume of contaminated soil around W-1 is estimated to be about 2,000 cubic yards. It is anticipated that some significant contaminated soil (greater than 100 ppm TPH) will be found under the existing 1,000-gallon fuel tank. That soil might extend from about 35 to 43 feet under the tank and extend out about 20 feet from the tank. That volume of soil is estimated to be about 350 cubic yards. In this deeper zone the Regional Water Quality Control Board might require removal of soil to a lower TPH concentration of about 10 ppm, or less. Considering the small size of the site and the need to have excavation slopes no steeper than 1 (horizontal) to 1 (vertical) there is a potential that the excavation would extend into L Street. Protection of L Street might require support of the slopes with retaining structures such as shoring or sheet piles. With a sloped excavation about 15,000 cubic yards of uncontaminated soil would be excavated and then replaced in the hole.

Excavated soil would be removed to an approved waste site, and the excavation would be refilled with clean soil. The excavated soil could also be stored on-site and treated by aeration and bioremediation prior to removal from the site. The relative costs of removal could then be reduced since the treated soil could possibly be disposed of at a Class III Landfill instead of a Class I facility. Considering the extra cost of shoring, removal and replacement of clean soil, and hauling to a waste site, the excavation option may not be appropriate for this site.

Soil Vapor Extraction

Petroleum contaminated soil above the saturated zone can be treated to remove volatile hydrocarbons by extraction of soil gas (the gas in the soil pores) using a vapor extraction system (VES). The extraction system consist of one or more small diameter extraction wells drilled into the vadose zone (unsaturated zone of soil) above the groundwater table. A vacuum pump is used to draw soil gas from the extraction wells. The removed vapors are either released into the air, if permitted, or a pumped through activated carbon filled canisters to remove the petroleum vapors prior to discharge to the atmosphere. The vacuum pump and carbon canisters would probably require about two parking spaces. The system might use two 55-gallon drums of activated carbon, for a small system, or two 1,000-pound canisters for a large system. Spent carbon is replaced with new carbon and the spent carbon is returned to the manufacturer for regeneration. Such a system might operate for at least one year.

Bioremediation of Groundwater

Contaminated groundwater can be treated by extraction and treatment in a closed loop bioremediation system. The system would consist of one or more groundwater extraction wells (the 4-inch on-site wells) and a grid of injection wells. Extracted water would be pumped to an above ground tank where oxygen, nutrients, and heat would be added to the water. This mixture would be metered into the injection wells to increase and maintain the in-situ bacteria and microorganism population necessary to remediate the soil and groundwater in the subsurface. This method might require substantial initial documentation before regulatory agency approval. At least one year might be required to complete remediation of the soil and groundwater. A second year might be needed to reach desired cleanup levels. This method is generally less expensive than the VES and other groundwater

extraction and treatment systems. However, permitting might require a pilot study before gaining approval, which would be time consuming.

Groundwater Extraction and Treatment

Extraction of petroleum contaminated groundwater and treatment is a proven technology that has been used for many years. The system is similar to that described above, but the groundwater is treated in a treatment plant on the surface. The two on-site 4-inch diameter wells could be used for extraction of groundwater. Groundwater would first be pumped through an oil/water separator and then through a canister of activated carbon prior to discharge to the sanitary sewer. The carbon would be replaced as necessary to achieve the allowable discharge concentrations. The cost of operations would depend upon the amount of petroleum product in the groundwater. Systems are available that can be brought to the site in a prefabricated unit on a trailer. The initial operations would extract much of the floating product which would be separated for disposal along with the used carbon canisters. As contaminant concentrations decrease the operations costs would decrease. An extraction and treatment system is expected to operate for at least one year, and then for another year until the cleanup concentrations are achieved. This method of groundwater treatment would be more easily permitted, and could be installed in less time than the bioremediation system.

EVIDENCE OF SITE HISTORY

Research has produced confirmation that a Mobil Service Station was located at 187 North L Street From 1951 to at least 1968. Copies of this information are contained in Appendix E. A "Ten Year Service Award" dated 1961 was found inside the building in 1972 after Arrow Rentals purchased the site. A City of Livermore Building Permit for an underground storage tank at 187 North L Street is dated December 2, 1960. Valley Mobil Service is listed in the October 1966 Pacific Telephone directory, and the 1968 Livermore City Directory.

The declaration of Mr. Michael Comer (Appendix F) documents Mr. Comer's observations of removal of the two remaining underground tanks in the spring of 1985. He states that he observed no noticeable odor of gasoline coming from the soils at the bottom of the tank pit, nor did the soil in the tank pit have a dark discolored look which is characteristic of a tank which has been leaking. He concluded that the two tanks had not been leaking.

The declaration of Mr. Gary R. Pinks (Appendix F) documents his observations of the gasoline spill by Petcock Petroleum on June 18, 1985. Mr. Pinks states that he was present at 187 North L Street when the delivery truck from Petcock Petroleum arrived to deliver an order of regular gasoline. Mr. Comer observed that the delivery truck driver had put the nozzle of the fill hose into the vapor well rather than the mouth of the tank. Mr. Comer states that the driver called Petcock Petroleum on this truck radio and reported he had dropped 600 gallons of gas into the dirt. Mr. Comer states that a day or so after the spill the Petcock delivery truck returned and delivered gasoline into the tank.

DISCUSSION

Since the costs of remediation are anticipated to be significant some discussion of the relative causes of contamination is appropriate. The distribution of gasoline contaminated soil indicates that most of the contaminated soil appears to be located beneath the location of piping from the pump island to the five former underground gasoline tanks, in operation for many years by Mobil Oil Company from 1951 until about 1968. The estimated relative volume of contaminated soil under pipes to the former Mobil tanks is estimated to be about 2,000 cubic yards, or about 85 percent of the estimated total contaminated soil. About 350 cubic yards of contaminated soil is estimated to underlie the existing tank and vapor well spill, or about 15 percent of the total contaminated soil. Therefore about 85 percent of the gasoline-contaminated soil appears to be a result of pipe leakage from 1951 to 1968, and about 15 percent appears to be a result of the 1985 spill into the vapor well. The calculated 1.8 g of tetraethyl lead and other lead isomers per gallon of gasoline in soil from Boring B-F supports this conclusion, since this is typical of a pre-1985 gasoline.

Considering that the laboratory tests have shown that the floating product in W-1 is a gasoline produced in 1985, based upon its lead content and other characteristics, and that the approximate 200- to 500-gallon volume is consistent with the reported Petcock Petroleum spill of 600 gallons in August of 1985, the floating product is the result of the Petcock spill. The remaining portion of the reported 600-gallon spill most likely remains in the soil and backfill under and around the existing tank. Since the soil contamination in W-1, under the piping to the former Mobil tanks, extends downward to the top of groundwater, some groundwater contamination might be the result of releases from the piping to the former Mobil Tanks. Most likely the more downgradient dissolved fraction of gasoline (BTEX) is

the result of an earlier release related to the soil contamination under the piping to the former Mobil tanks between 1951 to 1968.

The broad aerial extent of the groundwater contaminant plume downgradient of the former Mobil tanks is considered to be due to releases from piping before 1972. Considering the relatively low concentrations of gasoline in that plume as compared to the volume of floating gasoline from the 1985 spill, the greatest amount of gasoline floating and dissolved in the groundwater is from the 1985 spill. About 97 to 99 percent of the gasoline floating on and dissolved in the groundwater appears to be due to the 1985 spill. This is based on an estimated 200 to 500 gallons of floating product and a 100-foot diameter of contaminated groundwater with about 3-1/2 gallons of dissolved gasoline from the 1985 spill. About 1 to 3 percent of the gasoline in the groundwater could be due to releases from 1951 to 1968. This is based upon an estimated 7 gallons of gasoline in the broad groundwater plume.

The remediation effort to recover each of these releases from the groundwater will be dependent more upon the aerial extent of the contaminant plume. Assuming moderate recovery of about 1/2 gallon per minute for pumping of groundwater for treatment, about 1 to 2 years would be needed to extract the 1985 spill. Assuming the same recovery rate, more than 5 years would be needed to pump and treat groundwater from the broad plume. The time required for groundwater remediation could be reduced by addition of extraction wells to increase the rate of extraction of groundwater.

SUMMARY OF CONCLUSIONS

- 1. Based on this data, gasoline contamination in soil and groundwater at the site is the result of two separate sources: (1) a pipe leak from pipes connected to former Mobil tanks between 1951 and 1968, and (2) a sudden spill into a vapor monitoring well by Petcock Petroleum in 1985.
- Laboratory tests of samples of soil and groundwater for petroleum products indicates
 that soil contaminated with gasoline is located primarily on-site in the vicinity of well
 W-1 from a depth of about 15 to 43 feet, and near Boring B-1A from about 35 to 43
 feet.

- Floating gasoline found in well W-1 has a lead content consistent with a gasoline produced around 1985, and spilled by Petcock Petroleum into the vapor well in 1985.
- 4. The location of the floating product in W-1, about 40 feet downgradient from the reported Petcock Petroleum spills, appears consistent with the estimated present location for a spill at the vapor well in 1985. Floating product appears to be located on-site in the vicinity of W-1.
- The estimated volume of floating product from measured thickness and diameter is consistent with the reported 600-gallon spill by Petcock Petroleum at the vapor monitoring well.
- 6. Laboratory tests of groundwater from off-site wells W-C (upgradient), W-D and W-E (downgradient) show no significant detection of gasoline or BTEX. One ppb benzene in Well W-D is most likely a result of cross contamination during transport to the laboratory. Dissolved fractions of gasoline (BTEX) extend off-site in the downgradient direction towards W-D and W-E.
- 7. Laboratory tests of soil from a depth of 16 feet in Boring B-F shows the tetraethyl lead content of gasoline in this soil is consistent with a gasoline produced prior to 1985. The source of this gasoline is most likely a pipe leak from the former Mobil tanks prior to 1968.
- 8. About 85 percent of the soil contamination appears to be the result of leaks from piping to the former Mobil underground fuel tanks prior to 1972. About 15 percent of the soil contamination appears to be the result of the 1985 Petcock Petroleum spill.
- 9. About \$7\$ to 99 percent of the groundwater contaminant appears to be the result of the Petcock Petroleum spill, and about 1 to 3 percent appears to be the result of leaks from the former Mobil tanks, prior to 1968. Assuming a 1/2-gallon-per-minute recovery rate, about 1 to 2 years might be needed to extract the groundwater contamination from the 1985 spill. At the same extraction rate more than 5 years would be needed to extract the pre-1985 groundwater contamination.

10. While a number of options are available for remediation of soil and groundwater, the most feasible soil remediation method is soil vapor extraction, and the most feasible groundwater remediation method is groundwater extraction and treatment with carbon filters.

LIMITATIONS

The scope of this investigation is limited by time constraints, expense, and practicality. A limited number of samples were taken at locations on and off the site, and a limited number of chemical analyses were performed on those samples. Professional opinions concerning the presence of gasoline and petroleum products were developed based upon the resulting data. It would be prohibitively expensive and time consuming to sample all locations on and off site for all substances which are now, or in the future might be, considered hazardous. Therefore, WCC cannot be held responsible should the investigation fail to detect the presence or quantity of all hazardous substances at all locations on and off site in the study area.

REFERENCES

- Woodward-Clyde Consultants, 1988, consultants report entitled, "Phase I Environmental Assessment, Railroad Avenue Property, Livermore, California", dated December 27, 1988, prepared for Livermore Redevelopment Agency.
- Woodward-Clyde Consultants, 1989a, consultants report entitled, "Phase II Site Exploration, Railroad Avenue Property, Livermore, California", prepared for Livermore Redevelopment Agency.
- Woodward-Clyde Consultants, 1989b, consultants report entitled, "Phase III Environmental Assessment, 187 North L Street, Livermore, California", prepared for Livermore Redevelopment Agency.

Table 1 LABORATORY TEST RESULTS OF SOIL SAMPLES **
Arrow Rentals, 187 North L Street, Livermore,CA

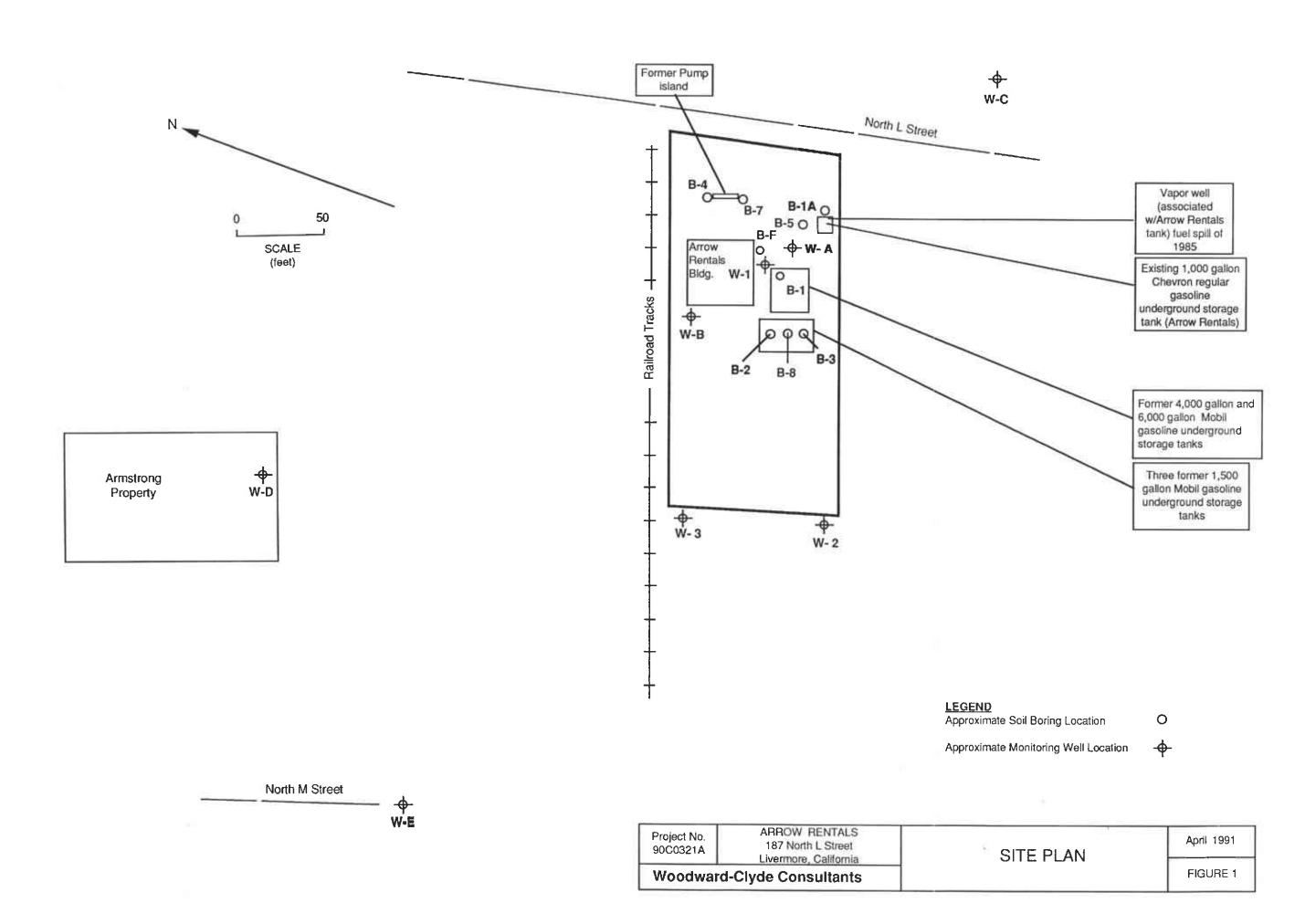
Sample No.	TPH Gas.	Organie Pb (pph)	Benzene (ppb)	Toluene (pph)	Ethylbenoene (ppb)	Xylenes (aph)	2-Mathyl Naphthalana	Naphthalene (ppm)	Phenol (ppm)	Dyes	Cd	Cr (ppm)	Cu (ppm)	NI (ppm)	V (ppm)	Zn (ppm)
B-1A-10			100.0		1	5.59.511	THE RESERVE OF THE PERSON NAMED IN COLUMN TO								1	1
B-1A-15	<10															
B-1A-20	<10	<5														
B-1A-30	<10	c 5								1						
B-1A-35	<10															
B-1A-40	390	< 5														
B-1A-45	54	<5														
B-1A-50	<10	< 5														
P-1	<0.5		<500	<500	<500	<500										
B-F-1,2	16	10*	2	25	30	34										
W-A-20	<1	<1	A18	320	240	210					0.2	2.3	12	64	0.3	1 1
W-A-30	2	<1	394	130	35	1200	<1	<1	<10		<.1	23	13	55	6.3	23
W- 4-3 5		< 1									0.6	3 1	15	6.9	9.1	15
W-A-40	19884		12000	37000	7500	27000				none						
W-B-25	<1	< 1									<.1	76	15	82	6.3	2 1
W-B-30							<1	<1	<1							
W- B-26	<1		898	260	110	70					0.2	22	18	74	7.1	1 4
W-C-20	<10		<10	<1	c1	<1				none						
W-C-40	c10															
W-D-25	<1		<10	<1	c1	<1				none						
W-D-40	<1		<10	<1	c1	<1										
W-E-20	c10															'
W-E-40	<10			1												

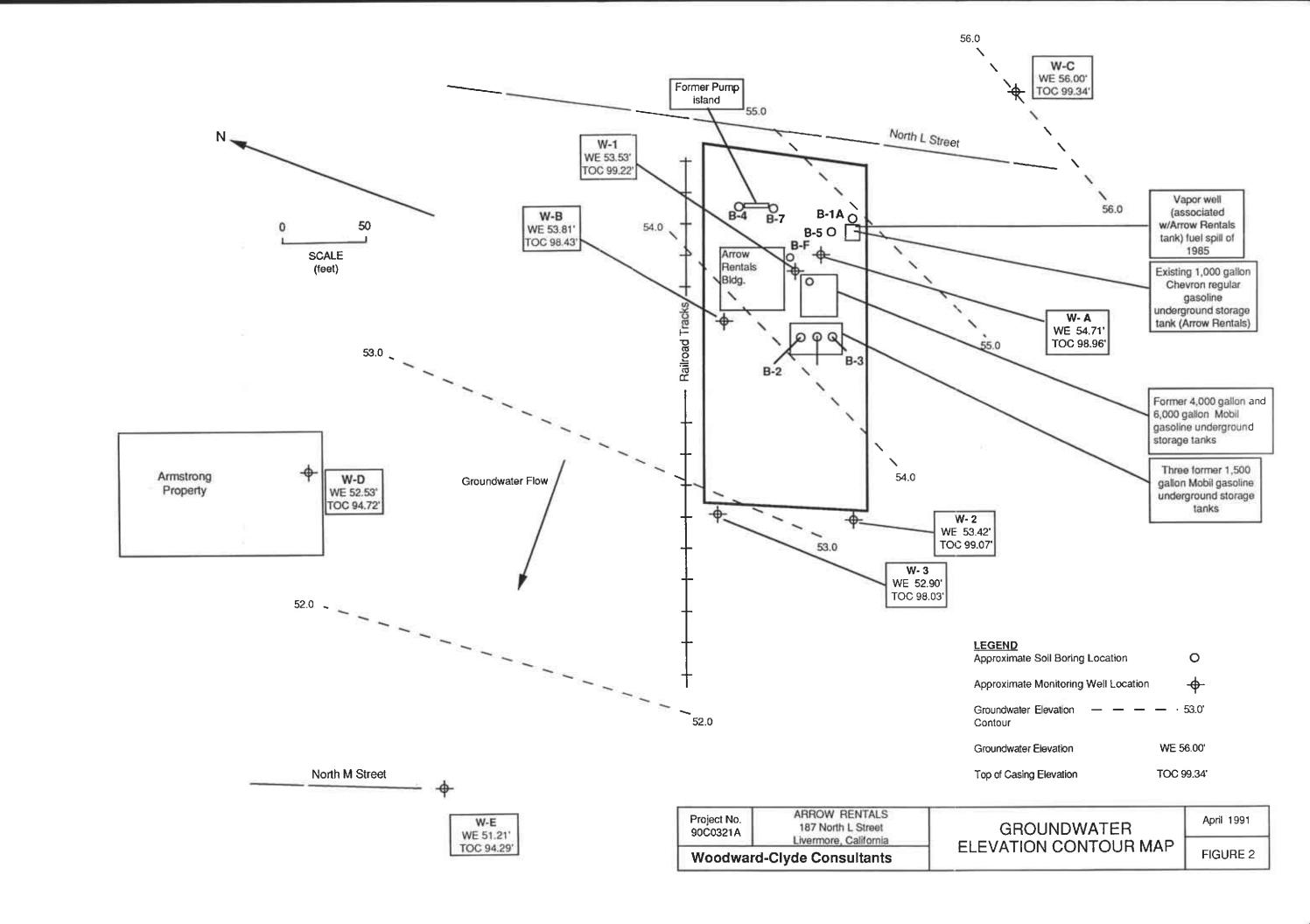
[&]quot;= Tetraethyl lead, ""= Previous laboratory tests summarized in Appendix C

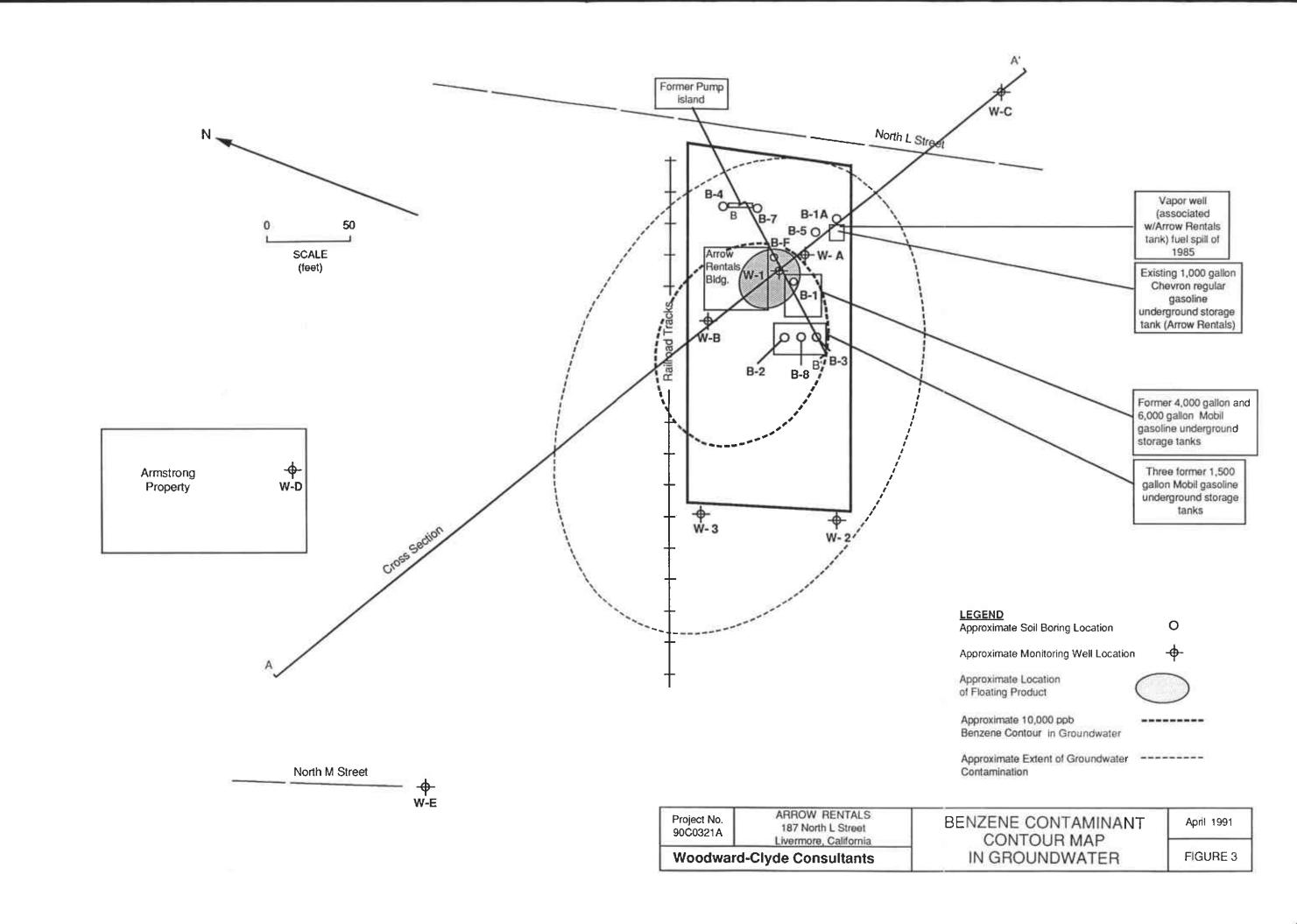
Table 2 LABORATORY TEST RESULTS OF WATER SAMPLES ***
Arrow Rentals, 187 North L Street, Livermore, CA

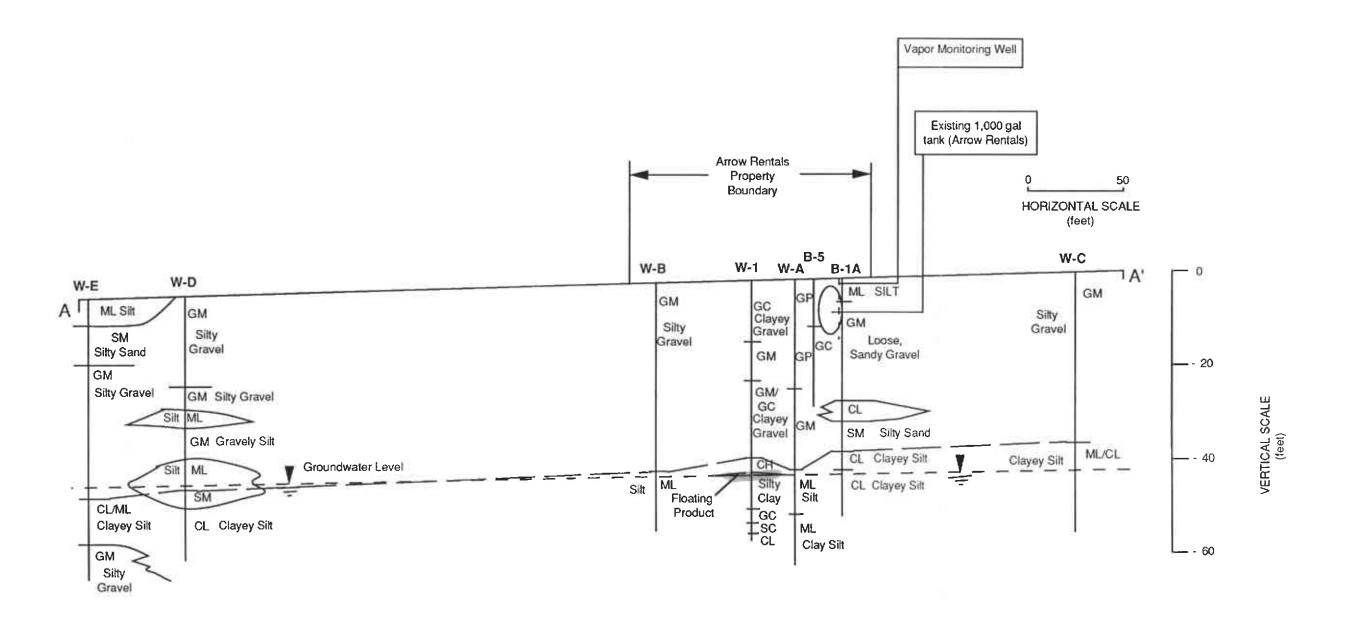
Number	Benzene (ppb)	Toluene (ppb)	Ethyl Benzene (ppb)	Xylenes (ppb)	TPH Gas. (ppm)	TPH Oles.	Org. Pb (ppm)	Phenol (ppm)	Naphth- alene (ppm)	2 Methyl Naphthalen (ppm)		Cr (ppm)	(ppm)	(bbw)	(bbw)	Zn (ppm)	Teirn Alky) Lend (g/gel)	Tetra Ethyl Loud %ofPh	Total Hydro Carbon ppm/(pob
W-1**	6.40%	4.70%	1.30%	5.10%				<1,000	<1,000	200							10.00		02=300
W-1 Dupe**	6.60%	4.70%	1.30%	5,10%				<1,000	<1,000	200									
W-A	6800	5500	620	3400	10	2.4		<0.1	<0.01	<0.01	<0.1	<0.1	<0.1	0.2	<0.1	<0.1			
W-A Dupe	6900	5600	820	6800							<0.1	<0.1	< 0.1	<0.1	<0.1	<0.1			
W-8	22000	7900	2000	4000	13	1.7		<0.1	<0.1	<0.1	<0.1	<0.1	<0.1	0.2	<0.1	<0.1			
W-B2*	21000	7300	1800	3700	21	1.6		<0.1	<0.1	<0.1	<0.1	<0.1	₹0.1	0.2	<0.1	<0.1			
w-c	<1	<1	<t< td=""><td><1</td><td><0.01</td><td><0.1</td><td></td><td><0.1</td><td><0.1</td><td><0.01</td><td><0.1</td><td><0.1</td><td><0.1</td><td><0.1</td><td><0.1</td><td><0.1</td><td></td><td></td><td></td></t<>	<1	<0.01	<0.1		<0.1	<0.1	<0.01	<0.1	<0.1	<0.1	<0.1	<0.1	<0.1			
W-D	1	2	1	1	0.1	€0.1		<0.1	<0.01	<0.01	<0.1	<0.1	<0.1	<0.1	<0.1	0.2			
W-E	<1	<1	<1	<1	<0.01	<0.1		<0.1	<0.01	<0.01	<0.1	<0.1	<0.1	<0.1	<0.1	<0.1			
W-EB	11	11	4	3	1.7	0.1		<0.1	>0.01	<0.01	<0.1	<0.1	<0.1	<0.1	<0.1	<0.1			
FT**	4.20%	8.90%	1.70%	6.10%			<50												
FT(Pres)**	4.20%	8.90%	1.70%	6.20%			<50								1				
W1-CH																	0.18g/ga	83%ofPb	
WA-CH		1					in.												7.6ppm
WB-CH																			2.4ppm
тв															1				(73ppb)

^{*-} Duplicate, **- Fuel Product, ***- Previous Laboratory Tests Summarized in Appendix C

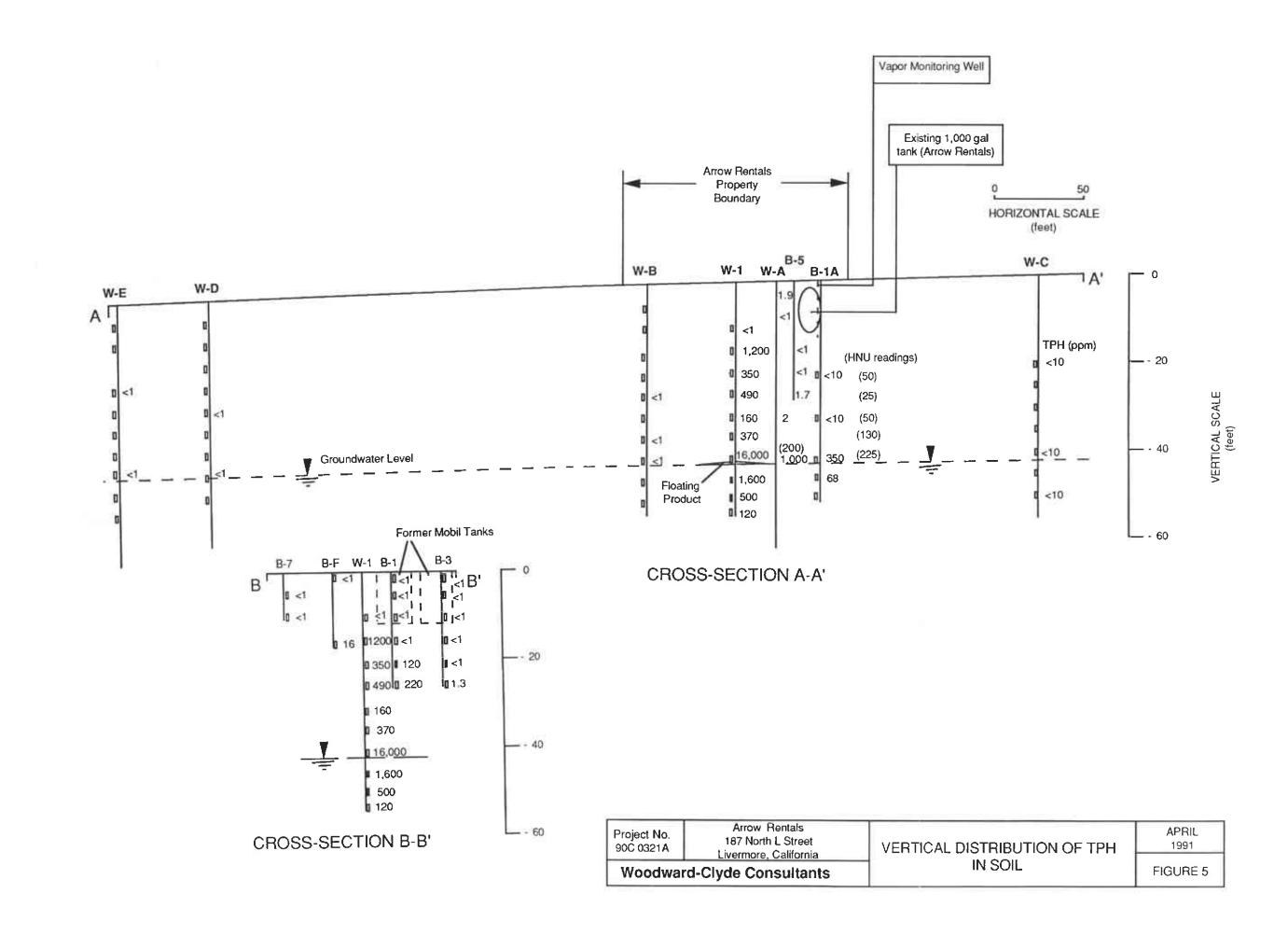


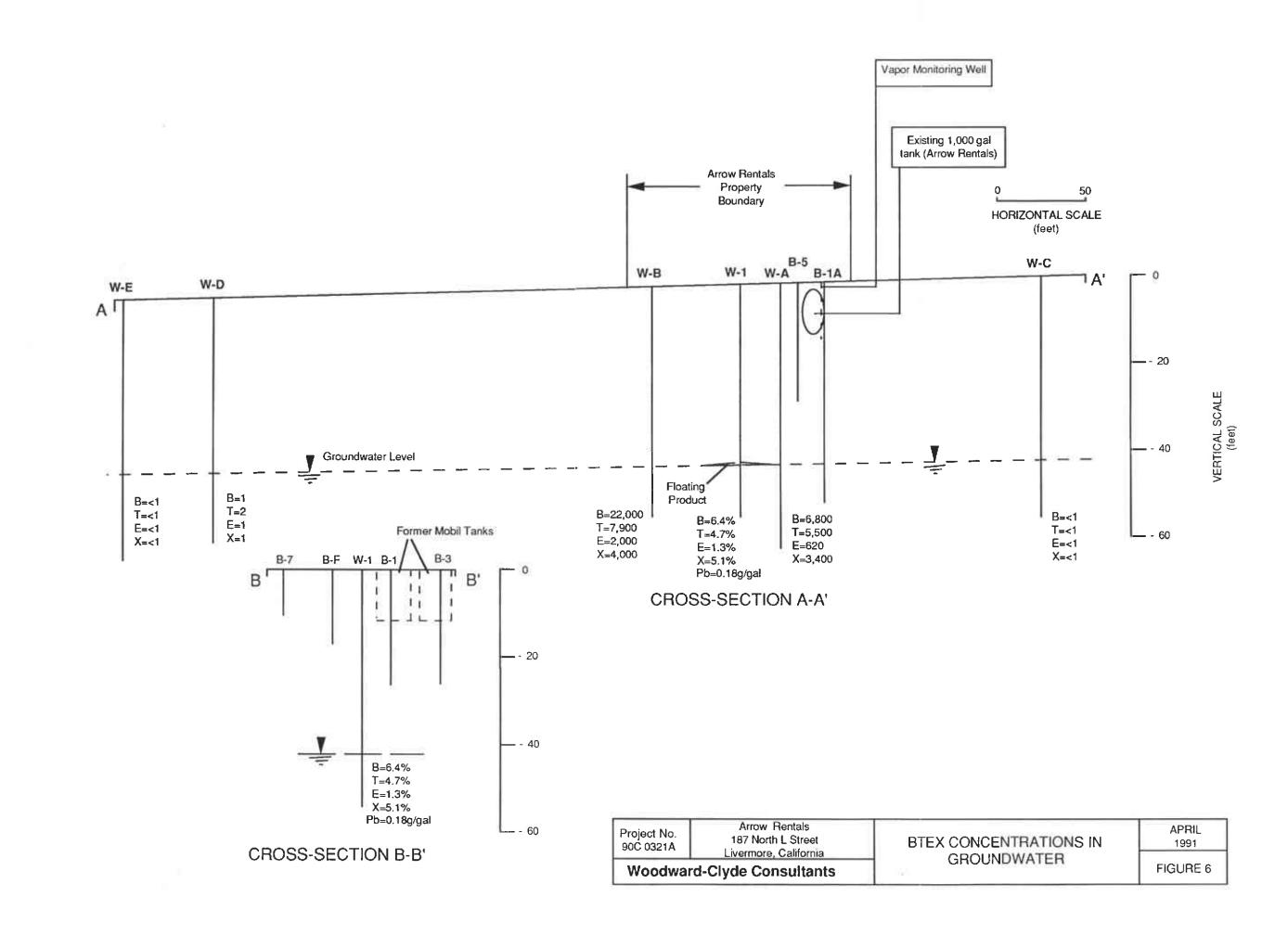


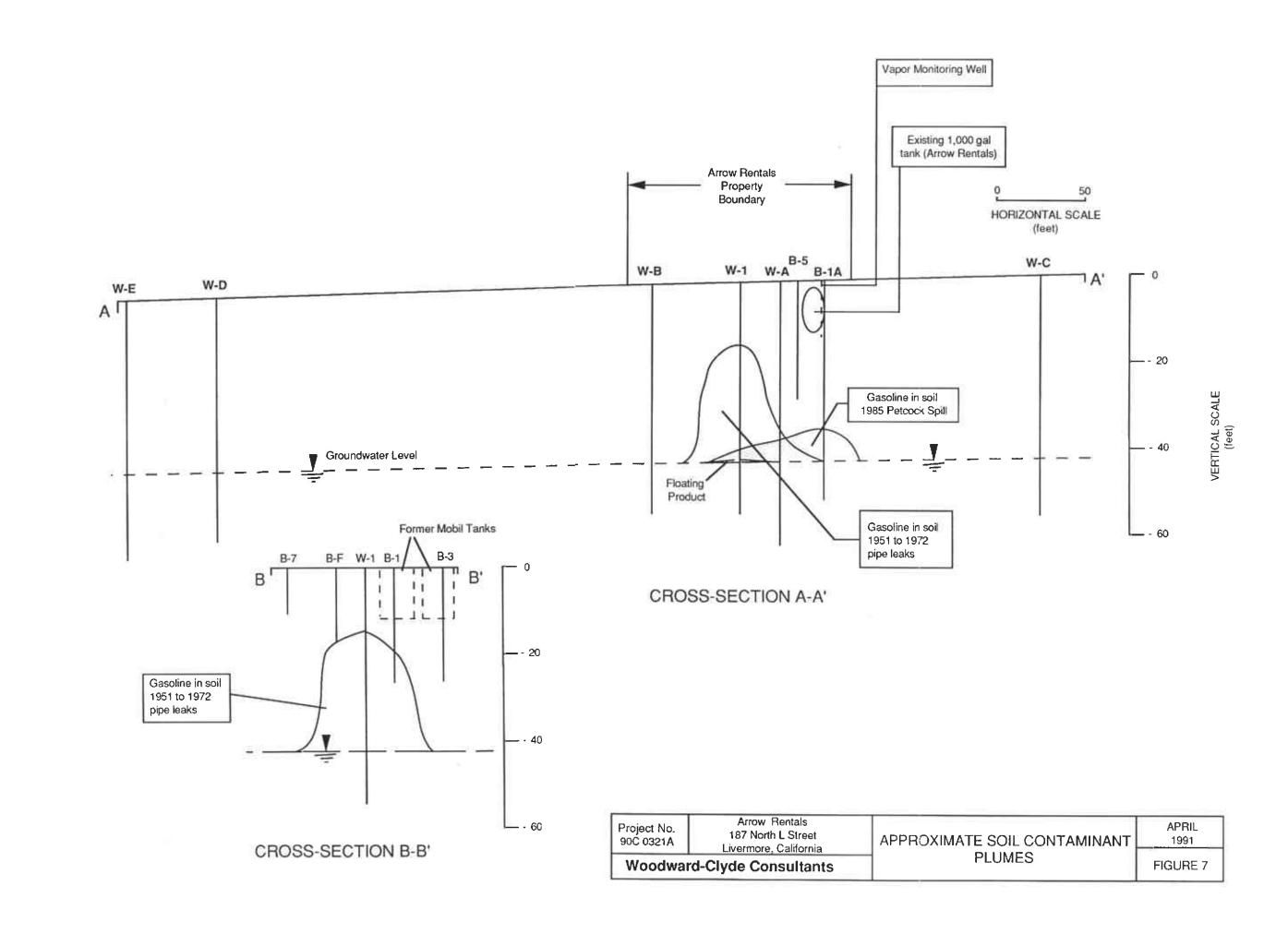




Project No. 90C 0321A	Arrow Rentals 187 North L Street Livermore, California	CROSS-SECTION A-A'	APRIL 1991
Woodwai	rd-Clyde Consultants		FIGURE 4







APPENDIX A TRACER RESEARCH REPORT



PREPARED FOR:

Woodward-Clyde Consultants 500 12th Street Oakland, California 94607-4014 (415) 874-3125

SHALLOW SOIL GAS INVESTIGATION **ARROW RENTALS** 187 NORTH "L" STREET LIVERMORE, CALIFORNIA

JULY 1990

SUBMITTED BY:

Tracer Research Corporation

190522S.REP 1-90-522-S



TABLE OF CONTENTS

INTRODUCTION 1
SHALLOW SOIL GAS INVESTIGATION - METHODOLOGY 2
EQUIPMENT 3
SAMPLING PROCEDURES 3
ANALYTICAL PROCEDURES 4
QUALITY ASSURANCE/QUALITY CONTROL PROCEDURES 5
RESULTS 6
CONCLUSIONS 7
A DDENINIV A
APPENDIX A ANALYTICAL DATA
APPENDIX B MAPS 9



INTRODUCTION

A shallow soil gas investigation was performed by Tracer Research Corporation (TRC) at Arrow Rentals located 187 North "L" Street in Livermore, California. The investigation was conducted on June 11 and 12, 1990 under contract to Woodward-Clyde Consultants. The purpose of the investigation was to characterize the nature and extent of volatile organic compounds (VOCs) present in the subsurface.

During this survey, a total of twenty-four soil gas samples were collected and analyzed in the field. The analytical equipment was calibrated for the following compounds:

benzene
toluene
ethylbenzene
xylenes
total hydrocarbons (THC)

Xylenes are reported as the total of the three xylene isomers and total hydrocarbons are reported as gasoline range constituents consisting of approximately C_4 - C_9 aliphatic, alicyclic and aromatic compounds.

The compounds in this suite were chosen as target compounds based on their suspected presence in the subsurface and amenability to soil gas detection. Soil gas samples were screened on the flame ionization detector (FID).



SHALLOW SOIL GAS INVESTIGATION - METHODOLOGY

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

Soil gas contaminant mapping helps to reduce the time and cost required to delineate underground contamination by volatile contaminants. The soil gas investigation does this by outlining the general areal extent of contamination. Conventional bore holes or observation wells are used to verify both the presence and extent of the subsurface contamination as indicated in the soil gas survey. In this manner, soil gas contaminant mapping can assist in determining the placement of monitoring wells. Thus, the likelihood



of drilling unnecessary monitoring wells is reduced. The soil gas survey is not intended to be a substitute for conventional methodology, but rather to enable conventional methods to be used efficiently.

EQUIPMENT

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

SAMPLING PROCEDURES

Sampling probes consisted of 7-14 foot lengths of 3/4 inch diameter hollow steel pipe that are fitted with detachable drive tips. Soil gas samples were collected by driving the steel probe to a depth of 8-10 feet below grade. Once inserted into the ground, the above-ground end of the sampling probes were fitted with a steel reducer and a length of polyethylene tubing leading to a vacuum pump. Gas flow is monitored by a vacuum gauge to insure that an adequate flow is obtained. To adequately purge the volume of air within the probe, 2 to 5 liters of gas is evacuated with a vacuum pump. During the soil gas evacuation, samples are collected in a glass syringe by inserting a syringe needle through a silicone rubber segment in the evacuation line and down into the steel probe. Ten milliliters of gas are collected for immediate analysis in the TRC analytical field van. Soil gas is subsampled (duplicate injections) in volumes ranging from 1 uL to 2 mL, depending on the VOC concentration at any particular location.

Sample probe vacuum, measured with a vacuum gauge, ranged from two to twenty-three inches Hg. Maximum pump vacuum was measured at twenty-six inches Hg.



ANALYTICAL PROCEDURES

One Varian 3300 gas chromatograph, equipped with a flame ionization detector (FID), was used for the sample analyses. The FID was used for the analysis of benzene, toluene, ethylbenzene, xylenes and total hydrocarbons. Separation of the FID compounds was achieved on a 6' by 1/8" OD packed column with OV-101 as the stationary phase. The column was contained in a temperature controlled oven and nitrogen was used as the carrier gas.

Hydrocarbon compounds detected in soil gas were identified by chromatographic retention time. Quantification of compounds was achieved by comparison of the detector response of the sample with the response measured for calibration standards (external standardization). Instrument calibration checks were run periodically throughout the day and system blanks were run at the beginning of the day to check for contamination in the soil gas sampling equipment. Air samples were also routinely analyzed to check for background levels in the atmosphere.

Detection limits for the compounds of interest are a function of the injection volume as well as the detector sensitivity for individual compounds. Thus, the detection limit varies with the sample size. Generally, the larger the injection size the greater the sensitivity. However, peaks for compounds of interest must be kept within the linear range of the analytical equipment. If any compound has a high concentration, it is necessary to use small injections, and in some cases to dilute the sample to keep it within linear range. This may cause decreased detection limits for other compounds in the analyses.

The detection limits range down to approximately 0.02 ug/L for compounds screened on the FID, depending on the conditions of the measurement, in particular, the sample size. If any component being analyzed is not detected, the detection limit for that compound in that analysis is given as a "less than" value (e.g. <0.02 ug/L). Detection limits obtained from GC analyses are calculated from the current response factor, the sample size, and the estimated minimum peak size (area) that would have been visible under the conditions of the measurement.



QUALITY ASSURANCE/QUALITY CONTROL PROCEDURES

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

- . Steel probes are used only once during the day and then washed with high pressure soap and hot water spray or steam-cleaned to eliminate the possibility of cross-contamination. Enough probes are carried on each van to avoid the need to reuse any during the day.
- Probe adaptors (TRC's patented design) are used to connect the sample probe to the vacuum pump. The adaptor is designed to eliminate the possibility of exposing the sample stream to any part of the adaptor. Associated tubing connecting the adaptor to the vacuum pump is replaced periodically as needed during the job to insure cleanliness and good fit. At the end of each day the adaptor is cleaned with soap and water and baked in the GC oven.
- Silicone tubing (which acts as a septum for the syringe needle) is replaced as needed to insure proper sealing around the syringe needle. This tubing does not directly contact soil gas samples.
- Glass syringes are usually used for only one sample per day and are washed and baked out at night. If they must be used twice, they are purged with carrier gas (nitrogen) and baked out between probe samplings.
- . Injector port septa through which soil gas samples are injected into the chromatograph are replaced on a daily basis to prevent possible gas leaks from the chromatographic column.



- . Analytical instruments are calibrated each day by analytical standards from Chem Service, Inc. Calibration checks are also run after approximately every five soil gas sampling locations.
- . Subsampling syringes are checked for contamination prior to sampling each day by injecting nitrogen carrier gas into the gas chromatograph.
- Prior to sampling each day, system blanks are run to check the sampling apparatus (probe, adaptor, 10 cc syringe) for contamination by drawing ambient air from above ground through the system and comparing the analysis to a concurrently sampled ambient air analysis.
- All sampling and subsampling syringes are decontaminated each day and no such equipment is reused before being decontaminated. Microliter size subsampling syringes are reused only after a nitrogen carrier gas blank is run to insure it is not contaminated by the previous sample.
- . Soil gas pumping is monitored by a vacuum gauge to insure that an adequate gas flow from the vadose zone is maintained. A reliable gas sample can be obtained if the negative pressure reading on the vacuum gauge is at least 2 inches Hg less than the maximum pressure of the pump.

RESULTS

Concentrations of hydrocarbons in the C_4 - C_9 range were detected slightly above background in the subsurface at the Arrow Rentals site in Livermore, California. A total of 24 soil gas samples and 6 ambient air samples were collected at this site. The total hydrocarbons detected in these samples were not attributable to benzene, toluene, ethyl benzene or xylenes as no significant levels of these specific compounds were detected in the soil gas. The total hydrocarbons reported were generally lighter, early eluting compound in the C_4 - C_9 range. The analytical data reported in micrograms per liter (ug/L) for this site is condensed in Appendix A and maps showing sampling locations and isoconcentration contours are included in Appendix B.



An isoconcentration contour map was generated for total hydrocarbons in the C4-C9 range (Figure 2). The highest level of total hydrocarbons was detected at sampling location SG-3 with a concentration of 2 ug/L. This location, along with sampling location SG-2, was contained within a 1 ug/L contour line that was open to the north. Sampling location SG-15 with a concentration of 1 ug/L was also contained within an open-ended contour at the north-east portion of the site. Several other sampling locations on the site were contained within 0.1 ug/L contour lines. These 0.1 ug/L contour line are somewhat speculative and border on being insignificant but the sake of completeness they are included. Additional sampling locations would be necessary to completely define the plumes depicted.

CONCLUSIONS

The soil gas survey conducted at the Arrow Rentals Site found slightly elevated levels of hydrocarbons (C_4 - C_9) in the subsurface. Several contour lines were generated at this site, depicting a small plume just above the level of significance. The highest concentrations of total hydrocarbons were detected in sampling location SG-3 located near the north corner of the office building. To further delineate and define the total hydrocarbon plume additional samples would be necessary.

Tracer Research Corporation

APPENDIX A: ANALYTICAL DATA

WOODWARD CLYDE CONSULTANTS/ARROW RENTALS/LIVERMORE, CALIFORNIA JOB#1-90-522-S

06/11/90

CONDENSED DATA

COMPENDE			ETHYL		
	BENZENE	TOLUENE	BENZENE	XYLENES	THC
SAMPLE	ug/l	ug/!	ug/l	ug/l	ug/l
AIR	< 0.04	< 0.05	< 0.05	< 0.07	< 0.05
SG1-10'	< 0.04	< 0.05	< 0.05	< 0.07	0.4
SG2-10'	< 0.08	<0.1	< 0.1	< 0.1	. 1
SG3-10°	< 0.04	< 0.05	< 0.05	< 0.07	2
SG4-10'	< 0.08	< 0.1	< 0.1	< 0.1	< 0.1
SG5-10'	< 0.08	<0.1	< 0.1	< 0.1	< 0.1
SG6-10'	< 0.04	< 0.05	< 0.05	< 0.07	0.6
AIR	< 0.04	< 0.05	< 0.05	< 0.07	2
SG7-10*	< 0.08	< 0.1	< 0.1	<0.1	< 0.1
SG8-10'	< 0.08	<0.1	< 0.1	<0.1	< 0.1
SG9-10'	< 0.08	<0.1	< 0.1	< 0.1	< 0.1
SG10-10'	< 0.08	<0.1	< 0.1	< 0.1	< 0.1
SG11-10'	< 0.08	<0.1	<0.1	<0.1	0.7
AIR	<0.04	< 0.05	< 0.05	< 0.07	0.3

Analyzed by: P. Bausman
Checked by: K. Proctor
Proofed by: X. Auglorides

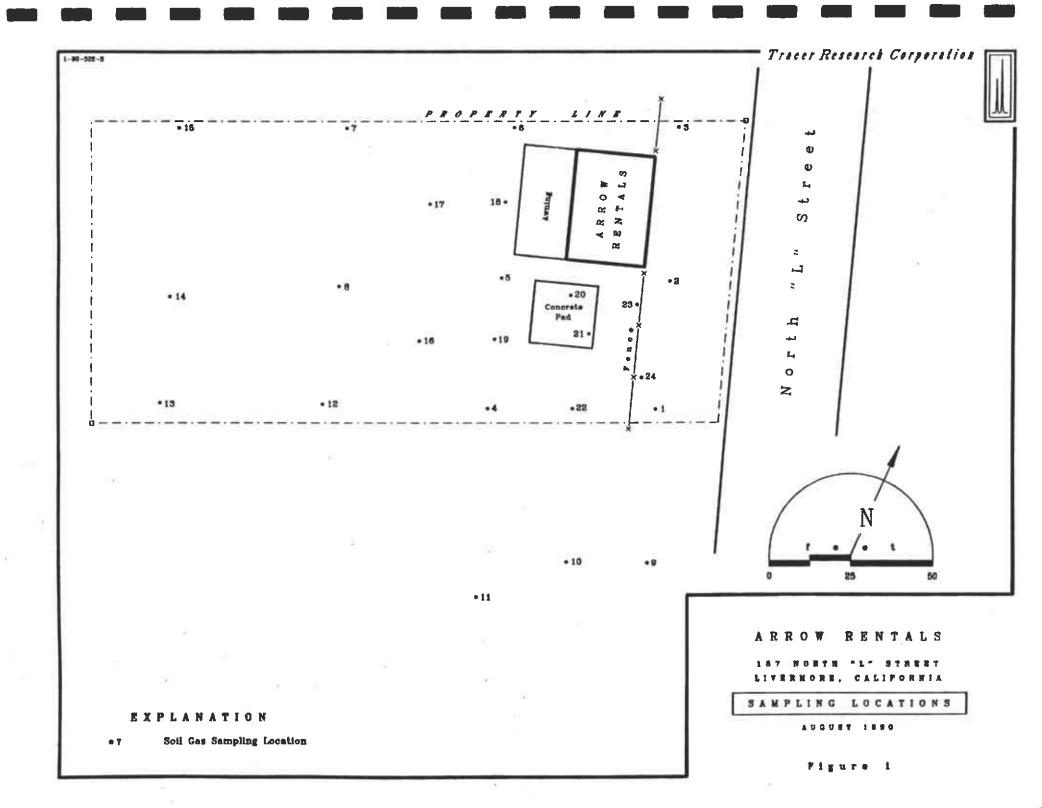
WOODWARD CLYDE CONSULTANTS/ARROW RENTALS/LIVERMORE, CALIFORNIA JOB#1-90-522-S

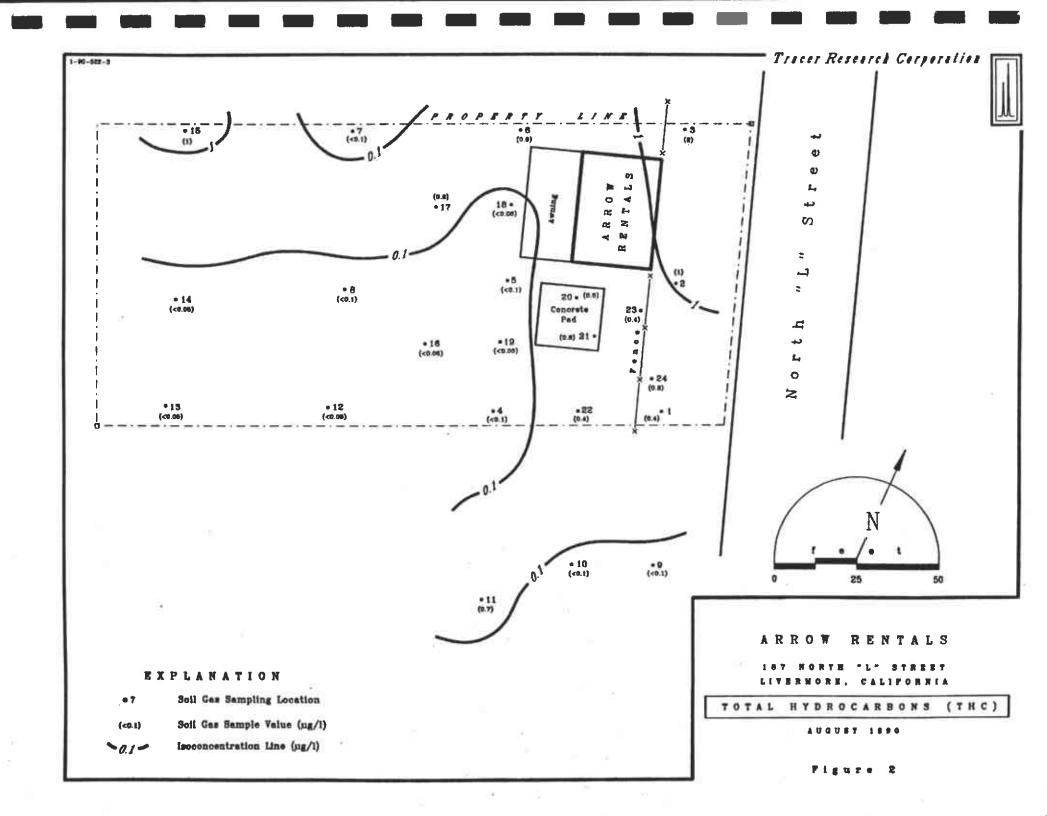
06/12/90 CONDENSED DATA

CONDENSE			ETHYL	XYLENES	THC
	BENZENE	TOLUENE	BENZENE		
SAMPLE	ug/l	ug/l	ug/l	ug/l	ug/l
SG12-10'	< 0.04	< 0.05	< 0.06	< 0.06	< 0.05
AIR	< 0.04	< 0.05	< 0.06	< 0.06	< 0.05
SG13-10'	< 0.04	< 0.05	< 0.06	< 0.06	< 0.05
SG14-10'	< 0.04	< 0.05	< 0.06	< 0.06	< 0.05
SG15-8'	< 0.04	< 0.05	< 0.06	< 0.06	1
SG16-10'	< 0.04	< 0.05	< 0.06	< 0.06	< 0.05
AIR	< 0.04	<0.05	< 0.06	< 0.06	< 0.05
SG17-10'	< 0.04	< 0.05	< 0.06	< 0.06	0.2
SG18-10'	< 0.04	< 0.05	< 0.06	< 0.06	< 0.05
SG19-10'	< 0.04	<0.05	< 0.06	< 0.06	< 0.05
SG20-10'	<0.04	< 0.05	< 0.06	< 0.06	0.5
SG21-10'	< 0.04	< 0.05	< 0.06	< 0.06	0.8
SG22-10'	< 0.04	<0.05	< 0.06	<0.06	0.4
SG23-10'	< 0.04	< 0.05	< 0.06	< 0.06	0.4
SG24-10'	< 0.04	< 0.05	< 0.06	< 0.06	0.2
AIR	< 0.04	< 0.05	< 0.06	< 0.06	< 0.05

Analyzed by: P. Bausman
Checked by: K. Proctor
Proofed by: A. Naslande







APPENDIX B BORING AND WELL LOGS

DAWPDOCSV90C0321.ALL\S 0611911455

Woodward-Clyde	Consultants
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PROJECT NAME ARROW RENTALS NO. 90C0321A

BORING	G LOCATI	ON MA between	on Mil-1 and ank							ELEVAT	ION A	ND DAT	UM				
DRILLING AGENCY Kwithaug DRILLER Rod Furlow Brian Vincent										DATE STARTED 7/12/90 -> 7/12/90							
-	IG EQUIP		Drill			Brian Vi	ricent		7	COMPLI	_		63 ft		SAMPLER 2 in.		
	IG METHO				DRILL B	IT			7	NO. OF		IST.			UNDIST.	в	
-			meter Schedule 40 P	vc			_		7	WATER LEVEL		IRST	50 f	ı.	COMPL.	24 HRS. 44,25	
TYPE O	F PERFO		otted PVC		FROM	57 -1/2	छ	42 6	FT	LOGGE	BY:				CHECKED B		
SIZE AM	ID TYPE (OF PACK Monterey	r Sand # 2/12	32	FROM	63	TO		뒤								
<u> </u>		NO. 1 Bentonite		111	FROM	40	TO		FT		Loit	Gruer	nberg				
TYPE O	FSEAL		4° agg - Grout/ to set Christy box		FROM	36 - 1/2	то	0	FT				-				
		Quick max	w ser unnsty DOX	نما		- 11E		Ť	_		_	SAM	PLES				
DEPTH (feet)			LITHOLOGIC DESCRIPTION					Well C	com	pletion em	DEPTH (feet)		BLOW. COUNTS	q		ARKS Loss, Odor, etc.)	
	Asoheli	Concrete - 4 inches						222									
										2.2	9	-					
-								222		2,2	- 54	1					
1		0	alla anciette e S.	_						2.2	5 -	1					
5 =	Cutting	s-carevel (GP), some	silt, much resistance	₽.			1			2,2	3	-		No sar	nple taken		
-								^^^		2,2,2	- 13	-					
-										2.2	1	1					
4			22					22.2		200	5	1					
10 -	Grades	to more silty						22.2	2		10 *		35 50/	Sampl HNU =	e: W-A-10 : 1 ppm		
													6				
1								222		222	19	-					
-		99						222		2.2.	19	1					
15-								22.2		222	15	\top	50/ 6	little re	covery		
- 5	3							22.2		224	1 2	_	°	no san		û.	
]								222		22	8	-					
-								222		2 0 0	2	+					
20 —	ŠILTY	, GRAVELLY SAND	(GM), gray brown, g	ravels	to 1/2 , v	rery		22.2			20 -		35 50/		le: W-A-20		
- 1	coars	e to coerse sand, tred	ce clay, wet, loose, or	dor.				222		2.22	8		6	mivu=	15 ppm		
]							1				3]					
											1 8	-					
25 —			C					1 4 41		* *	25		50/		e: W-A-25		
-										100	0		6	HNU =	110 ppm		
								222			1 3	1					
								1-2-24			1						
30 -											30	-	45		e: W-A-25		
-				36				[222	. 0		50/ 6	Very st	rong odor in l	porehole	
-								1-2-2			120						
3										222	3	-					
1 7								10,00		1,000			1	I .			

NO. 90C0321A **Woodward-Clyde Consultants** PROJECT NAME ARROW RENTALS SAMPLES LITHOLOGIC DESCRIPTION REMARKS (Drill Rate, Fluid Loss, Odor, etc.) DEPTH (feet) Well Completion Diagram 50/ Sample: W-A-35 HNU > 200 ppm SET (ML), brown with black patches, with some clay, some coarse sand, soft to medium stiff, moist. Sample : W-A-40 HNU = 250 ppm 40 5 9 20 Grades to homogenous brown, abundant clay, no sand, medium Sample: W-A-45 35 HNU = 150 ppm 30 35 50 -50 -Sample: W-A-50 HNU = 100 ppm Grades to saturated 60 60 Boring terminated at approximately 53 ft. below grade.



PROJECT NAME ARROW RENTALS NO. 9000321A

BORIN	IG LOCAT	ION 187 N. L Street, Livermore, Co	A						ELEVA	TION A	AND DAT	TUM					
DRILLI	NG AGEN	CY Kvilhaug		DRILLER	Mike C Joel Vir				DATE S			10/90		-	7/10/9	90	
DRILLI	NG EQUIP	MENT B 53 Mobile Drill							COMPL	ETIO	V	50		SAMPLER	2 in) ,	
DRILLI	NG METH	DD Hollow Stern Auger		DRILL B	ıτ				NO. OF		DIST.			UNDIST.	10		
SIZE A	ND TYPE (OF CASING None	- 17					\exists	WATER	1	FIRST	47 ft		COMPL.		24 HRS	
TYPE (OF PERFO	RATION None		FROM		то		FI	_					CHECKED	BY:		×
SIZE A	ND TYPE	OF PACK None		FROM		то		FT									
\vdash		NO.1 None	17	FROM		TO		ቨ		Lois	Gruent	erg					
TYPE	OF SEAL	NO.2 Grout	**	FROM	51-1/2	то	0	Fī									
\vdash			DXX					_	_		SAM	PLES					
DEPTH (feet)		LITHOLOGIC DESCRIPTION					Well (Com	pletion un	DEPTH		BLOW. COUNTS	ų	RE Odli Rate, Flu	MARK! id Loss		rtc.)
\vdash	Aspheli	Concrete - 4 inches.					\neg		Т		+	-					
5-	SANDY	s- SILT (ML), brown with orange, graveled , SILTLY GRAVEL (GM), brown-gray a moist, gravels to 1/2", loose.			very					5.		7 13 15		e: B-1-5 no respons	е		
10-	Grades	to more silt, some clay, slightly wet								10		10 19 23	Sample HNU=	e: B-1-10 1 0 ppm			
15 — - -	Grades	to moist with pockets of wet clay.								151		35 50 50	Sample HNU=1	e: B-1-15 no responsa		ĸ	
20 -	Noticeat	ole ador and sheen in packets of maistu	re.		E					20		20 40 36	Sample HNU=	e: B-1-20 50 ppm	(4)		
25 — -	CLAYE loose, i	Y SILT (CL), brown, gravels to 1/2", so moist to wet.	. — me very	— — coarse sa	and,	-	78			25 ·		20 35 35	Sample HNU= 2	: 8-1-25 15 ppm			
30 -	SILTY Si Irace cla	AND (SM), brown-gray, very coarse sar y, moist to wet, loose to medium dense.	id, grave	els to 1/2 "						30		12 17 27	Sample HNU = !	8-1-30 50 ppm			
-											-						

	Woodward-Clyde Consultants		F	ROJECT	T NAI	/E		ARRO!	W RENTALS NO. 80C0321A
DEPTH (feet)	LITHOLOGIC DESCRIPTION		W	ell Complet Diagram		DEPTH (feet)	SAM	BLOW-	REMARKS (Drill Rate, Fluid Loss, Odor, etc.)
35 -	CLAYEY, SANDY S&T (SM-CL), brown-gray with some orange, coarse aand, wet. Very easy drilling encountered.					35		5 12 5	Sample: B-1-35 HNU = 130 ppm
40 -	CLAYEY SILT (CL), hornogenous brown, noticeable hydrocarbon odor soft, wet.					40 -		4 5 5	Sample : 8-1-40 HNU =225 ppm
45 -	Grades to mottled with some, small, white and black pebbles, odor.	모				45		233	Sample: B-1-45 HNU= 180 ppm
50	Grades to gray with rare layer of coarse sand, odor, saturated.					50		10 15 17	Sample: 8-1-50 HNU= 110 ppm
55					21	\$5	W. Carlotte		Boring terminates at approximately 51-1/2 feet below grade.

BORING LOCAT	TION W.B.	ELEVATION AND DATUM								
DRILLING AGEN	-	DATE STARTED 7/12/90 - 7/13/90								
DRILLING EQUI	PMENT B 61 Mobile Drill		Brian V		COMPLE		55	SA	MPLER	
DRILLING METH		DRI	ILL BIT		NO. OF	DIST.		UN	(DIST	6
SIZE AND TYPE		0 PVC			WATER		48 ft.	cc	OMPL.	24 HRS 44.62
TYPE OF PERF		FFRC	XM 55	10 40 F	LOGGE	BY:		Cł	ECKED BY:	12
SIZE AND TYPE		FFK	1000	то <u>яг</u>	7	_				
	NO. 1 Bentonite Pellets	FFK		70 F		ois Gruen	berg			
TYPE OF SEAL	Grout/Quick mix to set	î PK	M.	το ₀ F	Ť		-			
	Christy box	l'and	30	ΤŤ	-	S	AMPLES			
(leest)	LITHOLOGIC DESCRIPTION			Well Co Diag	mpletion gram	DEPTH (leet)	BLOW- COUNTS	(Drill I	REMAR Rate, Fluid Lo	KS ss, Odor, etc.)
5 SILTY clay, d	GRAVELS (GM), brown to dark brown, amp, loose. Gravels 1/2 to 2 * des to more sitty, moist.		clay, moist			15	122 255 50 6 40 50/ 6	sampler HNU = 10 Little reco HNU= 15 Sample: W HNU = 2 p	e taken very, 2" grav ppm very ppm	relstuck in

	Woodward-Clyde Consultants		P	ROJECT	NAI	AE			W RENTALS NO. 9000321A
DEPTH (feet)	LITHOLOGIC DESCRIPTION		We	ill Completi Diagram	ion	DEPTH (feet)	SAMPL	COUNTS IN	REMARKS (Drill Rate, Fluid Loss, Odor, etc.)
35 —			1800		190	35-		50/ 6	Sample : W-B-35 HNU >20 ppm
40 -	SILT (ML), brown with black patterns, some clay, medium-low plasticity, moderately soft, damp to moist.					49-		8 12 15	Sample : W-B-40 HNU = 2 ppm
45 —	Grades to brown with green-yellow and gray, black patches, mottling	₩				45		17 19 26	Sample: W-8-45 HNU = 30 ppm
50 -						so_		5 25 40	Sample: W-B-50
55						55			Boring terminated at approximately 55 leet below grade.
60 -						80-			× 8

FIELD LOG OF BORING NO. W-B

SHEET 2

BORING	LOCAT	ION A	cross Arrow Rentals on Nor	th L St						ELEVAT	TION A	ND DA	NUTUM			
DRILLIN	IG AGEN	CY KVIII	naug		DRILLER	Mike C			П	DATE S	TARTE	D -	7/11/90			7/11/90
PALIN	G EQUIP	MENT E	53 Mobile Drill			41				COMPL	ETION	_	55 ft.		SAMPLER	2 in
RILLIN	G METH	00 F	tollow Stem Auger		DRILL B	IT				NO. OF	į D	IST.			UNDIST.	6
IZE AN	D TYPE (OF CASING	2 "-diameter Schedule 4	io PVC						WATER	_	IRST	47	ħ.	COMPL.	24 HRS. 43.34
YPE O	F PERFO	RATION	0.010 Slotted PVC		FROM	A in the second	70	45 10	FT	LOGGE	D BY:				CHECKED	
IZE AN	D TYPE	OF PACK	Monterey Sand #2	157	FROM	55	то	37 - 1/2	FI							
		NO. 1	Bentonita Pellets	11	FROM	37 - 1/2	то	35	FT		Lois (Gruer	berg			
YPE OF	FSEAL	NO. 2	Grout/ Quick mix to	2.0	FROM	35	TO	0	FI							
\neg		1	set Christy box	L^_^	1			Ť	_			SAI	WPLES			
Sec.			LITHOLOGIC DESCRIPTION						Com	pletion am	DEPTH (feet)		BLOW: COUNTS	(AARKS d Loss, Odor, etc.
\neg	Asphali	Concrete	- 5 inches 2 -diameter, silty, brown, dr				1	2.3					Т			
Ţ								2.2								
1										202	į	1				
-												1				
5 —	SILTY (GRAVELAG	RAVELLY SILT (GM) brow slightly damp, loose.	n, some	siit, some	very				222	5 -	⊢	+			
1	inzi gle i ui	LNS, UI Y 10	anginiy darip, loose.					22.2		200	Î	_		INO SAI	nple taken.	
1											1					
]]				
10 -											10 -	L		No sar	nple taken.	
								2.2		222						
-								200		22		╀	1			
-												1				
+											İ	1				
15-								2.2		2-2-	15~	T	1	No san	nple taken.	
1								-2.2		222		\vdash	1			390
1								200		200	ļį					
4												-				
20 -										1,74	20 -		20	Sample	e: W-C-20	
+													40 35	HNU=	no response	•
1						2	1				į					
]			£)							2.22]				
25 -			(1)					200			25 -	_	17	Samnle	: W-C-25	
1	Grades	to more s	ilt with trace clay, moderate	y stiff.						200			18 20		o response	
-										2.2	=					
-											1	1				
+										^^^	1					
30 -								2.3		2,22	30 -		20 35		: W-C-30 no response	
1				9.7				2.2		^^^	į		40	11140 =	in resputise	+ 1
4								3.3			1	1				
-								22.2		22.2	•	1				

Woodward-Clyde	Consultants
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PROJECT NAME _____ARROW RENTALS NO. __90C0321A

BORING LOCATION 1951 Chestnut, Livermore, CA (in backyard of residence of William Armstrong)									ELEVATION AND DATUM								
DRILLING AGENCY Kvilhaug DRILLER Rod Furlow Brian Vincent										DATE STARTED DATE FINISHED 7/12/90 - 7/12/90							
DRILLING	EQUIP	MENT B 61 Mobile Onli							COMPLE DEPTH	TION		57 - 1	/2 fL	SAMPLER	2 in		
DRILLING	METH	DD Hollow Stem Auger		DRILL B	IT			1	NO. OF		ST.			UNDIST.	6		
	_	OF CASING 4 *-diameter Schedule 40	PVC					1	WATER		RST	46 f	t,	COMPL.		IRS. 42.19	
TYPE OF	PERFO	RATION 0.010 Slotted PVC		FROM	57-12	TO	42	_	LOGGE	BY:				CHECKED	BY:	-	
SIZE AND	TYPE		121	FROM	57 - 1/2	то	39 - 1/2	FI									
		NO.1 Bentonite Pellets	11	FROM	34	то		FI		Lois G	iruenbe	erg					
TYPE OF	SEAL	NO.2 Grout		FROM	32	TO	0	FT									
$\neg \tau$	_		h-0-			Т	Ť	_			SAME	_		-			
DEPTH (feet)		LITHOLOGIC DESCRIPTION				1	Well C		DEPTH (feet)		BLOW- COUNTS	(MARKS id Loss, Od	KS oss, Odor, etc.)		
	Some 4	prass and dry, silty soil in backyard.				+	1000		1"2"2	25	-	8 8	-		_		
4							222		200	-							
-							2.2.2		-2.2	97							
+							2-2-		-2-2	-							
- 1							22.2		22.2				Samul	n met token			
5-	Cutting	s - SILTY (GM), brown, gravels to 3/8 *	dry.			1	2-2-		2.2.2	5-		50/ 6	HNU «	e not taken : 1 ppm			
1							222		222								
]							222		-2-2								
4							2222		22.2				l.				
10-	GRAV	/ELLY SILT (GM), brown with black and	oranoe	blotches.			22.2		200	10-		30	Little n	ecovery		12	
4	fractu	red gravels to 1/2", trace clay, moist, v	ery dens	€.			2.22		2-2-			25 28	HNU	c1 ppm			
1-						1	22.2		200							5	
+							2.22		2.22	1							
-							2.22		2.22	15-				2			
15-							222			15		25 50/		le: W-D-15 = 1 ppm			
]							22.22					6			(37		
]							222		222		1						
4							22.2		-2.2		1						
20 -	SILTŸ	GRAVEL (GM), brown, gravels to 1°, tra	ice day.	loose, m	pist, very		22.2		222	20 -	1	50/ 6		ie not taken			
	dense .				-		222		2.2		1_	Į"	HNU	1.5 ppm	80		
+							22.2		2.22		1						
7							2.2.2		22.2								
25						_	-2-2		2.2.2	25 -		12	Sampl	e: W-D-25			
	SILT	ML), brown with black patches, some o	isy, soft.						200		-	12	HNU.	1.75 ppm			
							222		2.2		-	1					
-									200		1						
+							22.2				1			10			
30 -	GRAV	ELLY SILT/SILTY GRAVEL(GM), brow	vn with o	ray rocks		-	222			30		50/		stuck in sam			
-	trace	clay, moist to wet, rock stuck in sample					222					6	130		30		
j							222				-						
							222		222		+						
								_	A A	RDIO	OFR	YRING	NO	Wb SI	ÆET 1	OF _	

Woodward-Clyde Consultants				PROJECT NAM					OW RENTALS NO. 90C0321A
(leet)	LITHOLOGIC DESCRIPTION		We	li Compli Diagram	etion 1	DEPTH (feet)	SAMF	BLOW.	REMARKS (Drll Rate, Fluid Loss, Odor, etc.)
35 -	SILT (ML), brown with bleck patches, some clay, moderately stiff.					36		12 15 20	Sample: W-D-35 HNU = 2 ppm
40 -	SANDY SILT (SM), brown, homeogenous, coarse sand, dense, wet Some sloughing occured from 40 to 34 feet, just above the sand pack, during drilling.					40		30 50/ 6	Sample :W-D-40 HNU = 1.5 ppm
5 - 5	CLAYEY SILT (CL), brown, medium plasticity, moderately soft, wet.	Ξ			100 VIN 100 VI	45-		12 15 22	Sample: W-A-45 HNU = 150 ppm
50 —						50			
55 -					1 M 12	55			2 0
1 00						eo –			Boring terminated at approximately 57-1/2 feet below grade.

Woodward-Clyde (Consultants
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PROJECT NAME __

ARROW RENTALS

NO 90C0321A

BORING LOCAT	TION End of M Street, Livermore, CA						ELEVA	TION	AND DAT	UM .				
DRILLING AGEN	DATE S	DATE STARTED 7/10/90 7/10/90												
DRILLING EQUI	PMENT B 53 Mobile Drill	COMPL	ETIO											
DRILLING METH	NO. OF		DIST.			UNDIST.	8							
SIZE AND TYPE OF CASING 2 *-diamter Schedule 40 PVC									FIRST	47 fL		COMPL	24 HRS. 43.08	
TYPE OF PERF	ORATION 0.010 Slotted PVC		FROM	60 -1/3	то	40 - 1/2 FT	LOGGE	D BY				CHECKED B	Y: 15	
SIZE AND TYPE	OF PACK Monterey Sand # 2/12		FROM	61	ТО	37 F		1 -'	. C-:	harr				
TYPE OF SEAL	NO.1 Bentonite Pellets	7	FROM	30	то	29 F	1	FOI	s Gruen	iberg				
	NO. 2 Sack, 3/4" agg Grout / Quick Seal to set Christy box		FROM	29	то	0 F		,	1	DI 50				
F.	LITHOLOGIC					W-80-		Į	-	PLES		REM	ARKS	
See the	DESCRIPTION					Well Cor Diag	LSILD LICIDATION	DEPTH		BLOW. COUNTS		(Drill Rule, Fluid Loss, Odor, etc.)		
5 — SILTY grave	ittings - SILT (ML), gravels to 1/2°, brown things - SILT (ML), gravels to 1/2°, brown strings - SILT (ML), prown and gray, some graves with black veins, fryable, loose. VELLY SILT/ SILTY GRAVEL (GM), brown and rocks.	vels to 1	/2°. qua	actured,				10		100 141 18 200 300 440 12 225 25 25 26 28 28 28 28 50/5	Sample HNU= Sample HNU= Sample	e: W-E-5 2 ppm e: W-E-10 2.5 ppm e: W-E-15 no response e: W-E-20 no response		

Woodward-Clyde Consultants					PROJECT NAMEARROW RENTALNO. 90003						
			_			SAM	PLES				
LITHOLOGIC DESCRIPTION		Well Completion Diagram		DEPTH (best)		SLOW- COUNTS	REMARKS (Drift Rate, Fluid Loss, Odor, etc.)				
				127	1						
Grades to wet. Very soft drilling encountered. Sloughing occured at approximately 37 to 30 feet during well installation.					36-		12 15 12	Sample: W-E-35 HNU < 1 ppm Little recovery			
CLAYEY SILT (ML-CL), brown, some medium sand, homogeneous, soft, moist					40 -		2 3 5	Sample : W-E-40 HNU < 1 ppm			
Grades to less sand, brown with faint black blebs.	후	18		5	45		5 8 8	Sample: W-E-45 HNU = no response			
More difficult drilling encountered. GRAVELS (GM), pebbles black/gray, saturated, some silt.					50	 - -	20 30 25	Sample not taken.			
					55	-					
					60	-					
	CLAYEY SILT (ML-CL), brown, some medium sand, homogeneous, soft, moist. Grades to less sand, brown with faint black blebs.	CLAYEY SILT (ML-CL), brown, some medium sand, homogeneous, soft, moist. Grades to less sand, brown with faint black blebs.	Grades to wet. Very soft drilling encountered. Sloughing occured at approximately 37 to 30 feet during well installation. CLAYEY SILT (ML-CL), brown, some medium sand, homogeneous, soft, moist. Grades to less sand, brown with faint black blebs.	Grades to wet. Very soft drilling encountered. Sloughing occured at approximately 37 to 30 feet during well installation. CLAYEY SILT (ML-CL), brown, some medium sand, homogeneous, soft, moist. Grades to less sand, brown with faint black blebs.	Grades to wet. Very soft drilling encountered. Sloughing occured at approximately 37 to 30 feet during well installation. CLAYEY SILT (ML-CL), brown, some medium sand, homogeneous. Soft, moist. Grades to less sand, brown with faint black blebs.	Grades to wet. Very soft drilling encountered. Soughing occured at approximately 37 to 30 feet during well installation. CLAYEY SILT (ML-CL), brown, some medium sand, homogeneous, soft, moist. Grades to less sand, brown with faint black biebs. A5. GRAVELS (GM), pebbles black/gray, saturated, some silt.	Crades to wet. Very soft drilling encountered. Stoughing occurred at approximately 37 to 30 feet during well installation. CLAYEY SILT (ML-CL), brown, some medium sand, homogeneous, soft, moist. Grades to less sand, brown with faint black blebs. More difficult drilling encountered. GRAVELS (GM), pebbles black/gray, saturated, some silt.	Grades to wet. Very soft drilling encountered. Southing sociated at approximately 37 to 30 feet during well installation. CLAYEY SILT (ML-CL), brown, some medium sand, homogeneous, soft, moist. Grades to less sand, brown with laint black biebs.			

Woo	odi	ward-Clyde Consultants						Project: Ar	tow l	Rent	als 90C0321A
Weli	Ni	umber and Location: Boring of at former	Arro	w Gas I	Pump			Elev. and [atur	n;	
Drilli	ng .	Agency: Arrow Rentals		Oriller:	Tony Sullins			Date Starte Date Comp			11/91
Drilli	ng	Equipment: Bobcat					_	Total Depti			12/91
		Method: flight auger		Drill bit:	8- inch			Sampler:			4inch brass liners
-		d Type of Casing: NA							First		Compl. 24 HRS
		Perforation: NA		From:	ft To:		ft	Level: No. of	Dist.	_	Undist.
_		Type of Pack: NA	9:1	From:			ft	Samples	Didi.	2	Ollolot.
		entonite Pellets NA		From:			ft	Logged by	:		Checked by:
	Ģı	rout NA		From:			ft	A. Ridle	у		
Ceptin (feet)		LITHOLOGIC DESCR	IPT	ION		TUHOLOGA		NITORING WELL INSTRUCTION	Sample	Blow	REMARKS
		Concrete payment- 4 inches thick					\vdash				
	Г	SILTY GRAVEL(GM) FILL					1			1	
1	L										
И.	L	CLAYEY GRAVEL (GC)								_	P-1 collected on
2	L								P-1	┖	1/22/91 from this
	H	-dark brown with sand, moist to 1" diameter	, with	gravel		GC	ı				location in brass
3	H	to v diamotor					ı				liner, no gasoline
	-										odor
4	H										
	Н						l				
5	Г						ı		1		
_							ı				
6											
7	L								1		
<i>'</i>	L										
8	L	12									
	L				18		1				
9	H										
	H	Difficult drilling, larger gr	avel '	to 4" dia	ameter		ı		l		2.4
10	H						l				
	H						1				1
11	r										_
	۲	Increasing clay content,	mois	τ			ı				no gasoline odor
12		1			91						Odor
	r										
13	r										
	Γ										
14											
						GC					
15		Boring terminated at 16 ' in CLA	YEY	GRAVE	L						DEtecto
16	L								1,2		B-F-1 and 2 in brass liners,
li '°							1		1		elight gasoling

APPENDIX C LABORATORY ANALYSES AND FIELD AQUIFER TESTS

0611911455

April 30, 1991

Mr. Peter McDonald, Suite 210 400 Main Street Pleasanton, Ca. 94566

Dear Mr. McDonald:

I have read the fuel characterization data report by Chevron Research and Technology Company dated October 16, 1990. The results of their analyses are most interesting and I would like to comment on some of their findings.

It was stated that the results of their chemical tests were not conclusive and some ambiguities in the data were seen. The conclusions were that the alkyl lead isomers were not present in the correct pattern, yet their was a possibility that a detergent additive utilized by Chevron may have been present.

The level of tetraalkyllead was found to be near 0.18 g/gal. and appeared to be similar to regular gasolines formulated after 1985. It was stated that Chevron formulated a regular fuel with 0.22 g/gal lead in 1985 and that 95% of the lead was in the form of tetraethyllead. The sample from the Arrow Rental Co. was found to be only 83% tetraethyllead, implying that the unknown fuel was not Chevron.

The numbers and quantitative data do not exactly match between the commercial product of the Richmond refinery and the unknown fuel. However, the numbers generated from the unknown fuel are very close, considering a weathered fuel, and one exposed to bacterial decomposition and water hydrolysis.

It also would be informative to ascertain what the pattern of the other alkyllead compounds were in the unknown Arrow Rental fuel. Has Chevron ever utilized others mixtures of alkyllead compounds (triethylmethyllead, diethyldimethyllead, etc.)? These could be the additives seen with the unknown sample.

Finally, from the data, analytical error, precision, and variability with the results obtained from standards are not presented for samples exposed to the environment. Therefore, the values of 95% and 83% appear very close, considering the analytical method and type of sample.

From the results of the Chevron laboratory, it appears that the fuel is a modern, regular gasoline, containing lead at concentrations similar to post 1985 refinery fuels. The analysis

utilizing gas chromatography with flame photo-ionization detectors may be adequate for these analyses, however, no precision or degree of accuracy was presented which may validate the numbers and comparison. From these findings, I am left with the conclusion that the unknown fuel from the Arrow Rental Co. is most probably Chevron which was accidentally poured down a vapor monitoring well at the site in 1985.

I hope that I have been some help with the review of these data. If you would require additional information or analytical testing, please call me at (415) 422-0903.

Sincerely, Andresen

Brian D. Andresen, Ph.D.

3. Identification of Tetraethyllead (TEL)

After the GC-MS mass chromatogram plots of m/z 208, 237, and 295 will highlight TEL and other lead isomers. The TEL profiles will be seen near naphthalene (m/z 128) in the mixture of fuel compounds. The mass chromatogram plots will clearly confirm regular fuel in the soil sample.

4. Semi-Quantitation of TEL

Once the mass chromatogram plots of TEL have been obtained the areas of each of the three ions is generated (by computer). The areas are summed and compared directly to the area of the total ionization plot.

Area (208 + 337 + 295) = Fraction % of TEL in the Fuel Area of the Total Ion Plot

Fractional % x 3784g/gallon fuel = g TEL/gallon

Discussion

All of the isolated fuel is carefully concentrated to avoid loses to the volatiles in the fuel. The choice of methylene chloride or carbon disulfide is important because these solvents are so very volatile, capable of extracting fuels from soil easily, and generate few competing fragment ions in the mass spectrometer.

The choice of mass chromatogram plots $(m/z\ 208,\ 237,\ and\ 295)$ are diagnostic of TEL in gasolines. No other fragment ions from other hydrocarbons generate these characteristic ions which must appear at exactly the same retention time.

The calculation of the amount of TEL in the final extract is semi-quantitative. If the soil is particularly rich in naturally occurring organic compounds, a blank soil sample should be analyzed under the identical conditions. The total ion plot area is then subtracted from the total ion plot are of the gasoline contaminated soil sample.

Soil samples, soaking with fuel, can have the TEL determined to within very close agreement to values obtained directly from raw fuel. The more the sample has to be concentrated, the more difficult it will be to accurately determine the amount of TEL/gallon. However, this approach will clearly demonstrate the difference between grams/gallons and fractional gram amounts of TEL/gallon in regular gasoline.

I hope that these procedures will aid in your investigations. They are reasonable easy to perform and have a good change of success when the samples are fresh.

BRIAN D. ANDRESEN

WORK ADDRESS:

Lawrence Livermore National Laboratory
University of California
Condensed Matter and Analytical
Sciences Division - Chemistry
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Livermore, California 94550
(415) 422-0903

HOME ADDRESS:

7694 Hillsdale Court
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(415) 846-2783

PERSONAL

Born: Reed City, Michigan (January 20, 1947).

PROFESSIONAL OBJECTIVES

To use the training obtained in synthetic organic chemistry, mass spectrometry, and computer assisted analysis of organic compounds in environmental, biomedical or related research areas. To design unique analytical equipment for trace organic analysis.

EXPERIENCE

- (1) <u>Assistant Professor</u> at the University of Florida, College of Pharmacy (1974-1979), responsible for teaching undergraduate and graduate students techniques in drug analysis, initiating research concerning the synthesis of drugs incorporating stable isotopes, and in identifying trace levels of drugs and biologically active compounds in man and in the marine environment.
- (2) <u>Assistant Director</u> of the University of Florida, College of Pharmacy's Analytical Toxicology Laboratory District IV, Licensed Director (Clinical Chemistry) Florida State Certified (No. 00067), (1974-1981).
- (3) Associate Professor (Tenured 1982) at the Ohio State University, College of Medicine, Department of Pharmacology. Courtesy Appointment in the Department of Obstetrics and Gynecology, College of Medicine. Director of Mass Spectrometry Facilities, College of Medicine (1979-1983).
- (4) <u>Director</u> of the National Reye's Syndrome Research Laboratory, at The Ohio State University College of Medicine (1980-1983).
- (5) National Institutes of Health (NIH) Grant Review Cadre Member for Cancer of the Pancreas and Large Bowel (1980-1984).

- (6) Perinatology Program in <u>Maternal-Fetal Medicine and</u>
 <u>Neonatology</u>, Associate, The Ohio State University, Department of Obstetrics, Gynecology and Neonatology, 1981-1983.
- (7) Courtesy Appointment: <u>Faculty Member in the Department of Obstetrics and Gynecology, School of Medicine</u>, The Ohio State University, 1981-1983.
- (8) Central Ohio Heart Association <u>Peer Review Cadre Member</u>, 1982-1983.
- (9) Senior Research Scientist, Environmental and Biomedical Sciences Divisions, Instrumentation and Measurement Sciences Section (1983-1988) at the Lawrence Livermore National Laboratory.
- (10) Senior Research Scientist, Condensed Matter and Analytical Sciences Division, Chemistry Special Projects (1988-present).
- (10) Drug Enforcement Administration Certified (No. PA0183296) as a Director of an Analytical Laboratory (1976-present).
- (11) Department of Energy, Subcommittee Member: Detection Sub-Group, "Research and Development to Counter the Narcotics Threat", part of the Intelligence Research and Development Council (IR&DC) under the direction of the CIA/ORD (1987-Present).

EDUCATION

- 1972-1974 Ph.D. obtained under Professor Klaus Biemann at Massachusetts Institute of Technology (MIT),
 Department of Chemistry. Thesis Title:
 "Deuterium Labeling in Drug Metabolism Studies by Mass Spectrometry".
- 1971-1972 Second Master's degree obtained at Woods Hole Oceanographic Institute (MIT-WHOI Joint Program) in the Chemical Oceanography Department. Thesis Title: "Identification of the Organic Components in Green River Shale".
- 1969-1971 First Master's degree obtained at Massachusetts Institute of Technology (MIT), Department of Chemistry in Analytical Chemistry.
- 1965-1969 Undergraduate Education <u>Florida State University</u>
 Major in Chemistry, minor in math, biology and oceanography (GPA 3.2/4.0 overall, 3.7/4.0 science).

INSTRUMENTATION

1. Mass Spectrometry - Proficiency in high resolution mass spectrometry using either a CEC-110B or Varian MAT 311A double focusing instruments. Experienced with both electronic and photographic recording capabilities and with computer control and calculations of high resolution data. Much experience with low

and high resolution mass spectrometry for generating and interpreting mass spectral data. Extensive use of combined gas chromatography-mass spectrometry-computer systems. Have been involved with the complete installation, maintenance, and upgrade of existing instruments including Varian-Finnigan, Hewlett-Packard, Dupont, Hatachi, Nuclide, and Nicolet. Hold several patents in GC-MS interfacing and new analytical equipment.

- 2. <u>Gas Chromatography</u> Proficiency in the use of all types of gas chromatographic equipment and techniques. Very familiar with packed and capillary column gas chromatographic-mass spectrometry techniques and interfacing.
- 3. High Pressure Liquid Chromatography Familiar with equipment and techniques necessary for the identification and quantitation of compounds in biological fluids. Have been responsible for an Altex and Spectro-Physics HPLC systems used to isolate and quantitate drugs in body fluids.
- 4. Computers Familiar with IBM 1800, PDP-11/34, HP-2200 computers and data handling equipment. Very familiar with microcomputers and techniques for data acquisition and reduction, having built a Z-80, 32K RAM, dual-disk drive system. Also have built a 635K, 10 megabit IBM-XT computer (with associated hardware and software) and taught courses related to high speed analog-to-digital data recording and manipulation.
- 5. Other Specialized Techniques Much research and teaching experience has been acquired in the areas of infrared, ultraviolet, and nuclear magnetic resonance instrumentation and spectral interpretation. Have had much experience with design and problem solving in electronics. Experienced with isolated organ perfusion apparatus and experimentation. Recently have gained much experience with fiber optics, lasers, and remote sensing utilizing optical fibers.

ORGANIC SYNTHESIS

Proficient in a wide variety of synthetic skills, having synthesized many natural products, drugs, drug metabolites, as well as useful intermediates. Work has also included the development of useful synthetic pathways for the preparation of isotopically labelled compounds for internal standards in quantitative analysis, and labelled drugs for feeding experiments in pharmacological studies of drug metabolism in animals and man. Recently have gained much experience in the synthesis of potent carcinogens and mutagens in overcooked food. Research work has also centered on the isolation, characterization and synthesis of pollutants in the San Francisco bay. This work includes chemicals isolated from seawater, ocean sediments and fish.

MEDICAL BACKGROUND

Analyses have been performed on the body fluids of over 1200 parients suffering from drug overdose. This work was performed

in Boston at MIT, at the University of Florida and at The Ohio State University to aid doctors with immediate body fluids analyses utilizing computer controlled gas chromatography and mass spectrometry to provide rapid and reliable information for the identification of the toxic substances in comatose patients. Quite often this work has included the analysis of and testimony concerning drugs, alcohol, and poisons. Major research interest has centered on the identification of biologically significant compounds in disease states and during pregnancy. Much work has been aimed at the complete analysis of amniotic fluids, spinal fluids, and the identification of abnormal compounds in the body fluids of Reye's Syndrome victims. The compound identifications have included both environmental compounds, drugs and metabolites, and new biochemicals of potential toxicity. Much experience has been obtained concerning animal surgeries with dogs, rats and baboons. In baboons, a considerable amount of work has been performed in the area of fetal pharmacology. In these surgical experiments, live baboon fetuses were removed, studied and returned to the uterus to be carried to term. Much experience has also been gained concerning complications of pregnancy in women.

GENERAL

Have worked on a variety of projects including the analysis of the Apollo Lunar samples for organic chemical content; anti-tumor compounds; fish migration and the chemical characterization of home stream water; crustacean communication, pheromones, and sex attractants between male and female lobsters; oil shale analysis by combined gas chromatography and mass spectrometry; pseudo-Jupiter soil analyses; the identification of various pollutants in river and ocean water (Chemical Oceanography); the identification of antibiotics from marine plants and animals; Fetal Alcohol Syndrome (FAS); the identification of chemicals indicative of distinct disease states (Reye's Syndrome, Maple Syrup Urine Disease); drug metabolism studies, including the identification of metabolites of beta-blockers, (acebutolol); the characterization of amniotic fluids by gas chromatography and mass spectrometry to identify chemicals indicative of the maturity and health of the developing fetus; and more recently the synthesis of mutagenic substances in over cooked food. addition, techniques have been developed to identify the accelerants in arson cases, or the identification of underground gasoline leaks utilizing mass spectrometry and pattern recognition.

Most mass spectrometry work has required the analysis of many different types of environmental and biological samples. These have included solid, liquid and gaseous samples. Central to this work has been the development of new procedures for the analysis of blood, urine, cerebral spinal fluids, serum, breath, bile, stomach contents, vitreous humor (from the eye), amniotic fluids, meconium, follicular fluids (from human ovaries), tissues (liver, brain, muscle, skin), and many varied samples of biological origin. All of these studies have presented many challenging

situations which require unique sample handling and derivatization techniques, yet allowed added experiences to be gained in the types of unusual compounds that may be encountered during any gas chromatography-mass spectrometry-computer analysis. New areas of interest have included computer profiling techniques for the comparisons of diseased and healthy body fluid and tissue samples by gas chromatography and mass spectrometry. With these new computerized profiling techniques we are now able to observe major and subtle changes in the body chemistry. This new area of research may be important for disease state diagnosis.

MASS SPECTROMETRY FACILITIES

In addition, much time has been spent in the development of mass spectrometry research groups and mass spectrometry facilities within the three different University systems (The University of Florida, The Ohio State University, and the University of California at Berkeley [Lawrence Livermore National Laboratory]). In this capacity, many students and laboratory personnel have been trained in the art of mass spectrometry, including instrument maintenance, electronics design, chemical analysis, and problem solving. This final point has been especially important. Mass spectrometry has been taught as a sophisticated tool, capable of solving problems which are often encountered in the analysis of submicrogram amounts of components within a complex biological matrix. Students who leave our group have a much greater appreciation for the capabilities and limitations of computer guided gas chromatography-mass spectrometry systems in analytical chemistry and a better understanding of its power when applied to environmental or biomedical problems.

NEW METHODS OF ANALYSIS

Recently new methods have been developed for sample introduction into a mass spectrometer. These include Electrophoresis-Mass Spectrometry (EPMS) and a new type of Moving Belt Interface for high performance liquid chromatography-mass spectrometry. Both designs are completely original and promise to give an added dimension to the mass analysis of complex mixtures containing polar or thermally labile molecules which have previously been most difficult to analyze by standard mass spectral techniques. From this work four new patent applications have been submitted.

The Electrophoresis-Mass Spectrometry interface has resulted in an IR-100 Award for 1986. This award recognizes the top 100 best research and development ideas which have been judged on the basis of their significance, uniqueness, and usefulness from a technical standpoint. The competition considered many thousands of applicants world wide.

New methods are currently being developed to review mass spectral data directly with the aid of an IBM PC. It is the goal of this research to utilize high speed microcomputers to manipulate mass spectral data for pattern recognition.

TO: Mr. Al Ridley

FROM: Brian D. Andresen

RE: Identification and semi-quantitation of tetraethyllead in

soils by extraction and GC-MS analysis

Enclosed is my procedure for the isolation of trace amounts of gasoline from soils. Although there is no EPA number for these analyses, they work well for a variety of sample types. The procedures are divided into three parts: (1) Following sample concentration, the GC-MS analysis is directed toward generating a recognizable gasoline profile. (2) Fragment ions associated with tetraethyllead (TEL) are highlighted and their areas determined. (3) A semi-quantitative treatment of the data is obtain an estimate of the original TEL in the regular gasoline calculated.

1. Fuel Isolation from Soils

Into a clear beaker (with aluminum foil top) is added 100 to 200 g of fresh soil which has revealed the characteristic smell of gasoline. The soil is broken into small pieces under a volume of methylene chloride (b.p. 40°C) or carbon disulfide (b.p. 46°C) which is sufficient to cover the soil sample. The mixture of solvent and soil is then placed in an ultrasonic bath for 15 minutes (avoiding excessive heat) to extract the fuel from the soil.

The beaker is removed from the ultrasonic bath and the fine sediments are allowed to settle (approximately 10 minutes). The clear solvent layer is decanted into a pear shaped flask and rotary evaporated carefully so that the solvent is removed slowly while cold. The extraction solvent is reduced to a volume of approximately 1.0 to 0.5 mL (never to dryness) and immediately capped and then analyzed by computer controlled gas chromatography-mass spectrometry.

2. Gas Chromatographic-Mass Spectrometric Analysis

Approximately 2 uL of the extracted sample is injected into a capillary GC-MS instrument. A concentrated enough sample should be injected to insure a strong fuel signature with a reasonably intense total ionization plot. A capillary column is not essential, however, it will allow the separation of the very volatile extraction solvent from the fuel. The GC conditions should be initiated near 40-50°C to insure separation of the fuel from the extraction solvent. The temperature program conditions should ramp high enough (250°C) to allow all gasoline components to elute during the GC-MS run. The mass spectrometer should record all mass spectral data (45-400 amu).

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- 1. K.J. Ng, B.D. Andresen, and J.R. Bianchine, "Capillary Gas Chromatographic-Mass Spectrometric Profiles of Organic Acids from Amniotic and Other Body Fluids", in <u>Reproductive Medicine Series</u>

 Marcel Dekker, Inc. Publishers).
- 2. K.J. Ng, B.D. Andresen, J.R. Bianchine, J.D. Tams, R.W. O'Shaugnessy, M. Nelson, F.P. Zuspan, and L.F. Stempel, "Variation in Hippuric Acid Concentration in Amniotic Fluid with Gestational Age", J. Obstetrics and Gynecology.
- 3. M. Alexander, M. Borrisud, B. Andresen, A. Staubus, S. Jacobs, S. Roemer, and J. Bianchine, "Acebutolol and its Metabolites Following Oral and Intravenous Administration in Man", Clin. Pharmacology and Therapeutics.
- 4. B.E. Watkins, M.G. Knize, C.J. Morris, B.D. Andresen, et. al., "Synthesis of derivatizes of the Cooked-Food Mutagens IQ, MeIQx, and PhIP as Haptenic Compounds", (Submitted Hetercycles 1987).

TEACHING ACTIVITIES

(Shared With Other Department Members at The Ohio State University)

- 1. Pharmacology 824 Psychopharmacology
- 2. Pharmacology 702 Practice of Pharmacology (Laboratory methods)
- 3. Pharmacology 700 Phase II Medical Students Required Pharmacology Course
- 4. Pharmacology 609 Molecular Pharmacology
- 5. Pharmacology 600 Introduction to Pharmacology
- 6. Pharmacology 400 Pharmacology for Clinical Chemistry Technologist

TEACHING ACTIVITIES (Total Responsibility)

- 1. Pharmacology 850 General Pharmacology Review (All Topics in Pharmacology)
- 2. Pharmacology 860 Chromatography and Instrumental Methods of Drug Assay (With Biomedical Mass Spectrometry)
- 3. Pharmacology 999 Graduate Student Supervision
- Analytical Pharmaceutical Procedures 445 (1979-1983 University of Florida)
- 5. Pharmaceutical Laboratory 447 (1979-1983 University of Florida)

GRADUATE STUDENTS (Under My Supervision)

- 1. Mrs. Judy Lubbers M.D. and Ph.D. Candidate
- 2. Mrs. Diane Moreno Ph.D. Candidate
- Mrs. Li-Shei Lin Master' Student (Awarded 1981)
- 4. Ms. Cynthia Beach Ph.D. Candidate (Awarded 1983)
- 5. Dr. Kwokei Ng Ph.D. Candidate (Awarded 1981)
- 6. Dr. Dennis Eneanya Ph.D. Candidate (Awarded 1981)
- 7. Dr. Ginn Wu Postdoctoral Fellow
- 8. Dr. Dale Sharp Postdoctoral Fellow
- 9. Dr. Matana Borrisud Ph.D. Candidate (Awarded 1983)
- 10. Mr. Harry Duran, Mr. Jason Chang, Ms. Denise Contos, Ph.D. Canadates (partially suported in my laboratory)
- 11. Dr. Michael Alexander -Ph.D. candidate (awarded 1983)

OTHER STUDENT SUPERVISION (The University of Florida)

- Dr. Frank Davis Ph.D. Student (awarded 1978)
- Mr. James Templeton Masters Student (Awarded 1977)
- Dr. Harry Panzik Postdoctoral Fellow

(Ph.D. THESIS and DISSERTATIONS DIRECTED)

- 1. "Applications of Synthetic Organic Chemicstry, Stable Isotope Labelling, and Mass Spectrometry to Drug Metabolism Studies", Dr. Frank Davis (1978).
- 2. "Gas Chromatographic-Mass Spectrometric Profiling of Organic Acids of Human Amniotic Fluid", Dr. Kwokei Ng (1981).
- 3. "Metabolic Study of Ritodrine With High Performance Liquid Chromatography", Dr. Matana Borrisud (1983).
- 4. "Metabolic Studies of Acebutolol by HPLC and GC-MS", Dr. Michael Alexander (1983).

MEDICAL STUDENTS (Receiving Awards and Financial Support While Under My Supervision)

- 1. Mrs. Mary Ann Gravinstein
- 2. Mr. Stephan Gravinstein
- 3. Mr. James Kotsonas
- 4. Ms. Barbara Bunn
- 5. Mr. Rick Kukukla
- 6. Mr. David Bray
- 7. Ms. Linda Weber

GRANTS AWARDED AS THE PRINCIPAL INVESTIGATOR AT THE OHIO STATE UNIVERSITY

- 1. The Ohio State University School of Medicine Starter Grant, \$5,000: "The Characterization of Unusual Compounds in the Urine of Reye's Syndrome Patients" (1980), Grant #540107.
- 2. The Ohio State University Graduate Research Program, \$5,400: "Acebutolol A Threat of Toxic Metabolites", (1980), Grant #221312.
- 3. The National Reye's Syndrome Foundation, \$21,000: "A Reye's Syndrome Analytical Facility" (1908-1983), Grant #713086.
- 4. The Central Ohio Heart Foundation, \$12,000: "The Impact of Metabolites of Acebutolol on Cardiofunction" (1980), Grant #713369.
- 5. The Stohler Isotope Chemical Co., \$5,000: "The Analysis and Synthesis of Labelled Compounds" (1980), Grant #536577.
- 6. The Central Ohio Heart Association, \$15,000: "The Potential Danger of In-Vivo Acebutolol Decomposition" (1981-1982).

GRANTS AWARDED AS CO-INVESTIGATOR AT THE OHIO STATE UNIVERSITY (Principal Investigator)

- 1. The National Institutes of Health, \$433,497: "Unsaturated Fatty Acids, Prostaglandins, and Diabetes;" (Dr. Howard Sprecher, 1981-1985).
- 2. Public Health Services, \$95,767: "Prostaglandin Control of Macrophage Anti-tumor Activity", (Dr. B.S. Zwilling, 1981-1985).
- 3. The National Institutes of Health, No. 669,218: "Stochastic Fourier Transform Ion Cyclotron Resonance Mass Spectrometry", (Dr. Allan Marshall, 1983-1986).

GRANTS AWARDED AT THE LAWRENCE LIVERMORE NATIONAL LABORATORY (As Principal Investigator)

1. The Institutional Research and Development Grants (LLNL), "Electrophoretic Mass Spectrometry", \$59,000 (1985).

- 2. The Institutional Research and Development Grants (LLNL), "Electrophoretic Mass Spectrometry", \$120,000 (1986).
- 3. The Institutional Research and Development Grants (LLNL), "High Performance Liquid Chromatography High Resolution Mass Spectrometry", \$110,000 (1985)
- 4. The Institutional Research and Development Grants (LLNL), "High Performance Liquid Chromatography High Resolution Mass Spectrometry", \$119,000 (1986).
- 5. Department of Energy, "Galileo Project", \$27,000 (1986-1989).
- 6. Department of Energy, "Very Small Mass Spectrometer (VSMS)", \$350,00 (1988-1990).

GRANTS AWARDED AT THE LAWRENCE LIVERMORE NATIONAL LABORATORY (As Co-investigator)

- 1. National Institutes of Health, "Mutagens From Cooking of Food", (Dr. Fred Hatch, \$3,800,000, 1981-1987).
- 2. National Institutes of Health, "Immunoassays for the Detection of Cooking-Food-Mutagens", (Dr. James Felton, \$724,821, 1985-1987).
- 3. National Institutes of Health, "Genotoxicity of Food Related to 1,2-Dicarbonyl Compounds in Cooked Food", (Dr. Robert Taylor, \$532,177, 1985-1987).
- 4. Oakridge National Laboratory, "Improved <u>In-Situ</u> Detection of Organic Compounds", (Dr. Fred Milanovich, \$210,00, (1986-1987).
- 5. Environmental Protection Agency, "Groundwater Monitoring By Spactroscopy With Optical Fibers and Optrodes", (Dr. Fred Milanovich, \$150,000, 1983-1986).
- 6. Department of Energy, "Subsurface Organic Chemical Analysis Using Remote Fiber Optic Technology", (Dr. Florence Harrison, \$50,000, 1985-1986).
- 7. Department of Energy, "Chemical Characterization and Analysis Techniques (CCAT-Organic)", R. Hawley-Fedder and B.D. Andresen (Co-Principal Investigators) \$385,000 (1986-1988).
- E. Department of Energy, "SKYBRIGHT", R. Hawley-Fedder, E. Raber, and B. Andresen, \$1,200,000 (1986-1989).
- 9. Various Government Agencies, "Unique Projects of Interest" (1989-1990), supporting funds.

COMMUNITY SERVICES

Miscellaneous seminars for various departments at The University of Florida.

Miscellaneous seminars for various departments at The Ohio State University.

Central Ohio Heart Chapter, Inc. (many community presentations concerning science in medicine, Columbus, Ohio Museum of Science and Technology, Continuous Video Tape).

Miscellaneous radio programs on drugs of abuse (AM 610, FM 96, and FM 98 Columbus, OH)

Reye's Syndrome National Laboratory presentations (many radio and television programs)

Reye's Syndrome - New York Times Newspaper (1981).

Ohio State Coronor's Association Annual Seminar Presentations (1980-1983).

New Horizons in Science Presentations - UPI and AP News Updates

North Florida District IV Medical Examiner Toxicology Laboratory Coronor's Association Annual Seminar Presentations - 1974-1979.

Livermore Police - Drug Taskforce.

AWARDS

- 1. <u>Director</u> Nation Reye's Syndrome Research Laboratory Awarded June 1981 at the National Press Club, Washington, D.C.
- 2. <u>Industrial Research (IR 100)</u> Awarded June 1987 at the Museum of Science and Industry, Chicago, IL. for the newly invented Electrophoresi-Mass Spectrometry Interface.

LECTURES

Lions Clubs Rotary Club National Telephone Companies American Associations of Clinical Chemists

MISCELLANEOUS

The Presendent's Club - Financial contributions to the Ohio State University Alumnae Fund.

Included in the American Men and Women of Science - 1982, Fifteenth Edition, page 134.

Contribution - National Baseball Hall of Fame - 1939 Centennial Coin.

CONSULTANT TO OR MASS SPECTROMETRY RELATED WORK WITH THE FOLLOWING GROUPS

- 1. Stohler Isotope Chemicals, Inc. Waltham, Massachusetts
- 2. Syntex Laboratories, Inc. Palo Alto, California
- 3. Adria Laboratories, Inc. Dublin, Ohio
- 4. Varian MAT Breman, Germany
- 5. Florida Crime Lab Tallahassee, Florida
- 6. Columbus Testing Laboratory, Inc. Columbus, Ohio
- 7. Isotope Labelling Company Whippany, New Jersey
- 8. Anchor Hocking Lancaster, Ohio
- 9. Childrens Hospital Columbus, Ohio
- 10. Sepragen, Corporation San Leandro, CA.
- 11. Baseline Environmental Consulting Oakland, Ca.
- 12. Homestake Minning Co. Davy McKee Corp.
- 13. ST&E, Inc. Livermore, Ca. (415) 449-8516.
- 14. Distributors Processing Inc. Porterville, CA, (209) 781-0297.
- 15. Midwest Physicans Network Minneapolis, MN, (612) 863-5613.
- 16. Fiber Chem., Inc. Albuquerque, NM, (505) 292-6901.
- 17. Alliance Technologies Corp. Bedford, Mass. (617) 275-9000.
- 18. Laser Cartrige Reconditioning Dublin, Ca. (415)846-9062.

CONSULTANT WITH THE FOLLOWING LAW FIRMS

- 1. Mr. Dan Conner and Associates Columbus, Ohio, (614) 464-2025
- 2. Mr. R. William Meeks and Associates Columbus, Ohio, (614) 228-3569.
- 3. Mr. Steve Larson and Associates Columbus, Ohio, (614) 224-0343.
- 4. Mr. Boyd Binning Columbus, Ohio, (614) 224-1979.
- 3. Mr. Richard Bush -Parkersburg, West Virginia, (304) 485-6438.
- Mr. Paul Cassidy and Associates Columbus, Ohio, (614) 228-3569.
- 7. Mr. Gary Dumm Columbus, Ohio, (614) 477-2593.
- S. Mr. James Hill Circleville, Ohio, (614) 477-2173.
- 9. Mr. Michael Hoddox and Mr. Ronald Lasmer Zanesville, Ohio, (614) 452-9358.
- 1... Mr. Rober Moses Columbus, Ohio, (614) 294-2678.
- 1 Mr. William Simonton Pennsboro, West Virginia, (304) 659-2966.
- 13. Mr. Rodney Wubdin Harrisville, West Virginia.
- 10. Mr. Larry Supp and Associates Morristown, New Jersey, (201) 267-5200.
- 14. Mr. Thomas Bell, Mr. Robert Addison and Associates Jackson, Mississippi, (601) 969-7607.
- 13. Mr. James R. Kingsley Circleville, Ohio, (614) 477-2546.
- 16. Mr. Joe A. Rosenfield Columbus, Ohio, (614) 469-9922.
- 1 Mr. Rodney C. Wincom Harrisville, West Virginia, (304) 643-4440.
- 10. Mr. Leonard Siegal Columbus, Ohio, (614) 866-4025.
- 19. Mr. Thomas Orvis Oakland, Ca. (415) 328-5510
- 2. Mr. Terry Broughton Dublin, Ca. (415) 828-4428
- 2... Mr. Harry Traback Livermore, Ca. (415) 447-7020
- 21 Mr. Terry Moore Cincinnati, Ohio (513) 961-6200
- 21. Mr. Michael Murry Oakland, Ca. (415) 451-4600

CONSULTANT FOR THE FOLLOWING TELEPHONE COMPANIES CONCERNING ENVIRONMENTAL CONTAMINATION

- 1. Pacific Bell (San Ramon, Ca.):
 Miss Dorthy (D.J.) Allen (415) 823-9820
 and Mr. E.J.(Ed) Koehler (415) 823-9821
- 2. Bell Atlantic (Arlington, Va.):
 Mr. Christopher E. Mandel, AIC (703) 974-3757
- 3. C & P Telephone Company of Virginia (Richmond, Va.):
 Mr. Ruben Burton, Jr., Mr. Gene Lane, and
 Mr. George Hamner, Legal Department (804) 772-3512
- 4. South Central Bell (Baton Rouge, Louisiana): Mr. Joe Pat Carnahan (504) 292-6465.
- 5. South Central Bell (Huntsville, Alabama):
 Mr. Mark A. Ray, Claims Office (205) 532-8678.
- 6. Mountain Bell, Station 29 (Albuquerque, New Mexico): Mr. Gill Quesada (505) 765-8113.
- 7. General Telephone of The Southwest (San Angelo, Texas): Mrs. Rosemary Davis - (915) 944-5335
- 8. Ohio Bell (Cleveland, Ohio):
 Mr. Al Richards (216) 822-42283
- 9. Bellcore (Piscataway, New Jersey)
 Mr. Robert Allen (201) 981-7825
- 10. Bell South Services (Birmingham, Alabama): Mr. Shep Holladay - (205) 321-8835
- 11. GTE (Fort Wayne, Indiana), Security Department: Mr. Mike Connell (219) 461-2653.
- 12. New Jersey Bell (Newark, N.J.):
 Mr. E.J. Carroll (210) 649-2206.
- 13. GTE (Carrolton, Texas) Security Dept.: Mr. Ben Day (214) 466-0738.
- 14. GET (Plano, Texas) Security Department:
 Mr. Bill Handcock (,
 Mr. Chris Schnaithman (214) 403-0404, and
 Mrs. Susan Freeman (214) 495-3112 x 3622.
- 15. South Central Bell Security Department: Mr. Jeffrey Morrison (318) 222-8596.

- CAPABILITIES -

I. GASOLINE ANALYSIS - IDENTIFICATION OF PETROLEUM PRODUCTS

- A. Fingerprinting and Pattern Recognition
- B. Lead Content
- C. Underground Tanks Leak Identification
- D. Asphalt Characterization
- E. Rocket, Jet, Missile Propulsion Fuel Analysis

II. IDENTIFICATION BY THE CHEMICAL MATCHING OF MATERIALS

- A. Paper, Metals, Plastics, Ink, Water, Oils, etc.
- B. Gun Powder Contact Memory
- C. Blood Residues Containing Unknown Substances
- D. Fingerprints and Smudges Containing Biological Materials
- E. Explosive Characterization
- F. Air and Gas Samples for Trace Level Analysis

III. TRACE ANALYSIS OF ORGANIC CHEMICAL RESIDUES

- A. On the Soles of Shoes, Paper, Leather, Cloths, etc.
- B. On Skin: The Hands for Trace Organic Residues
- C. In Breath: Nicotine, Marijuana, etc.
- D. In Hair Samples for Trace Organic Compounds of Interest

III. ARSON

- A. Identification of Acceleration
- B. Chemical Analysis of Materials
- C. Explosive Analysis Residues

T. BLOOD PROFILES

- A. Total Analysis of Blood Components Characterization of Individuals as to Health and Types of Foods Ingested
- B. Drugs in All Body Fluids (Blood, Urine, Saliva, etc.
- C. Adverse Drug Reactions and Pharmacology

I. URINE ANALYSIS

- A. Types of Foods Ingested and Overall Metabolic Health
- B. Drugs Analysis Rapid and Noninvasive

WII. TRACE ANALYSIS OF DRUG RESIDUES

- A. GC-MS-Computer Analysis: Confirmation and Quantitation
- B. Thin Layer Chromatography Rapid Screening
- C. Field Test for Marijuana and (THC) Residues
- D. Field Tests for Cocaine or Other Illicit Drugs
- E. Hair Analysis by GC-MS

VIII. INSTRUMENTATION AND OTHER SKILLS

- A. Mass Spectrometry Installation
- B. Gas Chromatography-Mass Spectrometry-Computer Design
- C. Synthetic Organic Chemistry
 - a. Design and Carry out Experiments
 - b. Prepare Target Compounds
 - Apparatus Glass Blowing
- D. Electronics Design and Construction
 - a. Generate Bread-board Layouts
 - b. Professional Quality Prototype
 - c. Construction and Fabrication
 - e. Graphics and Silk Screening
 - d. Circuit Board Design and Fabrication
 - e. Machining and Fabrication of Parts
- E. Recipes, Formulas, and Trade Secrets
 - a. Identification of Chemical Formulas
 - b. Chemical Code Breaking
 - b. Preparation of Trade Secret Recipes

CONTROLLED SUBSTANCES REGISTRATION CERTIFICATE UNITED STATES DEPARTMENT OF JUSTICE DRUG ENFORCEMENT ADMINISTRATION WASHINGTON, D.C. 20537

Tie Controlled Substances Act of 1970 reads in part as follows:

Sec. 304. (a) A registration pursuant to section 303 to manufacture, distribute, or dispense a controlled substance may be suspended or revoked by the Attorney General upon a finding that the registrant -

(1) has materially labified any application filed pursuant to or required by

(2) has been convicted of a telony under this little or title III or any other law of the United States, onot any State; relating to any substance defined in this title as a controlled substance; or

(3) has had his State license or registration suspended, revoked, or denied by competent State authority and is no longer authorized by State law to engage in the manufacturing, distribution; or dispensing of controlled DEA REGISTRATION NUMBER -

THIS REGISTRATION **EXPIRES**

PA0183296

06-30-90 SCHEDULES BUSINESS ACTIVITY

EXEMPT

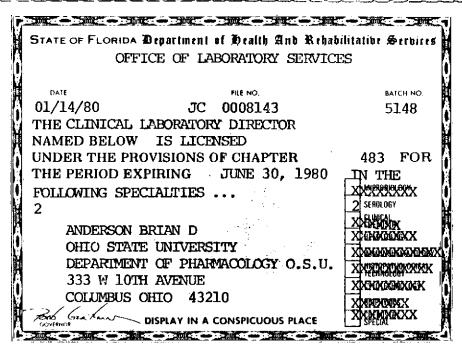
1.2.3.38.4.5 ANALYTICAL LAB

09-13-89

- DATE ISSUED

ANDRESEN. BRIAN D LAHRENCE LIVEMORE MATIONAL LAB PO BOX-808. L-310 LIVERMORE, CA

THIS CERTIFICATE IS NOT TRANSFERABLE ON CHANGE OF OWNERSHIP, CONTROL, LOCATION: OR BUSINESS



State of Florida
Department of Health And Rehabilitative Services
OFFICE OF LABORATORY SERVICES

ANDERSON BRIAN D

CLINICAL LABORATORY DIRECTOR

HAS PAID THE FEE REQUIRED BY CHAPTER FOR THE PERIOD EXPIRING JUNE 30, 1980

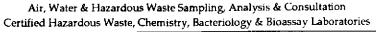
Dian Signature

PLEASE READ IMPORTANT
INFORMATION ON REVERSE

OFFICE OF LABORATION SERVICES POST OFFICE BOX 210

JACKSONVILLE FL 32231

AUDIT CONTROL NO.	FILE NO	BATCH NO.	FEE AMOUNT
035976	JC0008143	5148	\$30.00





141 Suburban Road 751 S. Kellogg, Suite A 1885 North Kelly Road 9333 Tech Center Dr., Ste. 800

2400 Cumberland Dr.

San Luis Obispo, CA 93401 • Goleta, CA 93117 •

Valparaiso, Indiana 46383

Goleta, CA 93117 Napa, CA 94558 Sacramento, CA 95826 • (916) 368-1333

(805) 543-2553

Fax (805) 967-4386
Fax (707) 226-1001

Fax (805) 543-2685

(916) 368-1333 • Fax (916) 362-2484 (219) 464-2389 • Fax (219) 462-2953

April 15, 1991

Mr. Albert Ridley Woodard-Clyde Consultants 500 12th St. Suite 100 Oakland, CA 93607

Re: Analysis of Soil for Gasoline and Tetraethyl Lead

Dear Mr. Ridley:

Coast-to-Coast Analytical Services analyzed sample H-01024-1 using the quantitative techniques specified by EPA 8270. A calibration curve using authentic tetraethyl lead supplied by Dupont Chemical Co. was used for quantification of tetraethyl lead (m/e 237) in mg/Kg of the sample matrix. The percent recoveries for the matrix spike and matrix spike duplicate were 64% and 50% respectively. The gasoline was not measured quantitatively by the semi-volatile GC/MS method. We did present the composition of tetraethyl lead in gasoline based on the non-quantitative semi-volatile method for gasoline as proscribed by the method supplied by Woodward-Clyde.

Quantitative measurement of gasoline was obtained using EPA method 5030 (Purge & Trap) with CAL DHS DRAFT TPH (modified) by GC/MS. Quantitation using the total ion current was based on an authentic gasoline standard. Matrix spike and matrix spike duplicate were 100% and 88%, respectively. The composition of tetraethyl lead in gasoline was calculated based on this more accurate and precise value for gasoline. The gasoline appears to be highly weathered and diminished in the lighter molecular weight hydrocarbons present in new gasoline. The additive methyl t-butylether was not detected in the sample.

Instrument and method blanks for the analytical batches containing sample H-01024-1 were free of tetraethyl lead and gasoline at and above the Practical Quantitation Limit. Instrument tuning criteria were met with each analytical batch run.

Respectfully submitted, Coast-to-Coast Analytical Services

Mary Havlicek, Ph.D., President



CLIENT: Albert Ridley

Woodward-Clyde Consultants

500 12th St. Ste. 100

Oakland, CA 94607

Air, Water & Hazardous Waste Sampling, Analysis & Consultation Certified Hazardous Waste, Chemistry, Bacteriology & Bioassay Laboratories

141 Suburban Road 751 S. Kellogg, Suite A 1885 North Kelly Road

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(707) 257-7211

Fax (707) 226-1001

(916) 368-1333 Fax (916) 362-2484 Fax (219) 462-2953 (219) 464-2389

Lab Number : H-1024-1

Project : 90C0321A-4000 Arrow

Rentals

: 03/27/91 Analyzed Analyzed by: DP

Method : GC/MS

REPORT OF ANALYTICAL RESULTS

Page 1 of 1

SAMPLE DESCRIPTION	MATRIX	SAMPLED BY	SAMPLED DATE RECEIVED			
B-F-1-16' & B-F-2-16'	Soil	Al Ridley	03/12/91		03/13/91	
CONSTITUENT		(CAS RN)	*PQL mg/Kg	RESUL mg/Kg		
TOTAL SEMIVOLATILE PETROLEUM HYDROCARBONS Total Petroleum Hydrocarbons (Gasoline)			5.	24.	1,2,3	
Tetraethyl Lead	-		0.005	0.010)	

CCAS is Certified by CA Department of Health Services: Laboratory #131 *RESULTS listed as 'ND' were not detected at or above the listed PQL (Practical Quantitation Limit)

- (1) Sample Preparation on 03/25/91 by RAD using Methylene Chloride in Sonic Bath.
- (2) The composition of Tetraethyl Lead in Gasoline is 1.2g/Gallon.
- (3) Quantitative values for gasoline are more accurate when extraction is done by EPA Method 5030 (Purge & Trap).

04/11/91 MSD5/ZC09A/#C26A MH/rdl/drc/elr HC25M550

Respectfully submitted,

COAST-TO-COAST ANALYTICAL SERVICES, INC.

Mary Havlicek, Ph.D.



141 Suburban Road 751 S. Kellogg, Suite A 1885 North Kelly Road

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Napa, CA 94558 Sacramento, CA 95826

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(219) 464-2389

Fax (805) 543-2685 Fax (805) 967-4386

Fax (219) 462-2953

Fax (707) 226-1001 (916) 368-1333 Fax (916) 362-2484

QC Batch ID: HC25M550

CLIENT: Coast-to-Coast Analytical Services

141 Suburban Rd. Ste. C-4 San Luis Obispo, CA 93401

Analyzed

: 03/27/91

Analyzed by: DP Method : GC/MS

INSTRUMENT BLANK

REPORT OF ANALYTICAL RESULTS

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Page 1 of 1

SAMPLE DESCRIPTION	MATRIX	SAMPLED BY	SAMPLED DATE RECEIVED			
INSTRUMENT BLANK	Soil					
CONSTITUENT		(CAS RN)	*PQL mg/Kg	RESULT mg/Kg	NOTE	
TOTAL SEMIVOLATILE PETROLEUM HYDROCARBONS Total Petroleum Hydrocarbons (Gasoline)			5.	ND		
Tetraethyl Lead	-		0.005	ND		

CCAS is Certified by CA Department of Health Services: Laboratory #131 *RESULTS listed as 'ND' were not detected at or above the listed PQL (Practical Quantitation Limit)

04/11/91 MSD5/ZC07A MH/erc H1024-1

Respectfully submitted, COAST-TO-COAST ANALYTICAL SERVICES, INC.

Denald R. Cortes, Group Leader

May Harlials
Mary Havlicek, Ph.D.



141 Suburban Road 751 S. Kellogg, Suite A 1885 North Kelly Road 9333 Tech Center Dr., Ste. 800

2400 Cumberland Dr.

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Fax (805) 543-2685 Fax (805) 967-4386

Fax (707) 226-1001 (916) 368-1333 Fax (916) 362-2484

Fax (219) 462-2953 Valparaiso, Indiana 46383 (219) 464-2389 QC Batch ID: HC25M550

CLIENT: Coast-to-Coast Analytical Services

141 Suburban Rd. Ste. C-4 San Luis Obispo, CA 93401

Analyzed

: 03/27/91

Analyzed by: DP

Method : GC/MS

METHOD BLANK

REPORT OF ANALYTICAL RESULTS

.

Page 1 of 1

SAMPLE DESCRIPTION	MATRIX	SAMPLED BY	SAMPLED DATE RECEIVED		
METHOD BLANK	Soil			-	
CONSTITUENT		(CAS RN)	*PQL mg/Kg	RESULT mg/Kg	NOTE
TOTAL SEMIVOLATILE PETROLEUM HYDROCARBONS Total Petroleum Hydrocarbons (Gasoline)			5.	ND	1
Tetraethyl Lead	•		0.005	ND	

CCAS is Certified by CA Department of Health Services: Laboratory #131 *RESULTS listed as 'ND' were not detected at or above the listed PQL (Practical Quantitation Limit) (1) Sample Preparation on 03/25/91 by RAD using Methylene Chloride in Sonic Bath.

04/11/91 MSD5/ZC08A MH/erc H1024-1

Respectfully submitted,

COAST-TO-COAST ANALYTICAL SERVICES, INC.

Donald R. Cortes, Group Leader

Mary Havlicek, Ph.D.



141 Suburban Road 751 S. Kellogg, Suite A 1885 North Kelly Road

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Sacramento, CA 95826

Valparaiso, Indiana 46383

(805) 543-2553 (805) 964-7838

Fax (805) 543-2685 Fax (805) 967-4386

(707) 257-7211 (916) 368-1333

(219) 464-2389

Fax (707) 226-1001 Fax (916) 362-2484 Fax (219) 462-2953

QC Batch ID: HC25M550

CLIENT: Coast-to-Coast Analytical Services

141 Suburban Rd. Ste. C-4 San Luis Obispo, CA 93401

Analyzed

: 03/27/91

Analyzed by: DP

: GC/MS Method

QC SPIKE

REPORT OF ANALYTICAL RESULTS

Page 1 of 1

SAMPLE DESCRIPTION MATRIX		SAMPLED B	Y	SAMPLED DA	IE RECE	RECEIVED	
QC SPIKE	Soil						
CONSTITUENT		*PQL mg/Kg	SPIKE AMOUNT	RESULT mg/Kg	%REC	NOTE	
TOTAL SEMIVOLATILE PETROLEUM HYDROCARBONS						1,2	
Tetraethyl Lead		0.005	0.50	0.32	64.		

CCAS is Certified by CA Department of Health Services: Laboratory #131 *RESULTS listed as 'ND' were not detected at or above the listed PQL (Practical Quantitation Limit)

(1) Sample Preparation on 03/25/91 by RAD using Methylene Chloride in Sonic Bath.

(2) Spike was in analyte-free soil.

04/11/91 MSD5/ZC10A MH/erc H1024-1

Respectfully submitted,

COAST-TO-COAST ANALYTICAL SERVICES, INC.

Mary Havlicek, Ph.D.



141 Suburban Road 751 S. Kellogg, Suite A 1885 North Kelly Road 9333 Tech Center Dr., Ste. 800

2400 Cumberland Dr.

San Luis Obispo, CA 93401

Goleta, CA 93117 Napa, CA 94558 Sacramento, CA 95826

Valparaiso, Indiana 46383

(805) 543-2553 (805) 964-7838 Fax (805) 543-2685 Fax (805) 967-4386

(219) 464-2389

(707) 257-7211 (916) 368-1333 Fax (707) 226-1001 Fax (916) 362-2484 Fax (219) 462-2953

QC Batch ID: HC25M550

CLIENT: Coast-to-Coast Analytical Services

141 Suburban Rd. Ste. C-4 San Luis Obispo, CA 93401

Analyzed

: 03/27/91

Analyzed by: DP Method : GC/MS

QC SPIKE

REPORT OF ANALYTICAL RESULTS

Page 1 of 1

SAMPLE DESCRIPTION	MATRIX	SAMPLED BY		SAMPLED DATE RECEIVED			
QC SPIKE DUPLICATE	Soi1						
CONSTITUENT		#PQL mg/Kg	SPIKE AMOUNT	RESULT mg/Kg	%REC	%DIFF	NOTE
TOTAL SEMIVOLATILE PETROLEUM HYDROCARBONS							1,2
Tetraethyl Lead	_	0.005	0.50	0.25	50.	14.	

CCAS is Certified by CA Department of Health Services: Laboratory #131 *RESULTS listed as 'ND' were not detected at or above the listed PQL (Practical Quantitation Limit)

(1) Sample Preparation on 03/25/91 by RAD using Methylene Chloride in Sonic Bath.

(2) Spike was in analyte-free soil.

04/11/91 MSD5/ZC11A MH/erc H1024-1

Respectfully submitted,

COAST-TO-COAST ANALYTICAL SERVICES, INC.

Donald R. Cortes, Group Leader

Mary Havlicek, Ph.D.

```
— Average RF
Calib File: CA_TEL::D5
                            Comp # 1
                                            ----- 1st Degree
Calib Date: 910410 20:15
                                                --- 2nd Degree
 omp: Tetraethyllead
         120-
         119-
         100-
          90-
          80-
          70-
Conc.
          60-
          50-
          40-
          30-
          20-
                                 Response Ratio
```

Compound # 1 Calib File: CA_TEL::D5

Compound: Tetraethyllead

Istd: 1,4-Dichlorobenzene-d4

File: >#C19A >#C20A >#C21A >#C22A >#C23A Conc: 20.00 40.00 60.00 80.00 100.00 Rf: .37935 .34402 .33459 .30955 .32967

Average of 5 Rfs: .33944 (7.55 % Rsd) Rx: .0000000 Ry: .0000000

1st Degree Equation: y = -.096006 + 3.219230(x)

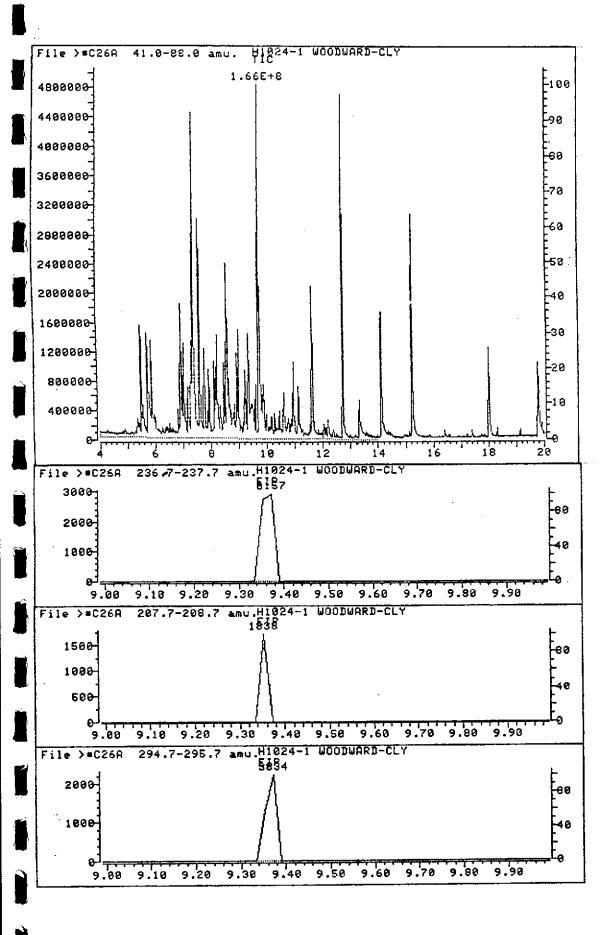
st Degree Corr Coef: .9970747

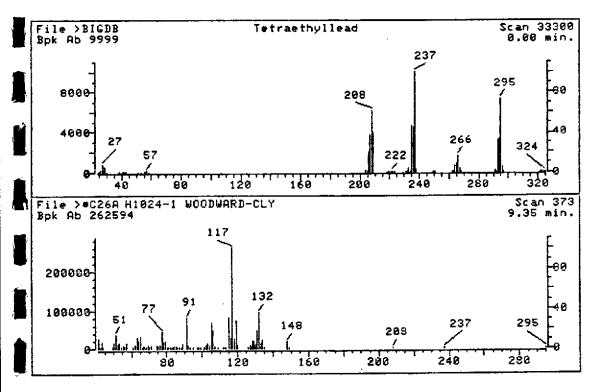
2nd Degree Equation: $y = -.231935 + 3.879870(x) + -.652235(x^2)$

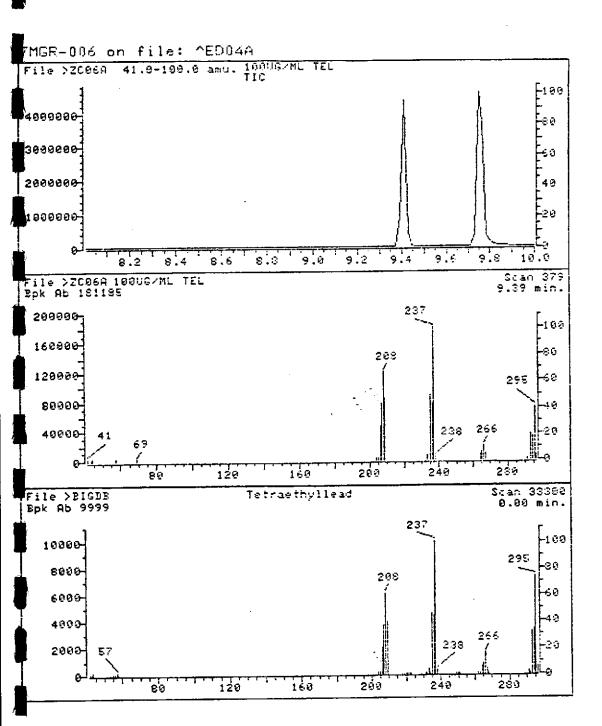
2nd Degree Corr Coef: .9978902

In the above equations:

Istd Conc for all calibration points is: 40.00









CLIENT: Albert Ridley

Air, Water & Hazardous Waste Sampling, Analysis & Consultation Certified Hazardous Waste, Chemistry, Bacteriology & Bioassay Laboratories

141 Suburban Road 751 S. Kellogg, Suite A 1885 North Kelly Road

9333 Tech Center Dr., Ste. 800 2400 Cumberland Dr.

Woodward-Clyde Consultants

500 12th St. Ste. 100

Oakland, CA 94607

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(916) 368-1333 • Fax (916) 362-2484 (219) 464-2389 • Fax (219) 462-2953

Lab Number : H-1024-1

Project

: 90C0321A-4000 Arrow

Rentals

Analyzed : 04/11/91

Analyzed by: DZ

Method : As Listed

REPORT OF ANALYTICAL RESULTS

Page 1 of 1

SAMPLE DESCRIPTION	MATRIX	SAMPLED BY	SAME	SAMPLED DATE RECEIVE			
B-F-1-16' & B-F-2-16'	Soil	Al Ridley	03	/12/91 0	3/13/91		
CONSTITUENT		(CAS RN)	*PQL mg/Kg	RESULT mg/Kg	NOTE		
FUEL FINGERPRINT ANALYSIS					1,2,3,4		
Benzene		(71432)	0.001	0.002			
Toluene		(108883)	0.001	0.025			
Ethylbenzene		(100411)	0.001	0.030			
Xylenes			0.001	0.34			
1,2-Dichloroethane (EDC)		(107062)	0.001	ND			
Ethylene Dibromide (EDB)		(106934)	0.001	ND			
Total Purgeable Petroleum Hydrocarbons	(Gasoline)		0.1	16.			
BTX as a percent of fuel				2.1			
Percent Surrogate Recovery				107.			

CCAS is Certified by CA Department of Health Services: Laboratory #131
*RESULTS listed as 'ND' were not detected at or above the listed PQL (Practical Quantitation Limit)

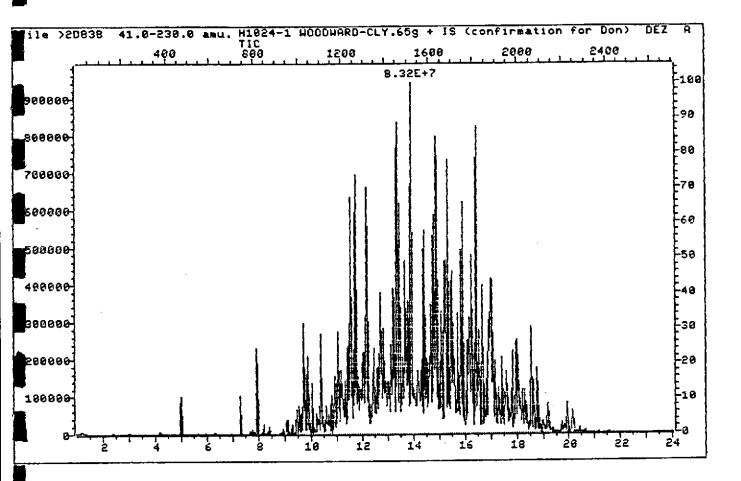
- (1) ANALYZED by CAL DHS DRAFT TPH (modified) and EPA 8240 (GC/MS)
- (2) EXTRACTED by EPA 5030 (purge-and-trap)
- (3) The composition of Tetraethyl Lead in Gasoline is 1.8g/Gallon.
- (4) Fuel Fingerprint analysis originally done on 4/1/91.

04/15/91 MSD2/2D83B MH/erc HD11M2 Respectfully submitted,

COAST-TO-COAST ANALYTICAL SERVICES, INC.

Frank R. Guenther, Ph.D., Technical Director

Mary Havlicek, Ph.D.





Air, Water & Hazardous Waste Sampling, Analysis & Consultation Certified Hazardous Waste, Chemistry, Bacteriology & Bioassay Laboratories

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(707) 257-7211 (916) 368-1333 (219) 464-2389

Fax (916) 362-2484 Fax (219) 462-2953

QC Batch ID: HD11M2

CLIENT: Coast-to-Coast Analytical Services

141 Suburban Rd. Ste. C-4 San Luis Obispo, CA 93401

Analyzed

: 04/11/91

Analyzed by: DZ

Method : As

: As Listed

METHOD BLANK

REPORT OF ANALYTICAL RESULTS

Page 1 of 1

SAMPLE DESCRIPTION	MATRIX	SAMPLED BY	SAM	RECEIVED	
METHOD BLANK	Soil				
CONSTITUENT		(CAS RN)	*PQL mg/Kg	RESULT mg/Kg	NOTE
FUEL FINGERPRINT ANALYSIS					1,2
Benzene		(71432)	0.001	ND	
Toluene	.··	(108883)	0.001	ND	
Ethylbenzene		(100411)	0.001	ND	
Xylenes			0.001	ND	
1,2-Dichloroethane (EDC)		(107062)	0.001	ND	
Ethylene Dibromide (EDB)		(106934)	0.001	ND	
Total Purgeable Petroleum Hydrocarbons	(Gasoline)		0.1	ND	
BTX as a percent of fuel				Not Appl.	
Percent Surrogate Recovery			110.		

CCAS is Certified by CA Department of Health Services: Laboratory #131
*RESULTS listed as 'ND' were not detected at or above the listed PQL (Practical Quantitation Limit)

- (1) ANALYZED by CAL DHS DRAFT TPH (modified) and EPA 8240 (GC/MS)
- (2) EXTRACTED by EPA 5030 (purge-and-trap)

04/15/91 MSD2/2D82B MH/erc H1024-1 Respectfully submitted,

COAST-TO-COAST ANALYTICAL SERVICES, INC.

Frank R. Guenther, Ph.D., Technical Director

Mary Havlicek, Ph.D.



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QC Batch ID: HD11M2

CLIENT: Coast-to-Coast Analytical Services

141 Suburban Rd. Ste. C-4 San Luis Obispo, CA 93401

Analyzed : 04/11/91

Analyzed by: DZ

Method : As Listed

QC SPIKE

REPORT OF ANALYTICAL RESULTS

Page 1 of 1

SAMPLE DESCRIPTION	MATRIX	SAMPLED B	Y	SAMPLED DA	IVED	
QC SPIKE	Soil					
CONSTITUENT		≉PQL mg/Kg	SPIKE AMOUNT	RESULT mg/Kg	%REC	NOTE
FUEL FINGERPRINT ANALYSIS						1,2
Benzene		0.001	0.003	0.003	100.	
Toluene		0.001	0.010	0.010	100.	
Ethylbenzene	•	0.001	0.002	0.002	100.	
Xylenes		0.001	0.010	0.010	100.	
1,2-Dichloroethane (EDC)		0.001		NS		
Ethylene Dibromide (EDB)		0.001		NS		
Total Purgeable Petroleum Hydrocarbons (Gasoline)	0.1	0.8	0.8	100.	
BTX as a percent of fuel	•		36.	32.		
Percent Surrogate Recovery			108.	110.		

CCAS is Certified by CA Department of Health Services: Laboratory #131

- * RESULTS listed as 'NS' were not spiked. PQL = Practical Quantitation Limit
- (1) ANALYZED by CAL DHS DRAFT TPH (modified) and EPA 8240 (GC/MS)
- (2) EXTRACTED by EPA 5030 (purge-and-trap)

04/15/91 MSD2/2D85B MH/erc H1024-1 Respectfully submitted,

COAST-TO-COAST ANALYTICAL SERVICES, INC.

Frank R. Guenther, Ph.D., Technical Director

in - proces

Mary Havlicek, Ph.D.



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QC Batch ID: HD11M2

CLIENT: Coast-to-Coast Analytical Services

141 Suburban Rd. Ste. C-4 San Luis Obispo, CA 93401

Analyzed : 04/11/91

Analyzed by: DZ

Method : As Listed

QC SPIKE

REPORT OF ANALYTICAL RESULTS

Page 1 of 1

SAMPLE DESCRIPTION	MATRIX	SAM	PLED BY	SAMP	LED DA	TE RECE	IVED
QC SPIKE DUPLICATE	Soil						
CONSTITUENT		*PQL mg/Kg	SPIKE AMOUNT	RESULT mg/Kg	%REC	%DIFF	NOTE
FUEL FINGERPRINT ANALYSIS							1,2
Benzene		0.001	0.003	0.003	100.	٥.	
Toluene		0.001	0.010	0.009	90.	10.	
Ethylbenzene	•	0.001	0.002	0.002	100.	0.	
Xylenes		0.001	0.010	0.010	100.	0.	
1,2-Dichloroethane (EDC)		0.001		NS			
Ethylene Dibromide (EDB)		0.001		NS			
Total Purgeable Petroleum Hydrocarbons ((Gasoline)	0.1	0.8	0.7	88.	13.	
BTX as a percent of fuel			36.	32.			
Percent Surrogate Recovery			108.	112.			

CCAS is Certified by CA Department of Health Services: Laboratory #131

04/15/91 MSD2/2D86B MH/erc H1024-1 Respectfully submitted,

COAST-TO-COAST ANALYTICAL SERVICES, INC.

Frank R. Guenther, Ph.D., Technical Director

Mary Havlicek, Ph.D.

^{*} RESULTS listed as 'NS' were not spiked. PQL = Practical Quantitation Limit

⁽¹⁾ ANALYZED by CAL DHS DRAFT TPH (modified) and EPA 8240 (GC/MS)

⁽²⁾ EXTRACTED by EPA 5030 (purge-and-trap)

Woodward-Clyde Consultants Chain of Custody Record 500 12th Street, Suite 100, Oakland, CA 94607-4041 (415) 893-3600 90C0321A-4000 PROJECT NO. ANALYSES APFOW Pental SAMPLEDS (Signature) REMARKS AI Ridler EPA Method 624 EPA Method 625 (Sample preservation, handling procedures, etc.) SAMPLE NUMBER DATE TIME BLUSS LINE 3/12/91 8:45 AM Brass Lines 3/12/91 8:48AM Sample From 15 depth 8"dia auger boxing

Dietrathy!

Lead test

By Special

Method

described separately. Combine samples as Needed for awalysis" ar See may H. about "special must special 2 - week TAT Bruss TOTAL NUMBER OF Linera CONTAINERS RECEIVED BY: OCO SE DECO RELINQUIŞHED,BY: DATE/TIME DATE/TIME RELINQUISHED BY : (Signature) (Signature) / 3/12 9:50 AM RECEIVED FOR LAB BY DATE/TIME COURIER : SHIPPED BY: (Signature) (Signature) (Signature) Federal Expres Butha Kulchach 3 1391

Attn: Albert Ridley

Chain of Custody # 910025

January 24, 1991

Albert Ridley Woodward-Clyde Consultants 500 12th Street; Suite #100 Oakland, CA 94607-4014

Dear Mr. Ridley:

Enclosed is the report for (Project ID 90C0321A) samples which were received at Woodward-Clyde Analytical Laboratory January 22, 1991.

The report consists of the following sections:

Sample Description ΙI Analysis Results

No problems were encountered with the analysis of your samples.

If you have any questions, please feel free to call.

Sincerely,

dward R. Morales

Lab Manager

Woodward-Clyde Consultants

COC# 910025

I SAMPLE DESCRIPTION

WCC LAB ID	SAMPLE		DATE		ANALYSIS
	10	MATRIX	SAMPLED	CONTAINERS	DESCRIPTION
910025-01-01	P-1(SOIL)	SOIL	01-22-91	4" BRASS LINER	TPH/BTEX

The samples were received under chain of custody, in good condition.

Woodward-Clyde Consultants

VOLATILE PETROLEUM HYDROCARBONS MODIFIED EPA METHOD 8015/5030

PROJECT NAME: ARROW RENTALS PROJECT NUMBER: 90C0321A COC# 910025

PROJECT NOMBER: 9000321A
PROJECT MANAGER: ALBERT RIDLEY

						DETECTION	
HCC	SAMPLE ID	MATRIX	COLLECTION DATE	EXTRACTION DATE	ANALYSIS DATE	LIMIT (mg/kg)	TPH (mg/kg)
METHOD BLANK	-	-	-	-	01-23-91	0.5	ND
910025-01-01	P-1(S0IL)	SOIL	01-22-91	-	01-23-91	0.5	ND

REC 1 REC 2 %RPD

Quantitated as Gasoline.

REVIEWED BY:

Woodward-Clyde Consultants

BENZENE, TOLUENE, ETHYLBENZENE, XYLENES DATA SHEET MODIFIED EPA METHOD 8020/5030

PROJECT NAME: ARROW RENTALS

COC# 910025

PROJECT NUMBER: 90C0321A
PROJECT MANAGER: ALBERT RIDLEY

						DETECTION			ETHYL	
WCC LAB ID	SAMPLE ID	MATR1X	COLLECTION DATE	EXTRACTION DATE	ANALYSIS DATE	LIMIT (ug/kg)	BENZENE (ug/kg)	TOLUENE (ug/kg)	BENZENE (ug/kg)	XYLENES (ug/kg)
METHOD BLANK 910025-01-01	- P-1(SOIL)	- \$01L	- 01-22-91	-	01-23-91 01-23-91	5 5	ND ND	ND ND	ND ND	ND ND

QUALITY ASSURANCE SUMMARY

REC 1	REC 2	%RPD
		-
95	108	13

REVIEWED BY:

7/0025

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DATE	TIME		SAMPLE NUM	MBER	General Mineral	Poority Po	EPA Metiod 624	EPA Method 625	EPA Method 608	# 2			Munber		ng procedur	_
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Chevron Real Estate Management Company

A Division of Chevron Industries, Inc. 225 Bush Street, San Francisco, California Mail Address PO Box 7137, San Francisco, CA 94120 7137

October 16, 1990

FUEL CHARACTERIZATION DATA LIVERMORE SITE

Mr. Albert P. Ridley, C.E.G. Woodward-Clyde Consultants 500 12th Street, Suite 100 Oakland, CA 94607-4014

Dear Al:

Attached please find results of samples that you submitted to Chevron Research and Technology Company (CRTC) for analysis. As we discussed, Chevron is providing assistance to Woodward-Clyde in the determination of the possible source of hydrocarbon contamination at 187 North L Street, Livermore. However, this assistance is not to be construed as an admission of liability by Chevron in relation to the incident that occurred at the above mentioned address.

It is my understanding that I will be shortly receiving a bill from CRTC for work completed. I will then pass this bill to Woodward-Clyde for payment.

If you have any questions, please give me a call at (415) 894-5427.

Very truly yours,

E. P. Johnke

Environmental Specialist

EPJ:oct Attachment

cc: J. P. Gorman

E. A. Harvey

CHEVRON RESEARCH AND TECHNOLOGY COMPANY ANALYTICAL SCIENCES UNIT PROJECT SUMMARY

Log No. 4527 Requested by E. P. Johnke
Date Initiated 08/23/90 Location 575 Market, Room 1882
Date Completed 10/04/90 Phone 894-5427
Charge Code TT52508

Project Description: Analyze one hydrocarbon sample, labeled W1-CH, and three water samples, labeled WA-CH, WB-CH, and TB, taken from Arrow Rentals property at 187 North L Street in Livermore, CA. The site was a Mobil gas station for over 30 years. Seven years ago a new gas tank was installed by Arrow Rentals. The tank has been supplied by a Chevron jobber, Petcock Petroleum, since then. In June of 1985, the Petcock delivery driver mistakenly poured gasoline down a vapor monitoring well at the site. Determine whether the hydrocarbon contamination comes from the old Mobil station or the 1985 accident.

<u>Result</u>: The results are not definitive, but suggest that the gasoline may have been produced in about 1985. The lead alkyl isomer pattern suggests Chevron did not produce the gasoline, while the detergent additive analysis suggests we may have.

W1-CH contains leaded regular grade gasoline. The tetraalkyllead content of the sample is 0.18 g/qal. The gasoline is no more than 10% evaporated, so that if we account for the evaporation, we calculate that the original lead level must have been no less than 0.16 g/gal. This level is typical of gasolines produced between late 1985 and 1988. If some of the lead has biodegraded or dissolved, the original level would be somewhat higher than 0.16. These processes occur slowly, however, and gasolines spilled before 1985 typically retain most of their lead, with lead levels of 0.5 to 3.0 g/gal. gasoline. The infrared spectrum of the gasoline shows that it is oxidized from exposure to air. It also has a rather weak absorbance at 1230 cm-1 that may be due to a polybutene-amine type detergent additive. Chevron used polybutene-amine-containing F310 detergent in leaded gasolines up until 1987. Mobil also uses a polybutene amine containing detergent. A dye analysis was not run because we have no record of what kind of dyes Chevron or Mobil have used in the past.

We have no chromatograms of Chevron June 1985 gasolines in our files. We do have a chromatogram of Chevron Supreme Unleaded produced at our Richmond refinery in late November 1985. There is nothing in this chromatogram to indicate that the W1-CH sample could not have come from the Richmond refinery. We also have an lead alkyl analysis of Chevron Regular Leaded gasoline produced at the Richmond refinery in November 1985. Chevron was adding 0.22 g/gal lead to its leaded regulars at that time. The November 1985 gasoline contained more than 95% of its lead as the tetraethyl isomer (TEL). The W1-CH gasoline contains only 83% of its lead as the tetraethyl isomer. This implies that the WL-CH gasoline is not Chevron's. However, it is possible that Chevron purchased an exchange gasoline from another supplier. Nick

Clark of Supply and Distribution (944-6212) should have the 1985 records on Livermore area exchange gasoline purchases.

The three water samples were extracted with carbon disulfide to determine the concentration of hydrocarbon present. The hydrocarbon in WA-CH is present at 7.6 uL/L (ppm), high enough to recognize it as dissolved or entrained gasoline. WA-CH contains 2.4 uL/L hydrocarbon, predominately the soluble aromatics from gasoline. TB hydrocarbons are present at 73 nL/L (ppb), and do not have a clear relationship to gasoline. None of the samples contain enough hydrocarbon for gasoline additive analysis.

<u>Analytical Approach</u>: The samples were analyzed by gas chromatography using a flame ionization detector (GC/FID) to determine hydrocarbon composition and an electron capture detector (GC/PID) to measure lead alkyl concentration. Water samples were extracted with carbon disulfide and analyzed by GC/FID. Detergent additive was analyzed by Sep-Pak extraction and infrared spectroscopy.

Reported by J. Kimberlin, E. A. Harvey

ZPJohnke
AWVerstuyft
THGouw
JKimberlin
EAHarvey
file

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122	14 00	W1-CH/a	_)						X			3	1
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122	12:44 h	NA-CH	(()									7	
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ENVIRONMENTAL CHEMISTS

Andrew John Friedman James E. Bruya, Ph.D. (206) 285-8282 3008-B 16th Avenue West Seattle, WA 98119 FAX: (206) 283-5044

August 28, 1990

Al Ridley, Project Leader Woodward Clyde Consultants 500 12th Street, Suite 100 Oakland, CA 94607-4041

Dear Mr. Ridley:

Enclosed are the results of the analyses of the samples submitted on August 4, 1990 from Project 90C0321A.

We appreciate this opportunity to be of service to you on this project. If you have any questions regarding this material, or if you just want to discuss any aspect of your projects, please do not hesitate to contact me.

Sincerely,

Andrew John Friedman, Chemist

AJF/fae

Enclosures

ENVIRONMENTAL CHEMISTS

Date of Report: August 28, 1990 Date Submitted: August 4, 1990

Project: 90C0321A

RESULTS OF ANALYSES OF THE WATER SAMPLES FOR BTX AND ETHYLBENZENE USING THE HEADSPACE METHOD Results Reported as ng/mL (ppb)

Sample #	Benzene	Toluene	Et-Benzene		<u>Vlene</u>
				m.p	Q
W-B	22,000	7,900	2,000	4,000	1,800
W-B2	21,000	7,300	1,800	3,700	1,700
W-EB	11	11	4	3	5
W-D	1	2	1	1	1
W-E	<1	<1	<1	<1	<1
W-C	<1	<1	<1	<1	<1
W-A	6,800	5,500	620	3,400	1,700
Travel Blank	7 a	37 a	14 a	43 a	22 a
Quality Assurance					
Method Blank	<1	<1	<1	<1	<1
W-A (Duplicate)	6,900	5,600	620	6,800	1,800
W-A (Matrix Spike) Spiked @ 1,000 ppb Percent Recovery	ъ	b	b	ъ	ъ
W-A (Matrix Spike Dupl: Spiked @ 1,000 ppb					
Percent Recovery	þ	þ	b	ъ	ъ

a - Contamination found is likely from the previous sample. This is supported by no confirmation of this analyte via second column confirmation analysis.

b - The amount spiked was insufficient to give meaningful recovery data.

ENVIRONMENTAL CHEMISTS

Date of Report: August 28, 1990 Date Submitted: August 4, 1990

Project: 90C0321A

RESULTS OF ANALYSES OF THE PRODUCT SAMPLES FOR BTX AND ETHYLBENZENE USING THE HEADSPACE METHOD Results Reported as Percent

	<u>Benzene</u>	<u>Toluene</u>	Et-Benzene	Xyl	<u>ene</u>
Sample #	-			m.p	Q
F-T	4.2 %	8.9 %	1.7 %	6.1 %	2.6 %
F-T (PRES)	4.2 %	8.9 %	1.7 %	6.2 %	2.6 %

ENVIRONMENTAL CHEMISTS

Date of Report: August 28, 1990 Date Submitted: August 4, 1990

Project: 90CO321A

RESULTS OF ANALYSES OF THE WATER SAMPLES FOR NONHALOGENATED ORGANICS BY EPA METHOD 8015 (GAS AND DIESEL) Results Reported as mg/L (ppm)

Sample #	Gasoline (ppm)	Diesel (ppm)
W-B	13	1.7
W-B2	21	1.6
W-EB	1.7	0.1
W-D	0.1	<0.1
W-E	<0.01	<0.1
W-C	<0.01	<0.1
W-A	10	2.4
Travel Blank	0.5	<0.1
Quality Assurance		
Method Blank	<0.01	<0.1
Tap Water (Matrix Spike) Spiked @ 10 ppm Percent Recovery	87%	107%
Tap Water		10/6
(Matrix Spike Duplicate) Spiked @ 10 ppm Percent Recovery	91%	74%

ENVIRONMENTAL CHEMISTS

Date of Report: August 28, 1990 Date Submitted: August 4, 1990

Project: 90C0321A

RESULTS OF ANALYSES OF THE WATER SAMPLES FOR FINGERPRINT CHARACTERIZATION BY CAPILLARY GAS CHROMATOGRAPHY

Sample #	GC Characterization
W-B	The gas chromatographic trace was indicative of a water soluble fraction of gasoline. This characterization is based on the presence of light hydrocarbons and monocyclic aromatic hydrocarbons (MCA) as the only compounds found in the sample.
₩-B2	The gas chromatographic trace was indicative of a water soluble fraction of gasoline. This characterization is based on the presence of light hydrocarbons and monocyclic aromatic hydrocarbons (MCA) as the only compounds found in the sample.
W-EB	The gas chromatographic trace was indicative of very low levels of the water soluble fraction of gasoline. This characterization is based on the presence of light hydrocarbons and MCA.
W-EB	The gas chromatographic trace contained only traces of highly volatile hydrocarbons, mostly butanes.
W-E	The gas chromatographic trace was indicative of no contamination.
W-C	The gas chromatographic trace was indicative of no contamination.
₩-A	The gas chromatographic trace was indicative of the water of a soluble fraction of gasoline. This characterization is based on the presence of light hydrocarbons and MCA.
Travel Blank	The gas chromatographic trace contained trace levels of benzene and toluene.

Nothing else was seen.

PRIEDMAN & DRUTA, INC.

ENVIRONMENTAL CHEMISTS

Date of Report: August 28, 1990 Date Submitted: August 4, 1990

Project: 90C0321A

RESULTS OF ANALYSES OF THE PRODUCT SAMPLES FOR FINGERPRINT CHARACTERIZATION BY CAPILLARY GAS CHROMATOGRAPHY

Sample #

GC Characterization

F-T

The gas chromatographic trace was indicative of a fresh gasoline. This characterization is based on a relatively typical pattern of peaks from $n\text{-}C_4$ to ca $n\text{-}C_{10}$ with a maximum at ca $n\text{-}C_7$ along with augmented levels of monocyclic aromatic hydrocarbons. Examination of the relative proportions of the four peaks we have found to be most conservative in gasolines gave abundance ratios of 1:1.17:0.861:0.0935. This did not match the W-1 or W-A-40 patterns. Analysis of the sample for organic lead gave <50 ppm.

F-T (PRES)

The gas chromatographic trace was indicative of a fresh gasoline. This characterization is based on a relatively typical pattern of peaks from $n\text{-}C_4$ to Ca $n\text{-}C_{10}$ with a maximum at ca $n\text{-}C_7$ along with augmented levels of monocyclic aromatic hydrocarbons. Examination of the relative proportions of the four peaks we have found to be most conservative in gasolines gave abundance ratios of 1:1.06:0.794:0.0876. This did not match the W-1 or W-A-40 patterns. Analysis of the sample for organic lead gave <50 ppm.

ENVIRONMENTAL CHEMISTS

Date of Report: August 28, 1990 Date Submitted: August 4, 1990

Project: 90CO321A

RESULTS OF ANALYSES OF THE WATER SAMPLES FOR PHENOL, NAPHTHALENE, AND 2-METHYL NAPHTHALENE Results Reported as mg/L (ppm)

Sample #	Phenol (ppm)	Naphthalene (ppm)	2-Methyl Naphthalene (ppm)
W-B	<0.1	<0.01	<0.01
W-B2	<0.1	<0.01	<0.01
W-EB	<0.1	<0.01	<0.01
W-D	<0.1	<0.01	<0.01
W-E	<0.1	<0.01	<0.01
M-C	<0.1	<0.01	<0.01
W-A	<0.1	<0.01	<0.01
Travel Blank	<0.1	<0.01	<0.01
Quality Assurance	<u>e</u>		
Method Blank (Tap Water)	<0.1	<0.01	0.02
Tap Water (Matrix Spike) Spiked @ 2.2 pp Percent Recover		120%	49%
Tap Water (Matrix Spike D Spiked @ 2.2 pp	m		
Percent Recover	У	120%	51%

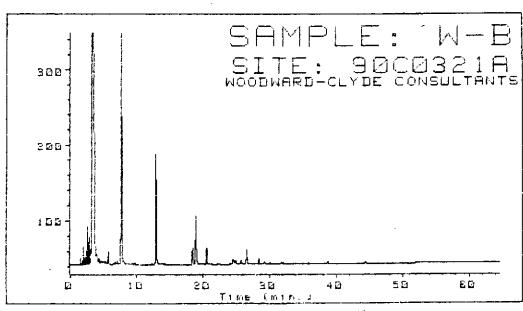
ENVIRONMENTAL CHEMISTS

Date of Report: August 28, 1990 Date Submitted: August 4, 1990

Project: 90C0321A

RESULTS OF ANALYSES OF THE WATER SAMPLES FOR SELECTED METALS BY ICP Results Reported as mg/L (ppm)

Sample #	Cadmium (ppm)	Chromium (ppm)	Copper (ppm)	Nickel (ppm)	Vanadium (ppm)	Zinc (ppm)
W-B	<0.1	<0.1	<0.1	0.2	<0.1	<0.1
W-B2	<0.1	<0.1	<0.1	0.2	<0.1	<0.1
W-EB	<0.1	<0.1	<0.1	<0.1	<0.1	<0.1
W-D	<0.1	<0.1	<0.1	<0.1	<0.1	0.2
W-E	<0.1	<0.1	<0.1	<0.1	<0.1	<0.1
W-C	<0.1	<0.1	<0.1	0.1	<0.1	<0.1
W-A	<0.1	<0.1	<0.1	0.2	<0.1	<0.1
Travel Blank	<0.1	<0.1	<0.1	<0.1	<0.1	<0.1
						1
Quality Assurance						
Method Blank	<0.1	<0.1	<0.1	<0.1	<0.1	<0.1
W-A (Duplicate)	<0.1	<0.1	<0.1	<0.1	<0.1	<0.1
W-A (Matrix Spike) Spiked @ 10 ppm Percent Recovery	81%	89%	91%	52%	77%	59%
W-A (Matrix Spike Dupli Spiked @ 10 ppm	cate)					
Percent Recovery	81%	93%	93%	58%	80%	60%



T: null.

Z: null.

Y: Sig. 2 of V3: NGC_A12A.D X: Sig. 2 of V3: NHC_A14A.D











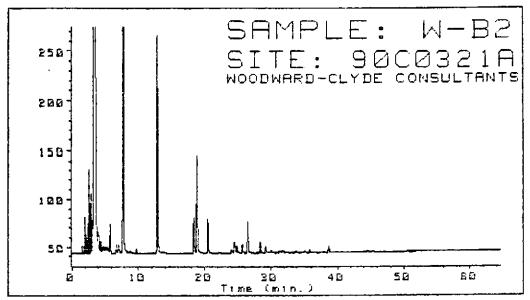










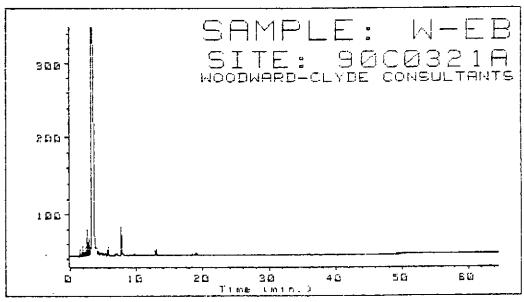


T: nall.

Z: Sig. 2 of VS:MGC_A12A.D Y: Sig. 2 of V3: NWC A14A.D

X: Sig. 2 of V3:MMC 8158.0





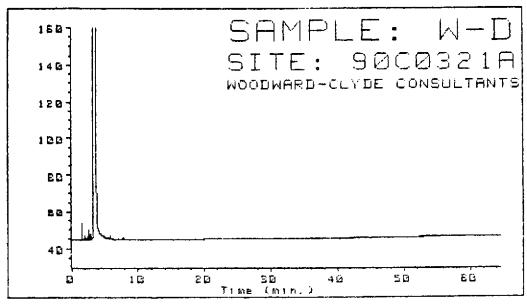
T: null. Z: null.

Y: null.

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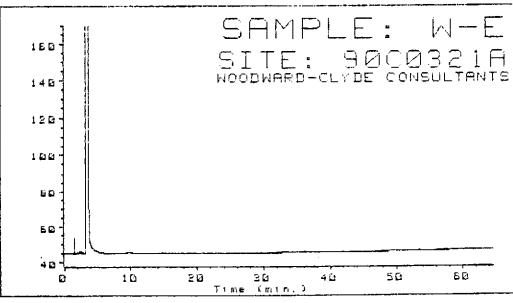
Z: mull.

Y: Sig. 2 of V3:WWC_A18A.D

X: Sig. 2 of Vs:MAC_HIZE.D



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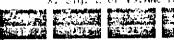
T: mull.

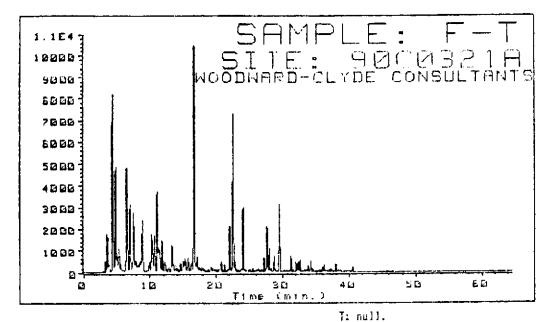
2: Sig. 2 of V3:WWC_A15A.D

Y: Sig. 2 of V3:WWC_A17A.D

X: Sig. 2 of V3:NWC A18A.D



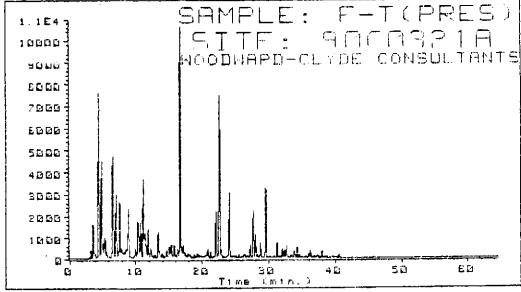




Z: noil.

Y: null.





T: mull.

Z: n#11.

Y: Sig. 2 of V3:WCA_A03A.D

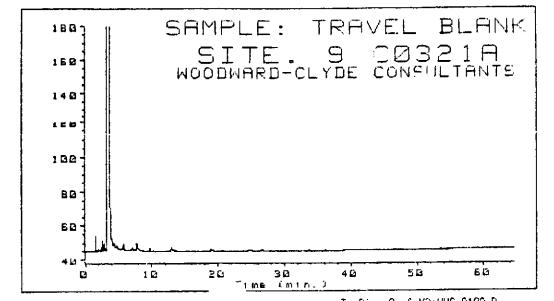
X: Sig. 2 of V3:WCA_R64A.D



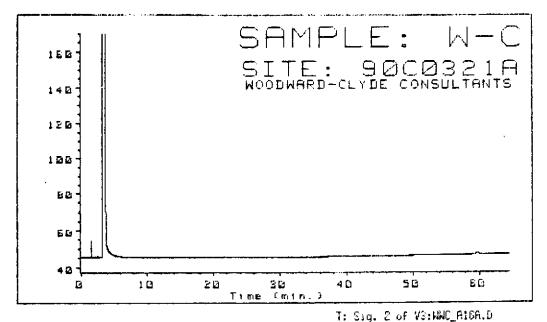




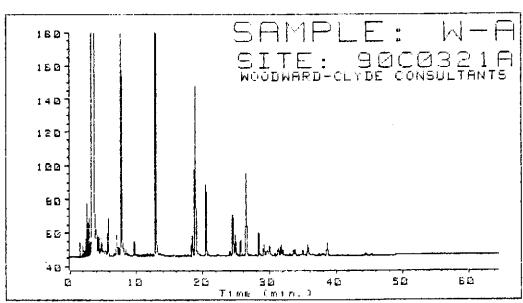








Z: Sig. 2 of V3:WHC_A17A.D
Y: Sig. 2 of V3:WHC_A18A.D
Y: Sig. 2 of V3:WHC_A18A.D
X: Sig. 2 of V3:WHC_A



T: Sig. 2 of V3:WWC_A17A.D Z: Sig. 2 of V3:WWC_A18A.D Y: Cig. 2 of V3:WWC_A19A.D

Y: Cig. 2 of V3: NNC A19A.D X: Sig. 2 of V3: NNC A20A.D

X: Sig. :





ENVIRONMENTAL CHEMISTS

Andrew John Friedman James E. Bruya, Ph.D. (206) 285-8282 3008-B 16th Avenue West Seattle, WA 98119 FAX: (206) 283-5044

September 28, 1990

Al Ridley, Project Leader Woodward-Clyde Consultants 500 12th Street, Suite 100 Oakland, CA 94607-4014

Dear Mr. Ridley:

Enclosed are the results of the analyses of the samples submitted on July 13, 1990 from Project 90C0321A.

We appreciate this opportunity to be of service to you on this project. If you have any questions regarding this material, or if you just want to discuss any aspect of your projects, please do not hesitate to contact me.

Sincerely,

Ishen Jh. Friedman, Chemist

AJF/fae

Enclosures

ENVIRONMENTAL CHEMISTS

Date of Report: September 28, 1990 Date Submitted: July 13, 1990

Project: 90C0321A

RESULTS OF ANALYSES OF THE SOIL SAMPLES FOR GASOLINE BY GC/FID (MODIFIED 8015) Results Reported as µg/g (ppm)

Sample #	Gasoline (ppm)
W-C-20	<10
W-C-40	<10
B-1-10	<10
B-1-15	<10
B-1-20	<10
B-1-30	<10
B-1-35	<10
B-1-40	350
B-1-45	54
B-1-50	<10
W-E-20	<10
W-E-40	<10
Quality Assurance	
Method Blank	<10
B-1-40 (Duplicate)	<10
B-1-40 (Matrix Spike) Spiked @ 500 ppm Percent Recovery	130%
B-1-40 (Matrix Spike Duplicate) Spiked @ 500 ppm Percent Recovery	97%

ENVIRONMENTAL CHEMISTS

Date of Report: September 28, 1990 Date Submitted: July 13, 1990 Project: 90C0321A

RESULTS OF ANALYSES OF THE SOIL SAMPLES FOR ORGANIC LEAD BY AA Results Reported as µg/g (ppm)

Sample #	Organic Lead (ppm)
B-1-20	<5
B-1-30	<5
B-1-40	<5
B-1-45	<5
Quality Assurance	
Method Blank	< 5
B-1-30 (Duplicate)	<5

ENVIRONMENTAL CHEMISTS

Date of Report: September 28, 1990

Date Submitted: July 13, 1990

Project: 90C0321A

RESULTS OF ANALYSES OF THE SOIL SAMPLES FOR FOR FINGERPRINT CHARACTERIZATION BY CAPILLARY GAS CHROMATOGRAPHY

Sample

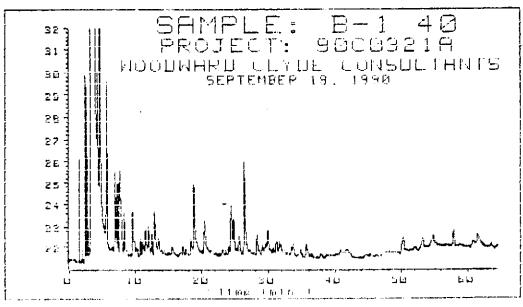
B-1 40

GC Characterization

The gas chromatographic trace was indicative of a low boiling petroleum product, such as gasoline. This characterization is based on the presence of a relatively typical suite of peaks present from ca $n-C_4$ to $n-C_{12}$ with a maximum at ca $n-C_6$. Augmented levels of benzene, toluene, ethylbenzene and the xylnes were seen which is common to most gasolines. Examining the four peaks we have found to be most conservative in gasolines we found the following relative abundances: 1:0.936:2.10:2.31. values are considerably different from samples analyzed previously from this project.

B-1 45

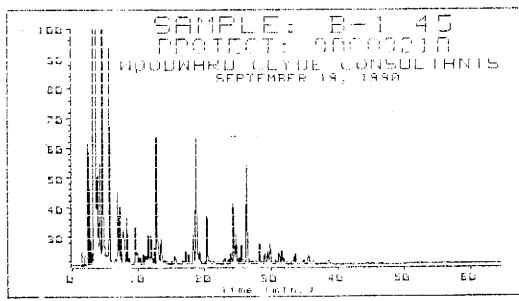
The gas chromatographic trace was indicative of a low boiling petroleum product, such as gasoline. This characterization is based on the presence of a relatively typical suite of peaks present from ca $n\text{-}\mathrm{C}_4$ to $n\text{-}\mathrm{C}_{12}$ with a maximum at ca $n\text{-}\mathrm{C}_6$. Augmented levels of benzene, toluene, ethylbenzene and the xylnes were seen which is common to most gasolines. Examining the four peaks we have found to be most conservative in gasolines we found the following relative abundances: 1:0.719:3.27:3.13. These values are similar to those from sample B-1 40 (above), though not a complete match, and different from other samples received from this project (W-1, W-A-40 or the F-T samples).



T: null. Z: nuli. Y: mull.

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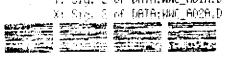


T: nall.

Z: null.

Y: Sig. 2 of DATA:WHC_ADIA.D

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ENVIRONMENTAL CHEMISTS

Andrew John Friedman James E. Bruya, Ph.D. (206) 285-8282

3008-B 16th Avenue West Seattle, WA 98119 FAX: (206) 283-5044

August 28, 1990

Al Ridley, Project Leader Woodward Clyde Consultants 500 12th Street, Suite 100 Oakland, CA 94607-4041

Dear Mr. Ridley:

Enclosed are the results of the analyses of the samples submitted on August 3, 1990 from Project 90C0321A-3000.

We appreciate this opportunity to be of service to you on this project. If you have any questions regarding this material, or if you just want to discuss any aspect of your projects, please do not hesitate to contact me.

Sincerely,

Andrew John Friedman, Chemist

AJF/fae

Enclosures

ENVIRONMENTAL CHEMISTS

Date of Report: August 28, 1990 Date Submitted: August 3, 1990

Project: 90C0321A-3000

RESULTS OF ANALYSES OF THE WATER SAMPLE FOR BTX AND ETHYLBENZENE USING THE HEADSPACE METHOD Results Reported as ng/mL (ppb)

	Benzene	Toluene	Et-Benzene	X	vlene
Sample #	-			m.p	<u>@</u> -
W-A	16,000ª	18,000ª	1,900ª	7,700ª	3,600ª

a - Value reported exceeded the calibration range established for the sample.

ENVIRONMENTAL CHEMISTS

Date of Report: August 28, 1990 Date Submitted: August 3, 1990

Project: 90C0321A-3000

RESULTS OF ANALYSES OF THE PRODUCT SAMPLE FOR BTX AND ETHYLBENZENE USING THE HEADSPACE METHOD Results Reported as ng/g (ppb)

	Benzene Toluene		Et-Benzene	<u>Xylene</u>		
Sample #				m.p	Q	
W-1	6.4%	4.7%	1.3%	5.1%	2.3%	
Quality Assurance						
W-1 (Duplicate)	6.6%	4.7%	1.3%	5.1%	2.3%	

ENVIRONMENTAL CHEMISTS

Date of Report: August 28, 1990 Date Submitted: August 3, 1990 Project: 90C0321A-3000

RESULTS OF ANALYSES OF THE WATER SAMPLE FOR PHENOL, NAPHTHALENE, AND 2-METHYL NAPHTHALENE Results Reported as mg/L (ppm)

Sample #	Phenol (ppm)	Naphthalene (ppm)	2-Methyl Naphthalene (ppm)
W-A	<0.5	<0.5	<0.05

ENVIRONMENTAL CHEMISTS

Date of Report: August 28, 1990 Date Submitted: August 3, 1990

Project: 90C0321A-3000

RESULTS OF ANALYSES OF THE PRODUCT SAMPLE FOR PHENOL, NAPHTHALENE, AND 2-METHYL NAPHTHALENE Results Reported as mg/L (ppm)

Sample #	Phenol (ppm)	Naphthalene (ppm)	2-Methyl Naphthalene (ppm)
W-1	<1,000	<1,000	200
Quality Assuranc	<u>e</u>		
Method Blank	<1,000	<1,000	<100
W-1 (Duplicate)	<1,000	<1,000	200
W-1 (Matrix Spike) Spiked @ 10,000 % Recovery	ppm b	150%	50%
W-1 (Matrix Spike Duplicate) Spiked @ 10,000 % Recovery	ppm b	150%	50%

 $[{]f b}$ - The analyte indicated was not added to the matrix spike sample.

ENVIRONMENTAL CHEMISTS

Date of Report: August 28, 1990 Date Submitted: August 3, 1990

Project: 90C0321A-3000

RESULTS OF ANALYSES OF THE WATER SAMPLE FOR FINGERPRINT CHARACTERIZATION BY CAPILLARY GAS CHROMATOGRAPHY

Sample #

GC Characterization

W-A

The gas chromatographic trace was indicative of a water soluble fraction of gasoline. This characterization is based on the presence of mostly monocyclic aromatic hydrocarbons (BTEX + C_3 - Benzenes).

ENVIRONMENTAL CHEMISTS

Date of Report: August 28, 1990 Date Submitted: August 3, 1990

Project: 90C0321A-3000

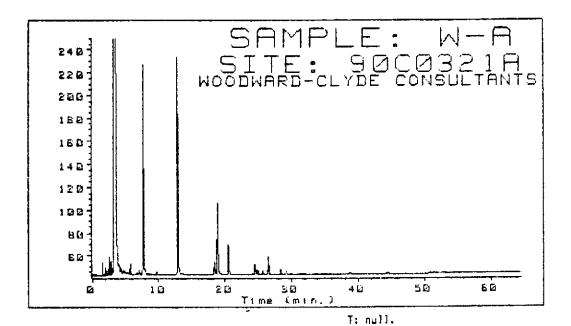
RESULTS OF ANALYSES OF THE PRODUCT SAMPLE FOR FINGERPRINT CHARACTERIZATION BY CAPILLARY GAS CHROMATOGRAPHY

Sample #

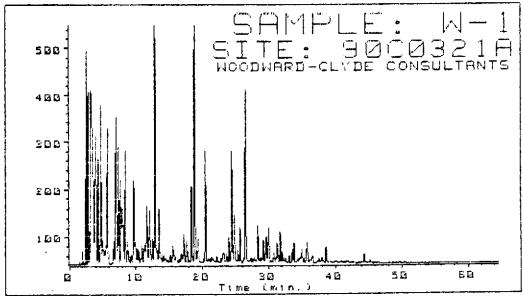
W-1

GC Characterization

The gas chromatographic trace was indicative of a fresh gasoline. This characterization is based on the presence of a relatively typical pattern of peaks present from ca n- C_4 to n- C_{12} with a maximum at ca n- C_7 , along with augmented levels of the monocyclic aromatic hydrocarbons. Examination of the relative proportions of the four peaks we have found to be most conservative in gasolines gave abundance ratios of 1:2.01:1.64:0.153. These did not give a good match to the F-T or W-A-40 samples. Analysis of the sample for organic lead gave a value of 100 ppm.



Z: null. Y: mull. $\mathsf{IDE}\mathsf{3}$ X: Sig. 2 of V3:HHC_A09A.D



T: null.

Z: null.

Y: Sig. 2 of V3:WWC_A10A.D

X: Sig. 2 of V3:WWC_A10A.D

[IE]





ENVIRONMENTAL CHEMISTS

Andrew John Friedman James E. Bruya, Ph.D. (206) 285-8282 3008-B 16th Avenue West Seattle, WA 98119 FAX: (206) 283-5044

August 28, 1990

Al Ridley, Project Leader Woodward Clyde Consultants 500 12th Street, Suite 100 Oakland, CA 94607-4041

Dear Mr. Ridley:

Enclosed are the results of the analyses of the samples submitted on July 13, 1990 from Project 90C0321A.

We appreciate this opportunity to be of service to you on this project. If you have any questions regarding this material, or if you just want to discuss any aspect of your projects, please do not hesitate to contact me.

Sincerely,

andrew John Friedman, Chemist

AJF/fae

Enclosures

ENVIRONMENTAL CHEMISTS

Date of Report: August 28, 1990 Date Submitted: July 13, 1990

Project: 90C0321A

RESULTS OF ANALYSES OF THE SOIL SAMPLES FOR ORGANIC LEAD BY AA Results Reported as $\mu g/g$ (ppm)

Sample #	Organic Lead (ppm)
W-A-20	<1
W-A-30	<1
W-A-35	<1
W-B-25	<1
W-B-35	<1
Quality Assurance W-A-20	
(Duplicate)	<1
W-A-20 (Matrix Spike) Spiked @ 2.5 ppm Percent Recovery	120%
W-A-20 (Matrix Spike Duplicate) Spiked @ 2.5 ppm Percent Recovery	120%
-	

ENVIRONMENTAL CHEMISTS

Date of Report: August 28, 1990 Date Submitted: July 13, 1990

Project: 90C0321A

RESULTS OF ANALYSES OF THE SOIL SAMPLES FOR DYE CHARACTERIZATION BY THIN LAYER CHROMATOGRAPHY

Sample #	GC Characterization
W-D-25	No dyes found.
W-A-40	No dyes found.
W-B-40	No dyes found.

ENVIRONMENTAL CHEMISTS

Date of Report: August 28, 1990 Date Submitted: July 13, 1990

Project: 90CO321A

RESULTS OF ANALYSES OF THE SOIL SAMPLES FOR FINGERPRINT CHARACTERIZATION BY CAPILLARY GAS CHROMATOGRAPHY

Sample #	GC Characterization
W-D-25	The gas chromatographic trace was not indicative of any hydrocarbon contamination.
W-D-40	The gas chromatographic trace was not indicative of any hydrocarbon contamination.
W-A-20	The gas chromatographic trace was not indicative of any hydrocarbon contamination.
W-A-30	The gas chromatographic trace was not indicative of any hydrocarbon contamination.
W-A-40	The gas chromatographic trace was indicative of weathered gasoline. This characterization is based on the presence of a relatively typical pattern of peaks present from ca n - C_4 to n - C_{12} with a maximum at n - C_8 . Only low levels of monocyclic aromatics were probably present. They were mostly solubilized or biodegraded. Examination of the relative proportions of the four peaks we have found to be most conservative in gasolines gave abundance ratios of 1:4.26:3.88:0.442. These did not give a good match to the F-T or W-1 samples.
W-A-50	The gas chromatographic trace was not indicative of any hydrocarbon contamination.
W-B-40	The gas chromatographic trace was not indicative of any hydrocarbon contamination.

ENVIRONMENTAL CHEMISTS

Date of Report: August 28, 1990 Date Submitted: July 13, 1990

Project: 90C0321A

RESULTS OF ANALYSES OF THE SOIL SAMPLES FOR GASOLINE BY GC/FID (MODIFIED 8015) Results Reported as μg/g (ppm)

Sample #	Gasoline (ppm)
W-D-25	<1
W-D-40	<1
W-A-20	<1
W-A-30	2
W-A-40	1,000
W-B-25	<1
W-B-35	<1
W-B-40	<1
Quality Assurance	
Method Blank	<1
W-A-40 (Duplicate)	640
W-A-40 (Matrix Spike) Spiked @ 100 ppm Percent Recovery	a
W-A-40 (Matrix Spike Duplicate) Spiked @ 100 ppm Percent Recovery	a

a - The amount spiked was insufficient to give meaningful recovery data.

ENVIRONMENTAL CHEMISTS

Date of Report: August 28, 1990 Date Submitted: July 13, 1990

Project: 90CO321A

RESULTS OF ANALYSES OF THE SOIL SAMPLES FOR BTX AND ETHYLBENZENE USING THE HEADSPACE METHOD Results Reported as ng/g (ppb)

	Benzene	<u>Toluene</u>	Et-Benzer		Xylene
Sample #				m.p	<u>o</u>
W-D-25	<10	<1	<1	<1	<1
W-D-40	<10	<1	<1	<1	<1
W-A-20	410	320	240	210	190
W-A-30	390	130	35	1,200	750
W-A-40	12,000	37,000	7,500	27,000	13,000
W-B-25	<10	<1	<1	<1	<1
W-B-35	690	260	110	70	35
W-B-40	<10	<1	<1	<1	<1
Quality Assurance					
Method Blank	<10	<1	<1	<1	<1
W-A-40 (Duplicate)	8,900	29,000	6,000	23,000	11,000
W-A-40 (Matrix Spike) Spiked @ 1,000 ppb Percent Recovery	a	a	a	a	a
W-A-40 (Matrix Spike Dupl Spiked @ 1,000 ppb Percent Recovery		a	a	a	· a

The amount spiked was insufficient to give meaningful recovery data.

ENVIRONMENTAL CHEMISTS

Date of Report: August 28, 1990 Date Submitted: July 13, 1990

Project: 90C0321A

RESULTS OF ANALYSES OF THE WATER SAMPLES FOR SELECTED METALS BY ICP Results Reported as mg/L (ppm)

Sample #	Cadmium (ppm)	Chromium (ppm)	Copper (ppm)	Nickel (ppm)	Vanadium (ppm)	Zinc (ppm)
W-A-20	0.2	2.3	12	64	0.3	11
W-A-30	<0.1	23	13	55	6.3	23
W-A-35	0.6	31	15	6.9	9.1	15
W-B-25	<0.1	76	15	82	6.3	21
W-B-35	0.2	22	18	74	7.1	14
Quality Assurance						
Method Blank	<0.1	<0.1	<0.1	<0.1	<0.1	<0.1
W-A-30 (Duplicate)	0.1	26	16	68	7.4	21
W-A-30 (Matrix Spike) Spiked @ 100 ppm Percent Recovery	55%	66%	70%	77%	a	60%
W-A-30 (Matrix Spike Dupl: Spiked @ 100 ppm	icate)					, v
Percent Recovery	54%	53%	64%	39%	a	56%

 $[{]f a}$ - The analyte indicated was not added to the matrix spike sample.

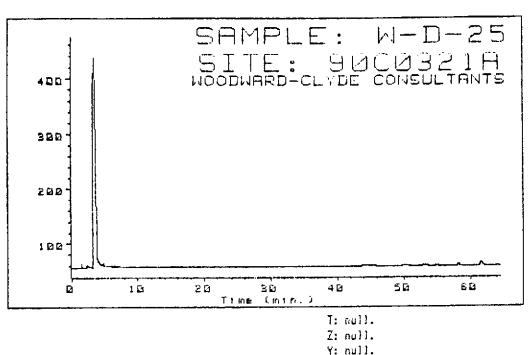
ENVIRONMENTAL CHEMISTS

Date of Report: August 28, 1990 Date Submitted: July 13, 1990

Project: 90C0321A

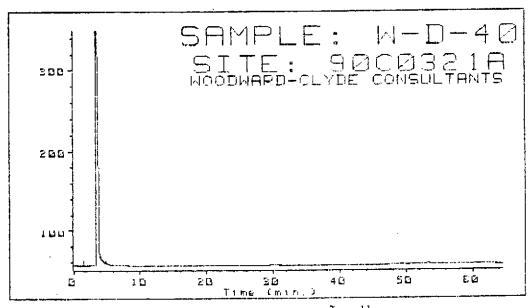
RESULTS OF ANALYSES OF THE SOIL SAMPLES FOR PHENOL, 2-MEHTYL NAPHTHALENE, AND NAPHTHALENE Results Reported as $\mu g/g$ (ppm)

Sample #	Phenol (ppm)	Naphthalene (ppm)	2-Methyl Naphthalene (ppm)
W-A-30	<10	<1	<1
W-B-30	<10	<1	<1
Quality Assurance			
Method Blank	<10	<1	2



Y: mull.





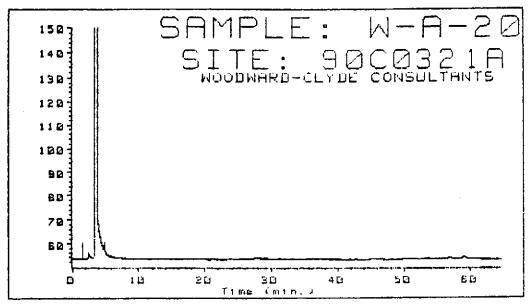
7: mull. Z: mull.

Y: Sig. 2 of V3:WOC_A01A.D

K: Sig. 2 of V3:WOC_A02A.D



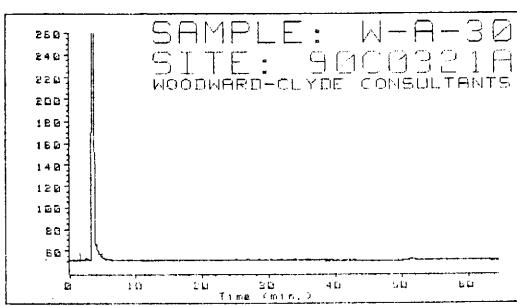
[[E]



T: null. Z: null.

Y: null.

X: Sig. 2 of V3:NOC_A03A.D [DE] is pare CHUNAI



T: null.

Z: null.

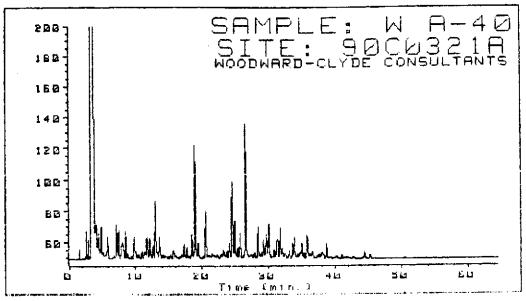
Y: Sig. 2 of V3:NOC_A03A.D

X: Sig. 2 of V3:WOC_A04A.D







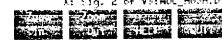


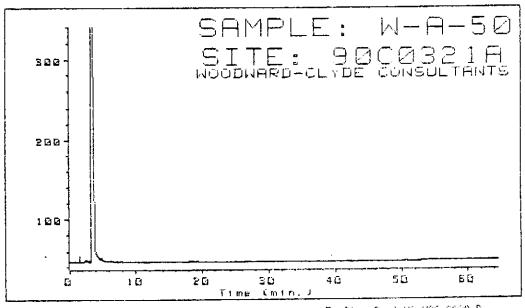
T: mull.

Z: Sig. 2 of V3:W0C_A03A.D

Y: Sig. 2 of V3:NDC_A04A.D X: Sig. 2 of V3:NDC_A05A.D







T: Sig. C of V3:NOC_AOSA.D

Z: Sig. 2 of V3:WOC_A04A.D

Y: Sig. 2 of V3:WOC_AO5A.D

X: Sig. 2 of V3:WOC_A06A.D

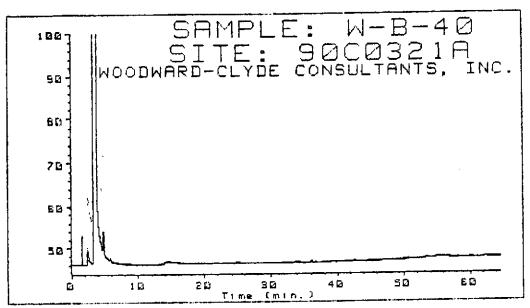












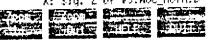
T: null.

Z: null.

Y: mull.

X: Sig. 2 of V3: NOC_A07H.D





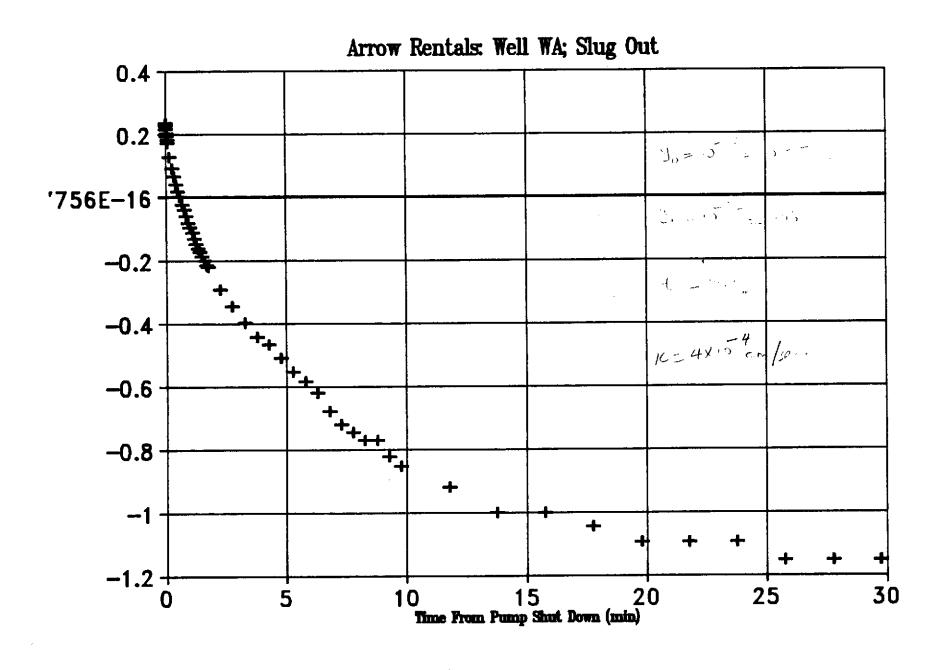


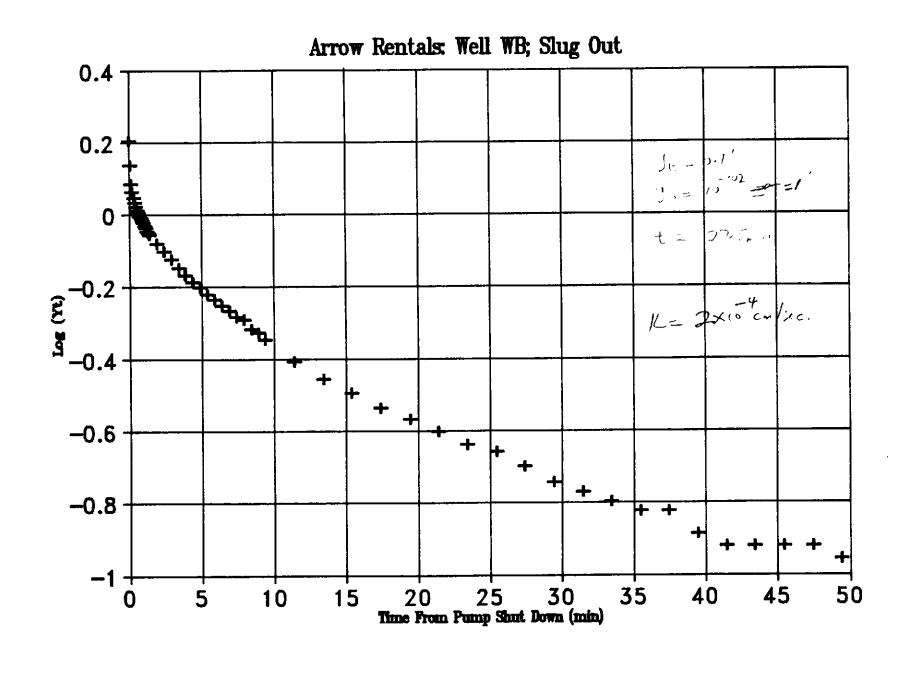


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Woodward-Clyde Consultants Chain of Custody Record 500 12th Street, Suite 100, Oakland, CA 94607-4041 (415) 893-3600 PROJECT NO. 90 CO 3 21 A **ANALYSES** Priority Pollutant Metals
EPA Method 624
EPA Method 625 Number of Containers SAMPLERS (Signature) REMARKS EPA Method 608 (Sample preservation, handling procedures, etc.) SAMPLE NUMBER TIME DATE 73 W-C-20 7/11 W-C= 25 74 W-C-30 -C - 35 7/10 ŹD જા 13-1-45 W. E-10 W-E-15 W- F-20 W- E-25 W- E-30 W- F - 40 9/0 TOTAL NUMBER OF CONTAINERS RECEIVED BY: DATE/TIME RELINQUISHED BY: DATE/TIME RECEIVED BY: RELINQUISHED BY : (Signature) (Signature) (Signature) (dis () muberg METHOD OF SHIPMENT : DATE/TIME RECEIVED FOR LAB BY : COURIER: SHIPPED BY : ICE Chest place (Signature) (Signature) (Signature)

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FIELD ACTIVITY REPORT ARROW RENTALS, LIVERMORE, CA 90C0321A-3000

FIELD PERSONEL: Jacki Lee

SUBCONTRACTOR: Frank Briggs, Datum Corporation

PURPOSE: To develop five new monitoring wells and obtain water levels on existing wells. Collect samples if gasoline is present.

GENERAL SUMMARY: Each new monitoring well received partial development. Additional pumping is still required for complete development.

WELL DEVELOPMENT PROCEDURES (see Table 1)

- 1. Initial water level and total depth were measured from the north rim of the well casing, surface completion was checked.
- 2. Bailed out as much silt deposit as possible.
- 3. Swab the screen interval for 10 to 15 minutes (for 4 inch wells only).
- 4. Bail well again after swabbing.
- 5. Pump well if possible until silt was minimal (for well W-D).

COMMENTS:

Well W-A: No floaters were present, however, strong gasoline odor was detected. A water sample was collected. Subcontractor was able to bail the well dry after 215 gallons were removed. Well was bailed again on the second day. High amount of silt is still present.

Well W-B: No floaters or odor were detected. No sample was collected. Subcontractor was able to bail the well dry after 70 gallons. Allowed a recharge time of 15 minutes. Only 10 gallons were removed before well went dry again. Amount of settled silt is about 50 ml.

Well W-C: No odor was detected. Originally started out with very thick silt. After removing about 3-4 gallons, viscosity was like water, however, still a high presence of silt. Subcontractor could not bail dry.

Well W-D: No odor was detected. Subcontractor could not get a 8 foot long bailer down the well. It would get stuck at various depths. However, a 3 foot long bailer fits down the well to total depth. This well was hand-bailed. Silt is present.

A total of 10 empty barrels were left at the site. All discharged groundwater is stored in labelled barrels. Total number of barrels used for development is 22 barrels.

Groundwater samples from wells W-A and W-1 were sent along with a travel blank to Seattle, WA. Samples from Well W-1 was pure product.

Jadrie, 10-31-50

Sanz was 2"4"

of product First

Day, Next day

Filled Built to

topu/ Fuel!

10-31-50

al

TABLE1. WELL DEVELOPMENT FOR ARROW RENTAL, LIVERMORE, CA.

		INITAL	INITIAL	TOTAL AMOUNT	TOTAL	TOTAL AMOUNT	COMMENTS
WELL I.D.	DATES OF	WATER	TOTAL	OF WATER	TIME OF	OF WATER	
Ì	DEVELOPMENT	LEVEL	DEPTH	BAILED	SWABBING	DISCHARGED	
		(FEET)	(FEET)	(GALLONS)	(minutes)	(BAILED AND PUMPED)	
W-A	7/26 & 7/27	44.25	57.99	165	12	250	GASOLINE ODOR
W-B	7/27	44.62	55.12	100	0	100	BAILED DRY
w-c	7/27	43.34	50.3	30	0	30	
W-D	7/26	42.19	58.65	250	10	550	
W-E	7/27	43.08	58.61	35	0	35	CFICOKED WELL
EXISTING WELLS							
W-1		45.69					PURE PRODUCT
W-2		45.65					GASOLINE ODOR
W-3		45,13					GASOLINE ODOR

HEALTH AND SAFETY COMPLIANCE AGREEMENT

I, the undersigned, have received a copy of the health and safety plan for the project identified below. I have read the plan, understand it, and agree to comply with all of the health and safety requirements therein. I understand that I may be prohibited from continuing work on the project for failing to comply.

I have __ have not __ (check one) been briefed by a project safety authority on the health and safety requirements of the project.

Project No	9000321A-3000	
Project Title		
Date of Plan _		
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Sample No. W-A	W	ATEF	SAI	MPL	E LOG		Samp	lle No	. W-A
									26/90
VV-1 3= 73,01					GNTAL of Driv			OKE	
collected rust/orance gasoline from top	Sample Loc Well Descri	ption: 4	~h 40	ngru	_ QTAX I.I	rway.			
	· •	nditions:	lear	Sues	; 2100	n Shad	b , 4	VAR	
0960	Observation	s / Commen	is: No	Float	lers, ho	wever	stro	ris a	dor-collected
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W-A									
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		narge:				asing Volum			
·	Method of				lo label		erret	<u> </u>	<u> </u>
	Number as	nd size of sar	mple conta	lners filled	. 7 bernel	S			
						Man	harand		e Consultants
	Collected	by:	<u>.</u>						Dakland, CA 94607-4014

Sample No.	₹ W	ATER	SAN	IPLE	LOG		Sample	e No.	w-B
W-B	Project Name	i An	eron 1-B	K61 bel	uthe hind sh		jale:	7/20	6/90/7/27/
quit @1515 on W-B. Going to W.A again	Well Descript	ion:4	" 59 //_	1 80	rskuo atas, r		o <u>r:</u> 1	L6),	ample
	Quality			Method to	Measure Wate	r Level : _	<u> </u>	XIN	
	Method of cla	eaning Pump	(Qailer)	_St20 2309	can Chra	and_	Ca	Morated	7/27/90
	Commonte	7A = 54	1.62+.	30 =	55.12				ied-lined iling dry
	Sampli			Water Le		d Start:	44.61	L	End:
	7/23/10	Discharge (gallons)	pH /2	Temp. (°C)	Specific Conductance (µmhos / cm)	Turbidity	Color	Odor	Comments
	1351	* 30 70	7. % 7.05	20	1099	#16 A	720	No_	Tismldult
	1503	100	7.0	23		nSE	"		soml feith
	Fotal Discr	arge:	100) 	s bbls	asing Votur	nes Remo	wed:	
	Method of Number ar	disposal of o	discharged	l water:l	:_None	· [
	Collected	by: J (u 10	97011		W000	tward	-Clyd ute 100, (415) 80	le Consultants Opkland, CA 94607-4014 3-3600

Sample No.	3					LOG				w-c
8-8:30 POWNTIME - GET CLEVIS TO FIT DOWN Z" Well.			:Ar ion:Q ion:Z ditions:Q / Comments	opport	fo Sampling	foggy A	Reala Doi	l, just	k.	bailer
		Pump Lines:		Now 1	Cleaned	o Measure Wall	Bailer Lines	:	New	
		pH Meter No. Specific Cond	luctance Met	er No.: _	_ PO 1.	7/36		Ca	litivated alityrated	7/26/go Am
		Sampli	ng rements	3	Water Le	vel (below MP) ng Point (MP):	al Start:	(3.3º	4_ Lim)	End:
		Develop Three 1 27/90	Discharge (gallons)	ρН	Temp. (°C)	Specific Conductance (µmhos / cm)	Turbidity	Color	Odor	Comments
		0847	2 2.0	7.90	20	1250 (000 2 930	#16# ••	BRN '.		Thick
		917	4 Y.O	+.72	20	900		**		stating to peak
		i	arge: 3	lischarge		1 barnel	Casing Volu	mes Remo	wed:	
			y Jlee							le Consultants Oakland, CA 94607-4014 3-3600

Sample No.					LOG		_		W-E
Start Briling @ 1345 - Mike (drapped of parrels (10 of them) total # of new burners = 20 bacrels.	Project No. : Project Name: Sample Location Well Description Weather Cond Observations / Observations / Ouality	90Ce Am on: On in: 2" kilons: 2 Commenta iler da Assura	0321, ow fi dead (mid qc ii Wel wan	ental ental ental ental die of shad is e sell.	t - MAS Rd) no rooked Method: R	t to through	D+ O+ Out, med I	M A M A barc	- 1/21 - Dev to Shop. wto Shop. Wy get able Bailer from W-
	Method of clea	aning Pump	Baile	Cleaned Stead 0230° 13 30 ° 5	n Clean 157 1752 19.61	ailer Lines:	Ca	New /	2/056 7+10 red-limal
	Samplii Measui	ng rements	}	Water Le	g Point (MP):	slan:	/3.0 x (A	/_ E	ind:
	7/27/90	Discharge (gallons)	pH	Temp. (°C)	Specific Conductance (µmhos / cm)	Turbidity	Color	Odor	Comments
	16040	30	7.49 7.38 7.34	21	399 899 880	1,	**	"	
	1653	35	7.29	21	040				
	Total Disch Method of Number an	disposal of o	35 lischarge	d water: _	l barre , Nona	asing Volum	nes Remo		
		J(Woo (tward	-Clyd	e Consultants Dakland, CA 94607-4014

Sample No.	W	ATEF	R SA	MPL	E LOG		Samp	ie No	· ω-D
1515-1525-Swah					-3000		Date: _	71.	26/90
Prnp C/Sque : 75 - 25	Sample Location: BEHIND PINK HOUSE, IN It'S BACKYOTO Well Description: I Schooper Weather Conditions: Hot-clear skies Observations / Comments:							kyord	
	Quality Assurance Sampling Method: None Method to Measure Water Level: Solinst								
	Method of cle	eaning Pum	Bailer	Cleaner Stea 02309	um Cka 157	Bailer Lines	: c	New allorated	7/24/90 Rd-lined
	Sampli Measu		s	Water L	evel (below MP) :	nt Start:	/2,19 :(N	eren)	End:
	Time	Discharge (gallons)	рН	Temp. (°C)	Specific Conductance (µmhos / cm)	Turbidity	Color	Odor	Comments
		/30				HIGH	BAN	?	140 mJ si7+
	1458	300	738	21	800	11	**	**	50 m/ sitt
	1638		7.46	21	800	710U MED	(† 	e,	min
	1648	550	7.%	21_	800	5783	"		
			Stri					<u></u> j	
	Total Discha Method of d Number and	isposal of d	550 Ilscharged nple conta	f water:	To 11 b	sing Volum			okeral)
	Woodward-Clyde Cons Soo 12th Street, Suite 100, Clahland, CA 9 (415) 893-3600						lakland, CA 94607-4014		







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From (Your Name), Please Print CK LCC Company WOODHARD-CLYDE C Street Address 500 12TH ST STE	HS 8	74 2 C 79 Department/Floor No Compar Exact St	pient's Name) Please Print A PPING / KECEW IN CAMAN + KIN Intel® Address (No Camor Daliver to P.O. 28 - 12 10 h Ave	ing lipt.	ppent's Phone Number (Very Important) 206) 295 9272 Department/Floor No.
CITY OAKLAND YOUR INTERNAL BILLING REFERENCE INFORMA	State ZIP Required TION (First 24 characters will appear on invoice.)	U.Y. CIV	(111LL) IF HOLD FOR PICK-UP, Street	State WA Print FEDEX Address Hore	78119
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(Check only one box) Priority Overnight Standard Overnight Service (Delivery by mexi business invenight business invenight) 11 VOUR PRICKAGING 51 16 FEDEX LETTER * 56 FEDEX LETTER * 12 FEDEX PAK * 52 FEDEX PAK * 13 FEDEX BOX 53 FEDEX BOX	DELIVERY AND SPECIAL HANDLING 1 HOLD FOR PICK-UP: If a in Rick Mile 2 DELIVER SATURDAY (I this Challe) 3 CHILD ANABORNOUS GOODS (I this Challe) 4 CONSTANT SURFLANCE SVC (CSS) 5 CONSTANT SURFLANCE SVC (CSS) 6 DRY ICE This	MCMARS in house will see on se	Use of this eithli constitutes in our current Service Gu sender's copy of this arbit We will not be responsible package, whether the resmodeling or miscillariation and arbitrorial charge, and claim Maximum amount Express Service Guide at Express for my loss motils sales, income interest profit damage whether direct or control of the program of the p	MIT OF LIABILITY you agreement to the service con- de, available upon request. See bi- lor information in excess of \$10 for any claim in excess of \$10 it of loss, demage, delay, non-de- non unless you declars a higher with document your actual loss for a limitations found in the current if pay Your inglit to recover from Inding informs when of the peckage, afterney seets, costs, and other for idental consequential, or springals.	Base Charges seck of OO per servery. Other 1 editors of charge Other 2 Other 2 Other 2
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Table 1
Organic Vapor Concentrations Measured with an HNu Photoionization Detector

Boring Number	Depth <u>feet</u>	Maximum HNU Reading* (ppm-HNu Units
B-1	2 5 10 15 20 25	0 0 0 0 5.6 68.5
B-2	2 5 10 15 20 25	0 0 0 0 0 32.6
B-3	2 5 10 15 20 25	0 0 0 0 3.1 1.2
B-4	2 5 10 15	0 0 0 0
B-5	2 5 10 15 20 25	0 0 0 0 0 16.3
B-6	2 5 10 15 20 25	0 0 0 0

^{*} Relative hydrocarbon levels

Table 2

<u>Summary of Laboratory Analyses of Soil Samples</u>

Total Petroleum Hydrocarbons and BTEX (EPA 8015/8020)

Boring Number	Depth (feet)	TPH gasoline (ppm)	TPH diesel (ppm)	Benzene ppm	Toluene ppm	Ethyl- benzene ppm	xylene ppm
8-1	2	ND	NT	ND	ND	ND	ND
	5	ND	NT	ND	ND	ND	ND
	10	ND	NT	ND	ND	ND	ND
	15	ND	2.3	ND	ND	ND	ND
	20	170	NT	2.1	1.4	0.22	1.5
	25	220	NT	ND	2.4	2.1	11.0
B-2	2 5 10 15 25	3.5 8.2 ND ND 1.7	NT NT NT 2.3 NT	ND ND ND ND ND	ND ND ND ND ND	ND ND ND ND	0.1 ND ND ND 0.13
B-3	2 5 10 15 20 25	ND ND ND ND ND	NT NT NT 2.6 NT NT	ND ND ND ND ND	ND ND ND ND ND	ND ND ND ND ND	ND ND ND ND ND
B-4	2	ND	NT	ND	ND	ND	ND
	5	ND	NT	ND	ND	ND	ND
	10	ND	NT	ND	ND	ND	ND
	15	ND	NT	ND	ND	ND	ND
B-5	2 5 10 15 20 25	ND 1.9 ND ND ND 1.7	NT NT NT NT ND NT	ND ND ND ND ND	ND ND ND ND ND	ND ND ND ND ND	ND ND ND ND ND
B-6	5	1.8	NT	ND	ND	ND	ND
	10	ND	NT	ND	ND	ND	ND
	15	ND	NT	ND	ND	ND	ND
	20	ND	NT	ND	ND	ND	ND
	25	ND	NT	ND	ND	ND	ND

ND = Non-Detected NT = Not-Tested

Woodward-Clyde Consultants

Table 2 (concluded) Summary of Soil Samples Analytical Laboratory Results

Volatile Organics (EPA 8240)

Boring Number	Depth <u>(feet)</u>	Benzene µg/kg	toluene ug/kg	Ethyl- benzene <u>ug/kg</u>	xylene µg/kg
B-1	25	3 80	7,100	6,400	52,000
B-2	25	ND	ND	ND	550

Semi-Volatile Organics (EPA 8240)

Boring	Depth	Component	Concentration
Number	(feet)		ug/kg
B-1	25	2-Methylnaphthalene 2-Naphthalene phenol	3,500 3,400 300

Metals Analyses 1940 Railroad Ave.

Boring	Depth	Cr	Cu	Ni	Pb	Hg	Zn
Number	(feet)	mg/kg	mg/kg	mg/kg	mg/kg	mg/kg	mg/kg
B-6	2	42	34	100	5.8	0.035	36
	5	43	16	100	4.4	0.059	35
(TTLC)*		(2500)	(2500)	(2000)	(1000)	(20)	(5000)
(STLC)+		(560)	(25)	(20)	(5)	(0.2)	(250)

^{*} TTLC = Total Threshold Limit Concentration (CAC, Title 22) + STLC = Soluble Threshold Limit Concentration (CAC, Title 22)

Table 1. ELEVATIONS OF MEASURING POINTS AND ELEVATION OF GROUNDWATER, 187 NORTH L STREET, LIVERMORE, CALIFORNIA

Well Number	Measuring Point Elevation (Project Datum, feet)	Depth to Groundwater (feet)	Elevation (feet) June 2, 1989
W-1	99.22	43.16	56.06
W-2	99.07	44.24	54.83
W-3	98.03	44.50	53.53

Note: Assumed temporary benchmark elevation 100 feet.

Table 2. SUMMARY OF SOIL SAMPLE ANALYTICAL RESULTS FOR MONITORING WELLS W-1, W-2 AND W-3 AND BORINGS B-7 AND B-8, 187 NORTH L STREET, LIVERMORE, CALIFORNIA

Well/ Boring Number	Sample Depth	High Boiling Point Hydrocarbons (Diesel) (ppm)	Low Boiling Point Hydrocarbons (Gasoline) (ppm)	Benzene (ppm)	Toluene (ppm)	Ethylbenzene (ppm)	Xylenes (ppm)
——— W-1	5'	NR	ND	ND	ND	ND	ND
M_1	10'	NR	ND	ND	ND	ND	ND
	15'	NR	1,200	ND	21	20	130
	20'	380	350	2.5	14	6.3	30
	25'	NR	490	3.5	24	9.4	46
	30'	NR	160	1.0	7.9	3.6	18
	35'	NR	370	2.4	20	8.2	40
	40'	1,500	16,000	220	1,100	340	1,500
	45'	NR	1,600	30	120	34	160
	50'	NR	2,500	28	200	59	270
	55 ¹	NR	120	3.2	10	2.7	13
W-2	5'	NR	1.2	ND	0.14	ND	ND
H-C	10'	NR	ND	ND	0.1	ND	ND
	15 '	NR	ND	ND	0.1	ND	ND
	20'	NR	ND	ND	ND	ND	ND
	25'	NR	ND	ND	ND	ND	ND
	30'	NR	ND	ND	ND	ND	ND
	35 '	NR	ND	ND	ND	ND	ND
	40'	NR NR	ND	ND	ND	ND	ND.
	451	ND	ND	ND	ND	ND	ND
	50'	NR	ND	ND	ND	N D	ND
W-3	5'	NR	ND	ND	DM	ND	ND
M-2	10'	NR.	ND	ND	ND	ND	ND
	15'	NR NR	ND	ND	ND	ND	ND
	20'	NR	ND	ND	ND	ND	ND
	25'	NR	ND	ND	ND	ND	ND
	30'	NR	ND	ND	ND	ND	ND
	35'	NR	ND	ND	ND	ND	ND
	401	ND ND	ND	ND	ND	ND	ND
	45'	NR NR	ND	ND	ND	ND	ND
	50'	₩Ŕ	12	0.06	ND	ND	ND

8810220A-T CON-3

Table 2. SUMMARY OF SOIL SAMPLE ANALYTICAL RESULTS FOR WELLS W-1, W-2 AND W-3 AND BORINGS B-7 AND B-8, 187 NORTH L STREET, LIVERMORE, CALIFORNIA

Well/ Boring Number	Sample Depth	High Boiling Point Hydrocarbons (Diesel) (ppm)	Low Boiling Point Hydrocarbons (Gasoline) (ppm)	Benzene (ppm)	Toluene (ppm)	Ethylbenzene (ppm)	Xylenes (ppm)
B-7	5'	ND	ND	ND	ND	ND	ND
	10'	NR	ND	ND	ND	ND	ND
B- 8	5'	NR	ND	ND	DM	ND	ND
	10'	ND	ND	ND	DM	ND	ND
Detecti	on Limits:	1.0	1.0	0.05	0.1	0.1	0.1

ND = Not Detected

NR = Analysis Not Run

Table 3. SUMMARY OF GROUNDWATER SAMPLE ANALYTICAL RESULTS, NOVEMBER 1988, 187 NORTH L STREET, LIVERMORE, CALIFORNIA

		Micrograms	per Lite	r (µg/L)		
Well Number	High Boiling Point Hydrocarbon (Diesel)	Low Boiling Point Hydrocarbon (Gasoline)	ية Benzene	Toluene	Ethyl Benzene	Xylenes
W-1	300,000	210,000	29,000	30,000	5,400	24,000
W-2	ND	360	6.7	2.1	0.47	1.3
W-3	2,200	11,000	290	120	150	140
Detection Limits:	n 50.0	30.0	0.3	0.3	0.3	0.3
State or Drinkin Water I (MCLS)	ng		1.0	2,000	680	1750
State Drinkin Water Levels			0.7	100	680	620

ND = Not Detected

APPENDIX D TANK INTEGRITY TEST

D:\WPDOC\$\tag{90}00321.ALL\tag{55}

Data Chart for Tank System Tightness Test

PAINT	Annow	Jan Mal	187 100	+	Mariative	Ca.l.
VNER Property 🗹	Name		Address		adnistive	Telephone
· ·	Name		Address "SZT	_ v		1
ERATOR	Accident S	(Sert 37)	Address -			Telephone
	Annuel	Test				
ASON FOR				· · · · · · · · · · · · · · · · · · ·		
plain fuily)				3 - 4	· 0 - · +> 1	
- PEAUERTER	Trong Su	ui vs	Title	Company or	WENTS!	Date
HO REQUESTED ST AND WHEN	Name .					Talaghona
			Address Brand/Supplier	Grade	Approx. Age	Steel/fibergines
	Identify by Direction	Capacity	P. Hoods	Grec.	<u> इञ्चलक</u>	18teel
ANK INVOLVED	South-East	1,000				
aa addiijonai iln es						
r manifolded tanks						Pumps
		Covet	FILE	Yen ta	Sighones	Pumps
NSTALLATION	Location			1112"		Suche
ATA	L85	ا مدستاهام	<u>-</u>	1, 6	Mone	Sucilon, Remote.
	The latest	Cancrale, Black Top.	Size, Titellit miske, Drop tubes, Remote Fills	žirė, Manifolded	Which Works?	Make II known
·	Notify inside History Rear of station, etc.	Earth, etc.	(DBS), Remote 7 min		le the water over the tars	k?
INDERGROUND WATER	Depth to the Water table -	1804			□ Y# ☑ Nº	455-1900
ILL-UP	Terias to be fined			TOUY S	Namě	Telephone
ARRANGEMENTS	Eatra product to "top of!"	and the lank tester. How	and who to provide? Constit			
ARRANGEMENTS		and run lank reguer. How	and with it is seen.			Telephone
ARRANGEMENTS	Terminal or other content for notice or inquity	and run tank regter. From			Name	Telephoria
	Terminal or other contact				Name	Yelephone
CONTRACTOR.	Terminal or other contact				Name	Yelephoria
CONTRACTOR, MECHANICS, any other contractor	Terminal or other contact				Name	Yelophone
CONTRACTOR, MECHANICS, any other contractor	Terminal or other contact				Name	Yelephone
CONTRACTOR, MECHANICS, any other contractor involved	Terminal or other contact				Name	Yelophone
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27.	Sensor Calibration 16601/16600		PRE	STATIC SSURE		IME MEASUREMEN RECORD TO DDI GA		34. TEMPERATURE COMPENSATION USE FACTOR (a)			38, NET VOLUME CHANGING EACH READING	39. -CCUMUR ATED CHANGE	
	LOG OF TEST PROCEDURES		LOWING										
28. DATE	Record details of setting up and running test: (Use full	29. Reading	Standpipe in the	Nes.	32. Produ Grad		33, Product Replaced I-1	35. Thermal	36 Change Higher •	Computation (c) *(a) *	Temperature Adjustment Volume Minus	At Low Level compute	
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14. ARROW DENTAL 187	No. and Street D	Lucarone C		of Test
15. TANK TO TEST 15a. BR	EF DIAGRAM OF TANK FIELD	16. CAPACITY	From	:
South - East Lat		Nominal Capacity 1000 Gallons	Charte supplied with	
Bernell and Grade	Sheel	By most accomin especify chart melleble Gallo	Dow	Trad Gellera
17. FILL-UP FOR TEST			Gallors	na. Floreting
Stick Water Bences Solding Filt-up G 14" Gallons	York Observator	inedity.	FULL TOUR	+ 10
18. SPECIAL CONDITIONS AND PROCEDURES TO TEST THIS	TARK United in tent. United	a) being tested with LVLLT	water	<u> </u>
Sour determine applicable. Church below and record precedure to log (27).	High water table in tank	accivation	Total Preduct	1046
Cles maximum allowable test paralises for all tests. Four present rule dags riot apply to deutstreated tests.	19. TANK MEASUREMENTS TSTT ASSEMBLY	Cala	21. VAPOR RECOVERY SYSTEM	*
Complete arctice below: 1. In four pound min required? Yes No	Softers of tenk to probe any	30 to	24b. COEFFICIENT OF EXPANSION RECIPROCAL METHOD	_
2. Heinlight to 12" seems droven brofficien del trients.	20. EXTENSION HOSE SETT	14 m	Type of Product	
3. Processes at Section of Lank 2.75. P.94.	Extend hose on suction table 8" or store below lank top	٠	Temperature in Tank After Circulation Temperature of Sample	=1.4
4. Pressure at lap of tents 1.58 P.S.L.	"If Fit pipe axtends above grade, use to 22. Thermal-Sensor randing after cir-		Oliferation (+/-)	4 1,
Depth of build 45 in	23. Objics per "F fix range of suspects	Between	Observed APL Gravity	
Tank din. 455 in		ON (Complete after circulation)	Fold quantity in Reciprocal fail to the (16 or 17)	70 25565. Volume change in this task per *F
Weler trible	Observed A.P.L. Gravity	_	term amon file on 193	Yeansler to Linu 29s.
Mater Level	†êydroumter employed	=1.0	24c. FOR TESTING WITH WATE	R ass Table CAG
than a monitor well in the truk	Consected A.P I. Gravity	h:/0.	Vinter Temperature efter Circulation Table C	NA.
The above conditions are to be used for thy soil conditions to	Conflictant of Expendion for Implied Product From Table B	MA	Contricional of Walter Table 13	
establish a positive pressure advantage, or when using the four posses? Trute to compensate by the presence of subsurface wells; in the tank time.	Yransier COE to Line 25b.	<u></u>	Added Surfactant? The No Tru	
Sinter to HEFA. 30, Sections 2-3.24 and 2-7.2 and the tank in manufactorer regarding allowable system test presented.	25. (a) 104Ce Total quentity in full tresh (18 or 17)	Coefficient of duporation but		gallone

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APPENDIX E HISTORICAL EVIDENCE OF MOBIL SERVICE STATION

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EVIDENCE OF EXISTENCE OF MOBIL SERVICE STATION AT 187 NORTH "L" STREET IN LIVERMORE, CALIFORNIA CIRCA 1951-1968

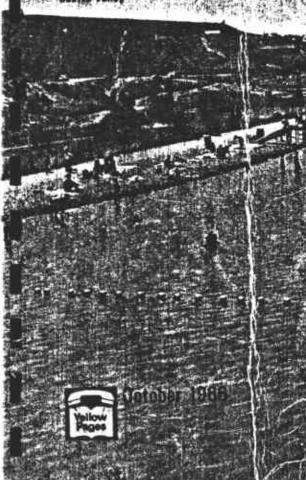
- 1. Pacific Telephone Directory, October 1966, Page 507
- 2. 1968 Livermore City Directory, Page 53
- 3. City of Livermore Building Permit to Socony-Mobile Co. For Underground Storage Tank, December 2, 1960
- 4. Mobil Oil Company, Ten Year Service Award to John Bowersox, 1961



SOUTHERN ALAMEDA County

Area Code 415

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1968
City
Directory

INCLUDES

- Alphabetical directory of business concerns and private citizens
- Complete classified business directory
- Street and Avenue guide:

 a list of householders and occupants of office buildings and businesses with telephones
- Reverse telephone directory: a list of telephone numbers in numerical order
- Rural information of the surrounding area.

AND

Miscellaneous information about the city and county

Published by

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162 H. 6th Aver, Eugene, Oregon 97401

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Goodwill Industries Inc Stark's Bargain House 2136 Railroad av 447-1973 4 Thriff Shop 1111 E Stanley Blvd. . in A 447~47030

SECRETARIAL SERVICE

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PECK'S SUBLAND SERVICE 160 ROLMES

PHILLIPS 66

Batteries

Tune-Ups 54

Complete Car Service

Free Pick Up & Delivery

PHONE 443-2944

2688-1st St.

OWNER

PHIL CORZINE

PECK SUNLAN SERVICE

Motor Tune-Up Expert

160 Hotmes

Rincon Richfield 899 Rincon Av. 447-01698 Sanchez Rick Chevron Station 1334-1st, 447-05480 Sexton's Phillips 66 Service Station 2680-1st Spencer's Gene Richfield Service 286 Livermore Av. / 447-98584 Valley Mobil Service 187 North Louis Vineyard 76 Union Oil Service Station 900 447-98070 AV respectively restricted his reserve Williams Ira Shell Service \$18 S Livermore

SEWER CONTRACTORS

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BAILEY, TOM-PLUMBING, HEATIN SHEET METAL 2888 RAILROAD

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Eracet No. 'L' Street	OWNER

24 HOURS NOTICE REQUIRED FOR INSPECTIONS 449-4007 between 9:00 a.m. - 4:30 p.m. BUILDING INSPECTION FECORD OF LIVERMORE Type of Fuel Storage BLOCK on TRACT: LOT: Building: Value: 1500. INSPECTOR DATE INSPECTIONS CONTRACTUR PERMITS DATE/NO. 12/18/80 YARDS & SETBACKS BUILDING 31206 FOUNDATION OWNER/Builder SUBFLOOR/Under flr. PLUMBING/SAS FRAME SLECTRICE. ELECTRICAL MECHANILIA PLUMBING MECHANICAL. SEWE-SAS TEST PG&E NUTTE SEWER ENCROACHMEN" INSULATION SHEETRUCK POOL PEL-IP WITE 200F POOL PRE-TIEN FIREPLACE/STOLE POOL PRE-PLASTER FINAL

North "L" St.

STREET :

NUMBER

OWNER:

Arrow Rentals

ohn owersox

We are pleased you have completed Ten Years as a Mobil Lealer

By your willingness to render professional services to the motorist, during this time, you have earned this special recognition for business achievement from your customers, from your community and from all of us at

Mobil Oil Company 51

1961

APPENDIX F DECLARATIONS OF MICHAEL COMER AND GARY PINKS

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DECLARATION OF GARY R. PINKS

My name is Gary R. Pinks.

I have personal knowledge of all the facts stated in this declaration and, if called as a witness, I could and would testify to them under oath.

I am the Manager at Arrow Rentals at 187 North "L" Street in Livermore, California. On or about June 18, 1985 I was so employed and on duty at Arrow Rentals when the Petcock Petroleum driver arrived to deliver an order of regular gasoline. Our usual order was for 500 gallons of regular grade gasoline.

The driver has to check in with me to get the key to the underground storage tank. On this day, it was a different driver from the driver who usually came from Petcock Petroleum. On this day, the Petcock driver was a shorter, older guy about 50 years old. I gave him the key to the 1000 gallon tank (as shown on attached diagram).

He was outside filling for what seemed like a long time. I stepped outside to see what was happening and saw the he had the nozzle from the gasoline truck in the vapor well rather than the mouth of the tank. I called his attention to the mistake he was making.

He pulled the nozzle out of the vapor well and put it back on the truck. He then immediately called the Petcock Petroleum office on his truck radio and said he was at Arrow Rentals and had just dropped 600 gallons of gas into the dirt. I was standing by his truck and heard his office advise him to get off the radio and to call from the phone inside. One of my employees, Robert Coleman, was also standing by when the radio conversation took place. The Petcock driver did make a phone call from inside Arrow Rentals and then he left.

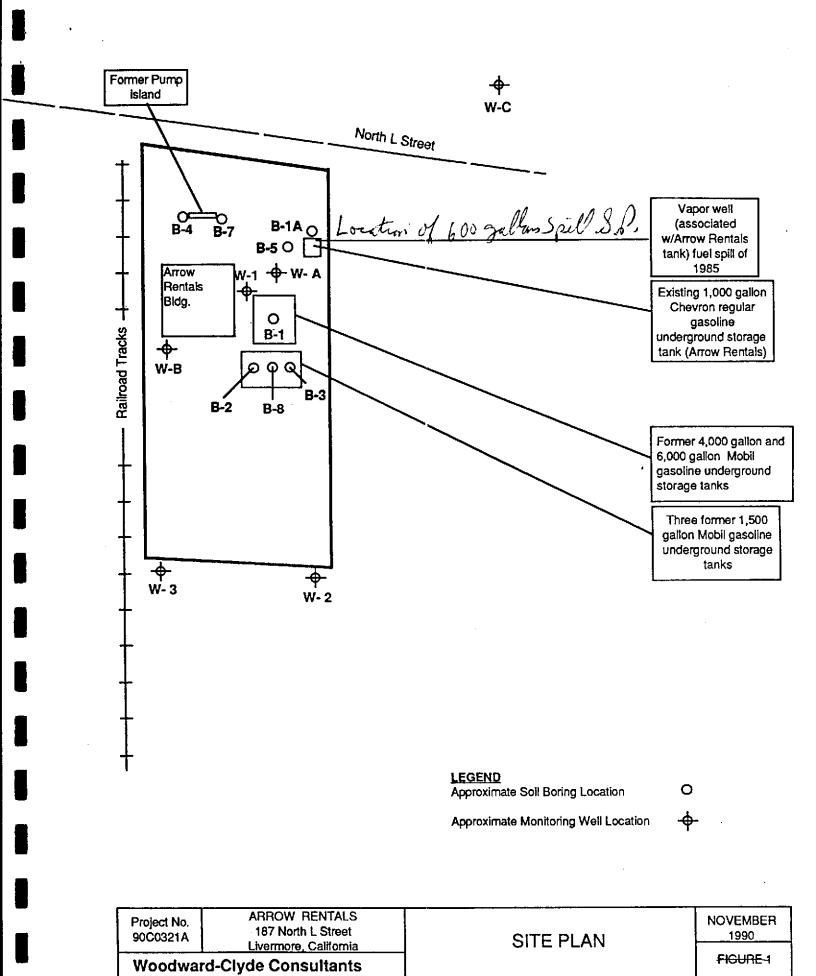
A day or so after the spill, the regular driver for Petcock showed up and delivered the gasoline we had originally ordered.

I declare under penany, ng statements are true and correct.

Dated this 12 day of February, 1991 at Luling.

Show the statement of the stateme I declare under penalty of perjury under the laws of the State of California that the foregoing statements are true and correct.

California.



DECLARATION OF MICHAEL P. COMER

My name is Michael P. Comer.

I have personal knowledge of all the facts stated in this declaration and if called as a witness, I could and would testify competently to them under oath.

I am the owner of Comer Petroleum Equipment. In my business, I frequently assist people in the removal of old underground storage tanks.

In the spring of 1985 I assisted Tony Sullins in the removal of two underground petroleum tanks (one approximately 4000 gallons and one approximately 6000 gallons) at Arrow Rentals in Livermore (as shown on attached map). When we were disconnecting plumbing going to the tank and the plumbing associated with the fill boxes, there was an odor of gasoline which is normal.

When we pulled the tanks out of the ground there was no noticeable odor of gasoline coming from the soils at the bottom of the tank pit. Nor did the soil in the tank pit have the dark discolored look which is characteristic of a tank which has been leaking underground. Based upon my experience with tank removal, I concluded that the two underground storage tanks which we removed from Arrow Rentals that day had not been leaking.

I declare under penalty of perjury under the laws of the State of California that the foregoing is true and correct.

Dated this 3/13/91 day of February, 1991 at Researches.

Michael P. Comer

