ENVINCYMENTAL PROTECTION

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#### QUARTERLY GROUNDWATER MONITORING REPORT First Quarter 1995

JOE SIO CHEVROLET 914-916 San Pablo Avenue Albany, California STID-3808

February 28, 1995

# Prepared for:

MS. FLORENCE ANN CONNORS
Executor for the Estate of Josephine A. Dibble
1658 Del Dayo Drive
Carmichael, California 95608

### Prepared by:

#### BURLINGTON ENVIRONMENTAL INC.

A Philip Environmental Company 5901 Christie Avenue, Suite 501 Emeryville, California 94608

SIO101/12104

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Ms. Florence Ann Connors Executor for the Estate of Josephine A. Dibble 1658 Del Dayo Drive Carmichael, California 95608

Subject: QUARTERLY GROUNDWATER MONITORING REPORT

First Quarter 1995 Joe Sio Chevrolet

914-916 San Pablo Avenue, Albany, California

Dear Ms. Connors:

Burlington Environmental Inc. (Burlington) is pleased to submit the following quarterly monitoring report for Joe Sio Chevrolet, located at 914-916 San Pablo Avenue in Albany, California (see Figure 1, Site Location Map). The groundwater monitoring and sampling was conducted by Burlington in January 1995.

#### **BACKGROUND**

Two 550-gallon underground storage tanks (USTs) were removed from the site on March 20, 1989 by Petroleum Engineering, Inc. One UST contained gasoline and was located under the sidewalk between the former building and San Pablo Avenue. The other UST contained waste oil and was located adjacent to the southwest corner of the former building (see Figure 2, Site Plan). Soil samples collected from beneath the former gasoline UST contained concentrations of total petroleum hydrocarbons (TPH) ranging between 270 and 1,300 milligrams per kilogram (mg/kg). As a result of the TPH in the soil samples from beneath the former gasoline UST, Alameda County Department of Environmental Health (ACDEH) requested that additional excavation be conducted in the vicinity of the former gasoline UST, and groundwater monitoring wells be installed and sampled to determine groundwater quality, flow direction, and gradient.

On July 24 and 25, 1991, Aqua Terra Technologies (ATT) of Walnut Creek, California, installed three groundwater monitoring wells (MW-1, MW-2, and MW-3) at the site (see Figure 2, Site Plan). The three groundwater monitoring wells were developed on July 31, 1991 and sampled on August 7, 1991. Elevated concentrations of TPH and benzene, toluene, ethylbenzene and total xylenes (BTEX) were found in the groundwater sample collected from well MW-1 (see Table 1). At the time the wells were sampled,

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Ms. Florence Ann Connors February 28, 1995 Page 2

ATT determined that the groundwater flow direction was to the west-northwest with an approximate hydraulic gradient of 0.01 feet/foot.

In a letter dated November 9, 1993, Ms. Juliet Shin (ACDEH) directed that quarterly groundwater monitoring be resumed at the site. In April 1994, Burlington received authority to proceed with quarterly groundwater monitoring at the site.

#### **MONITORING ACTIVITIES**

The first quarter 1995 monitoring event was conducted on January 17, 1995. In each well, the depth to groundwater and the presence or absence of phase-separated hydrocarbons (PSH) were determined. Groundwater samples were collected and analyzed according to U. S. Environmental Protection Agency (EPA) guidelines to determine the concentrations of TPH as gasoline (TPHg), BTEX, and total lead. In addition, groundwater from monitoring well MW-3 was analyzed for cadmium, chromium, zinc, and nickel. The groundwater sample from well MW-3 to be analyzed for the five metals was filtered during sampling with a 0.45 µm filter.

The ground water sample from well MW-2 was also analyzed for halogenated volatile organics in response to matrix interference observed during the 3rd and 4th quarter 1994 monitoring events, per the request of Ms. Juliet Shin (ACDEH) in her November 2, 1994 correspondence to you.

The monitoring and sampling procedures are presented in Appendix A. Field data sheets are presented in Appendix B. Western Environmental Science & Technology, located in Davis, California, performed the analysis. The analytical results and detection limits are presented in Table 1 (Groundwater Analytical Data)

#### RESULTS

The groundwater elevation in the monitoring wells beneath the site on January 17, 1995 ranged from 32.64 to 34.00 feet above mean sea level (see Table 2, Groundwater Elevation Data). A contour map of these data is presented in Figure 3 (Groundwater Elevation Contours). The approximate groundwater flow direction is to the east with an approximate hydraulic gradient of 0.007 feet/foot. The reversal in groundwater flow direction from previous monitoring events could be a result of the intensive rain that fell at the site during the sampling period. This flow direction reversal is believed to be temporary.

The results of the chemical analyses are presented in Table 1. No PSH were detected in any of the groundwater monitoring wells. No detectable concentrations of TPHg or BTEX were found in the groundwater samples collected from well MW-2 or MW-3. However, elevated concentrations of petroleum hydrocarbons were found in the

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Ms. Florence Ann Connors February 28, 1995 Page 3

groundwater sample collected from well MW-1, including 600 micrograms per liter (μg/l) of TPHg and 250 μg/l of benzene. The groundwater sample collected from well MW-3 contained trace levels of chromium and zinc (see Table 1). No detectable concentrations of lead were found.

Elevated concentrations of tetrachloroethene (PCE) were found in the groundwater sample from well MW-2, at 100 µg/l. In addition, trace concentrations of chloromethane, carbon tetrachloride, trichloroethene, and cis-1,2-dichloroethene were found in the sample. Although the PCE level exceeds the California maximum contaminant level (MCL) of 5 µg/l, there is no known source of PCE onsite, and an offsite source for the halogenated hydrocarbons is likely.

Chain-of-custody documentation and certified analytical results are presented in Appendix C. Purge and rinsate water was stored on the site in 55-gallon drums. The drums were labeled by the field sampling technician. Purge and rinsate water disposal will be arranged in the future.

#### CONCLUSIONS

One year of quarterly groundwater monitoring has been completed at this time. The recurrence of TPHg and BTEX in the groundwater samples collected from well MW-1 suggests that the groundwater below the former gasoline UST continues to be impacted. Burlington recommends that a Corrective Action Feasibility Study be prepared at this time to evaluate the most feasible corrective action for the site. Burlington further recommends that the quarterly groundwater monitoring be temporarily discontinued while the Feasibility Study is being prepared and reviewed.

Burlington appreciates the opportunity to provide you with quality consulting and environmental services. Please feel free to contact me if I can provide further assistance.

Sincerely,

BURLINGTON ENVIRONMENTAL INC.

David C. Tight, R.G. No. 4603

Investigation/Remediation Manager

Ms. Florence Ann Connors February 28, 1995 Page 4

#### Attachments:

Figure 1 - Site Location Map

Figure 2 - Site Plan

Figure 3 - Groundwater Elevation Contours

Table 1 - Groundwater Analytical Data

Table 1A- Halogenated/Volatile Organics Analytical Data

Table 2 - Groundwater Elevation Data

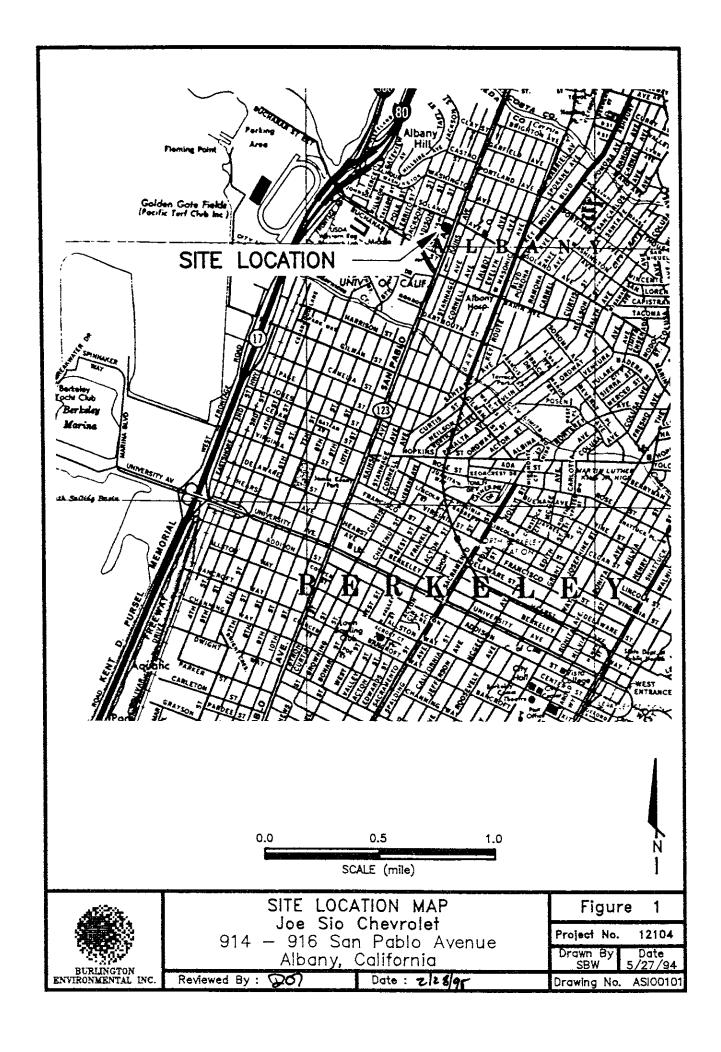
Appendix A - Groundwater Sampling and Analysis Procedures

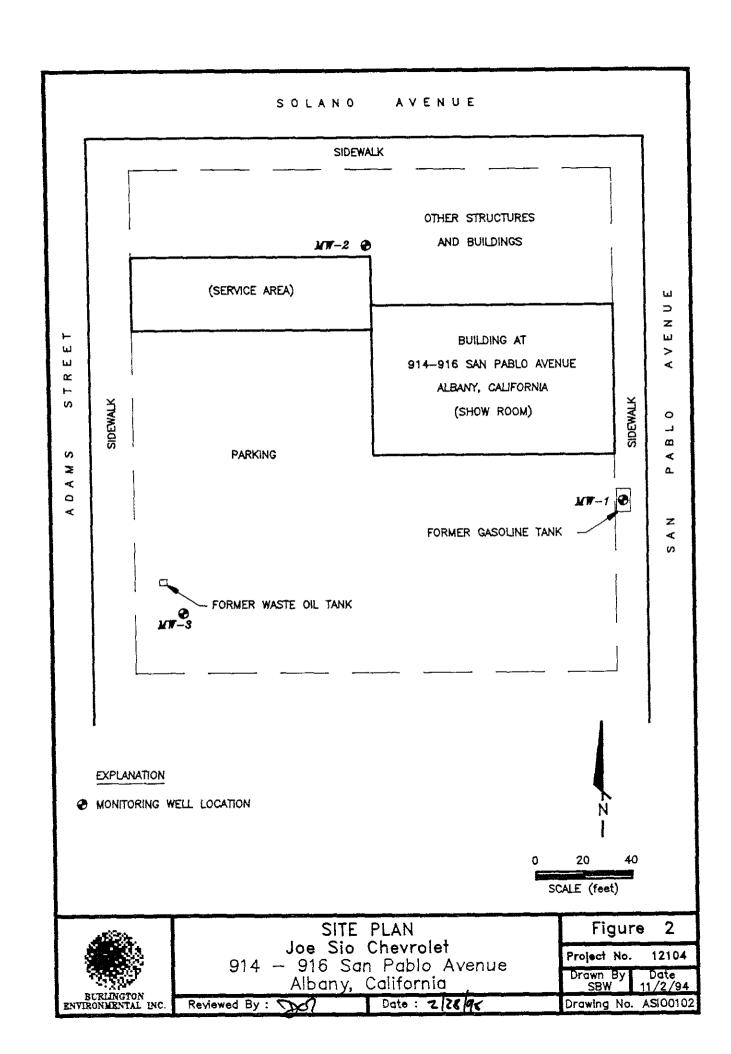
Appendix B - Water Sample Field Data Sheets

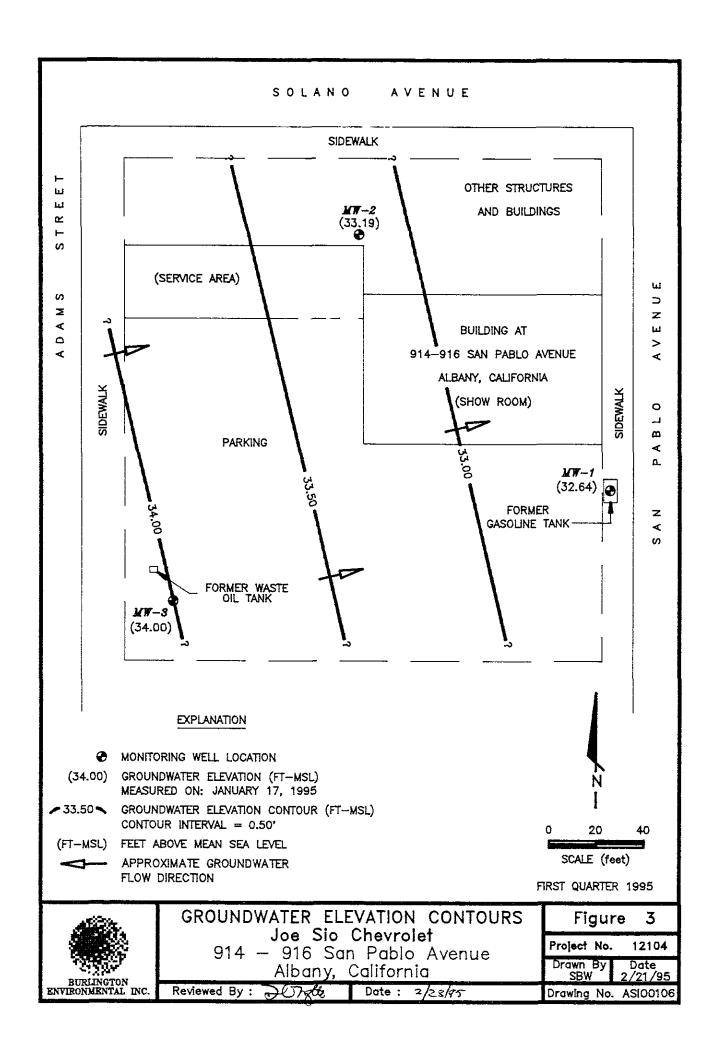
Appendix C - Chain-of-Custody Records and Certified Analytical Reports

cc: Ms. Juliet Shin (ACDEH)

FIGURES 1-3







TABLES 1 - 2

# TABLE 1 GROUNDWATER ANALYTICAL DATA

Joe Sio Chevrolet 914-916 San Pablo Avenue, Albany, California 50 PPB wester stel.

			TPH			Ethyl-	Total	Total Oll				*******	Zinc
	B	Sample	Gasoline	Benzene	Toluene	benzene	Xylenes	and Greese	Cadmlum	Chromium	Lend	Nickei (mg/i)	(mg/l)
Monitoring	Date	Sample No.	(ug/l)	(ug/l)	(ug/l)	(ug/l)	(ug/i)	(mg/l)	(mg/l)	(mg/l)	(mg/l)	AA	AA
Vell No.	Sampled	nalytical Method:	8015m	602	602	602	602	9070	AA	AA	AA		
	ETH M	talytical incinos.		<del></del>			_						
Groundwater A	inalvaes:							NA	NA	NA	NA	NA	NA
MW-1	8/7/91	MW-1	110	16	2	0.7	15		NA.	NA.	0.0093	NA	NA
(*1** - 1	4/16/94	MW01-041594	2,500	880	22	79	47	NA	NA NA	NA.	0.0059	NA	NA
	7/14/94	MW01-071494	470	110	22	21	87	NA	NA NA	NA.	0.008	NA	NA
	10/14/94	MW01-101494	380	86	17	24	77	NA		NA.	0.0096	NA	NA
	1/17/95	MW01-011795	600	250	11	5.3	66	NA	NA	WA	0.0000		
						ND4 +0 F01	ND(<0.50)	NA	NA	NA	NA	NA	NA
MW-2	8/7/91	WM-5	NA(<60)	ND(<0.50)	ND(<0.50)	ND(<0.50)	ND(<0.50)	NA NA	NA.	NA	0.022	NA	NA
	4/16/94	MW02-041494	ND(<50)	ND(<0.30)	ND(<0.30)	ND(<0.30)	0.71 *	NA NA	NA.	NA	0.023	NA	NA
	7/14/94	MW02-071494	ND(<50)	ND(<0.30) *	0.73 *	ND(<0.30) *		NA NA	NA.	NA	0.021	NA	NA
	10/14/94	MW02-101494	ND(<50) *	ND(<0.30) *	ND(<0.30) *	ND(<0.30) *	ND(<0.50) *	NA NA	NA.	NA	0.031	NA	NA
	1/17/95	MW02-011795	ND(<50) *	ND(<0.30) *	ND(<0.30) *	ND(<0.30) *	ND(<0.50) *	NA	147	1421			
						ND1 -0 501	ND(<0.50)	ND(<5)	NA	NA	NA	NA	NA
MW-3	8/7/91	WM-3	NA(<50)	NO(<0 50)	ND( < 0.50)	ND(<0.50)		NA NA	0.012	0.25	0.22	0.34	0.49
	4/15/94	MW03-041594	ND(< 50)	ND(<0.30)	ND(<0.30)	ND(<0.30)	ND(<0.50)	NA NA	NA	NA	NA	NA	NA
	4/15/94 d	DW01-041494	ND(<50)	ND(<0.30)	ND(<0.30)	ND(<0.30)	ND(<0.50)	NA.	0.017	0.55	0.22	0.73	0.84
	7/14/94	MW03-071494	ND(<50)	ND(<0.30)	ND(<0.30)	ND(<0.30)	0.50	NA NA	NA.	NA.	NA	NA	NA
	7/14/94 d	DW01-071494	ND(<50)	ND(<0.30)	ND(<0.30)	ND(<0.30)	0.53		0.019	0.64	0.14	0.86	0.90
	10/14/94	MW03-101494	ND(<50)	ND(<0.30)	ND(< 0.30)	ND(<0.30)	ND(<0.60)	NA	NA	NA.	NA	NA	NA
	10/14/94 d	DW01-101494	ND(< 50)	ND(<0.30)	ND(<0.30)	ND(<0.30)	ND(<0.50)	NA		0.0088	ND(<0.003)	ND(<0.015)	0.022
	1/17/95	MW03-011795	ND(<50)	ND(<0.30)	ND(<0.30)	ND(<0.30)	ND(<0.50)	NA	ND(<0.004)	NA	NA NA	NA.	NA
	1/17/95 d	DW03-011796	ND(< 50)	ND(<0.30)	ND(<0.30)	ND(<0 30)	ND(<0.60)	NA	NA	NA	140		
Binsate Analys					107 40 30)	ND(<0.30)	ND(<0.50)	NA	NA	NA	NA	NA	
	4/15/94	RS01-041594	ND(<60)	ND(<0.30)	ND(<0.30) 0.33	ND(<0.30)	0.65	NA	NA	NA	NA	NA	
	7/14/94	RS01-071494	ND{<50}	ND(<0.30)			ND(<0.50)	NA.	NA	NA	NA	NA	
	10/14/94	RS01-101494	ND(<50)	ND(<0.30)	ND(<0.30)	ND(<0.30)	ND(<0.50)	NA.	NA	NA	NA	NA	NA
	1/17/95	RS01-011795	ND(<50)	ND(<0.30)	ND(<0.30)	ND(<0.30)	ND( < 0.50)	140					
												NA	NA
Trip Blank Ans		TB01-041594	ND(<50)	ND(<0.30)	ND(<0.30)	ND(<0.30)	ND(<0.50)	NA	NA	NA	NA	NA NA	
	4/15/94	TB01-071494	ND(<50)	ND(<0.30)	ND(<0.30)	ND(<0.30)	ND(<0 50)	NA	NA	NA	NA		
	7/14/94	TB01-101494	ND(<50)	ND(<0.30)	ND(<0.30)	ND(<0.30)	ND(<0.50)	NA	NA	NA	NA	NA	
	10/14/94		ND(<50)	ND(<0.30)	ND(<0.30)	ND(<0.30)	ND(<0 50)	NA	NA	NA	NA	NA	NA
	1/17/95	TB01-011795	(4D) < 00)	MD( < 0.30)	115( < 0.00)	.,_,						<del> </del>	
DOINIVING MA	ATER STANDARD	os:								0.25	0.05		. 5
	ximum Contamina			1	-	680	1750	-	0.01	0.05	0.05	-	•

Results above detection limit are bolded for emphasis.

An external standard quantitation was used on this sample due to matrix interference

Analyte found in method blank

d Duplicate sample

mg/l Milligrams per liter (parts per million)

NA Not analy≥ed

ND Concentration below detection limit presented in parentheses

ug/l Micrograms per liter (parts per billion)

#### TABLE 1A HALOGENATED/VOLATILE ORGANICS ANALYTICAL DATA

#### Joe Slo Chevrolet 914-916 San Pablo Avenue, Albany, California

Monitoring Well No.	Date Sampled EPA An	Sample No. alytical Method:	Chloro- methene 601	Carbon Tetra- chloride 601	Trichloro- ethene 601	cls-1,2- Dichloro ethene 601	Tetrachloro- ethene 601
MW-2	1/17/95	MW02-011795	0.94	0.98	0.58	0.51	100
	TER STANDAR		-	0.5	5	6	5

Results above detection limit are bolded for emphasis.

All results presented in micrograms per liter (ug/l)

601 analytes not listed are all below method detection limits

# TABLE 2 GROUNDWATER ELEVATION DATA

Joe Sio Chevrolet 914-916 San Pablo Avenue, Albany, California

		Total	TOC	Depth to	Water
Monitoring	Date	Depth	Elevation	Water	Elevation
Well No.	Measured	(ft-BTOC)	(ft-MSL)	(ft-BTOC)	(ft-MSL)
			40.04	10.40	32.12
MW-1	8/7/91	NM	42.61	10.49	
	8/12/91	NM	42.61	10.37	32.24
	4/15/94	29.80	42.61	10.60	32.01
	7/14/94	29.70	42.61	10.55	32.06
	10/14/94	29.75	42.61	10.88	31.73
	1/17/95	29.75	42.61	9.97	32.64
MW-2	8/7/91	NM	42.73	11.64	31.09
	8/12/91	NM	42.73	11.69	31.04
	4/15/94	26.88	42.73	10.16	32.57
	7/14/94	26.85	42.73	10.91	31.82
	10/14/94	26.88	42.73	12.10	30.63
	1/17/95	26.87	42.73	9.54	33.19
MW-3	8/7/91	NM	39,44	8.94	30.50
	8/12/91	NM	39.44	8.94	30.50
	4/15/94	25.58	39.44	7.68	31.76
	7/14/94	25.62	39.44	8.40	31.04
	10/14/94	25.61	39.44	9.31	30.13
	1/17/95	25.79	39.44	5.44	34.00

Water levels measured on 8/7/91 and 8/12/91 by Aqua Terra Technologies (ATT) of Walnut Creek, California.

TOC elevations obtained from survey data provided in the ATT Groundwater Monitoring Report dated 11/11/91.

ft-BTOC Feet below top of casing ft-MSL Feet above mean sea level

NM Not measured TOC Top of casing

### APPENDIX A

Groundwater Sampling and Analysis Procedures



#### APPENDIX A

#### Groundwater Sampling and Analysis Procedures

#### INTRODUCTION

The sampling and analysis procedures for water-quality monitoring programs are contained in this Appendix. These procedures ensure that consistent and reproducible sampling methods are used, proper analytical methods are applied, analytical results are accurate, precise, and complete, and the overall objectives of the monitoring program are achieved.

#### SAMPLE COLLECTION

Sample collection procedures include equipment cleaning, water-level and total well-depth measurements, and well purging and sampling.

# **Equipment Cleaning**

Sample bottles, caps, and septa were precleaned and provided by a DHS-certified laboratory. All sampling containers were used only once and discarded after analysis was complete.

Before starting the sampling event, all equipment to be placed in the well or come in contact with groundwater was disassembled and cleaned thoroughly with detergent water, then steam cleaned with tap water, and rinsed with distilled water. Any parts that may absorb contaminants, such as plastic pump valves or bladders, were cleaned as described above or replaced.

During the sampling event all equipment used in the well was washed with detergent, steam-cleaned, and rinsed with distilled water before purging or sampling the next well. The rinsate water was contained for temporary storage in 55-gallon drums and disposal

will be arranged by the client. The 55-gallon drums were stored onsite and labeled by the field technician.

### Quality Assurance Samples

A trip blank was analyzed to insure contamination did not result from travel exposure.

# WATER-LEVEL, FLOATING-HYDROCARBON, AND TOTAL WELL-DEPTH MEASUREMENTS

Before purging and sampling, the depth to water, floating hydrocarbon thickness, and the well total depth were measured using an oil water interface probe and an electric sounder. The electric sounder, manufactured by Slope-Indicator, Inc., is a transistorized instrument that uses a reel-mounted, two conductor, coaxial cable that connects the control panel to the sensor. Cable markings are stamped at 1-foot intervals. An engineers rule was used to measure the depths to the closest 0.01 foot. The water level was measured by lowering the sensor into the monitoring well. A low current circuit is completed when the sensor contacts the water, which serves as a conductor. The current is amplified and fed across an indicator light and audible buzzer, signaling when water has been contacted. A sensitivity control compensates for very saline or conductive water. The oil water interface probe signals with a solid sound when it contacts phase-separated hydrocarbons. When the probe detects water, the sound changes to a beeping sound.

No phase-separated hydrocarbons were detected in any of the monitoring wells. When PSH is detected at greater than 1/32-inch in thickness, a sample is not collected.

All liquid measurements were recorded to the nearest 0.01 foot in the field logbook. The groundwater elevation at each monitoring well was calculated by subtracting the measured depth to water from the surveyed well-casing elevation. Well total depth was then measured by lowering the sensor to the bottom of the well. Well total depth, used to calculate purge volumes and to determine whether the well screen is partially obstructed by silt, was recorded to the nearest 0.01 foot in the field log book.



#### WELL PURGING

Before sampling, standing water in the casing was purged from the monitoring wells using a PVC hand bailer. Samples were collected from the monitoring wells after a minimum of four casing volumes had been evacuated or the pH, electrical conductivity, and temperature had stabilized. In the case that the monitoring well was purged until dry, the well was allowed to recover to within 80% of its static water level and sampled.

The pH, electrical conductivity, and temperature meter were calibrated each day before beginning field activities. After every well volume of groundwater removed from the monitoring well, field measurements were taken. The data is presented on the water sample field data sheets. The calibration was checked once each day to verify meter performance. All field meter calibrations were recorded in the field log book.

Groundwater generated from well-purging operations were contained for temporary storage in 55-gallon drums. All drums were labeled and stored onsite. The sampler recorded on the drum label for each drum generated:

- drum content (i.e., groundwater)
- source (i.e., well identification code)
- date generated
- client contact
- project number
- name of sampler.

The purge water will be disposed of by the client.



#### WELL SAMPLING

A Teflon bailer was used for well sampling. Glass bottles of at least 40 milliliters volume and fitted with Teflon-lined septa were used in sampling for volatile organics. These bottles were filled completely to prevent air from remaining in the bottle. A positive meniscus forms when the bottle is completely full. A convex Teflon septum is placed over the meniscus to eliminate air. After capping, the bottle was inverted and tapped to verify that it did not contain air bubbles. The sample containers for other parameters were filled, and capped.

## SAMPLE HANDLING AND DOCUMENTATION

The following section specifies the procedures and documentation used during sample handling.

#### Sample Handling

All sample containers were labeled immediately following sample collection. Samples were kept cool with ice cubes until received by the laboratory. At the time of sampling, each sample was logged on a chain-of-custody record which accompanied the sample to the Western Environmental, Science, and Technology.

### Sample Documentation

The following procedures were used during sampling and analysis to provide chain-of-custody control during sample handling from collection through storage. Sample documentation included:

- field log books to document sampling activities in the field
- · labels to identify individual samples; and

 chain-of-custody record sheets for documenting possession and transfer of samples.

#### Field Log Book

In the field, the sampler recorded on the Water Sample Field Data Sheet for each sample collected:

- project number
- client name
- location
- name of sampler
- · date and time
- pertinent well data (e.g., casing diameter, depth to water, well depth
- calculated and actual purge volumes
- · purging equipment used
- sampling equipment used
- appearance of each sample (e.g., color, turbidity, sediment)
- results of field analyses (i.e., temperature, pH, electrical conductivity)
- general comments

The field logbooks were signed by the sampler.



#### Labels

Sample labels contained:

- project number
- sample number (i.e., well designation)
- sampler's initials
- · date and time of collection
- type of preservative used (if any)

# Sampling and Analysis Chain-of-Custody Record

The Sampling and Analysis Chain-of-Custody record, initiated at the time of sampling, contains, but is not limited to, the well number, sample type, analytical request, date of sampling, and the name of the sampler. The record sheet was signed, timed, and dated by the sampler when transferring the samples. The number of custodians in the chain of possessions were kept to a minimum. A copy of the Sampling and Analysis Chain-of-Custody record is included in Appendix C.

# APPENDIX B Water Sample Field Data Sheets

#### FIELD REPORT WATER LEVEL / FLOATING PRODUCT SURVEY

PROJECT NO.: 12104

LOCATION: 914 SAN PABLO ANE, ALBANY

SAMPLER: QLATAB TIME AND DATE OF SYSTEM START-UP: NA

ILLE SIA CHENKOLET

DATE: 1-194

STATION NO.:	be sid chevi	roce i	DATE: J.T	7.75		<del></del>	-70
WELL ID	YOTAL DEPTH (Feet)	WELL DIAMETER (fn)	DEPTH TO WATER (Feet)	DEPTH TO FLOATING PRODUCT (Feet)	FLOATING PRODUCT THICKNESS (Feet)	TIME	COMMENTS
mw-1	29.75	2	9.97		-	0925	
m.+Z	26.87	2	9.54	_		0909	
MW-1 MW-2 MW-3	25.79	2.	5.44		-	0902	
							· · · · · · · · · · · · · · · · · · ·

# WATER DATA SHEET

PROJECT NO.: 12104	SAMPLE ID .: MWOI · OII 7	193
LOCATION: 914 SAN PABLO AUF., ALBANY	DATE: 1.17.95	
STATION NO.: 58-101	WELL/SAMPLE	
SAMPLER: D.LAMB	POINT DESIGNATION: MW	-1
SAMPLING DEVELOPING	BAILING FLOATING P	RODUCT
Casing Diameter: Screened Int. (ft.): 18		vi. (gal.): 3.36
2 inch X 3 inch Initial DTW (ft.):9.	97 @ 092.5 Calc. Purge Vol	(gal.): 13.45
4 inch Initial TD (ft.): 29:75	Final DTW (ft.):	13.84 @ 0948
other Water Column Height	(ft.): 19.78 Final TD (ft.):_	29.76
TD (Actual) (ft.): 30 80 % Recovery (ft.):_	13.93 Product Balled	(gal.):
FIEL	D MEASUREMENTS	
691.) (units) (dog 0933 3.5 6.92 6 0938 7.0 6.57 6 0942 10.5 6.48 6	EC. (umhos/cm) 6.9 6.4 x10 2 6.13 6.10 x 10 2 2.4 6.23 x 10 2 2.9 6.26 x 18 2	COLOR DTW (Fldry)  GEN / BEN  YEllow / BEN- VELLOW) BEN
9147 13.5 6.45 6		
Odor? SliGHT		
Actual Purge Vol. (gal.): 13.5		
PURGE METHOD:	SAMPLE METHOD:	
Bailer (Tefion)	۷	∠ Bailer (Teflon)
Baller (PVC)		Bailer (PVC) Dedicated Bailer
Well Wizard Dedicated Baller		Other
Other	<del></del>	
REMARKS: MWS1.011795 SAMPLES	0 2020 ON 1.	17.95.
WEATHER: SUNNY Gol, & 50.		

# WATER DATA SHEET

PROJECT NO.: 12104	SAMPLE IO.:	MW02. 0	11795	
LOCATION: 914 SAN PAND ANE., ALBANY	DATE:	1.17.93	5	
STATION NO.: 50.181	WELL/SAMPL			
SAMPLER: D.LAMB	-	NATION: M	1W-Z	
SAMPLING DEVELOPING	BAIUI	NG FLOATING	PRODUCT	
Casing Diameter: Screened Int. (ft.): 8 -	28_		/ol. (gal.): 2.95	-
3 inch Initial DTW (ft.): 9.54	<u>e 0909</u>	Calc. Purge Vo	(4°65) (6° - 1.5) ol. (gal.):	
4 Inch			9.48 @	1045
Casing Elev. (ft.): Water Column Height (ft	u): <u>17.33</u>	Final TD (ft.):_	26.88	
TD (Actual) (ft.): 28 80 % Recovery (ft.): 1	3.01	Product Balled		
FIELD	MEASUREMEN	ITS		
TIME VOLUME PH TEM	IP.	E.C.	COLOR	0774
1032 3 7.13 56. 1037 6 7.02 58	9 6	mhos/om) .02 x/0 <sup>2</sup>	YF/kir/BPN.	DTW (I day)
1041 9 6.94 60.		$\frac{.20 \text{ x/0}^2}{.53 \text{ x/0}^2}$	XEllow BLW.	
1044 12 6.91 60.	A	51 x102	yellow 18W.	
				<del></del>
Odor? NaNE				<del></del>
Actual Purge Vol. (gal.): 1Z				
PURGE METHOD:	SAME	PLE METHOD:		
Baller (Teflon)Baller (PVC)	<b>.</b>		(Bailer (Teffon)	
Well Wizard			Baller (PVC) Dedicated Baller	
Dedicated Bailer Other			Other	
REMARKS: MWOZ.01795 SAMPLES	<i>a</i>		1	
- Sitts shipes	e 1100	<u> </u>	1.17.9.5	<del></del>
WEATHER: SWAY, CLEP, 250.				

# WATER DATA SHEET

PROJECT NO.:	12184		SAMPL		10 1 · 011 103 · 011	795 () 1795	opeican	<b>E</b>
LOCATION: 9	14 San Abd	OANE, ALBA			7.95			
STATION NO.:	ठांठ १०।	<b>-</b>	WELL/	SAMPLE				
SAMPLER:	D.LAMB	_	POINT	DESIGNAT	10N: <u>/1</u>	W-3	<del></del>	
X SAMPL	ING	DEVELOPIA	ig 🗀	BAILING	FLOATING F	PRODUCT		
Casing Diameter			L): 7-27		17) 57 = .3	ol. (gal.): 3.4 si (4° = .65) (6	1.5	
3 inch 4 inch	-	Initial DTW (ft.):	5.44 @	10% C	sic. Purge Vo	il. (gal.): <u>13-9</u>	<u> 4</u>	
6 Inch	<del>-</del>	Initial TD (ft.):	25.79	_ FI	nal DTW (ft.):	<u>5.5</u> 2	21	129
Casing Elev. (fi	L):	Water Column	Height (ft.): <u>20</u> .	<i>35</i> fi	nal TD (ft.):_	25.78	<del></del>	
TO (Actual) (ft.)	: 27	80 % Recovery	(ft.): <u>9.51</u>	Pr	oduct Balled	(gal.): g	<del></del>	
			FIELD MEASU	rement <b>s</b>				
TIME 1113	VOLUME	pH (units) 6.84	TEMP. (degrees F) 57.4	E. (umb: 2.14	.C. ≈/an) ~ √ 10	COLOR	_	DTW @doy)
1119	7.0	7.74	59.9	2.2	¥10°	YEIN!	BEN.	
<u> 1124</u> 1128	14.0	7.62	612	2.12 2.16	- 40 <sup>2</sup>	VEllaw/	-	, <del></del>
					<u> Ale</u>	Y CHOW /	prov.	
Odor?	NONE							
Actual Purge	vol. (gal.): 14	<u>.                                    </u>						
PURGE METH	iOD:			SAMPI F	E METHOO:			
	Baller (Teflon)	•		•		XBaller (Tello	n)	
<u>ـــ</u>	Baller (PVC)					Baler (PVC)		
	_Well Wizard Dedicated Ba	1~				Dedicated 8	afer	
	Other				_	_Other		-
REMARKS:	MW03.	011795	SAME	<b>e</b>	1145	ON	1.17	95
	DW01.	811795	Francis .	a	1210	3N	1.77	75
<del></del>								
						<del></del>		
WEATHER:	Sury,	900 1, 0	= 510.					

## APPENDIX C

Chain-of-Custody Records and Certified Analytical Data



Sample Log 11124

Sample: MM01-011795

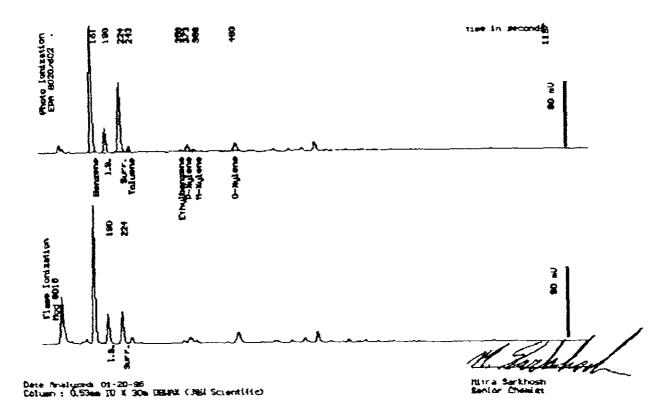
Prom : Project # 12104 (Joe Sio Chevrolet)

Sampled: 01/17/95

Dilution: 1:3 QC Batch: 4111M

Matrix : Water

Parameter	(MRL) wg/L	Measured Value was
Benzene	(.75)	250
Toluene	(.75)	11
Ethylbenzene	(.75)	5.3
Total Xylenes	(1.3)	56
TPH as Gasoline	(130)	600
Surrogate Recover	Y	98 %





Sample Log 11124 11124~4

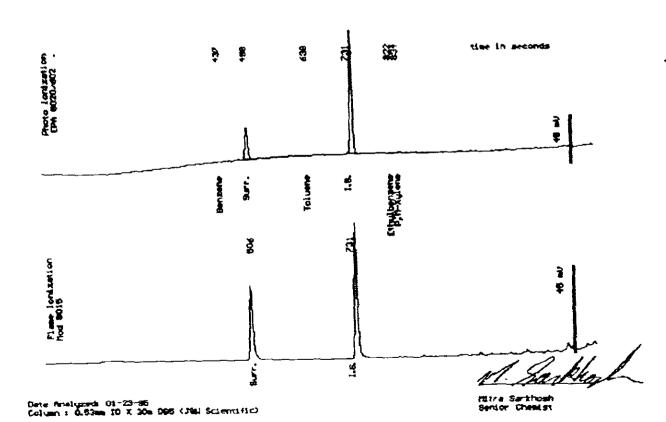
Sample: NW02-011795

From : Project # 12104 (Joe Sio Chevrolet)

Sampled: 01/17/95 Dilution: 1:1 QC Batch : 6138K

Matrix : Water

Parameter	(MRL) was	Measured Value w/L
Benzene Toluene Ethylbenzene Total Xylenes TPH as Gasoline	(.30) (.30) (.30) (.50) (50)	<.30 <.30 <.30 <.50 <50
Surrogate Recover * External standa interference.	y rdization was used d	101 * % ue to matrix





January 24, 1995 Sample Log 11124

Sample: HW02-011795

From : Project # 12104 (Joe Sio Chevrolet)

 Sampled : 01/17/95
 Received : 01/17/95

 Matrix : Water
 Analyzed : 01/23/95

#### 601 - Balogenated Volatile Organics

out management to the comment		Heasured	
Parameter	(MRL) wg/L	Value w/c	Flag
Chloromethane	(0.50)	.94	
Chloroethane	(1.0)	< 1.0	
Vinyl Chloride	(0.50)	<0.50	
Bromomethane	(1.0)	< 1.0	
Trichlorofluoromethane	(0.50)	<0.50	
1,1-Dichloroethene	(0.50)	<0.50	
Dichloromethane	(0.50)	<0.50	
t-1,2-Dichloroethene	(0.50)	<0.50	
1,1-Dichloroethane	(0.50)	<0.50	
Chloroform	(0.50)	<0.50	
1,1,1-Trichloroethane	(0.50)	<0.50	
1,2-Dichloroethane	(0.50)	<0.50	
Carbon Tetrachloride	(0.50)	.98	
1,2-Dichloropropane	(0.50)	<0.50	
Trichloroethene	(0.50)	.58	
Bromodichloromethane	(0.50)	<0.50	
c-1,2-Dichloroethene	(0.50)	.51	
c-1,3-Dichloropropene	(0.50)	<0.50	
t-1,3-Dichloropropene	(0.50)	<0.50	
1,1,2-Trichloroethane	(0.50)	<0.50	
Tetrachloroethene	(0.50)	100	
Dibromochloromethane	(0.50)	<0.50	
Chlorobenzene	(0.50)	<0.50	
Bromoform	(0.50)	<0.50	
1,1,2,2-Tetrachloroethane	(0.50)	<0.50	
1,4-Dichlorobenzene	(0.50)	<0.50	
1,3-Dichlorobenzene	(0.50)	<0.50	
1,2-Dichlorobenzene	(0.50)	<0.50	
2-Chlorotoluene (Surrogate)		93 %	

Joseph Life Search Constat



January 20, 1995 Sample Log 11124

From: Project # 12104 (Joe Sio Chervolet)

Date Sampled: 01/17/95

Matrix: Water

Date Received 01/17/95

Units: (mg/L)

# Total Lead by GFAA by SW-846 Method 7421

WEST ID	Sample ID	Result	MRL	Date Digested	Date Analyzed
11124-3	MW01-611795	0.0096	0.003	01/19/95	01/20/95
11124-4	MYY9Z-011795	0.931	0.003	01/19/95	01/20/95

MRL = Method Reporting Limit

Michelle L. Anderson inorganics Supervisor Sample: MM03-011795

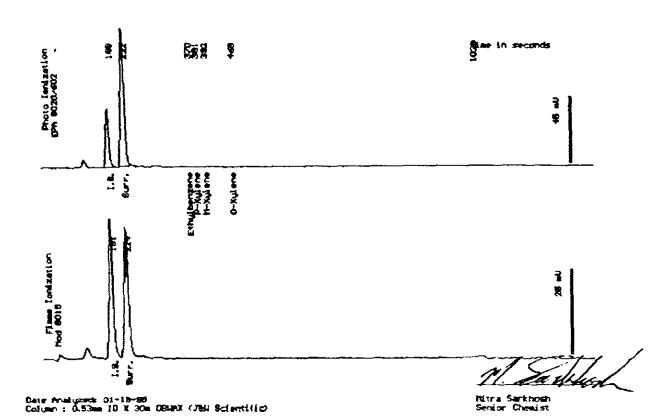
Prom : Project # 12104 (Joe Sio Chevrolet)

Sampled: 01/17/95

Dilution: 1:1 QC Batch: 2112c

Matrix : Water

Parameter	(MRL) was	Weasured Value w/L
Benzene	(.30)	<.30
Toluene	(.30)	<.30
Ethylbenzene	(.30)	<.30
Total Xylenes	(.50)	<.50
TPH as Gasoline	(50)	<50
Surrogate Recover	Y	108





Sample Log 11124 13124-6

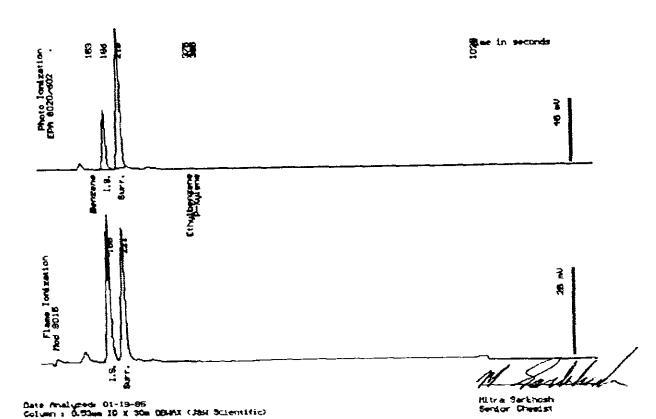
Sample: DW01-011795

From : Project # 12104 (Joe Sio Chevrolet)
Sampled : 01/17/95

QC Batch : 2112c Dilution: 1:1

Matrix : Water

Parameter	(MRL) was/L	Measured Value wa
Benzene Toluene Ethylbenzene	(.30) (.30) (.30)	<.30 <.30 <.30 <.50
Total Xylenes TPH as Gasoline Surrogate Recover	(.50) (50) Y	<50 107 <b>\$</b>





January 24, 1995 Sample Log 11124-5

Sample: MW03-911795

From Project # 12104 (Joe Sio Chevrolet)

Matrix: Water

Date Sampled: 01/17/95 Date Received: 01/17/95

Units (mg/L)

# Dissolved Metals Analyses by ICP and GFAA by SW-844 6 LUFT: "Waste Oil" Notals

Analyte	Result	MRL	EPA Method	Date Digested	Date Analyzed
Cadmium (Cd)	<0.004	0 004	6010	01/23/95	01/24/95
Chromium (Cr)	0.0086	0.007	6010	01/23/95	01/24/95
Lead (Pb)	<0.003	0.003	7421	01/19/95	01/20/95
Nickel (Ni)	<0.015	0.015	6010	01/23/95	01/24/95
Zinc (Zn)	0.022	0.010	6010	01/23/95	01/24/95

MRL = Method Reporting Limit

Michelle L. Anderson Inorganics Supervisor



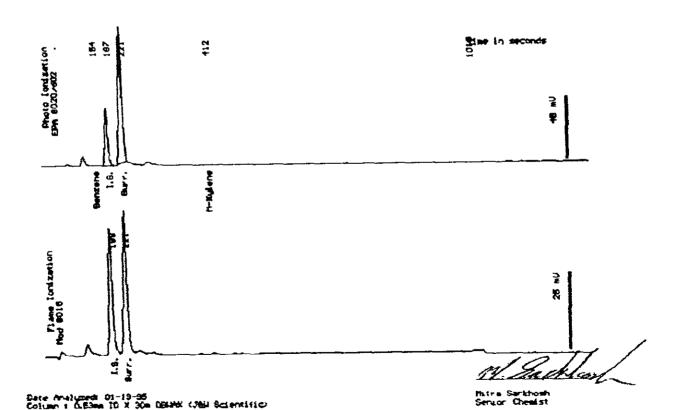
Sample Log 11124 11124-2

Sample: RS01-011795

From: Project # 12104 (Joe Sio Chevrolet)
Sampled: 01/17/95
Dilution: 1:1 QC Batch: 21 QC Batch : 2112c

Matrix : Water

Parameter	(MRL) vg/L	Measured Value w/L
Benzene	(.30)	<.30
Toluène	(.30) (.30)	<.30
Ethylbenzene	(.30)	<.30
Total Xylenes	(.50)	<.50
TPH as Gasoline	(50)	<50
Surrogate Recover	y	105





Sample Log 11124

Sample: TB01-011795

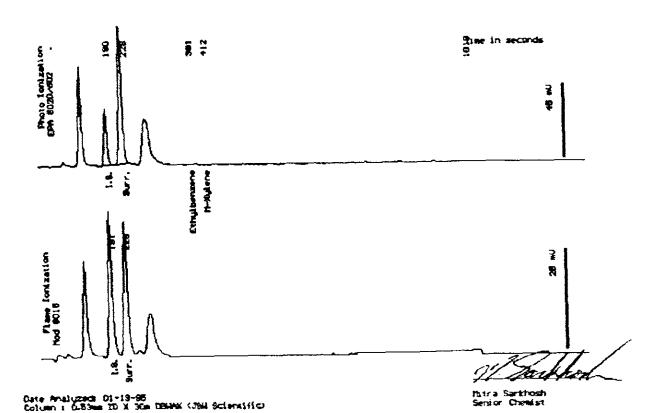
From : Project # 12104 (Joe Sio Chevrolet)

Sampled: 01/17/95

Dilution: 1:1 QC Batch: 2112c

Matrix : Water

Toluene Ethylbenzene Total Xylenes	(MRL) was	Measured Value w/L
Benzene Toluene Ethylbenzene Total Xylenes TPH as Gasoline	(.30) (.30) (.30) (.50) (50)	<.30 <.30 <.30 <.50 <50
Surrogate Recover	y	112 %





January 20, 1995

### Metals QC Report for Sample Log 11124

From . Project # 12104 (Joe Sio Chervolet)

Matrix: Water

Sample Spiked for MS/MSD: 11124-5

Units: (mg/L)

#### Method Blank

Anatyse	Result	MRL	EPA Method	Date Digested	Date Analyzed
Lead (Pb)	<0.003	0.003	7421	01/19/95	01/20/95

MRL = Method Reporting Limit

Laboratory Control Sample (LCS)

Analyte	% Recovery	EPA Method	Date Digested	Date Analyzed
Lead (Pb)	98	7421	01/19/95	01/20/95

LCS Limits are 85 - 115%.

#### **Matrix Spikes**

Analyte	M8 % Recov	MSD % Recov	RPD	EPA Method	Date Digested	Date Analyzed
Lead (Pb)	94	96	4	7421	01/19/95	01/20/95

MS = Matrix Spike MSD = Matrix Spike Duplicate RPD = Relative Percent Difference Spike Recovery Limits for Matrix Spikes are 75 - 125%. The RPD Limits are  $\pm$  20%.

Michelle L. Anderson Inorganics Supervisor



January 24, 1995

# Metals QC Report for Sample Log 11124 (cont'd)

From: Project # 12104 (Jos Sio Chevrolet) Sample Spiked for MS/MSD: 11124-5

#### Matrix Spikes

Analyte	MS % Recov	M8D % Recov	RPD	EPA Method	Date Digested	Date Analyzed
Cadmium (Cd)		115	2	6010	01/23/95	01/24/95
Chromium (Cr)		98	0	6010	01/23/95	01/24/95
Lead (Pb)		98	4	7421	01/19/95	01/20/95
Nickel (Ni)		99	1	6010	01/23/95	01/24/95
Zinc (Zn)		107	1	6010	01/23/95	01/24/95

MS = Matrix Spike MSD = Matrix Spike Duplicate RPD = Relative Percent Difference Spike Recovery Limits for Matrix Spikes are 75 - 125%. RPD Limits are ± 20%.

Michelle L. Anderson Inorganics Supervisor



January 24, 1995

## Metals QC Report for Sample Log 11124

From: Project # 12104 (Joe Sio Chevrolet)

Matrix: Water

Units: (mg/L)

#### Method Blank

Analyte	Result	MRL	EPA Method	Date Digested	Analyzed
Cadmium (Cd) Chromium (Cr) Lead (Pb) Nickel (Ni) Zinc (Zn)	<0.004 <0.007 <0.003 <0.015 <0.010	0.004 0.007 0.003 0.015 0.010	6010 5010 7421 6010 6010	01/23/95 01/23/95 01/19/95 01/23/95	01/24/95 01/24/95 01/20/95 01/24/95

MRL = Method Reporting Limit

Laboratory Control Sample (LCS)

Analyte	% Recovery	EPA Method	Date Digested	Date Analyzed
Cadmium (Cd) Chromium (Cr) Lead (Pb) Nickel (Ni) Zinc (Zn)	112	6010	01/23/95	01/24/95
	102	6010	01/23/95	01/24/95
	98	7421	01/19/95	01/20/95
	104	6010	01/23/95	01/24/95
	109	6010	01/23/95	01/24/95

LCS Limits are 85 - 115%.

Michelle L. Anderson

Inorganics Supervisor

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