Ultramar

ALCO HAZMAT

Ultramar Inc.

P.O. Box 466 525 W. Third Street Hanford, CA 93232-0466 (209) 582-0241 94 FEB 17 PM 3: 52

Telecopy: 209-584-6113 Credit & Wholesale 209-583-3330 Administrative 209-583-3302 Information Services 209-583-3358 Accounting

check paser when I

February 15, 1994

34-54'

Ms. Eva Chu Department of Environmental Health Alameda County Health Care Services 80 Swan Way, Room 200 Oakland, CA 94612

SUBJECT:

BEACON STATION NO. 604, 1619 FIRST STREET, LIVERMORE,

CALIFORNIA

Dear Ms. Chu:

Enclosed is a copy of the Ground-Water Monitoring Report, First Quarter 1994 for the above-referenced Ultramar facility. Also included is a copy of the Quarterly Status Report which describes the work completed this quarter and the work anticipated to be completed next quarter.

Evidently, CalTrans had misplaced the Ultramar encroachment permit application and has just recently found the application. It is hoped that the encroachment permit will be approved by the end of February. The City encroachment permit has been approved. It is anticipated that the additional wells can be installed in March.

Please call if you have any questions regarding this site.

Sincerely,

ULTRAMAR INC.

Terrence A. Fox

Tenene A. 24

Senior Project Manager

Marketing Environmental Department

cc:

Alameda County Local Coordinator, San Francisco Bay Region,

RWQCB



Ultramar

Ultramar Inc.

P.O. Box 466 525 W. Third Street Hanford, CA 93232-0466 (209) 582-0241

Telecopy: 209-584-6113 Credit & Wholesale 209-583-3330 Administrative 209-583-3302 Information Services 209-583-3358 Accounting

ENVIRONMENTAL PROJECT QUARTERLY STATUS REPORT

DATE REPORT SUBMITTED: February 15, 1994

QUARTER ENDING: March 31, 1994

SERVICE STATION NO.: 604

ADDRESS: 1619 First Street, Livermore, CA

COUNTY: Alameda

ULTRAMAR CONTACT: Terrence A. Fox

TEL. NO: 209-583-5545

BACKGROUND:

November 1992, three underground storage tanks were In Hydrocarbons were detected and the excavation was removed. extended to a depth of 27 feet in the southwest corner. Hydrocarbons were detected in the sample collected from the base of the overexcavation.

In May and June 1993, three monitoring wells (MW-1 through MW-3), three vapor wells (VW-1 through VW-3), and one boring. the soil plume has been defined.

The site has been placed on a quarterly monitoring program.

SUMMARY OF THIS QUARTER'S ACTIVITIES:

Awaiting on approval of encroachment permits. Performed quarterly monitoring on January 13, 1994.

RESULT OF QUARTERLY MONITORING:

Monitoring data indicates that the benzene concentrations decreased in MW-1 from 4,700 ppb to 1,300 ppb and in MW-3 from 57 ppb to 2.6 ppb. The benzene concentration increased in MW-2 from 17,000 ppb to 20,000 ppb.

PROPOSED ACTIVITY OR WORK FOR NEXT QUARTER:

ACTIVITY

ESTIMATED COMPLETION DATE

Continue quarterly monitoring program.

Install the additional monitoring wells.

March 15, 1994





Beacon Station 604 Page 2

RESULT OF QUARTERLY MONITORING:

Monitoring data indicates that the benzene concentrations decreased in MW-1 from 8,000 ppb to 4,700 ppb and in MW-2 from 19,000 ppb to 17,000 ppb. The benzene concentration increased in MW-3 from 8.2 ppb to 57 ppb.

PROPOSED ACTIVITY OR WORK FOR NEXT QUARTER:

<u>ACTIVITY</u>

ESTIMATED COMPLETION DATE

Continue quarterly monitoring program.

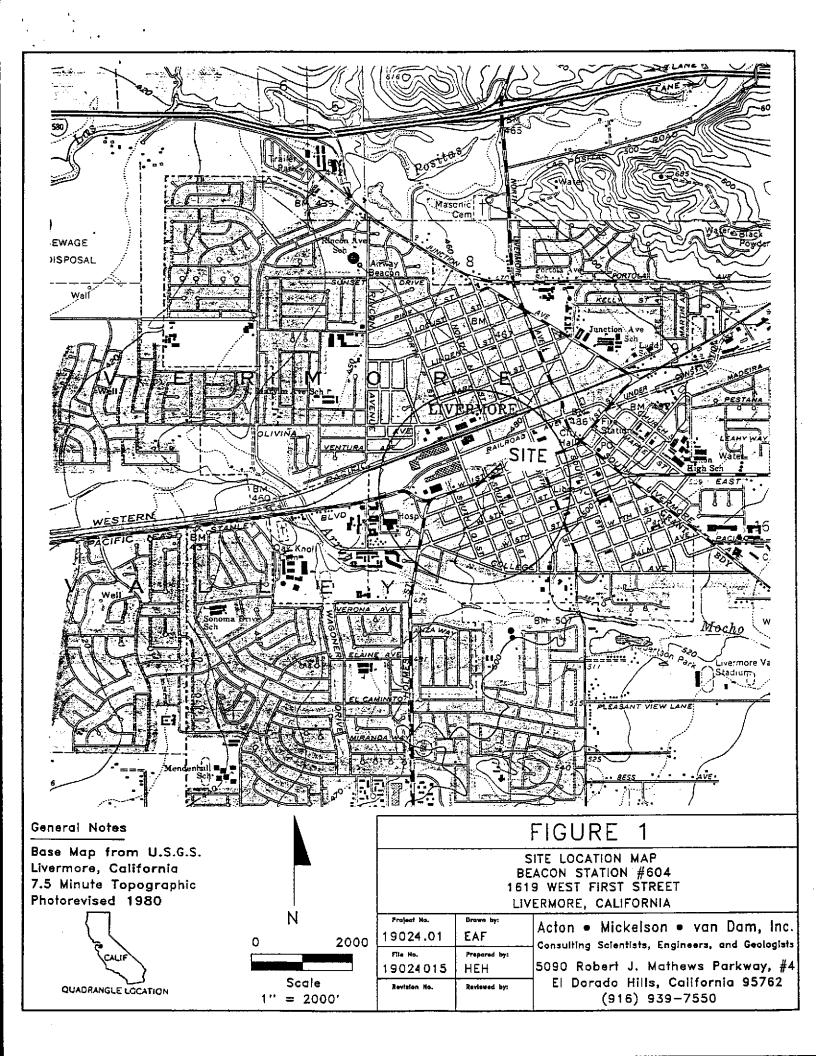
Install the additional monitoring wells.

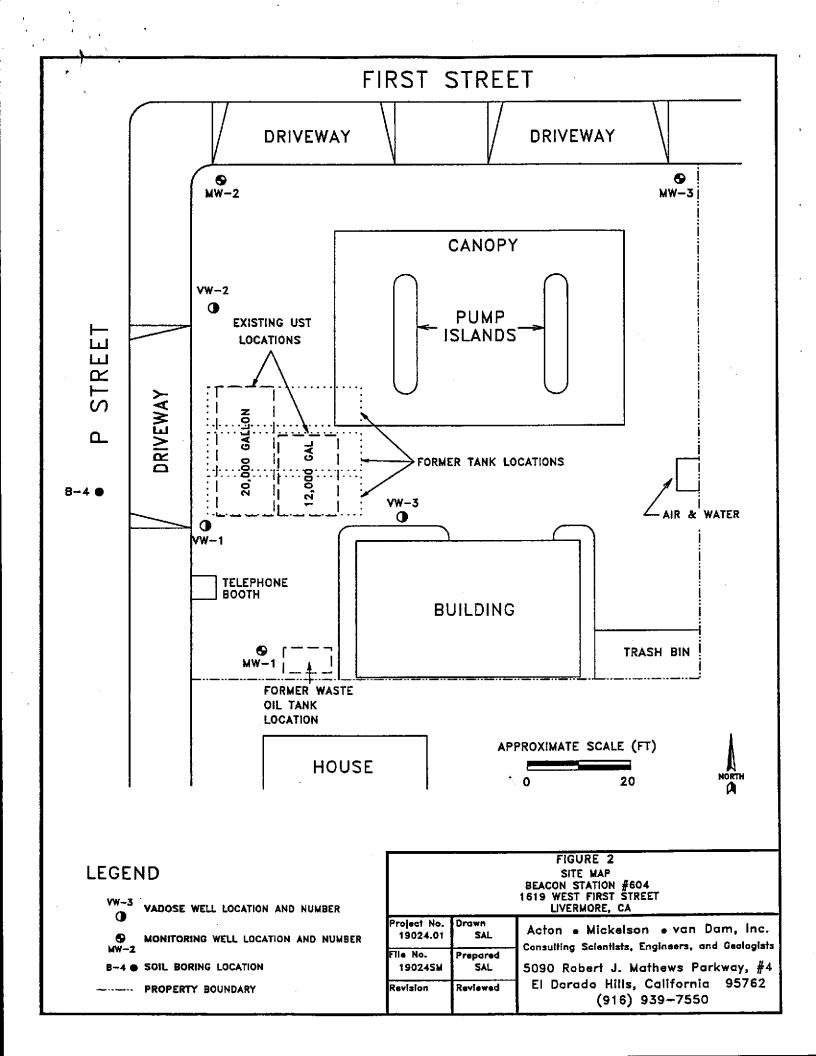
January 31, 1994

Perform vapor extraction and groundwater pumping tests. February 28, 1994

Submit a PAR

March 31, 1994





Consulting Scientists, Engineers, and Geologists

February 9, 1994

Mr. Terrence A. Fox Ultramar Inc. 525 West Third Street Hanford, California 93230

19024.04

Subject:

Ground Water Monitoring Report, First Quarter 1994

Beacon Station #604--1619 West First Street, Livermore, California

Dear Mr. Fox:

Acton • Mickelson • van Dam, Inc. (AMV), has been authorized to continue a hydrogeologic investigation of ground water conditions at Beacon Station #604, located at 1619 West First Street, Livermore, California (Figures 1 and 2). The investigation is intended to assess the distribution of petroleum hydrocarbon constituents in the ground water beneath the site. This letter report summarizes the results of ground water sampling conducted on January 13, 1994. The procedures used to purge and sample monitoring wells and measure water levels are described in Enclosure A.

Ground Water Level Measurements, Hydraulic Gradient, and Flow Direction

Depth to ground water was measured in monitoring wells MW-1 through MW-3. Depth to ground water ranged from 33.95 to 35.85 feet below top of casing. Ground water level measurements from this sampling event, as well as previous ground water depth measurements, are presented in Table 1. Ground water levels increased an average of approximately 7.6 feet between October 1993 and January 1994. The inferred direction of ground water flow was toward the northwest (Figure 3) which is consistent with previous monitoring events. Gradient was calculated to be approximately 0.03 foot per foot.

Ground Water Sample Analytical Results

Ground water samples were collected from monitoring wells MW-1 through MW-3 on January 13, 1994, using the procedures outlined in Enclosure A. Field observations and ground water sampling documentation are presented in Enclosure B. Ground water samples were submitted to a state-certified laboratory for analysis of benzene, toluene, ethylbenzene, xylenes,

Mr. Terrence A. Fox February 9, 1994 Page 2

and total petroleum hydrocarbons as gasoline (TPHg). Ground water sample analytical results from this sampling event and previous events are compiled in Table 2. Copies of the certified laboratory analytical reports from this sampling event are presented in Enclosure C.

Discussion

Benzene concentrations in ground water ranged from 20,000 μ g/l in the sample collected from monitoring well MW-2 to 2.6 μ g/l in a sample collected from monitoring well MW-3. Compared to previous monitoring events, benzene concentrations in water samples collected on January 13, 1994, decreased in monitoring wells MW-1 and MW-3 and increased slightly in MW-2. Benzene concentrations reported from the January 13, 1994, ground water sample analytical results are presented on Figure 4.

Future Work

Quarterly ground water monitoring will continue at the site. The next sampling event is scheduled for April 1994. AMV is currently in the process of acquiring encroachment and monitoring well permits to install additional off-site monitoring wells.

Remarks

The opinions and conclusions contained in this letter report represent our professional opinions. These opinions are based on currently available information and were developed in accordance with currently accepted hydrogeologic and engineering practices at this time. Other than this, no warranty is implied or intended.

AMV recommends that a copy of this quarterly monitoring report be forwarded to the following:

Ms. Eva Chu
Department of Environmental Health
Alameda County Health Care Services
80 Swan Way, Room 200
Oakland, California 94612

Mr. Cecil Fox
California Regional Water Quality Control Board,
San Francisco Bay Region
2101 Webster Street, Room 500
Oakland, California 94612

ACTON • MICKELSON • van DAM, INC.

hrpt049.mj

Mr. Terrence A. Fox February 9, 1994 Page 3

If you have any questions, please call the undersigned at (916) 939-7550.

Sincerely,

ACTON • MICKELSON • van DAM, INC.

William G. Rocha, Jr.

Staff Engineer

TAD:DAvD:mjd

Enclosures

Sale d. van Jam

Dale A. van Dam, R.G.

California Registered Geologist #4632

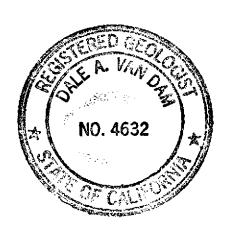


TABLE 1

WATER ELEVATION DATA

Beacon Station #604 1619 West First Street, Livermore, CA

Monitoring Well	Date	Top of Riser	Depth of Water (feet)	Ground Water Elevation (feet)	Physical Observation
MW-1	06-01-93 06-22-93 10-06-93 01-13-94	100.00	37.50 38.46 42.22 34.52	62.50 61.54 57.78 65.48	No Product No Product No Product No Product
MW-2	06-01-93 06-22-93 10-06-93 01-13-94	98.68	38.02 39.07 43.72 35.85	60.66 59.61 54.96 62.83	No Product No Product No Product No Product
MW-3	06-01-93 06-22-93 10-06-93 01-13-94	97.08	36.18 37.11 41.15 33.95	61.90 61.97 55.93 63.13	No Product No Product No Product No Product

NOTE: Monitoring well elevations were surveyed relative to an arbitrary benchmark at the top of the casing of monitoring well MW-1 with an assumed elevation of 100.00 feet.

TABLE 2

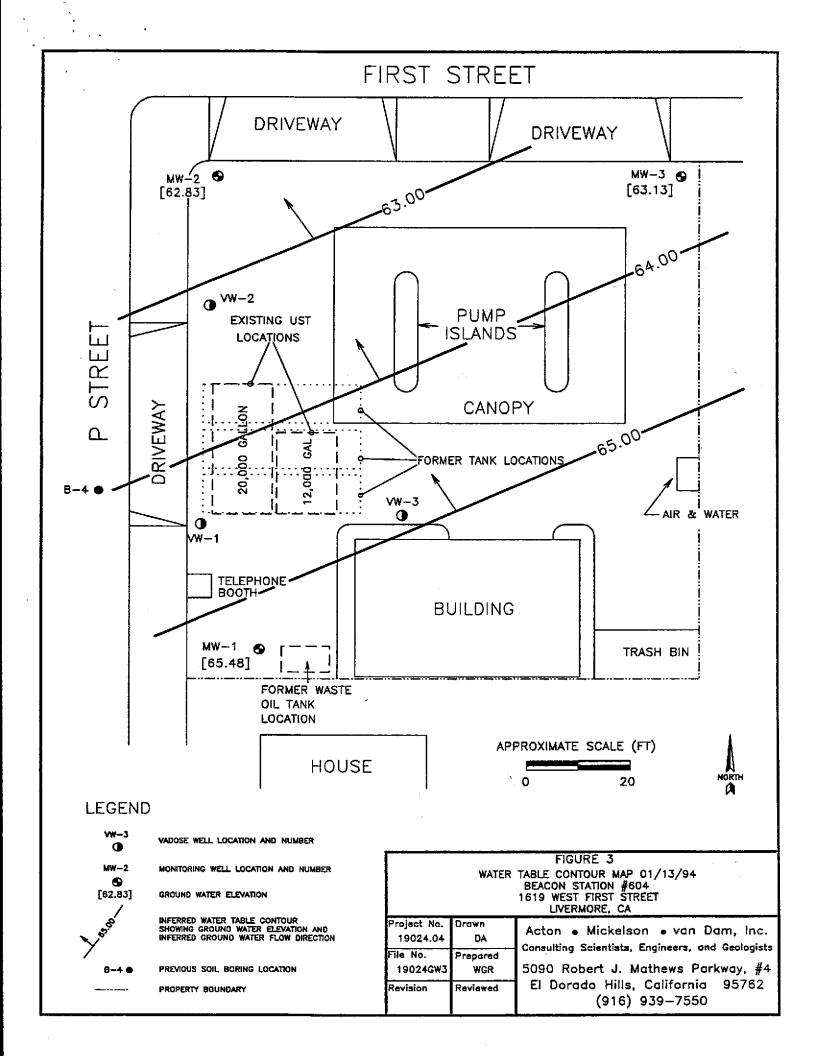
GROUND WATER SAMPLE ANALYTICAL RESULTS

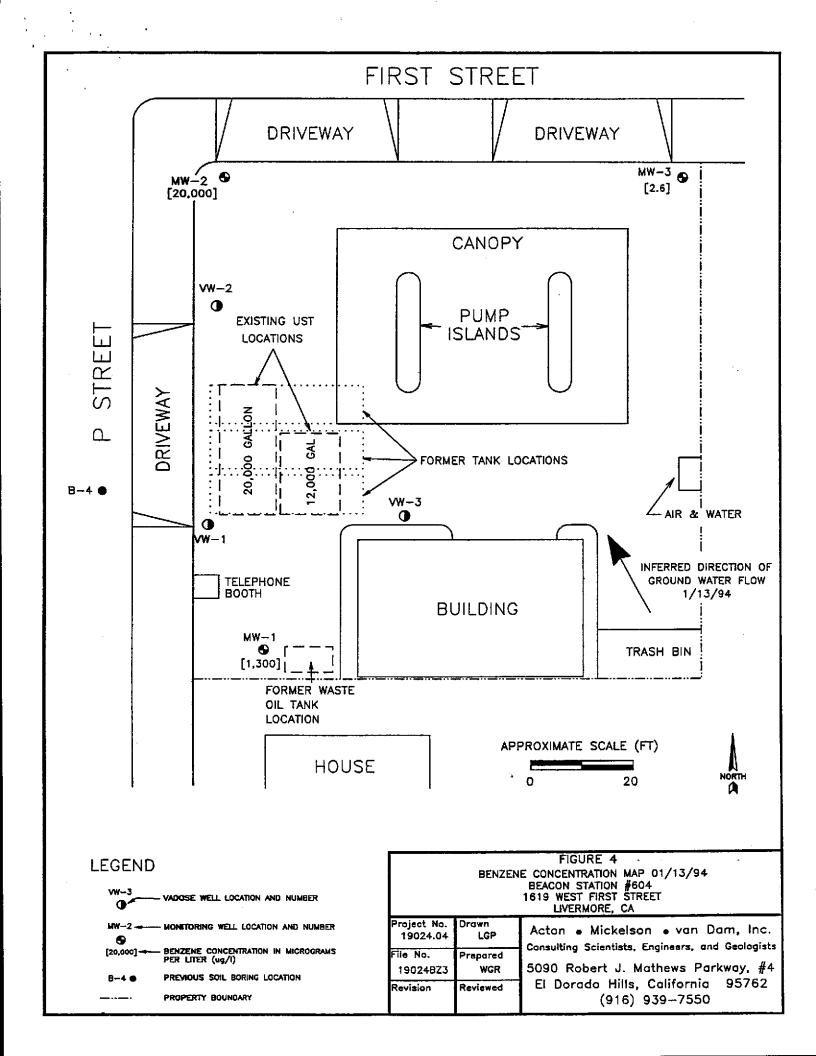
Beacon Station #604

1619 West First Street, Livermore, CA Concentrations in micrograms per liter $(\mu g/l)$

Monitoring Well	Date Sampled	Benzene	Toluene	Ethylbenzene	Total Xylenes	TPHg
MW-1	06-01-93	2,200	400	< 50	4,900	27,000
	06-22-93	8,000	10,000	260	10,000	87,000
	10-06-93	4,700	6,500	740	5,300	40,000
	01-13-94	1,300	950	110	850	9,400
MW-2	06-01-93	20,000	21,000	3,300	18,000	170,000
	06-22-93	19,000	22,000	3,500	18,000	160,000
	10-06-93	17,000	17,000	3,000	15,000	110,000
	01-13-94	20,000	19,000	2,300	14,000	93,000
MW-3	06-01-93	4.6	< 0.50	< 0.50	1.9	270
11211	06-22-93	8.2	< 0.50	< 0.50	0.72	160
	10-06-93	57	110	24	120	740
	01-13-94	2.6	0.67	0.78	4.2	83

^aTPHg = total petroleum hydrocarbons as gasoline.





ENCLOSURE A SAMPLING TECHNIQUES

ENCLOSURE A

SAMPLING TECHNIQUES

Proper sampling techniques were followed to assure that samples represented actual field conditions and that samples were labeled, preserved, and transported properly to retain sample integrity. This exhibit describes procedures followed by Acton • Mickelson • van Dam, Inc. (AMV), during collection of samples of subsurface soil and ground water. Sampling guidance documents from the American Society of Testing and Materials (ASTM), U.S. Environmental Protection Agency (EPA), and California Department of Health Services (DHS) were followed for all sampling procedures. Actual sampling procedures employed were based on field conditions and may differ from those described here.

1.0 WATER LEVEL AND LIQUID-PHASE HYDROCARBON (LPH) THICKNESS MEASUREMENTS AND GROUND WATER SAMPLING

1.1 Water Level and LPH Thickness Measurements

The static water level and/or LPH thickness in each well was measured prior to purging or sampling.

The depth to water/product was measured using an electronic interface probe. The wire of the interface probe is marked at 0.01 foot intervals. One tone is emitted from the interface probe if LPH is encountered; another tone for water. The wire of the interface probe was lowered slowly until LPH or water was encountered. At this point, the mark on the interface wire opposite the permanent reference point on the top of the well casing was read to the nearest 0.01 foot and recorded. If the first encountered substance was LPH, the probe was lowered until the tone corresponding to water was emitted. This depth was also recorded. The difference between the two depths corresponds to the LPH thickness. The interface probe was rinsed in deionized water between measurements in different wells.

A permanent reference point was marked on the well casings. The permanent reference point on the well casings was surveyed to a common reference point. All well casing riser elevations are known to within 0.01 foot.

Prior to well development, a disposable bailer was used to collect a sample of LPH, if present in a well, for subjective analysis. The sample was collected by gently lowering the bailer approximately one-half the bailer length past the air/LPH interface. The appearance (color, opacity, "freshness") was described and noted on field notes.

If LPH was encountered in the well, it was removed by bailing or pumping and the approximate volume of LPH removed was recorded. LPH thickness was then remeasured. If LPH was still present, the thickness was recorded and the well was not sampled. If LPH was not present, the well was developed, purged, and sampled as described below.

1.2 Well Evacuation and Development

After the static water level in a well was determined and prior to collection of a ground water sample, stagnant water was removed from the well casing and the surrounding gravel pack by bailing, pumping, or with a vacuum truck. At least three casing volumes of water were removed from each well from which a sample was collected. The volume of water in the casing was determined from the known elevation of the water surface, the well bottom elevation (as measured when the well is installed), and the well diameter.

If the well was bailed or pumped during purging, samples were collected and field analyzed for pH, temperature, and specific conductance. The well was considered stabilized when repeated readings of the following parameters were within the ranges indicated as follows:

• Specific conductance ± 10 percent of the reading range

pH ±0.1 pH unit
 Temperature ±0.5° C.

After stabilization, and after at least three well volumes were evacuated, a sample was collected for analysis. The field container used for well stabilization measurements, and the pH, temperature, and conductivity probes were rinsed between wells with deionized water.

All purge water was containerized and properly handled and documented for disposal. If the containers were stored on site, a label specifying the date of purging, source, and the known or suspected nature of the contents was affixed to each container.

1.3 Sample Collection, Preservation, and Handling

After purging, a new polyethylene disposable bailer was used to collect samples for analysis. The bailer was attached to a new disposable rope and lowered slowly into the water to avoid agitation of the collected sample. Containers for volatile organics analyses were filled completely so no airspace remained in the vial after sealing.

All sample containers were prewashed and prepared at the analyzing laboratory in accordance with quality assurance/quality control protocols of the laboratory. Only sample containers appropriate for the intended analyses were used.

After sample collection, the samples were placed into coolers with ice packs. Internal temperature of the cooler was maintained at approximately 4 degrees Celsius. Samples were kept in coolers during transport to the analyzing laboratory.

2.0 DECONTAMINATION AND DISPOSAL PROCEDURES

2.1 Equipment Decontamination

Sampling equipment was decontaminated as follows:

- 1. Prior to individual sample collection, any sampling device was cleaned in a TSP solution and rinsed twice in clean water. Any visible soil residue was removed.
- 2. Water sampling containers were cleaned and prepared by the respective analytical laboratories.
- 3. Field monitoring equipment (pH, conductivity, or temperature probes) was rinsed with clean water prior to use and between samples.

3.0 FIELD MEASUREMENTS

Field data were collected during various sampling and monitoring activities; this section describes routine procedures followed by personnel performing field measurements. The methods presented below are intended to ensure that field measurements are consistent and reproducible when performed by various individuals.

3.1 Conductivity, Temperature, and pH

Specific conductance, water temperature, and pH measurements were made when a water sample was collected. Regardless of the sample collection method, a representative water sample was placed in a transfer bottle used solely for field parameter determinations. A conventional pH meter with a combination electrode or equivalent was used for field-specific conductance measurements. Temperature measurements were performed using standard thermometers or equivalent temperature meters. Combination instruments capable of measuring two or all three of the parameters may have also been used.

All instruments were calibrated in accordance with manufacturer methods. The values for conductivity standards and pH buffers used in calibration were recorded daily in a field notebook. All probes were thoroughly cleaned and rinsed with fresh water prior to any measurements, in accordance with Section 3.1.

4.0 SAMPLE CUSTODY

This section describes standard operating procedures for sample custody and custody documentation. Sample custody procedures were followed through sample collection, transfer, analysis, and ultimate disposal. The purpose of these procedures is to assure that (1) the integrity of samples was maintained during their collection, transportation, and storage prior to analysis and (2) post-analysis sample material was properly disposed of. Sample custody is divided into field procedures and laboratory procedures, as described below.

4.1 Field Custody Procedures

Sample quantities, types, and locations were determined before the actual fieldwork commenced. As few people as possible handled samples. The field sampler was personally responsible for the care and custody of the collected samples until they were properly transferred.

4.1.1 Field Documentation

Each sample was labeled and sealed properly immediately after collection. Sample identification documents were carefully prepared so that identification and chain-of-custody records could be maintained and sample disposition could be controlled. Forms were filled out with waterproof ink. The following sample identification documents were utilized.

- Sample labels
- Field notebook
- Chain-of-custody forms

4.1.2 Sample Labels

Sample labels provide identification of samples. Preprinted sample labels were provided. Where necessary, the label was protected from water and solvents with clean label-protection tape. Each label contained the following information:

- Name of collector
- Date and time of collection
- Place of collection
- AMV project number
- Sample number
- Preservative (if any)

4.1.3 Field Notebook

Information pertinent to a field survey, measurements, and/or sampling were recorded in a bound notebook. Entries in the notebook may have included the following:

- Name and title of author, date and time of entry, and physical/environmental conditions during field activity.
- Location of sampling or measurement activity.
- Name(s) and title(s) of field crew.
- Type of sampled or measured media (e.g., soil, ground water, air, etc.)
- Sample collection or measurement method(s).
- Number and volume of sample(s) taken.
- Description of sampling point(s).
- Description of measuring reference points.
- Date and time of collection or measurement.
- Sample identification number(s).
- Sample preservative (if any).
- Sample distribution (e.g., laboratory).
- Field observations/comments.
- Field measurements data (pH, etc.).

4.1.4 Chain-of-Custody Record

A chain-of-custody record was filled out for and accompanied every sample and every shipment of samples to the analytical laboratories in order to establish the documentation necessary to trace sample possession from the time of collection. The record contained the following information:

- Sample or station number or sample I.D.
- Signature of collector, sampler, or recorder.
- Date and time of collection.
- Place of collection.
- Sample type.
- Signatures of persons involved in the chain of possession.
- Inclusive dates of possession.

The laboratory portion of the form was completed by laboratory personnel and contains the following information:

- Name of person receiving the sample.
- Laboratory sample number.
- Date and time of sample receipt.
- Analyses requested.
- Sample condition and temperature.

4.1.5 Sample Transfer and Shipment

Samples were always accompanied by a chain-of-custody record. When transferring samples, the individuals relinquishing and receiving the samples signed, dated, and noted the time on the chain-of-custody record. Samples were packaged properly for shipment and dispatched to the

appropriate laboratory for analysis. The chain-of-custody record accompanied each shipment. The method of shipment, courier name(s), and other pertinent information was entered in the chain-of-custody record.

4.2 Laboratory Custody Procedures

A designated sample custodian accepted custody of the shipped samples and verified that the information on the sample label matched that on the chain-of-custody record. Information regarding method of delivery and sample conditions was also checked on the chain-of-custody record. The custodian then entered the appropriate data into the laboratory sample tracking system. The laboratory custodian may have used the sample number on the sample label or may have assigned a unique laboratory number to each sample. The custodian then transferred the sample(s) to the proper analyst(s) or stored the sample(s) in the appropriate secure area.

Laboratory personnel are responsible for the care and custody of samples from the time they are received until the sample is exhausted. Once at the laboratory, the samples are handled in accordance with <u>U.S. Environmental Protection Agency SW-846</u>, <u>Test Methods for Evaluating Solid Waste Physical/Chemical Methods</u>, <u>Third Edition</u>, for the intended analyses. All data sheets, chromatographs, and laboratory records were filed as part of the permanent documentation.

4.3 Corrections to Documentation

Original data recorded in field notebooks, chain-of-custody records, and other forms were written in ink. These documents were not altered, destroyed, or discarded, even if they were illegible or contained inaccuracies that required a replacement document.

If an error was made or found on a document, the individual making the corrections did so by crossing a single line through the error, entering the correct information, and initialing and dating the change. The erroneous information was obliterated. Any subsequent error(s) discovered on a document were corrected. All corrections were initialed and dated.

4.4 Sample Storage and Disposal

Samples and extracts were retained by the analytical laboratory for 60 days after a written report was issued by the laboratory. Unless notified by the program manager, excess or unused samples were disposed of by the laboratory in an appropriate manner consistent with applicable government regulations.

ENCLOSURE B FIELD NOTES

ACTON • MICKELSON • vnn DAM, INC.

GROUND WATER LEVEL DATA

Project N	ame_ <i>13e</i>	acon 6	04				Project Number 19024.04				
Date_ /-,	13-94		Field Crew	6N:			Project Number 19029.04 Measuring Device Interface Probe and Number				
Well No.	Time	Depth to Product (feet)	Depth to Ground Water (feet)	Product Thickness (feet)	Reference Elevation (feet)	Ground Water Elevation (feet)	Physical Observations/Comments				
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ACTON • MICKELSON • van DAM, INC.

SAMPLING/DEVELOPMENT INFORMATION

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ACTON . MICKELSON . VII DAM, INC.

SAMPLING/DEVELOPMENT INFORMATION

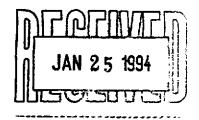
Sampling/Devel	opment Point	mw-3	3	Project Name	Beacon 609	£.
Sample I.D				Project No Work Order Data /-/ Field Crew_	3-94	
Number of bores evacuated before	tole volumes ore sampling:	33.95 4 casi'ns		•		inches AM/PM
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ENCLOSURE C GROUND WATER SAMPLE ANALYTICAL RESULTS



January 20, 1994 Sample Log 8379

William Rocha Acton, Mickelson & van Dam 5090 Robert J. Matthews Pkwy El Dorado Hills, CA 95762



Subject: Analytical Results for 3 Water Samples

Identified as: Project # 19024.04 (Beacon 604)

Received: 01/14/94

Dear Mr. Rocha:

Analysis of the sample(s) referenced above has been completed. This report is written to confirm results communicated on January 20, 1994 and describes procedures used to analyze the samples.

Sample(s) were received in 40-milliliter glass vials sealed with TFE lined septae and plastic screw-caps. Each sample was transported and received under documented chain of custody and stored at 4 degrees C until analysis was performed.

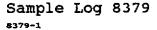
Sample(s) were analyzed using the following method(s):

"BTEX" (EPA Method 602/Purge-and-Trap)
"TPH as Gasoline" (Modified EPA Method 8015/Purge-and-Trap)

Please refer to the following table(s) for summarized analytical results and contact us at 916-757-4650 if you have questions regarding procedures or results. The chain-of-custody document is enclosed.

Approved by:

Senior Chemist





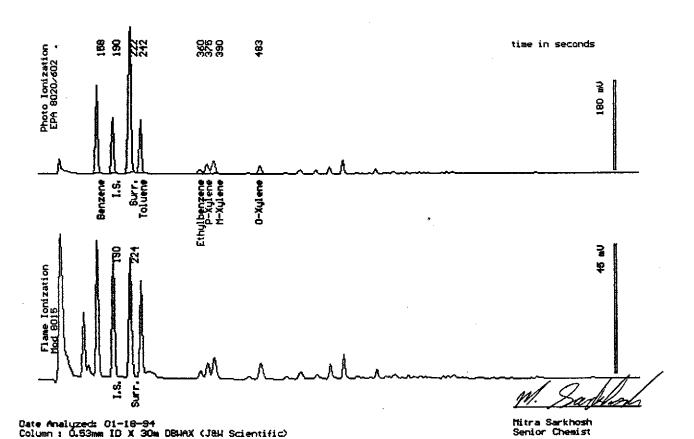
Sample: MW-1

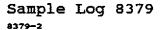
From : Project # 19024.04 (Beacon 604)

Sampled: 01/13/94 Dilution: 1:50 QC Batch: 4060m

Matrix : Water

Parameter	(MRL) ug/t	Measured Value ug/L
Benzene	(25)	1300
Toluene	(25)	950
Ethylbenzene	(25)	110
Total Xylenes	(25)	850
TPH as Gasoline	(2500)	9400
Surrogate Recovery	7	94 %







Sample: MW-2

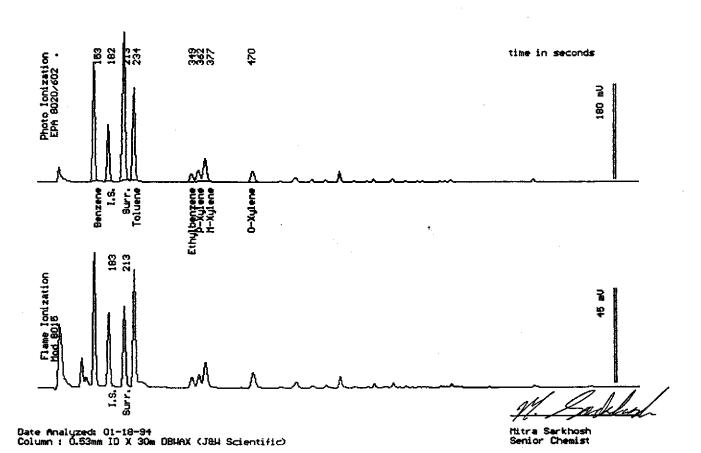
From : Project # 19024.04 (Beacon 604)

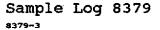
Sampled: 01/13/94 Dilution: 1:500

Dilution: 1:500 QC Batch: 4060m

Matrix : Water

Parameter	(MRL) ug/L	Measured Value ug/L
Benzene Toluene	(250) (250)	20000 19000
Ethylbenzene	(250)	2300
Total Xylenes	(250)	14000
TPH as Gasoline	(25000)	93000
Surrogate Recovery	, ·	95 %







Sample: MW-3

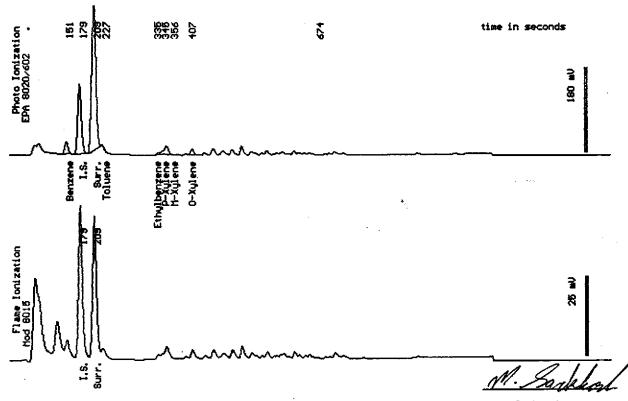
From : Project # 19024.04 (Beacon 604)

Sampled: 01/13/94

Dilution: 1:1 QC Batch: 2046A

Matrix : Water

Parameter	(MRL) ug/L	Measured Value ug/L
Benzene	(.50)	2.6
Toluene	(.50)	.67
Ethylbenzene	(.50)	.78
Total Xylenes	(.50)	4.2
TPH as Gasoline	(50)	83
Surrogate Recovery	7	98 %



Date Analyzed: 01-15-94 Column : 0.53mm ID X 30m DBWAX (J&W Scientific)

Mitra Sarkhosh Senior Chemist



Ultramar Inc.CHAIN OF CUSTODY REPORT

BEACON

Beacon Station No.	Sampler (Print	Sampler (Print Name)			ANALYSES				Date	Form No.	
604	William	Rochen Jr.		\vdash	<u> </u>	NALYS	ES	+	1-14-94	/ of /	
Project No.	Sampler (Signa	ature)		7							į
19029.04	William	KaMA			<u>6</u>			Ders			
Project Location 1619 First Street Livermore, CA	Affiliation AMU, J	Enc.			(gasoline) (diesel)			No. of Containers	Standar	d TAT	-
Sample No./Identification	Date	Time	Lab No.	BTE	HT H			No.	REMA		
$m\omega-1$	1-13-94	1555		x	У			3	40m/ Von	s, Presv. 1	HCL
mw-Z		1445		Х	X			3)		
MW-3		1335		X	<u> </u>			3	(
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William Roch M. , AMV, Inc	1-14-91	11:07 =	Florit Cu	1	/	/ "	1.1	1X	/	1/14/9/11	1:07
Relinquished by: (Signature/Alfiliation)	Date		ed by (Signatur	e/Ai	filiat	on)	// ·				ime
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Relinquisted by Signature Affiliation	Date	Time Receiv	ed by: (Signatur	re/Af	filiati	on)				Date T	ime
total Carlo	1/14/94	12:05	Mur	<u> </u>		1E)	37			14/901	1205
Report Vo:	1	₽⁄ill to:	ULTRAMAF 525 West T			et				, ,	
Bill Rocha AMU, Inc.			Hanford, CA			-0 Y	,				\$
MINIO INC.			Attention:		, ,-	<u> </u>					
WHITE: Return to Client with Report	YELLOW: Labo	oratory Copy	PINK: Origin	nator	Cor	у				32-8003 1	/90