

## BLAINE TECH SERVICES INC.

1370 TULLY RD., SUITE 505 SAN JOSE, CA 95122 (408) 995-5535

compiled June 13, 1988

Chevron USA, Inc. 2410 Camino Ramon San Ramon, CA 94583

Attn: Bill Barnard

SITE:

CHEVRON SERVICE STATION NO. 1026 3701 BROADWAY OAKLAND, CALIFORNIA

PROJECT:

FULL SERVICE STATION DEMOLITION WITH REMOVAL OF ALL ABOVE GROUND AND SUBSURFACE INSTALLATIONS

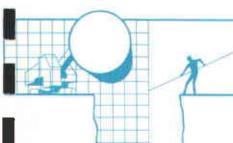
ALL SAMPLING THROUGH MAY 4, 1988

#### CUMULATIVE FORMAT SAMPLING REPORT 88126-C-2

This Cumulative Sampling Report includes the following:

	DATE	REPORT #	ACTIVITY
SECTION ONE	4/22/88	88113-M-1	Tank removal sampling
SECTION TWO	5/4/88	88125-C-2	Additional excavation sampling in the waste oil pit. Product line and stockpile sampling.

The Cumulative Report format presents all the environmental sampling and documentation activities performed by our firm during this project. The reports are presented in chronological order. Each sampling event is presented as a Section within the Cumulative Sampling Report. SECTION ONE contains the earliest work, SECTION TWO contains the next subsequent work, with additional Sections being added as more sampling is performed. The most recent information is always contained in the last Section of a Cumulative Report.



## BLAINE TECH SERVICES INC.

1370 TULLY RD., SUITE 505 SAN JOSE, CA 95122 (408) 995-5535

June 28, 1988

Chevron USA, Inc. 2 Annabel Lane Suite 200 San Ramon, CA 94583

Attn: Bill Barnard

SITE: CHEVRON SERVICE STATION NO. 1026 3701 BROADWAY OAKLAND, CALIFORNIA

PROJECT:
FULL SERVICE STATION DEMOLITION
WITH REMOVAL OF ALL ABOVE GROUND
AND SUBSURFACE INSTALLATIONS

SAMPLED ON: APRIL 22, 1988

#### TANK REMOVAL SAMPLING REPORT 88113-M-1

This report describes the initial environmental sampling and documentation performed by our firm on this project. In addition to the Sampling Report text, itself, supporting documents are provided as attachments. These include the chain of custody and the certified analytical laboratory report. All these documents should be kept together and preserved as a file of interrelated records which, together, comprise the documentation of the work performed at the site. If additional work is required, then a cumulative report format will be followed so that all information on the various phases of work at the site can be easily reviewed.

#### THE SCOPE OF WORK

In accordance with your request, field personnel were dispatched to the site to observe the tank removal, collect samples, arrange the proper analyses of the samples and maintain adequate documentation resulting in the issuance of a formal Sampling Report. The collection of environmental samples was

performed in accordance with the requirements of the State Water Resources Control Board, Regional Water Quality Control Board, and the specific directions of the Local Implementing Agency (LIA) inspector present at the site at the time of removal.

#### EXECUTION OF THE TANK REMOVAL SAMPLING

Personnel from our office were present at the subject site for the tank removal on April 22, 1988.

TANK I.D.	TANK SIZE IN GALLONS	TANK CONTENT	MATERIAL OF CONSTRUCTION	INSPECTION FOUND
A	10,000	GASOLINE	FIBERGLASS	NO HOLES
B C	10,000 10,000	GASOLINE GASOLINE	FIBERGLASS FIBERGLASS	NO HOLES
D	6,000	RECOVERY	FIBERGLASS	NO HOLES
WO	1,000	WASTE OIL	FIBERGLASS	NO HOLES

The subject site is located within the overall jurisdiction of the Regional Water Quality Control Board -- San Francisco Bay Region. Initial inspection and evaluation of the site is customarily conducted by the local implementing agency (LIA), which in this case was the Oakland Fire Department. The local implementing agency was represented by Ariel Bryant, who was present to observe the tank removal and sampling.

In accordance with the local regulations and the field judgement of the LIA representative, a detailed inspection was conducted in which the tanks were visually inspected and likely failure points were probed with small pointed metal examination tools. No holes were observed in any of the tanks.

Tanks designated A, B, C, and WO shared a common excavation. The portion of the tank pit that contained tanks A, B, and C was found to have standing water at a depth of approximately 14' below grade. Subsequent to the removal of the tanks, approximately 1/8" of free-floating product was observed on the water. Approximately 2,800 gallons of liquid were removed from the excavation prior to the collection of capillary zone samples. These samples were taken from the three walls of the excavation that were safely accessible to the excavation equipment. Due to the large number of installed monitoring wells on the property, the LIA representative determined that the procurement of a water sample was not necessary.

Tank WO was of a smaller diameter than the other tanks in the pit. As a result, the impression in the backfill material left by this tank remained above the water standing in the deeper portion of the excavation. As this area of the excavation remained dry, standard interface zone sampling procedures were followed for the collection of soil samples beneath this tank.

Tank D was used to store recovery products. It was noted that a water line had been leaking into the pit area for some time. The water was standing at a level of approximately 3' below grade at the time the tanks were exposed. After the removal of the tank and the backfill material, the water level dropped to 13' below grade.

During the removal of Tank D, the tank was damaged by the excavation equipment. This damage resulted in the release of recovery product into the water standing in the main portion of the excavation. The addition of the recovery product to the water already in the excavation resulted in approximately 1/4" of free-floating product on the surface of the water. Therefore, approximately 700 gallons of liquid was removed from the excavation prior to the collection of two capillary zone samples from the walls of the pit.

Information on the exact location of a particular sampling point will be found on the UNDERGROUND STORAGE TANK REMOVAL AND SAMPLING LOG, and the TABLE OF SAMPLING LOCATIONS AND ANALYTICAL RESULTS. Additional information on the exact method of sample collection will be found in the SAMPLING METHODOLOGY section of this report.

After completion of the field work, the sample containers were delivered to Anametrix Incorporated in San Jose, California. Anametrix Incorporated is a California Department of Health Services certified Hazardous Materials Testing Laboratory and is listed as DOHS HMTL #151. It was requested that the analytical procedures used for these analyses be those specified by the Regional Water Quality Control Board -- San Francisco Bay Region. The methods are defined in attachments to the San Francisco RWQCB (Region 2) publication, <u>Guidelines For Addressing Fuel Leaks</u>.

### DIAGRAMS & TABLES

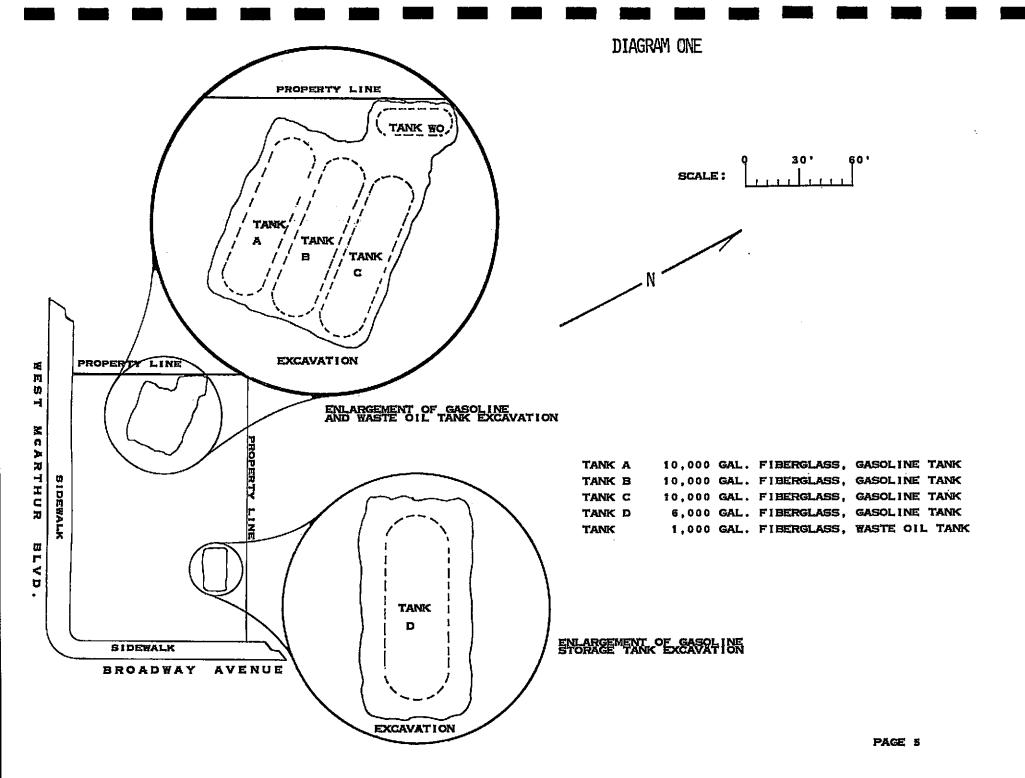
Graphic and Tabular presentation of all samples

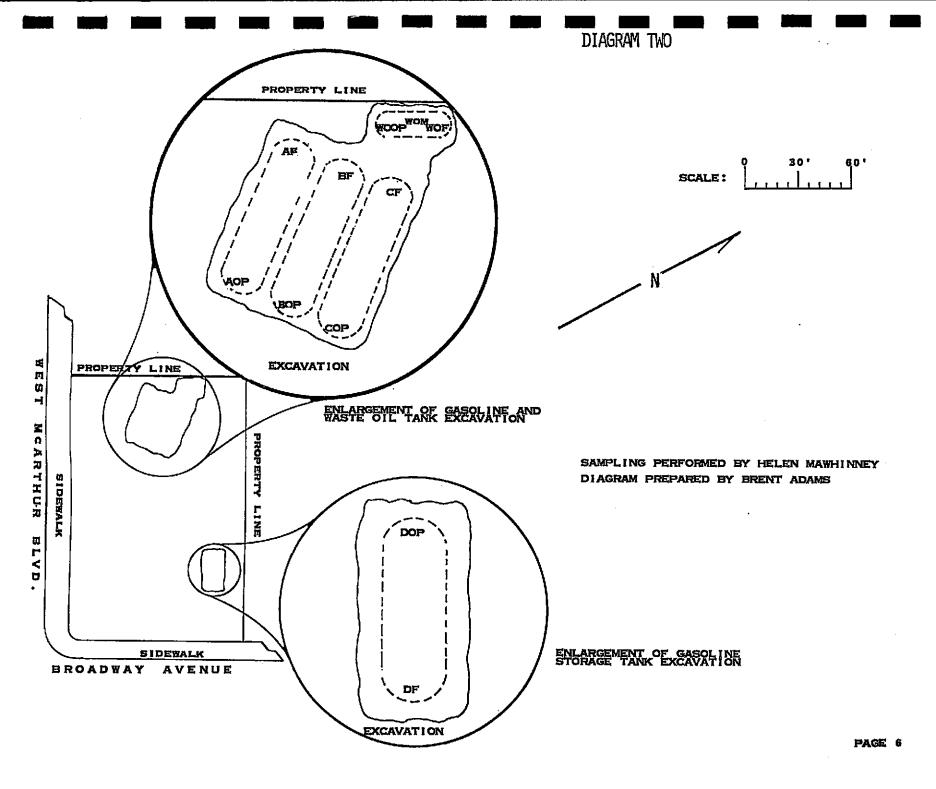
The TABLE OF SAMPLING LOCATIONS AND ANALYTICAL RESULTS relies on a system of designated SAMPLING AREAS which are specific locations on the site which correspond to the fuel dispensing fixtures that were originally in place there. Briefly, the tanks are assigned a letter and a suffix which comprise a descriptive identification. Even when the tanks have been removed, all samples taken in that area of the site are referenced to that particular SAMPLING AREA which can be cross referenced to construction blueprints, permit drawings and similar documents.

DIAGRAM ONE shows the location of key features of the site including streets, property lines, and the location of underground storage tanks. The diagram shows that each tank has been assigned an arbitrary letter designation (Tank A, Tank B etc.). This simplifies communications concerning a particular tank by providing a nomenclature that does not rely on descriptions by size and tank contents.

DIAGRAM TWO shows a finer level of detail. A descriptive suffix is added to the tank identification letter, in order to designate a SAMPLING AREA at that particular end of that tank. For example, Tank A is given an F suffix to indicate the fill pipe end and AF is used to define the area in which samples are taken. The opposite end of the tank from the fill pipe is given an OP suffix, and that SAMPLING AREA is, thereafter, referred to as Aop. The approximate midpoint of the tank is given an M suffix if a sample is taken from that location. The location of stockpiled material and the individual sampling points is also shown on Diagram Two.

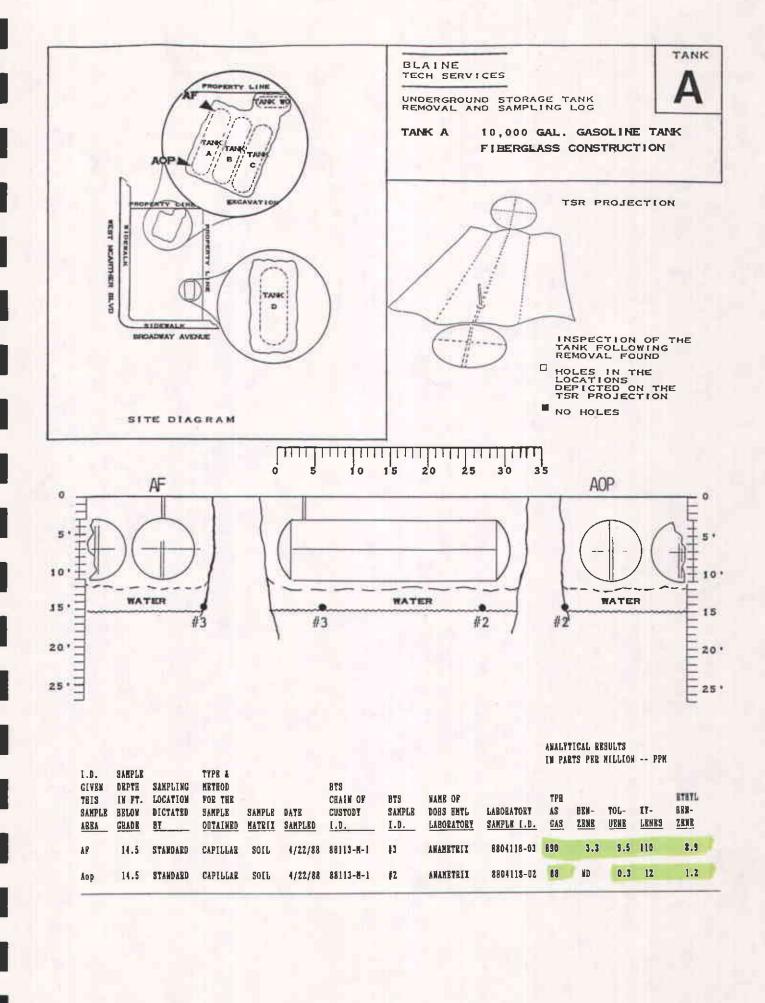
The relationship between a given tank and its sample collection points is more precisely illustrated in the UNDERGROUND STORAGE TANK REMOVAL AND SAMPLING LOGS. A log sheet is filled out for each of the tanks that was removed from the site. The log sheets contain end views and cross sections of each tank which graphically depict the SAMPLING AREAS at each end of that tank. Whenever holes are discovered in a tank during either removal or a post-removal inspection, the location of the holes is indicated on the TSR projection.

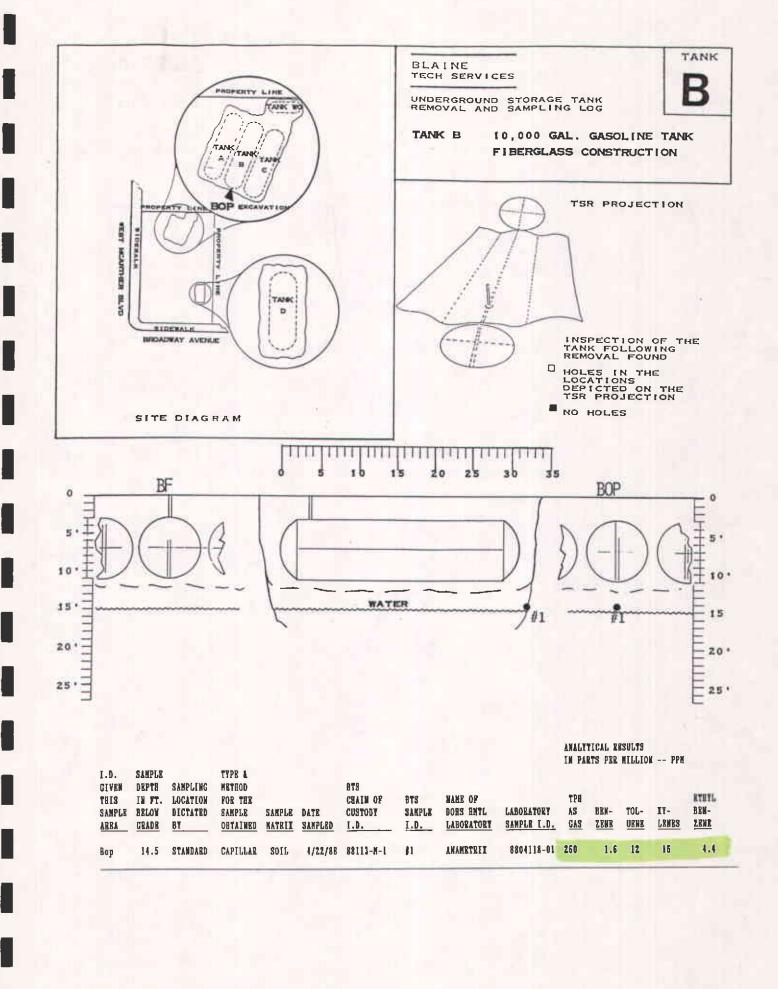


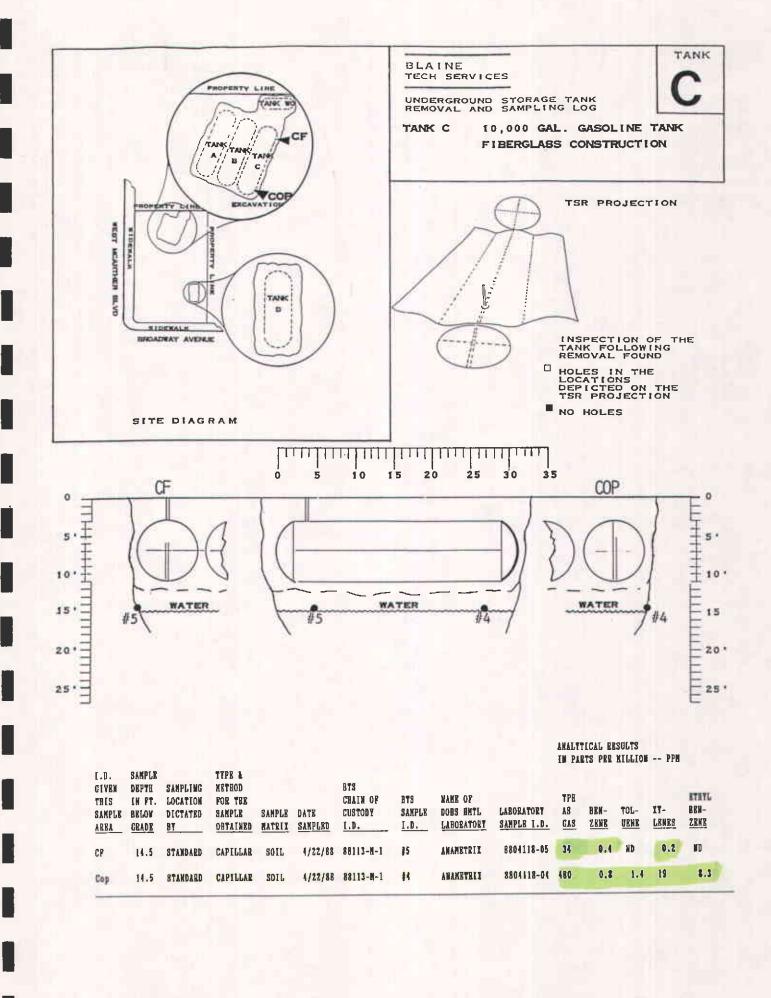


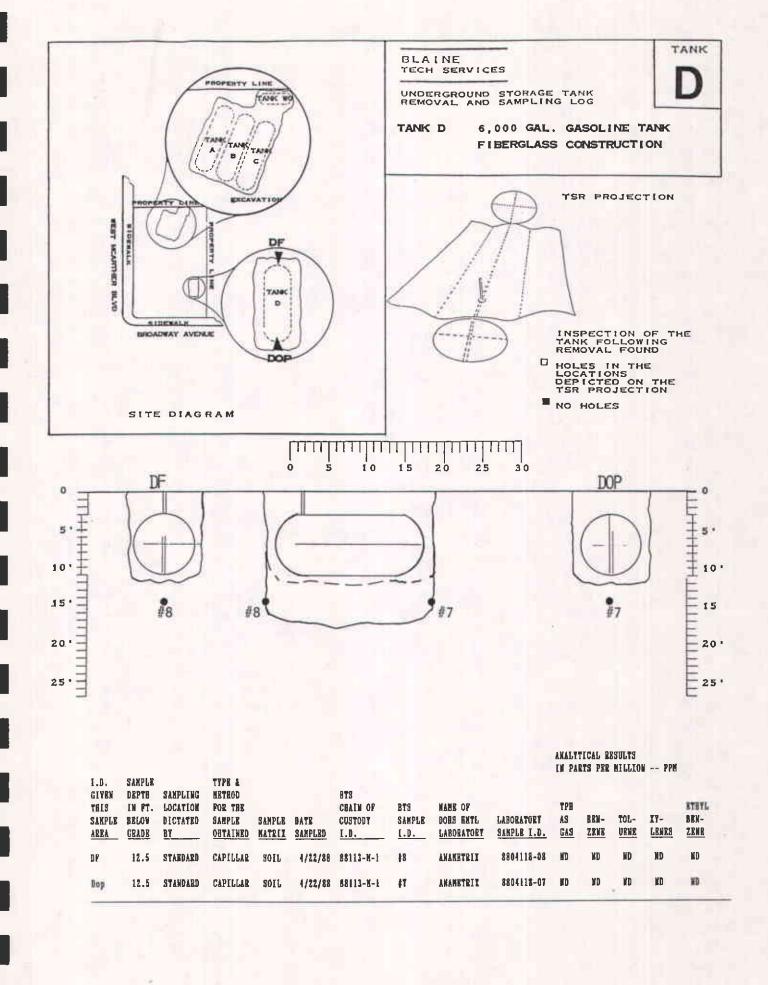
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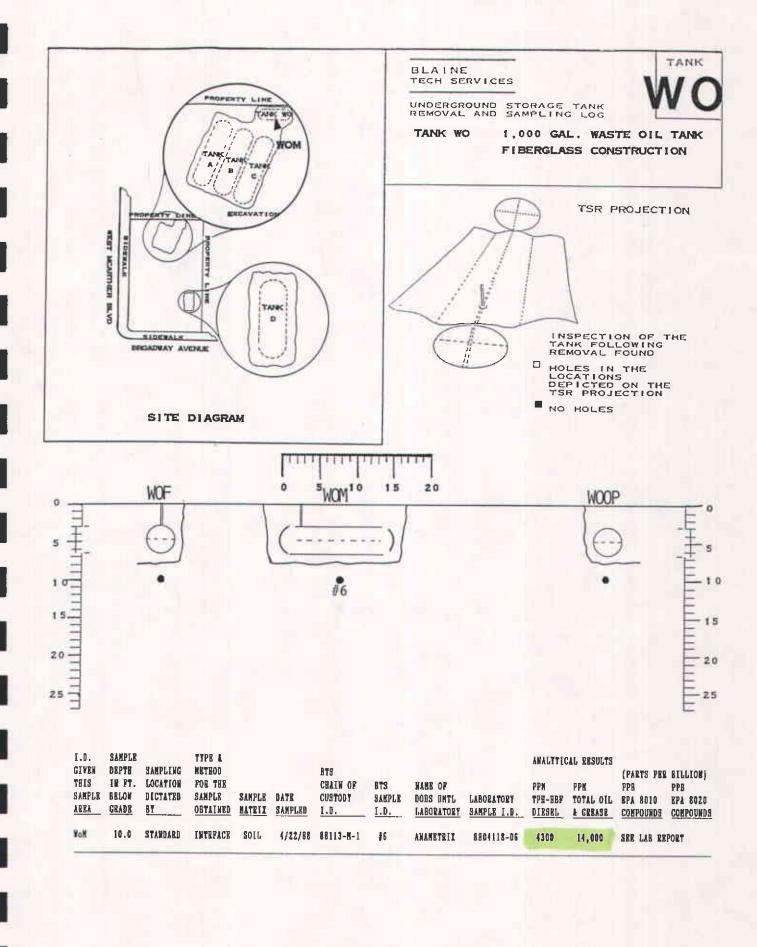
Rop Bop CP	14.5	STANDARD STANDARD STANDARD	CAPILLAR  CAPILLAR  CAPILLAR	SOIL	4/22/88	88113-M-1 88113-M-1 88113-M-1	#2 #1 #5	ANAMETRII ANAMETRII ANAMETRII	8804118-02 8804118-01 8804118-05	260	0.4	12	12 16 0,2	1.2 4.4 ND
Cop )P Oop	12.5	STANDARD STANDARD STANDARD	CAPILLAR CAPILLAR CAPILLAR	SOIL	4/22/88	88113-N-1 88113-N-1 88113-N-1	#4 #8	ANAMETRIX  ANAMETRIX  ANAMETRIX	8804118-04 8804118-08 8804118-07	ND ND	MD MD	ND T-4	ND ND	ND ND











#### PROCEDURES

Methodologies and Conventions

#### GENERAL PRACTICES WITHIN A MULTIPLE AGENCY HIERARCHY

U.S. Environmental Protection Agency standards serve as the foundation for all field sampling operations performed by our firm. The EPA SW 846 is the primary publication from which procedures are derived, though there are additional EPA sources such as training films and verbal communications. Sampling related to underground storage tanks and tank related threats to groundwater are governed by the California Water Resources Control Board and their Regional Water Quality Control Boards. While some aspects of field and laboratory work may be delegated to the California Department of Health Services, the CWRCB and the nine Regional Water Quality Control Boards establish the general and specific criteria for sampling performed in connection with underground storage tanks. This is done through the publication of guidance documents, the issuance of memoranda and verbal announcements.

Other agencies, such as Air Pollution Control Districts, may require additional samples, but these are usually in addition to samples required by the RWQCB. Local implementing agency (LIA) inspectors are frequently present during the tank removal phase of a project and either direct or request that samples be taken according to RWQCB specifications. Additional samples may, and frequently are, taken at the request of the LIA inspector.

Based on field conditions directly observable by the LIA inspector, our field personnel may be asked to collect samples that are tailored to the specific situation and which the inspector judges will provide substantial information about the site. Quite often these directions or suggestions coincide with the sampling areas established by the RWQCB as the proper collection points for samples which will be used as the Primary Criteria for a Regulatory Agency Determination on whether additional exploration or remediation will be required at a particular site. Similarly, there are instances when the LIA inspector's judgements do not coincide with Board specifications.

Two common examples of this are as follows:

- 1. A local implementing agency inspector notes that soil dug up from the correct RWQCB interface sampling point is relatively clean, but observes that there is quite obviously contaminated backfill underlying the center of the tank. The inspector directs that the contaminated backfill should be taken instead of the clean interface soil so as to provide information about the "worst case" conditions within the tank pit.
- 2. The soil at the specified interface sampling depth is found to be slightly contaminated, but much less so than the soil only a few inches above. Noting the relatively dense soil, the local implementing agency inspector decides not to have the interface soil sampled and has the backhoe dig deeper to see

if the contamination diminishes to acceptable levels. This exploration saves the property owner the cost of running two samples at that location, and enables the inspector to directly observe the condition of the deeper soil.

In both examples, different material is collected in lieu of a standard RWQCB interface sample. Further, the material collected is substantially different from what would have been obtained by taking representative soil at the Board specified sampling location. Note that both of these samples were taken at the direction of the local implementing agency inspector who was present at the site and elected to select alternative sampling locations. Note too, that these alternative samples may provide more information about the site than standard Board specified samples. However, as the LIA elected samples do not accurately reflect soil conditions at the sampling points specified by the RWQCB, the decision making process may be hampered.

Clearly there is no advantage in limiting the ability of the regulator in the field to make prudent judgements. Likewise, regulatory personnel and consultants who will review the reports without benefit of having been present at the site need to know that the samples taken were not obtained at the standard locations. A simple resolution to these situations is a brief notation indicating that the sampling was elective rather than in accordance with a standard Board specification. These notations appear in the third column of the TABLE OF SAMPLING LOCATIONS AND ANALYTICAL RESULTS. By referring to the notations in column three and four in the TABLE, any party reviewing the report should be able to determine if something other than Board standard samples were obtained, and when variant sampling was performed, clarify whether it was elected by the LIA inspector, elected by our field personnel, or the result of some physical condition at the site that made it impossible to obtain material from the correct sampling location.

### SAMPLING METHODOLOGIES USED ON THIS PROJECT

CAPILLARY ZONE SOIL SAMPLE: The capillary zone is the soil horizon immediately above the surface of standing groundwater into which moisture is drawn by capillary action. Capillary zone sampling is most often requested in open pit and open trench situations where lost petroleum products are evident or suspected. In these cases, it is reasoned that a sample of the capillary zone will demonstrate whether or not fuel has been drawn up into the soil above the groundwater and, thereby, provide a rough indication of the volume and duration of the lost fuel condition.

Engineers of the Region 2 RWQCB staff have specified the correct sampling area as being from zero to six inches above the surface of the standing perched water and no more than twelve inches back into the native soil from the lateral backfill/native soil interface.

There are two weaknesses which tend to invalidate capillary zone sampling on the basis of inconsistent results. First, is the difficulty encountered in locating the true surface of the perched water above which the capillary zone resides. The removal of the tank and backfill material tends to artificially lower the water in the immediate vicinity of the tank pit below the true standing water level and mislead observers attempting to evaluate where the capillary zone is located. Second, the zone itself is a narrow horizon which is bordered on the top and bottom by soil which would not be expected to contain nearly the concentration of fuel hydrocarbons as the capillary zone proper. Collecting the correct material is complicated by conditions at the site which usually consist of a broad excavation, with vertical walls descending into a water filled pit. Because of these conditions, direct approach to the sampling area is difficult, dangerous, or impossible.

Assuming that the true and original surface of the perched water can be determined, samples can be safely obtained by one of the following methods. The backhoe bucket can be used to dig up a segment of the pit wall that contains the capillary zone and bring it up for inspection and sampling. An alternative method is to use sections of light weight drill rod and a drive shoe which contains a brass sample liner. This train can be extended across the pit, positioned, and used to drive an undisturbed soil sample.

STANDARD RWQCB INTERFACE SAMPLE: The tank removal sampling followed the standard protocol for obtaining interface samples. These samples fall into the category of samples which are known to be of primary concern to the interested regulatory agencies for determining if additional action will be required at a site and the methodology has been closely defined in State and RWQCB publications, supplements, and presentations. These specify both the acceptable depth and lateral situation of sample collection points. In accordance with these specifications, sample collection is executed as close as possible to the center line (longitudinal axis) of the tank and on a vertical axis with the fill pipe. A corresponding location is also found at the opposite end of the tank whenever standard interface samples are being collected.

Briefly, the method consists of digging up native soil from directly below the fill pipe and the corresponding opposite end of the tank and obtaining a sample from the backfill/native soil interface or a short distance below the interface. In the case of tanks less than 1,000 gallons in capacity, only one sample from beneath the tank is required. A short distance has been defined by Region 2 Board engineers as not greater than twenty-four inches below the backfill/native soil interface and is generally taken to be one foot below the the backfill/native soil interface. This soil is brought up in the backhoe bucket. A shovel or trowel is used to cut away surface soil and backfill material which may have been included in the bucket, and the sample is taken by pushing or driving a brass sample liner into the newly exposed soil from the designated depth and location. Additional clarifications by Region 2 Board engineers have indicated that when there is an obvious difference in the relative contamination of soil brought up from the interface depth, then it is the relatively more contaminated soil that should be selected for inclusion in the sample.

### SAMPLE CONTAINERS

Our firm uses new sample containers of the type specified by either EPA or the RWQCB for the collection of samples at sites where underground storage tanks are involved. Water samples are contained in 40 ml volatile organic analysis vials (VOAs) when analysis for gasoline and similar light volatile compounds is intended. These containers are prepared according to EPA SW 846 and will contain a small amount of preservative when the analysis is for TPH as gasoline or EPA 602. Vials intended for EPA 601 analysis and EPA 624 GCMS procedures are not preserved. Closure is accomplished with an open headed (syringe accessible) plastic screw cap brought down on top of a Teflon faced septum which is used to seal the sample without headspace.

Water samples intended for semivolatile and nonvolatile analysis such as total oil and grease (TOG) and diesel (TPH HBF) are collected and transported in properly prepared new glass liter bottles. Dark amber glass is used in the manufacture of these bottles to reduce any adverse effect on the sample by sunlight. Antimicrobial preservative may be added to the sample liquid if a prolonged holding time is expected prior to analysis. Closure is accomplished with a heavy plastic screw cap.

Soil samples for volatile, semivolatile and nonvolatile analyses are all collected in properly prepared new brass liners which are 2 inches in diameter by 4 inches in length. Closure is accomplished with press fit plastic end caps which are fitted to the open ends of brass tube after a sheet of aluminum foil is wrapped over the exposed sample material. A noncontributing/nonsubtractive tape is wrapped completely around the joint areas where the plastic caps meet the outer wall of the brass tube. No preservative other than cold storage is used on samples captured in sample containers of this type.

#### SAMPLE HANDLING PROCEDURES

Solid sample material is captured by advancing the liner into the soil. This may be done by pushing the liner into soft soils or by containing the liner in a drive shoe which can be advanced and then retracted by means of a slide hammer. The open ends of the sample liner are covered with aluminum foil and plastic end caps. Excess aluminum foil is removed and the edge of the plastic end caps is tightly sealed against the outer surface of the brass liner with an unbroken wrap made with a tape which has been tested to confirm that it does not contribute compounds that would be detected in the type of analyses intended for the sample contained inside of the brass liner. The brass liner is then labeled with the appropriate identification numbers which specify the sampling activity designation number, sample collection area, depth etc. that apply to that particular sample. The sample liner is then placed in an ice chest which contains pre-frozen blocks of an inert ice substitute such a Blue Ice or Super Ice.

Water samples are collected in any of several appropriate devices such as bailers, Coliwasas, Middleburg sampling pumps etc. which are described in detail only as warranted by their employment at a given site. Sample liquid is decanted into new sample containers in a manner which reduces the loss of volatile constituents and follows the applicable EPA procedures for handling volatile organic and semi-volatile compounds. Only two variations from the EPA methods are generally employed. First, preservative is added to the sample container prior to addition of the sample liquid. This method was pioneered by Stoner Laboratories in 1982 and subsequently adopted by laboratories and environmental consulting firms as a practical means of reducing the time that a liquid is allowed to aerate prior to closure of the sampling container. Second, because tests have shown that the preservative readily mixes with sample liquid, glass stirring rods are not used to agitate the sample/preservative mixture.

#### SAMPLE DESIGNATIONS

All samples containers are identified with both an activity number and a discrete sample identification. Please note that the activity number is the number that appears on our chain of custody. It is roughly equivalent to a job number, but applies only to work done on a particular day of the year rather than spanning several days as an actual activity often does. This is followed by the sample I.D. number which is usually a simple number such as #1, #2, #3.

#### CHAIN OF CUSTODY

Samples are continuously maintained in either a chilled ice chest, refrigerator, or freezer from the time of collection until acceptance by the State certified Hazardous Materials Testing Laboratory selected to perform the analytical procedures. If the samples are taken charge of by a different party (such as another person from our office, a courier etc.) prior to being delivered to the laboratory, appropriate release and acceptance records are made on the chain of custody (time, date, and signature of person releasing the samples followed by the time, date and signature of the person accepting custody of the samples).

### LABORATORY IDENTIFICATION NUMBERS

Following receipt of the samples and completion of the Chain of Custody form, the laboratory then assigns their own identification numbers to the samples. Different laboratories use different numbering systems and, according to their own internal conventions, may or may not assign sequential numbers to samples which are placed on temporary "hold", pending the results of other analyses. Laboratory identification numbers (if assigned and available) are included in the TABLE, and will be found on the certified analytical report by the analytical laboratory.

#### CERTIFIED ANALYTICAL REPORT

The certified analytical report (CAR) generated by the laboratory is the official document in which they issue their findings. The Results of Analyses section of the TABLE OF SAMPLING LOCATIONS AND ANALYTICAL RESULTS should correspond exactly with the laboratory's CAR. Any discrepancy between analytical values should be decided in favor of the CAR, for while it may, itself, be in error with regard to a particular number, the CAR remains the recognized document until such time as it is amended with a corrected report.

The certified analytical report should also be reviewed when samples are taken from below waste oil tanks as any detection of the EPA halogenated and purgeable aromatic compounds may be grounds for requiring further action. Also the TABLE OF SAMPLING LOCATIONS AND ANALYTICAL RESULTS is insufficiently spacious to allow anything more that a simple listing of the detected compounds. The TABLE does not include such information as the detection limits at which other compounds were not detected. The full text of the laboratory report will be found in Section Four of this report.

#### GENERAL ADVISORY ON POSITIVE RESULTS

Though our firm specializes in sampling, monitoring and documentation, rather than interpretation and remediation, we have been asked by the engineering staff of the Regional Water Quality Control Board to include in our reports an advisory section outlining the general type of additional actions which may be required when contamination is found. This advisory is not intended to characterize conditions at this particular site or replace the services of a consulting firm specializing in the investigation, characterization and remediation of such conditions as may exist. Rather, it is intended to advise you that such additional actions may be required even though some time may elapse before you are contacted by one of the interested regulatory agencies.

In Region 2 (which is regulated by the San Francisco Regional Water Quality Control Board,) the thresholds are readily defined in the Board's publication, <u>Guidelines For Addressing Fuel Leaks</u>. According to this document, soil which has less than 100 parts per million total petroleum fuel hydrocarbon (TPH) contamination does not generally require immediate additional action. Board engineers emphasize that this does not mean that some action might not be required in the future. Still, the site is assigned a low priority unless it is situated in an area of high hydrogeologic concern.

The detection of more than 100 ppm TPH in the native soil beneath a tank is generally considered grounds for requiring an additional investigation in the form of soil borings and installation of at least one groundwater monitoring well followed by periodic monitoring. The detection of 1000 ppm TPH is usually viewed by the Board as an unacceptable level of fuel saturation which will mandate excavation of the effected ground down to the furthest practicable reach of conventional excavating machinery followed by soil

borings and installation of groundwater monitoring wells.

Other regions use different standards for determining when a groundwater investigation will be required. For example benzene is often used in lieu of TPH. Even very low levels of benzene are often seen as grounds for requiring a subsurface investigation. This criteria may be relaxed or stiffened depending on the location of the site in relation to different groundwater systems, the depth to water, type of soil, and the concentrations of benzene involved.

The above standards apply only to fuels. When samples taken in connection with a waste oil tank or a solvent tank are found to contain even small amounts of any of the EPA priority pollutants (such as TCE, PCE, DCE etc. which are detected by EPA methods 8010, 8020, and 8240) more stringent standards are often applied. In these cases, soil borings and monitoring well installation may be required if there is any detectable amount of any of the EPA priority pollutant compounds.

When contaminants are found to have reached the water underlying a site, the Board customarily requires that additional work be undertaken in order to define the extent of the contamination.

#### REPORTAGE

Submission to the Regional Water Quality Control Board and the local implementing agency should include copies of the sampling report, the chain of custody, and the certified analytical report issued by the Hazardous Materials Testing Laboratory. The property owner should attach a cover letter and submit all documents together in a package.

The following addresses have been listed here for your convenience:

Water Quality Control Board San Francisco Bay Region 1111 Jackson Street Room 6040 Oakland, CA 94607 ATTN: Greg Zentner

Alameda County Health Hazardous Materials Management 420 27th Street Oakland, Ca 94612 ATTN: Storm Goranson

Oakland Fire Prevention Bureau One City Hall Plaza Oakland, CA 94612 ATTN: Ariel Bryant Please call if we can be of any further assistance.

Jordichard C. Blaine

RCB/ral

attachments: supporting documents

### LABORATORY RESULTS

Supporting documents

This section contains the following:

CHAIN OF CUSTODY ANAMETRIX ANALYTICAL REPORTS TABLE OF SAMPLING LOCATIONS AND ANALYTICAL RESULTS

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from	• :	am/pm	-88 -> to				/PM -88
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# ANAMETRIX, INC.

ENVIRONMENTAL • ANALYTICAL CHEMISTRY
2754 AJELLO DRIVE • SAN JOSE, CA 95111 • (408) 629-1132

April 26, 1988
Work Order Number 8804118
Date Received 04/22/88
Release No. 18
Station No. 1026
3701 Broadway, Oakland
Blaine Tech Project No.88113M1

Bill Barnard Chevron U.S.A., Inc. 2 Annabel Lane Suite 200 San Ramon, CA 94583

Eight soil samples were received for analysis of volatiles by GC/MS, total hydrocarbons as gasoline and diesel by GC, and total oil and grease by gravimetric analysis, using the following method(s):

ANAMETRIX I.D.	SAMPLE I.D.	METHOD(S)
8804118-01	9-1026 #1	8015/8020
-02	" #2	11
-03	11 #3	11
-04	11 #4	11
-05	" #5	u
-06	" <b>#</b> 6	8240/8015/503E
-07	" #7	8015/8020
-08	n #8	. 11

#### RESULTS

See enclosed data sheets, Pages 3 thru 11.

### EXTRA COMPOUNDS

See enclosed data sheet, Page 12.

### QUALITY ASSURANCE REPORTS

See enclosed data sheets, Pages 13 thru 14.

## ANAMETRIX, INC.

## ENVIRONMENTAL • ANALYTICAL CHEMISTRY 2754 AIELLO DRIVE • SAN JOSE. CA 95111 • (408) 629-1132

If there is any more that we can do, please give us a call. Thank you for using ANAMETRIX, INC.

Sincerely,

Burt Sutherland

Laboratory Manager

BWS/km

c/c: Blaine Tech Services

1370 Tully Rd., Ste. 505

San Jose, CA. 95122

### ORGANIC ANALYSIS DATA SHEET - EPA METHOD 624/8240 ANAMETRIX, INC. (408) 629-1132

Sample I.D. : 9-1026 #6 Anametrix I.D. : 8804118-06

Matrix : SOIL Analyst : ACL Date sampled : O4-22-88 Supervisor : pG

Date sampled: 04-22-88 Supervisor: pG Date analyzed: 04-22-88 Date released: 04-26-88

Dilution 1000 Instrument ID F1

10(10)		instrument in	: F1
CAS #	Compound Name	Reporting Limit (vg/Kg)	Amount Found (ug/Kg)
74-87-3	* Chloromethane	1 10000	BRL
75-01-4	* Vinyl Chloride	10000	BRL
74-83-9	* Bromomethane	10000	BRL
75-00-3	* Chloroethane	10000	BRL
75-69-4	* Trichlorofluoromethane	5000	BRL
75-35-4	* 1,1-Dichloroethene	5000	BRL
76-13-1	# Trichlorotrifluoroethane	5000	BRL
67-64-1	**Acetone	20000	ERL
75-15-0	**Carbondisulfide	5000	BRL
75-09-2	* Methylene Chloride	5000	BRL
156-60-5	* Trans-1,2-Dichloroethene	5000	BRL
75-34-3	* 1,1-Dichloroethane	5000	BRL
78-93-3	**2-Butanone	20000	BRL
156-59-2	* Cis-1,2-Dichloroethene	5000	BRL
67-66-3	* Chloroform	5000	BRL
71-55-6	* 1,1,1-Trichloroethane	5000	BRL
56-23-5	* Carbon Tetrachloride	5000	BRL
71-43-2	* Benzene	5000	BRL
107-06-2	* 1,2-Dichloroethane	5000	BRL
79-01-6	* Trichloroethene	5000	
78-87-5			BRL
75-27-4	1,2-Dichloropropane	5000	BRL
110-75-8	* Bromodichloromethane	5000	BRL
108-05-4	* 2-Chloroethylvinylether	5000	DRL
	**Vinyl Acetate	10000	BRL
10061-02-6	* Trans-1,3-Dichloropropene	5000	BRL
108-10-1	**4-Methyl-2-Pentanone	10000	BRL
108-88-3	* Toluene	5000	26000
10061-01-5	* cis-1,3-Dichloropropene	5000	BRL
79-00-5	* 1,1,2-Trichloroethane	5000	BRL
127-18-4	* Tetrachloroethene	5000	BRL
591-78-6	**2-Hexanone	10000	BRL
124-48-1	* Dibromochloromethane	5000	BRL
108-90-7	* Chlorobenzene	5000	BRL
100-41-4	* Ethylbenzene	5000	10000
1330-20-7	**Total Xylenes	5000	€8000
100-42-5	**Styrene	5000	BRL
75-25-2	* Bromoform	5000	BRL
79-34-5	* 1,1,2,2-Tetrachloroethane	5000	BRL
541-73-1	* 1,3-Dichlorobenzene	5000	BRL
106-46-7	* 1,4-Dichlorobenzene	5000	BRL
95-50-1	* 1,2-Dichlorobenzene	5000	BRL
CAS #	Surrogate Compounds	Limits	% Recovery
17060-07-0	1,2-Dichloroethane-d4	84-132%	103%
2037-26-5	Toluene-d8	76-131%	103%
460-00-4	p-Bromofluorobenzene	74-116%	93%

<sup>\*</sup> A Method 624 priority pollutant compound (Federal Register, 10/26/84)

<sup>\*\*</sup> A compound on the U.S. EPA CLP Hazardous Substance List (HSL) # A compound added by Anametrix, Inc. BRL: Below reporting limit.

 Sample I.D.: 9-1026 #1
 Anametrix I.D.: 8804118-01

 Matrix: SOIL
 Analyst
 : m/2

 Date sampled: 04-22-88
 Supervisor
 : 875

 Date anl. TVH: 04-23-88
 Date released
 : 04-26-88

Date ext. TEH: NA Date ext. TOG : NA Date anl. TEH: NA Date anl. TOG : NA

CAS #	Compound Name	Reporting Limit (ug/kg)	Amount Found (ug/kg)
71-43-2	Benzene	200	1600
108-88-3	Toluene	200	12000
100-41-4	Ethylbenzene	200	4400
1330-20-7	Total Xylenes	200	16000
	TVH as Gasoline	5000	260000

BRL - Below reporting limit.

TVH - Total Volatile Hydrocarbons is determined by modified EPA 8015 with either headspace or purge and trap.

TEH - Total Extractable Hydrocarbons is determined by modified EPA 8015 with direct injection.

TOG - Total Oil & Grease is determined by Standard Method 503E.

BTEX- Benzene, Toluene, Ethylbenzene, and Total Xylenes are determined by modified EPA 8020.

Sample I.D. : 9-1026 #2 : SOIL Matrix Date sampled: 04-22-88

Date an1. TVH: 04-23-88

Date ext. TEH: NA Date anl. TEH: NA

Anametrix I.D. 8804118-02

Analyst - mh Supervisor : 825

Date released : 04-26-88

Date ext. TOG : NA Date anl. TOG : NA

CAS #	Compound Name	Reporting Limit (ug/kg)	Amount Found (ug/kg)
71-43-2	Benzene	200	BRL
108-88-3	Toluene	200	300
100-41-4	Ethylbenzene	200	1200
1330-20-7	Total Xylenes	200	12000
	TVN as Gasoline	5000	88000

BRL - Below reporting limit.

TVH - Total Volatile Hydrocarbons is determined by modified EPA 8015 with either headspace or purge and trap.

TEH - Total Extractable Hydrocarbons is determined by modified EPA 8015 with direct injection.

TOG - Total Oil & Grease is determined by Standard Method 503E.

BTEX- Benzene, Toluene, Ethylbenzene, and Total Kylenes are determined by modified EPA 8020.

Sample I.D. : 9-1026 #3

Anametrix I.D. : 8804118-03
Analyst : And

Matrix : SOIL
Date sampled : 04-22-88
Date anl. TVH: 04-23-88

Supervisor : Fig Date released : 04-26-88

Date ext. TEH: NA Date anl. TEH: NA

Date ext. TOG : NA Date anl. TOG : NA

CAS #	Compound Name	Reporting Limit (ug/kg)	Amount Found (ug/kg)
71-43-2	Benzene	200	3300
108-88-3	Toluene	200	9500
100-41-4	Ethylbenzene	200	8900
1330-20-7	Total Xylenes	200	110000
	TVH as Gasoline	5000	890000

BRL - Below reporting limit.

TVH - Total Volatile Hydrocarbons is determined by modified EPA 8015 with either headspace or purge and trap.

TEH - Total Extractable Hydrocarbons is determined by modified EPA 8015 with direct injection.

TOG - Total Oil & Grease is determined by Standard Method 503E.

BTEX- Benzene, Toluene, Ethylbenzene, and Total Xylenes are determined by modified EPA 8020.

Anametrix I.D. : 8804118-04 Sample I.D. # 9-1026 #4 Analyst SOIL my Date sampled: 04-22-88 Supervisor 5 FW Date anl. TVH: 04-23-88 Date released 04-26-88 Date ext. TOG : NA Date ext. TEH: NA AN : Date anl. TEH: NA Date anl. TOG

CAS #	Compound Name	Reporting Limit (ug/kg)	Amount Found (ug/kg)
71-43-2	Benzene	200	008
08-88-3	Toluene	200	1400
100-41-4	Ethylbenzene	200	8300
1330-20-7	Total Xylenes	200	19000
	TVH as Gasoline	5000	480000

BRL - Below reporting limit.

TVH - Total Volatile Hydrocarbons is determined by modified EPA 8015 with either headspace or purge and trap.

TEH - Total Extractable Hydrocarbons is determined by modified EPA 8015 with direct injection.

TOG - Total Oil & Grease is determined by Standard Method 503E.

BTEX- Benzene, Toluene, Ethylbenzene, and Total Xylenes are determined by modified EPA 8020.

Date anl. TOG : NA

 Sample I.D.: 9-1026 #5
 Anametrix I.D.: 8804118-05

 Matrix: SOIL
 Analyst
 Implies the sample of the sampl

Reporting Amount Found Limit CAS # Compound Name (ug/kg) (ug/kg) ------|71-43-2 | Benzene 400 200 BRL BRL 108-88-3 | Toluene 200 1100-41-4 | Ethylbenzene 200 1330-20-7 |Total Xylenes 200 200 |TVH as Gasoline 5000 34000

BRL - Below reporting limit.

Date anl. TEH: NA

TVH - Total Volatile Hydrocarbons is determined by modified EPA 8015 with either headspace or purge and trap.

TEH - Total Extractable Hydrocarbons is determined by modified EPA 8015 with direct injection.

TOG - Total Oil & Grease is determined by Standard Method 503E.

BTEX- Benzene, Toluene, Ethylbenzene, and Total Xylenes are determined by modified EPA 8020.

Anametrix I.D. : 8804118-06 Sample I.D. : 9-1026 #6 : SOIL Analyst Matrix : mh Supervisor Date sampled: 04-22-88 2 Tas Date anl. TVH: NA Date released 04-26-88 Date ext. TEH: 04-22-88 Date ext. TOG : 04-25-88 Date an1. TEH: 04-25-88 Date anl. TOG : 04-25-88

CAS #	Compound Name	Reporting Limit (ug/kg)	Amount Found (ug/kg)
	H as Diesel tal Oil & Grease	10000	4300000 14000000

BRL - Below reporting limit.

TVH - Total Volatile Hydrocarbons is determined by modified EPA 8015 with either headspace or purge and trap.

TEH - Total Extractable Hydrocarbons is determined by modified EPA 8015 with direct injection.

TOG - Total Oil & Grease is determined by Standard Method 503E.

BTEX- Benzene, Toluene, Ethylbenzene, and Total Xylenes are determined by modified EPA 8020.

Sample I.D. : 9-1026 #7 Anametrix I.D. : 8804118-07

Matrix : SOIL Analyst : Mr.
Date sampled: 04-22-88 Supervisor : Mr.
Date anl. TVH: 04-22-88 Date released : 04-28

Date anl. TVH: 04-22-88

Date ext. TEH: NA

Date ext. TOG : NA

Date anl. TEH: NA

Date anl. TOG : NA

     CAS #	Compound Name	Reporting Limit (ug/kg)	Amount Found (ug/kg)	1
171-43-2	Benzene	200	BRL	
108-88-3	Toluene	200	BRL	i
100-41-4	Ethylbenzene	200	BRL	ĺ
1330-20-7	Total Xylenes	j 200	BRL	i
1	TVH as Gasoline	5000	BRL	İ
i	i	Ì		İ
į		1	1	

BRL - Below reporting limit.

TVH - Total Volatile Hydrocarbons is determined by modified EPA 8015 with either headspace or purge and trap.

TEH - Total Extractable Hydrocarbons is determined by modified EPA 8015 with direct injection.

TOG - Total Oil & Grease is determined by Standard Method 503E.

BTEX- Benzene, Toluene, Ethylbenzene, and Total Xylenes are determined by modified EPA 8020.

Sample I.D. : 9-1026 #8 Anametrix I.D. : 8804118-08

Matrix : SOIL Analyst : The Date sampled : 04-22-88 Supervisor : The Supervisor : The Date sampled : The Dat

Date anl. TVH: 04-22-88 Date released : 04-26-88 Date ext. TEH: NA Date ext. TOG : NA

Date ext. TEH: NA Date ext. TOG : NA Date anl. TEH: NA Date anl. TOG : NA

	Compound Name	Reporting	Amount
		Limit	Found
CAS #		(ug/kg)	(ug/kg)
71-43-2	Benzene	200	BRL
108-88-3	Toluene	200	BRL
100-41-4	Ethylbenzene	200	BRL
1330-20-7	Total Xylenes  TVH as Gasoline	200 5000	BRL BRL

BRL - Below reporting limit.

TVH - Total Volatile Hydrocarbons is determined by modified EPA 8015 with either headspace or purge and trap.

TEH - Total Extractable Hydrocarbons is determined by modified EPA 8015 with direct injection.

TOG - Total Oil & Grease is determined by Standard Method 503E.

BTEX- Benzene, Toluene, Ethylbenzene, and Total Xylenes are determined by modified EPA 8020.

# ORGANICS ANALYSIS DATA SHEET - TENTATIVELY IDENTIFIED COMPOUNDS ANAMETRIX, INC. (408) 629-1132

Sample I.D. : 9-1026 #6

Anametrix I.D.

: 8804118-06

Matrix : SOIL
Date Sampled : 04-22-88

Analyst TC
Supervisor BDS
Date Released 04-26-88

Analyzed VOA : 04-22-88 Dilution VOA : 1:1000

Analyzed SV : NA Dilution SV : NA

1	CAS #	  Scan#	Volatile Fraction Compound Name	Det.    Limit   ppb	Found
1 1 1 2 1 3 1 5 1 5 1 5 1 5 1 5 1 5 1 5 1 5 1 5	505-22-6 620-14-4 95-63-6 1120-21-4 1758-88-9	1123 1209 1346	decane 1-ethyl-3-methylbenzene 1,2,4-trimethylbenzene undecane 2-ethyl-1,4-dimethylbenzene	5000  5000  5000  5000  5000  5000  5000  5000	5000  15000  20000  5000  5000
	CAS #	  Scan#	Semivolatile Fraction Compound Name	Det.  Limit   ppb	Found
12345578901234567890				10 10 10 10 10 10 10 10 10 10 10 10 10 1	

Tentatively identified compounds are significant chromatographic peaks (TICs) other than priority pollutants. TIC spectra are compared with entries in the National Bureau of Standards mass spectral library. Identification is made by following US EPA guidelines and acceptance criteria. TICs are quantitated by using the area of the nearest internal standard and assuming a response factor of one (1). Values calculated are ESTIMATES ONLY.

## TOTAL VOLATILE HYDROCARBON MATRIX SPIKE REPORT EPA METHOD 8015

Sample I.D. : 9-1026 #7

Matrix : SOIL
Date sampled : 4-22-88
Date analyzed : 4-23-88

Anametrix I.D. : 8804118-

Analyst : mt Supervisor : &

Date Released : 4-25-88

COMPOUND	SPIKE AMT. (ug/Kg)	MS (ug/Kg)	%REC MS	MSD (ug/Kg)	%REC MSD	RPD	%RE LIMIT
Gasoline	10000	8600	86%	12000	120%	33%	50-:

<sup>\*</sup> Limits established by Anametrix, Inc.

## ORGANIC ANALYSIS DATA SHEET - EPA METHOD 624/8240

ANAMETRIX, INC. (408) 629-1132

Sample I.D. : METHOD BLANK Anametrix I.D. : 1CB0422V001

Matrix : SOIL Analyst Supervisor : Date sampled : NA Supervisor : DOS

Date analyzed: 04-22-88 Date released : 04-26-88

Dilution : NONE Instrument ID : F1

CAS #	Compound Name	Reporting Limit (ug/Kg)	Amount Found (ug/Kg)		
74-87-3	* Chloromethane	10	BRL		
75-01-4	* Vinyl Chloride	10	BRL		
74-83-9	* Bromomethane	10	BRL		
75-00-3	* Chloroethane	10	BRL		
75-69-4	* Trichlorofluoromethane	5	BRL		
75-35-4	* 1,1-Dichloroethene	5	BRL		
76-13-1	# Trichlorotrifluoroethane	5	BRL		
67-64-1	**Acetone	20	20		
75-15-0	**Carbondisulfide	5	BRL		
75-09-2	* Methylene Chloride	5	1		
156-60-5	* Trans-1,2-Dichloroethene	5	BRL		
75-34-3	* 1,1-Dichloroethane	5	BRL		
78-93-3	**2-Butanone	20	BRL		
156-59-2	* Cis-1,2-Dichloroethene	5	BRL		
67-66-3	* Chloroform	5	BRL		
71-55-6	* 1,1,1-Trichloroethane	5	BRL		
56-23-5	* Carbon Tetrachloride	5	BRL		
71-43-2	* Benzene	5	BRL		
107-06-2	* 1,2-Dichloroethane	5	BRL		
79-01-6	* Trichloroethene	5	BRL		
78-87-5	* 1,2-Dichloropropane	5	BRL		
75-27-4	* Bromodichloromethane	5	BRL		
110-75-8	* 2-Chloroethylvinylether	5	BRL		
108-05-4	**Vinyl Acetate	10	BRL		
10061-02-6	* Trans-1,3-Dichloropropene	5	BRL		
108-10-1	**4-Methyl-2-Pentanone	10	BRL		
108-88-3	* Toluene	5	BRL		
10061-01-5	* cis-1,3-Dichloropropene	5	BRL		
79-00-5	* 1,1,2-Trichloroethane	5	BRL		
127-18-4	* Tetrachloroethene	5	BRL		
591-78-6	**2-Hexanone	10	BRL		
124-48-1	* Dibromochloromethane	5	BRL		
108-90-7	* Chlorobenzene	5	1		
100-41-4	* Ethylbenzene	5	BRL		
1330-20-7	**Total Xylenes		BRL		
100-42-5	**Styrene	5 5	BRL BRL		
75-25-2	* Bromoform	5			
79-34-5		5	BRL BRL		
541-73-1	* 1,1,2,2-Tetrachloroethane	5	BRL		
106-46-7	* 1,3-Dichlorobenzene		BRL		
95-50-1	* 1,4-Dichlorobenzene  * 1,2-Dichlorobenzene	5 5	BRL		
CAS #	Surrogate Compounds	Limits	% Recovery		
17060-07-0	1,2-Dichloroethane-d4	84-132%	93%		
2037-26-5	Toluene-d8	76-131%	100%		
460-00-4	p-Bromofluorobenzene	74-116%	104%		

<sup>\*</sup> A Method 624 priority pollutant compound (Federal Register, 10/26/84)

<sup>\*\*</sup> A compound on the U.S. EPA CLP Hazardous Substance List (HSL)
# A compound added by Anametrix, Inc. BRL: Below reporting limit.

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## TABLE OF SAMPLING LOCATIONS AND ANALYTICAL RESULTS

ANALYTICAL RESULTS
IN PARTS PER MILLION -- PPM

Í.D.	SAMPLE		TYPE 4							IN PARTS PER HILLION PPH				
GIVEN THIS SAMPLE ARBA	DEPTH IN FT. BELOW GRADE	SAMPLING LOCATION DICTATED BY	METHOD FOR THE SAMPLE OBTAINED	SAMPLE MATRIX	DATR SAMPLED	BTS CHAIN OF CUSTODY I.D.	BTS SAMPLE 1.D.	NAME OF DOBS HATL LABORATORY	LABORATORY SAMPLE 1.D.	TPH A8 <u>GAS</u>	BBN- Zene	TOL- UENR	IY-	BTHYL, BBN- 2BNB
AP	14.5	STANDARD	CAPILLAR	SOIL	4/22/88	88113-M-1	‡3	ANAMETRIX	8804118-03	890	3.3	9.5	110	8.9
Aop .	14.5	STANDARD	CAPILLAR	SOIL	4/22/88	88113-M-1	#2	AWAMETRIX	8804118-02	88	ND	0.3	12	1.2
Вор	14.5	STANDARD	CAPILLAR	SOIL	4/22/88	88113-M-1	<b>#</b> 1	ANAMETRIX	8804118-01	260	1.6	12	16	4,4
CF	14.5	STANDARD	CAPILLAR	SOIL	4/22/88	88113-M-1	<b>‡</b> 5	ANAMETRIX	8804118-05	34	0.4	ND	0.2	ND
Cop	14.5	STANDARD	CAPILLAR	SOIL	4/22/88	88113-M-1	H	ANANETRIX	8804118-04	480	0.8	1.4	19	8.3
DP	12.5	STANDARD	CAPILLAR	SOIL	4/22/88	88113-M-1	<b>‡</b> 8	ANAKETELY	8804118-08	ND	ND	ND	ND	ND
Dop	12.5	STANDARD	CAPILLAR	SOIL	4/22/88	88113-K-1	<b>\$</b> 7	ANAMETRIX	8804118-07	ND	ND	ND	ND	ND
I.D. GIVEN THIS SAMPLE ARBA	SAMPLE DEPTH IN FT. BELOW GRADE	SAMPLING LOCATION DICTATED BY	TYPE & METHOD FOR THE SAMPLE OBTAINED	SAMPLE MATRIX	DATR Sampled	BTS CHAIN OF CUSTODY I.D.	BTS SAKPLB I.D.	NAME OF DOES ENTL LABORATORY	LABORATORY SAMPLE I.D.	ANALYTII PPM TPH-BBP DIRSEL	PPN TOTAL A GRE	OIL	(PARTS PE PPB BPA 8240 COKPOUNDS	R BILLION)
Voll	10.0	STANDARD	INTERACE	SOIL	4/22/88	88113-M-1	<b>‡</b> 6	ANAWETRIX	8804118-06	4300	14,0	00	SBB LAB B	EPORT