BASELINE

No 116

ENVIRONMENTAL CONSULTING

24 September 2001 98381-B0

Ms. Ann E. Johnston COBLENTZ, PATCH, DUFFY, & BASS, LLP 222 Kearny Street, 7th Floor San Francisco, CA 94108-4510

Subject: Report on Passive Gas Sampling and Work Plan for Additional Investigation, 6623 San Pablo Avenue, Oakland, California

Dear Ann:

This letter documents recent subsurface investigation activities at 6623 San Pablo Avenue in Oakland (Figure 1). In accordance with a BASELINE work plan dated 31 May 2001, approved by the Alameda County Health Agency (County) in a letter 6 June 2001, BASELINE conducted a passive soil gas survey at the project site.

A passive soil gas survey can be an effective method for identification of source areas of chemical releases and delineation of plumes of volatile and semi-volatile compounds in soil and groundwater. In comparison with active soil gas surveys (where soil gas is actively pumped from the subsurface), passive soil gas surveys are effective in low permeability environments. The conditions at the project site, including low permeability surface soils, promoted the selection of a passive soil gas survey survey.

Field Activities

On 25 June 2001, a geologist from BASELINE installed 31 GORE-SORBERSM modules in a grid pattern at the project site (Figure 2). The modules contain a chemically inert expanded polytetrafluoroethylene membrane (GORE-TEX). The membrane is microporous and hydrophobic, which facilitates the efficient absorption of soil gas. One-inch borings were made with an electrically-powered hammer drill to depths ranging from 2.5 to 3.0 feet. Upon completion of each boring, the GORE-SORBERSM module and attached retrieval cord were set into the boring with a stainless steel insertion rod. The rod was removed from the boring. A one and one half-inch cork (to which the retrieval cord was attached) was placed in the top of the boring to minimize the potential for surface runoff entering the boring. In addition, each boring was temporarily sealed with a small amount of hydraulic cement (placed over the cork) to minimize the potential for tampering. Following use at each probe location, the drill bit and insertion rod were cleaned with a solution of trisodium phosphate and water and then rinsed

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with deionized water. The boring at each probe location was backfilled with cement grout after probe retrieval.

As recommended by the manufacturer, the modules were retrieved by a BASELINE geologist on 9 July 2001, 14 days after installation. Immediately upon retrieval, the modules were placed in labeled glass jars. The sealed jars were placed in a cooled container along with three travel blanks. The modules were shipped under chain-of-custody to W.L. Gore & Associates, Inc. (Gore) laboratory in Elkton, Maryland.

At the Gore laboratory, each module (including travel blanks) was treated with thermal desorption to liberate volatile and semi-volatile compounds and analyzed using gas chromatography. The analyses provide a measure of the total mass of each analyte that was collected on and desorbed from each of the modules. A report presenting a description of the testing methodology and the results of the analysis of the modules is enclosed as Attachment A. The report includes a color graphical presentation of the distribution (contouring) of the total mass of benzene, total BTEX, and gasoline-range petroleum hydrocarbons adsorbed to the modules. It should be noted that the colors presented in the contouring represent a standardized number of contour intervals between the highest and lowest detected values for the individual compounds. Therefore, the intervals (and corresponding colors) are not consistent for each compound.

Results and Conclusions

The results of the passive soil gas sampling survey are shown on Figures A-1 through A-3 in Attachment A (and the raw data are tabulated in Attachment A). The mapped spatial patterns showed fairly well defined soil gas plumes for benzene (Figure A-1), total BTEX (Figure A-2), and gasoline-range petroleum hydrocarbons (Figure A-3). MTBE was identified at only two locations (362835 and 362839), and therefore was not contoured for graphical presentation. It should be noted that the positive results for MTBE coincided with benzene and total BTEX soil gas "hotspots."

The results provide empirical data regarding the on-site locations where soil gas containing elevated levels of volatile organic hydrocarbons reach the surface. These data are useful when evaluating the potential exposure pathway of volatilization of VOCs to indoor air (which has been identified as a potential route of exposure at the site). In addition, it provides information on the potential extent of contaminants in the underlying soil and groundwater.

In general, there appears to be a correlation between VOCs identified in soil and groundwater as part of past sampling activities and the results of the soil gas survey. The soil sample collected from KB-1 (Table 1) in 1996 represents the highest benzene level identified in soils at the site to date (13 mg/kg) and is near the location of a "hotspot" identified by the passive soil howers.

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> gas sampling survey (located approximately 100 feet north of the restaurant building, as shown on Figure A-1). However, it does not appear that a precise correlation exists. For example, past sampling at location MW-3A indicates that benzene is present at elevated levels in soil (0.66 mg/kg in January 1999; Table 1) and groundwater (0.26 mg/L in January 2001; Table 2), but appears to be non-detect ($< 0.030 \ \mu g$) for soil gas. MTBE is also issue.

> The color contour image representing the distribution of benzene soil gas (which is expected to be the primary contaminant of concern in the risk assessment analysis that will be completed at a later date) defines the extent of near-surface benzene soil gas in the northwestern, northern, and eastern portions of the site (Figure A-1). In addition, the relative mass of benzene in nearsurface soil gas appears to decrease significantly to the south and southwest (Figure A-1).

own durand White for the past soil borings or monitoring wells has been installed at this location, and therefore data regarding the potential worst-case location (for volatilization of benzene to the surface have been identified by collection. indicates that a previously unexplored "hotspot" of gasoline-range hydrocarbons may be present near the southwest property boundary.

Recommendations (Work Plan)

Install two monitoring wells in the approximate locations shown on Figure 2. Proposed well MW-4A would be installed to characterize soil and groundwater quality in the vicinity of the benzene soil gas hotspot (Figure A-1). Proposed monitoring well MW-5A would be installed to investigate soil and groundwater quality conditions near the western property boundary where the soil gas survey indicated elevated levels of gasoline-range hydrocarbons (Figure A-3) and slightly elevated benzene levels. The purpose of installation and sampling of these wells is to characterize the potential worst-case conditions for benzene and gasoline-range petroleum hydrocarbons in soil and groundwater at the site.

The borings would be advanced using a direct-push rig and 2.5-inch casing. Each boring would be logged (continuous) in accordance with the Unified Soil Classification system. Up to three soil samples would be collected for analysis at each location, depending on field conditions encountered. All sample-driving equipment would be decontaminated between borings by steam-cleaning or washing with trisodium phosphate and rinsed with deionized water.

Each soil sample would be labeled, sealed with teflon film, plastic caps, and silicone tape. All samples would be stored in a cooled container and transported under chain-of-custody

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procedures to a state-certified analytical laboratory. Each sample would be analyzed for TPH as gasoline, BTEX, and MTBE. In addition, select soil samples would be analyzed for total organic carbon, moisture content, and bulk density to support analysis of potential residual health risks using the Risk-Based Corrective Action (RBCA) protocol.

The wells would be constructed with 3/4-inch PVC casing and screen (0.010 slot). The screen would be wrapped with a pre-fabricated filter pack. The location of the screened interval would depend on field conditions, but would be installed to account for shallow groundwater fluctuations, and would not exceed ten feet in length. The upper five feet of the boring would be hand-augered (six inches in diameter) to allow for utility clearance and installation of a sanitary seal. All work associated with the well installation would be supervised by a Registered Geologist. The top of casing of the new monitoring wells would be surveyed by a licensed surveyor to determine elevations to within 0.01 foot. Not sooner than 48 hours after completion, the wells would be developed by surging and pumping until relatively clear water was produced.

The newly installed wells would be sampled on a semi-annual basis, similar to the current approved sampling schedule for the existing wells at the site. Prior to sampling, groundwater levels would be measured (and the presence or absence of floating product determined, and, if present, the thickness measured) in each of the new wells using a dualinterface probe. To the extent practicable, the wells would be purged until physical parameters measured in the field (pH, EC, and temperature) had stabilized. Samples would then be collected directly from the flowing purge water¹ in glassware provided by the laboratory. Experience with the existing wells at the site indicates that the upper water-bearing zone in some locations is extremely low-yield, rendering traditional "purge until stabilized" methods infeasible.

All excess soil cores and wastewater (decontamination, development, and purge water) would be disposed of off-site at an approved facility.

Prior to initiation of well installation, appropriate permits would be acquired from the Alameda County Public Works Agency. All on-site activities would be conducted under a site-specific health and safety plan.

¹ BASELINE uses a peristaltic pump and new disposable tubing on each well. Purge water never comes into contact with the components of the pumping system, only with the disposable tubing, minimizing the potential for cross-contamination between wells.



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> A report would be prepared for submittal to the County that summarizes field activities and analytical results and provides recommendations for additional activities, as appropriate.

One copy of this report should be submitted to Mr. Barney Chan of the Alameda County Health Care Services Agency. Should you have any questions, or require additional clarification, please contact us at your convenience.

Sincerely,

Bruce Abelli-Amen Bruce Abelli-Amen

Project Manager

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Yane Nordhav Principal Reg. Geologist No. 4009

BAA:YN:cr Enclosure

Ms. Helen Loreto, McDonald's Corporation cc:

REGIONAL LOCATION

Figure 1



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6623 San Pablo Avenue Oakland, California



SITE PLAN

Figure 2

1



Legend

Location of Geophysical Anomally (two borings installed within the bounds of mapped anomally to confirm that no UST is present) Soil Boring Location (Kleinfelder) B-1 MW-2A \oplus Monitoring Well Location (BASELINE) Approximate Passive Soil Gas Sampling Location ο MW-4A 🔶 Proposed Monitoring Well Location 0 30 Feet 6623 San Pablo Avenue BASELINE

Oakland, California

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ATTACHMENT A

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Result of Passive Soil Gas Survey



W. L. GORE & ASSOCIATES, INC.

100 CHESAPEAKE BLVD., P.O. BOX 10 • ELKTON, MARYLAND 21922-0010 • PHONE: 410/392-7600 FAX: 410/506-4780

> GORE-SORBER® EXPLORATION SURVEY GORE-SORBER® SCREENING SURVEY

1 of 6

GORE-SORBER[®] Screening Survey Final Report

6623 San Pablo Ave. Oakland, CA

July 25, 2001

Prepared For: Baseline Environmental Service 101 H Street, Suite L Petaluma, CA 94952

W.L. Gore & Associates, Inc.

Written/Submitted by: Jim Whetzel, Project Manager

Reviewed/Approved by: Ray Fenstermacher Project Manager

Analytical Data Reviewed by: Jim E. Whetzel, Chemist

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GORE-SORBER[®] Screening Survey **Final Report**

REPORT DATE: July 25, 2001

AUTHOR: JW

SITE INFORMATION

Site Reference: 6623 San Pablo Ave., Oakland, CA Customer Purchase Order Number: 98381-80 Gore Site Code: BVN Gore Production Order Number: 10732467

FIELD PROCEDURES

Modules shipped: 34 Installation Date(s): 6/25/01 Field work performed by: Baseline Environmental Service

Retrieval date(s): 7/9/01 # Modules Retrieved: 31 # Modules Lost in Field: 0 # Modules Installed: 31

Exposure Time: 14 [days] # Trip Blanks Returned: 3 # Unused Modules Returned: 0

Date/Time Received by Gore: 7/11/01 @ 7:30 AM Chain of Custody Form attached: $\sqrt{}$ Chain of Custody discrepancies: None Comments: Modules 362857, 362858, 362859 were identified as trip blanks.

By: MM

GORE-SORBER[®] Screening Survey Final Report

ANALYTICAL PROCEDURES

W.L. Gore & Associates' Screening Module Laboratory operates under the guidelines of its Quality Assurance Manual, Operating Procedures and Methods. The quality assurance program is consistent with Good Laboratory Practices (GLP) and ISO Guide 25, "General Requirements for the Competence of Calibration and Testing Laboratories", third edition, 1990.

Instrumentation consists of state of the art gas chromatographs equipped with mass selective detectors, coupled with automated thermal desorption units. Sample preparation simply involves cutting the tip off the bottom of the sample module and transferring one or more exposed sorbent containers (sorbers, each containing 40mg of a suitable granular adsorbent) to a thermal desorption tube for analysis. Sorbers remain clean and protected from dirt, soil, and ground water by the insertion/retrieval cord, and require no further sample preparation.

Analytical Method Quality Assurance:

The analytical method employed is a modified EPA method 8260/8270. Before each run sequence, two instrument blanks, a sorber containing 5μ g BFB (Bromofluorobenzene), and a method blank are analyzed. The BFB mass spectra must meet the criteria set forth in the method before samples can be analyzed. A method blank and a sorber containing BFB is also analyzed after every 30 samples and/or trip blanks. Standards containing the selected target compounds at three calibration levels of 5, 20, and 50μ g are analyzed at the beginning of each run. The criterion for each target compound is less than 35% RSD (relative standard deviation). If this criterion is not met for any target compound, the analyst has the option of generating second- or third-order standard curves, as appropriate. A second-source reference standard, at a level of 10μ g per target compound, is analyzed after every ten samples and/or trip blanks, and at the end of the run sequence. Positive identification of target compounds is determined by 1) the presence of the target ion and at least two secondary ions; 2) retention time versus reference standard; and, 3) the analyst's judgment.

NOTE: All data have been archived. Any replicate sorbers not used in the initial analysis will be discarded fifteen (15) days from the date of analysis.

Laboratory analysis: thermal desorption, gas chromatography, mass selective detection Instrument ID: #3 Chemist: JW/WW Compounds/mixtures requested: Gore Fuel Hydrocarbons (A2) Deviations from Standard Method: None

Comments: Soil vapor analytes and abbreviations are tabulated in the Data Table Key (page 6).

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DATA TABULATION

CONTOUR MAPS ENCLOSED: Three (3) B-sized color contour maps LIST OF MAPS ENCLOSED:

- Benzene, Toluene, Ethyl benzene, and total Xylenes (BTEX)
- Gasoline Range Petroleum Hydrocarbons (GRPH)
- Benzene (BENZ)

NOTE: All data values presented in Appendix A represent masses of compound(s) desorbed from the GORE-SORBER Screening Modules received and analyzed by W.L. Gore & Associates, Inc., as identified in the Chain of Custody (Appendix A). The measurement traceability and instrument performance are reproducible and accurate for the measurement process documented. Semi-quantitation of the compound mass is based on either a single-level (QA Level 1) or three-level (QA Level 2) standard calibration.

General Comments:

- This survey reports soil gas mass levels present in the vapor phase. Vapors are subject to a variety of attenuation factors during migration away from the source concentration to the module. Thus, mass levels reported from the module will often be less than concentrations reported in soil and groundwater matrix data. In most instances, the soil gas masses reported on the modules compare favorably with concentrations reported in the soil or groundwater (e.g., where soil gas levels are reported at greater levels relative to other sampled locations on the site, matrix data should reveal the same pattern, and vice versa). However, due to a variety of factors, a perfect comparison between matrix data and soil gas levels can rarely be achieved.
- Soil gas signals reported by this method cannot be identified specifically to soil adsorbed, groundwater, and/or free-product contamination. The soil gas signal reported from each module can evolve from all of these sources. Differentiation between soil and groundwater contamination can only be achieved with prior knowledge of the site history (i.e., the site is known to have groundwater contamination only).
- QA/QC trip blank modules were provided to document potential exposures that were not part of the soil gas signal of interest (i.e., impact during module shipment, installation and retrieval, and storage). The trip blanks are identically manufactured and packaged soil gas modules to those modules placed in the subsurface. However, the trip blanks remain unopened during all phases of the soil gas survey. Levels reported on the trip blanks may indicate potential impact to modules other than the contaminant source of interest.

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• Unresolved peak envelopes (UPEs) are represented as a series of compound peaks clustered together around a central gas chromatograph elution time in the total ion chromatogram. Typically, UPEs are indicative of complex fluid mixtures that are present in the subsurface. UPEs observed early in the chromatogram are considered to indicate the presence of more volatile fluids, while UPEs observed later in the chromatogram may indicate the presence of less volatile fluids. Multiple UPEs may indicate the presence of multiple complex fluids.

Project Specific Comments:

- The minimum (gray) contour level, for each mapped analyte or group of analytes, was set at the maximum blank level observed or the method detection limit, whichever was greater. When target compounds are summed together (i.e., BTEX), the contour minimum is arbitrarily set at 0.02 µg or the maximum blank level, whichever is greater. The maximum contour level was set at the maximum value observed.
- Stacked total ion chromatograms (TIC's) are included in Appendix A. The six-digit serial number of each module is incorporated into the TIC identification (e.g.: <u>123456</u>S.D represents module #<u>123456</u>).
- No target compounds were detected on the trip blanks and/or the method blanks. Thus, target analyte levels reported for the field-installed modules that exceed trip and method blank levels, and the analyte method detection limit, have a high probability of originating from on-site sources.
- The mapped spatial patterns showed fairly well defined soil gas plumes.
- If the objective of the soil gas survey was to delineate the nature and extent of the contamination, then additional soil gas sampling is recommended in those areas where the color contours appear to extend into unsampled areas. Subsequent sampling events can be combined with the data from this event and mapped together to provide greater coverage.

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KEY TO DATA TABLE 6623 San Pablo Ave., Oakland, CA

UNITS	
μg	micrograms (per sorber), reported for compounds
MDL	method detection limit
bdl	below detection limit
nd	non-detect
ANALYTES	
GRPH	gasoline range petroleum hydrocarbons
DRPH	diesel range petroleum hydrocarbons
BTEX	combined masses of benzene, toluene, ethylbenzene and total xylenes
	(Gasoline Range Aromatics)
BENZ	benzene
TOL	toluene
EtBENZ	ethylbenzene
mpXYL	m-, p-xylene
oXYL	o-xylene
C11,C13&C15	combined masses of undecane, tridecane, and pentadecane (C11+C13+C15) (Diesel Range Alkanes)
UNDEC	undecane
TRIDEC	tridecane
PENTADEC	pentadecane
TMBs	combined masses of 1,3,5-trimethylbenzene and 1,2,4-trimethylbenzene
135TMB	1,3,5-trimethylbenzene
124TMB	1,2,4-trimethylbenzene
NAPH&2-MN	combined masses of naphthalene and 2-methyl naphthalene
NAPH	naphthalene
2MeNAPH	2-methyl naphthalene
MTBE	methyl t-butyl ether
OCT	octane
BLANKS	
TBn	unexposed trip blanks, travels with the exposed modules
method blank	QA/QC module, documents analytical conditions during analysis

APPENDIX A:

CHAIN OF CUSTODY DATA TABLE STACKED TOTAL ION CHROMATOGRAMS COLOR CONTOUR MAPS

GORE-SORBER[®] Screening Survey Chain of Custody

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Clearing Technologies	
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GORE-SORBER @ Screening Survey is a registered service mark of W.L. Gore & Associates, Inc.

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	07/19/01	362820	1.92	1.12	0.00	nd	nd	nd	nd	nđ	0.00	bdl	bdl
	07/19/01	362821	0.01	0.03	0.00	nd	nd	nđ	nd	nd	0.00	nd	nd
	07/19/01	362822	0.03	0.02	0.04	nd	nd	nd	0.02	0.02	0.00	nd	nd
	07/19/01	362823	0.00	0.03	0.00	nd	nd	nđ	nd	nd	0.00	nd	bn Dn
	07/19/01	362824	0.02	0.01	0.00	nd	nd	nd	nd	nd	0.00	nd	nd
	07/19/01	362825	1200.83	్ 28.81	9.85	2.27	2.38	2.71	1.52	0.97	0.35	0.23	0.10
	07/19/01	362826	101.04	3.02	1.07	0.67	0.13	0.04	0.13	0.10	0.06	0.06	bdl
ļ	07/19/01	362827	7.27	0.04	0.00	nd	nd	nd	nd	nd	0.00	nd	nd
	07/19/01	362828	0.01	0.01	0.00	nd	nd	nd	nd	nd	0.00	nd	nd
	07/19/01	362835	618.09	<u>*</u> 8.67	98.64	83.47	2.84	5.79	5.82	0.72	0.16	0.16	bdl
ļ	07/19/01	362836	356.83	1.52	0.28	0.15	0.06	bdi	0.04	0.03	0.00	bdl	nd
ļ	07/19/01	362837	0.06	0.01	0.00	nd	nd	nd	nd	bdl	0.00	nd	nd
	07/19/01	362838	nd	0.02	0.00	nd	nd	nd	nd	nd	0.00	nd	nđ
ļ	07/19/01	362839	345.93	12.29	94.20	85.39	2.29	2.91	3.19	0.42	0.50	0.33	0.08
	07/19/01	362840	3.85	0.07	0.30	0.08	0.06	nd	0.09	0.07	0.00	nd	nd
┟	07/20/01	362841	1.87	0.01	0.04	nd	nd	nd	0.02	0.02	0.00	nď	nd
ļ	07/20/01	362842	385.02	2.11	0.35	0.10	0.09	0.02	0.09	0.05	0.03	0.03	nd
	07/20/01	362843	0.02	0.02	0.00	nd	nd	лd	nd	nd	0.00	nd	nd
-	07/20/01	362844	702.53	2.16	249.63	228.00	4.58	12.03	4.75	0.27	0.08	0.08	лd
ļ	07/20/01	362845	106.47	0.42	0.83	0.40	0.16	0.04	0.14	0.09	0.00	nd	nd
ļ	07/20/01	362846	nd	0.02	0.00	nd	nd	nd	nd	nd	0.00	nd	nd
1	07/20/01	362847	214.91	1.43	3.21	2.58	0.26	0.08	0.22	0.07	0.06	0.06	nd
-1	07/20/01	362848	118.40	2.06	3.93	3.51	0.09	0.20	0.10	0.03	0.16	0.13	0.03
	07/20/01	362849	450.59	1.69	236.16	221.36	1.78	10.96	1.90	0.16	0.07	0.07	nd
	07/20/01	362850	290.84	1.83	1.22	0.71	0.22	0.06	0.16	0.07	0.04	0.04	bdl
	07/20/01	362851	0.27	0.09	0.02	nd	nd	nd	nd	0.02	0.00	nd	nd
	07/20/011	302052	563.06	19.05	0.34	0.03	0.03	0.04	0.14	0.10	0.12	0.07	0.02
ļ	07/20/01	362853	45.00	2.64	0.47	0.20	0.10	0.03	0.08	0.06	0.04	0.04	nd
ļ	07/20/01	362854	22.67	5.36	0.15	0.08	0.04	nd	0.03	bdl	0.04	0.04	bdl
	07/20/01	362855	10.55	19.90	0.25	0.10	0.06	nd	0.05	0.04	0.57	0.10	0.14
	07/20/01	362856	0.02	0.02	0.00	nd	nd	nd	nd	nd	0.00	nd	nd
		000057					ļ						
	07/19/01	362857	0.06	0.02	0.00	nd	nd	nd	nd	nd	0.00	nd	nd

No mdl is available for summed combinations of analytes. In summed columns (eg., BTEX), the reported values should be considered ESTIMATED if any of the individual compounds were reported as bdl.

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I DATE	SAMPLE					· · · · · · · · · · · · · · · · · · ·				······································	·····	
ANALYZED		GRPH, ug	DRPH, ug	BTEX, ug	BENZ, ug	TOL, ug	EtBENZ, ug	mpXYL. ua	oXYL. ua	C11 C13 &C15 un	UNDEC up	
	MDL=				0.03	0.03	0.02	0.02	0.02	· · · · · · · · · · · · · · · · · · ·		TRIDEC, ug
07/20/04	262060	0.05		0.00		0.00	0.02	0.02	0.02		0.03	0.02
0//20/01	302008	0.05	0.01	0,00	nd	nd nd	nd	nd	nd	0.00	nd	nd
07/20/01	362859	0.04	0.01	0.00	nd	nd	nd	nd		0.00		114
								110		0.00	na	. nd
07/40/04	mode of the		0.00									
0//19/01	method blank	na	0.00	0.00	nd	nd]	nd	nd	nd	0.00	nd	nd
07/20/01	method blank	ndi	nd	0.00	nd	nd	nd	hn	ba	0.00		110
									110	0.00	na	na
	Marine	4000.00										
ļ	Maximum	1200.83	28.81	249.63	228.00	4.58	12.03	5.82	0.97	0.57	0.33	0.14
}	Standard Dev.	283.37	6.98	63.50	58.45	1.09	3.03	1 / 3	0 22	0.14	0.00	
	Moan	178.07	2 60	22.64	20.00	0.40		1,40	0.22	0.14	0.08	0.03
	Moun	170.97	3.09	22.01	20.29	0.49	1.13	0.60	0.11	0.07	0.05	0.01

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SAMPLE									
NAME	PENTADEC, ug	TMBs, ug	124TMB, ug	135TMB, ug	NAPH&2-MN, ug	NAPH, ug	2MeNAPH, ug	MTBE, ug	OCT, ug
MDL=	0.02		0.02	0.02		0.02	0.02	0.48	0.10
362820	bdl	0.00	bdl	nd	0.00	nd	nd	bdi	nd
362821	nd	0.00	nd	nd	0.00	nd	nd	nd	nd
362822	nd	0.00	nd	nd	0.00	nđ	nd	nd	nd
362823	nd	0.00	nd	nd	0.00	nd	nd	nd	nd
362824	nd	0.00	nd	nd	0.00	nd	nd	bdi	nd
362825	0.02	5.22	2.81	2.41	3.55	1.12	2.43	bdl	5.38
362826	bdł	0.06	0.03	0.03	0.03	0.03	bdl	nd	0.21
362827	nd	0.00	nd	nd	0.00	nd	nd	nd	nd
362828	nd	0.00	nd	nd	0.00	nd	nd	nd	nd
362835	กป	5.87	3.75	2.12	0.29	0.29	bdl	40.55	1.24
362836	nd	0.00	nd	nd	0.00	nd	nd	bdl	0.40
362837	nd	0.00	nd	nd	0.00	nď	nd	nd	nd
362838	nd	0.00	nd	nd	0.00	nd	nd	nd	nd
362839	0.09	7.14	4.57	2.57	0.49	0.46	0.03	2.74	0.70
362840	nd	0.00	nd	nd	0.00	nd	nd	nd	nd
362841	nd	0.00	nd	nd	0.00	nd	nd	nd	nd
362842	nd	0.00	bdl	bdl	0.00	nd	nd	bdl	0.65
362843	nd	0.00	nd	nd	0.00	nd	nd	nd	nd
362844	nd	2.70	1.21	1.49	0.22	0.22	bdl	bdl	2.20
362845	nd	0.04	0.02	0.02	0.00	nd	nd	bdl	0.17
362846	nd	0.00	nd	nd	0.00	nd	nd	nd	nd
362847	nd	0.22	0.10	0.12	0.00	nd	nd	bdl	0.37
362848	bdl	0.65	0.14	0.51	0.15	0.15	bdl	bdl	0.20
362849	nd	3.35	2.27	1.08	0.07	0.07	nd	bdl	0.62
362850	bdl	0.15	0.10	0.05	0.00	nd	nd	bd	0.39
362851	nd	0.00	nd	nd	0.00	nd	nd	nd	лd
362852	0.03	0.05	0.03	0.02	0.02	0.02	bdl	лd	0.50
362853	nd	0.13	0.08	0.05	0.00	bdl	nd	nd	0.11
362854	bdl	0.15	0.04	0.11	0.00	nd	nd	bdl	nd
362855	0.33	0.10	0.07	0.03	0.08	0.02	0.06	bdl	nd
362856	nd	0.00	nd	nd	0.00	nd	nd	nd	nd
362857	nd	0.00	nd	nd	0.00	nd	nd	nd	nd

No mdl is available for summed combinations of analytes. In summed columns (eg., BTEX), the reported values should be considered

ESTIMATED if any of the individual compounds were reported as bdl.

I SAMPLE									
NAME	PENTADEC, ug	TMBs, ug	124TMB, ug	135TMB, ug	NAPH&2-MN, ug	NAPH, ua	2MeNAPH, ug	MTBE un	OCT un
MDL=	0.02		0.02	0.02		0.02	0.02	0.48	0 10
362889	nd	0.00	nd	nd	0.00	nd	bu	nd	nd
362859	nd	0.00	nd	nd	0.00	nd	nd	nd	nd
method blank	nd	0.00	nd	nd	0.00	nd	nd	nd	nd
method blank	nd	0.00	nd	nd	0.00	nd	nd	nd	nd
Maximum	0.33	7.14	4.57	2.57	3 55	1 1 2	2 / 2	40.55	E 20
Standard Dev.	0.06	1.92	1.18	0.75	0.64	0.22	0.44	7.00	2.38
Mean	0.02	0.83	0.49	0.34	0.16	0.08	0.08	1.41	0.42

7/24/01 Page: 4 of 4 No mdl is available for summed combinations of analytes. In summed columns (eg., BTEX), the reported values should be considered ESTIMATED if any of the individual compounds were reported as bdl.

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