

January 31, 2002

Mr. Barney Chan Alameda County Health Care Services Agency Department of Environmental Health 1131 Harbor Bay Parkway, 2nd Floor Alameda, California 94502

Dear Mr. Chan:

Please find enclosed for your review the report "Site Investigation and Screening Level Risk Assessment, Grey & Reynolds Development Site, Embarcadero Cove, Oakland." This report is submitted in response to your letter dated October 3, 2001, in which you request further delineation of the TPH contamination discovered by Baseline Environmental Consultants in a Phase II Investigation dated August 2001.

Your prompt review and comments on the report would be greatly appreciated. If you have any questions, please do not hesitate to contact me at (510) 627-1184.

Sincerely,

Douglas H. Herman

Associate Port Environmental Scientist

Cc w/encl: Tom Bender

Barbara Szudy Betty Graham

Cc w/o encl: Adrienne La Pierre

Phil Fitzwater Yane Nordhav

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SITE INVESTIGATION AND SCREENING-LEVEL RISK ASSESSMENT REPORT

Gray & Reynolds Development Site Embarcadero Cove 1275 Embarcadero Oakland, California

January 16, 2002

Prepared for:

Port of Oakland c/o Douglas P. Herman EH&SC Department 530 Water Street Oakland, California 94607

Prepared by:

Iris-Cambria Environmental, J.V.

1615 Broadway, Suite 1003 Oakland, California 94612

1144 65th Street, Suite B Oakland, California 94602

Robert W. Schultz, R.G.

Senior Project Geologist

Adrienne LaPierre Principal Toxicologist

Cambria Environmental • 1144 65th Street, Ste. B • Oakland, CA 94608 • t. 510.420.0700 • f. 510.420.9170

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1.0 INTRODUCTION

Iris-Cambria Environmental, J.V. (Iris-Cambria), has prepared this Site Investigation and Screening-Level Risk Assessment Report for the property located at 1275 Embarcadero, Oakland, California (Site) on behalf of the Port of Oakland (Port) to facilitate redevelopment of the property for commercial use.

1.1 Purpose

The primary objective of these investigations is to further characterize soil and groundwater quality beneath the Site. A subsurface environmental site assessment was performed to assess potential ramifications of chemicals in the subsurface for Site redevelopment.

1.2 Scope of Work

Multiple investigations have been completed at the Site to characterize site conditions. Completed investigations are listed below in chronological order:

- Grab sampling of soil and groundwater was completed to provide hydrogeological and analytical data for the Site and to guide the selection of locations for groundwater monitoring wells;
- Geophysical surveys were utilized to investigate the lateral extent of subsurface concrete foundation(s);
- Test pits were excavated at the Site in areas where the geophysical surveys yielded anomalies including areas that might have contained possible USTs;.
- Monitoring wells were installed, developed, and sampled;
- A conduit survey was completed;
- The tidal influence on groundwater levels was assessed and used to understand whether a local wooden bulkhead might be important in controlling the local flow of groundwater;
- Sanborn maps were reviewed to help confirm the historical location of the former UST and to help identify potential former Site activities that could have released chemicals at the Site;
 and
- A screening-level risk assessment was completed for the Site to evaluate whether chemicals
 detected at the Site exceed risk-based screening levels (RBSLs) for the projected future site
 uses.

2.0 SITE DESCRIPTION

The Site is shown on Figure 1 and is located at 1275 Embarcadero, Oakland, California. Current Site surface features include a parking lot and a vacant former restaurant. The Port owns the Site, and Gray & Reynolds Properties, Inc. (Gray & Reynolds) is proposing commercial redevelopment for the Site. Redevelopment will include a 3 or 4-story commercial building to be located within the footprint of the former restaurant, a parking lot, and landscaping.

2.1 Background

Past investigations at the Site included the March 14, 2001 Draft Review of Existing Site Conditions and Environmental Risk Evaluation completed by Henshaw Associates, Inc., (Henshaw) and the subsequent April 23, 2001 Soil and Groundwater Sampling and Analysis Workplan (Workplan) also completed by Henshaw. The Site was discussed as an "Adjoining Property" in the July 26, 1994 Phase I Environmental Site Assessment report (ESA) prepared by the Port for the Marriott Hotel Development Site (Port of Oakland, 1994). In the 1994 ESA, the Port reported that an underground storage tank (UST) and associated pump, several small wood frame sheds, and an incinerator had been removed from the Site in 1970. The April 23, 2001 Workplan identified the storage sheds, the UST, and a cooling tower. According to Henshaw, the structure was initially misidentified during aerial photograph review. Henshaw later confirmed that the structure was a cooling tower (Brathwaite, 2002). Subsequently, Baseline Environmental Consulting (Baseline) implemented Henshaw's April 23, 2001 Workplan.

Baseline completed four (4) borings to groundwater (SB-1, SB-1A, SB-2, and SB-3), and shallow borings where refusal was encountered (SB-1B, SB-1C, SB-2C, and SB-2D). Baseline completed six shallow borings (RN-A1, RN-A2, RN-A3, RN-A4, RN-B1, and RN-B2) to assess soil and groundwater conditions beneath the Site (Baseline, 2001). Baseline's field effort included soil sampling at randomly selected locations ("RN" series) and soil and grab groundwater sampling at source-specific locations ("SB" series.) Baseline's sampling locations are shown on Figure 2. Baseline collected and analyzed soil and groundwater samples from the area near the former UST and pump (borings SB-1, SB-1A, and SB-1B) and from the area between the former UST and the Bay (SB-2 and SB-2C). Baseline detected the following compounds in the former UST area: total petroleum hydrocarbons (TPH) as gasoline (TPHg), TPH as diesel (TPHd), naphthalene, 2-methylnaphthalene, benzene, toluene, ethylbenzene, and xylenes (BTEX). Baseline also identified bis(2-ethylhexly)phalate, commonly the result of laboratory cross contamination, in a sample from the former UST area. No samples from the former UST area were analyzed for TPH as motor oil (TPHmo). Baseline analyzed soil and groundwater samples from the area between the former UST and the Bay for TPHg, TPHd, and BTEX. Only TPHd was detected in these areas.

Soil and grab groundwater samples were collected from boring SB-3 adjacent to the former cooling tower; samples were analyzed for hexavalent chromium only. No hexavalent chromium was identified above the reporting limit in soil or groundwater from boring SB-3.

Soil and groundwater chemical data from all investigations are presented in Tables 1 through 4 for soils and Tables 5 through 7 for groundwater. These tables include data from Baseline's investigation of the Site and data from Iris-Cambria investigations.

2.2 Hydrogeology

2.2.1 Regional Hydrogeology

The Site is located within the East Bay Plain Groundwater Basin and is very near (<1/4 mile) to the boundary between the Oakland Sub-Area and the Central Sub-Area (California Regional Water Quality Control Board, San Francisco Bay Region, 1999). The Oakland Sub-Area contains a series of westward-dipping alluvial fan deposits, typically 300 to 700 feet thick. The Central Sub-Area is underlain by the Younger Bay Mud (silty clay and fine-grained facies). Merritt Sands that are exposed at the ground surface in areas of west Oakland, extend through sections of both sub-areas. The Merritt Sand is stratigraphically deeper than the Younger Bay Mud, and occurs in part, as interfingered channel fills. The Merritt Sand typically ranges from 10 to 50 feet in thickness. The Younger Bay Mud thins in the region of the Site, and the deposit terminates at, or near, the Site. West of the Site, beneath the Bay, the Younger Bay Mud thickens to over 20 feet in thickness, and the Merritt Sand is present beneath the Younger Bay Mud (Goldman, 1969). Holocene deposits and developing soils are present either at the ground surface or beneath manmade fills.

2.2.2 Site Lithology

Soils beneath the Site generally consist of gravelly fill to approximately 2 feet below ground surface (ft bgs) (0 to 2 ft bgs). In many areas investigated at the Site, there exists a layer of concrete and asphalt that extends to approximately 3 ft bgs (2 to 3 ft bgs). The concrete and asphalt layer appears to be a remnant of former foundation(s). Beneath the concrete and asphalt, approximately 2 feet of clayey, silty, sandy and/or gravel fill (3 to 5 ft bgs) was encountered during drilling. Soils from 5 ft bgs to approximately 8 ft bgs principally consisted of silty clay and clayey silt. In all borings except MW-4 (the furthest boring from the Bay), primarily clayey silts were detected between 8 and the total explored depth of 15 ft bgs. Boring logs from this investigation are presented in Appendix A.

The investigated soils were principally comprised of sequences of gravelly fill and remnants of old foundations, overlying lower permeability clays and silts. The lower permeability clays and silts may be Younger Bay Muds and/or dredge fill. The clays and silts potentially overly Merritt Sands or they may overly interlayered Holocene sediments and soils.

2.2.3 Depth to Groundwater at the Site

Depth to first encountered groundwater at the Site is between approximately 5 and 8 ft. Depth to groundwater is greatest in the north-northeastern portion of the Site (wells MW-1 and MW-4) and less in wells nearer the Bay (MW-3 and MW-2). First encountered groundwater at the Site appears to be unconfined.

2,2,4 Site Groundwater Flow Direction

Inconsistent with the regional flow pattern, groundwater at the Site is calculated to flow to the northeast. This finding is based on five (5) rounds of groundwater level monitoring in October and December 2001 in all four monitoring wells at the Site (MW-1, MW-2, MW-3 and MW-4). Figure 3 illustrates groundwater levels measured on October 19, 2001 and shows the groundwater elevation decreasing to the northeast. The hydraulic gradient is mapped on Figure 3 to the northeast with a magnitude of 0.02.

Groundwater flow direction at the Site as calculated from multiple rounds of water level measurements is inconsistent with general regional groundwater flow patterns expected for this area. Regional flow is generally to the southwest towards the Bay (California Regional Water Quality Control Board, San Francisco Bay Region, 1999). A northeastern flow direction at the Site would suggest that local hydraulic influences are controlling the direction of flow. Review of boring logs for the Site indicate that well MW-2 is screened both through the targeted clayey silts and 6 inches into an underlying higher permeability sand. This well construction detail could affect groundwater elevation in well MW-2. Despite this consideration, the groundwater flow direction calculated for the Site using wells MW-1, MW-3, and MW-4, only, remains northeastward. Discussion of local potential hydraulic influences such as utility conduits is presented in Section 3.8.4 of this report. Lastly, flow direction at the Site is to be investigated further with the installation of an additional monitoring well in the northeastern portion of the Site.

3.0 SITE INVESTIGATION

Iris-Cambria's investigations at the Site included soil and groundwater grab sampling, investigation of a buried foundation found to exist beneath a portion of the Site, excavation of test pits, installation and sampling of monitoring wells, completion of a conduit survey, evaluation of a buried wooden bulkhead, a Sanborn map review and a limited tidal study. Discussion of each phase of study is presented below.

3.1 Soil and Groundwater Grab Sampling

On August 30, 2001, Iris-Cambria collected soil and grab groundwater samples from six borings completed at the Site. These borings are numbered SB-A, SB-B, SB-C, SB-D, SB-E and SB-F and their respective locations are illustrated on Figure 2. All borings were advanced to groundwater with the exception of boring SB-C. Boring SB-C was terminated at 4 ft bgs because subsurface obstacles were encountered in the borehole. Boring logs and permits are presented in Appendices A and B, respectively.

Soil and groundwater samples collected from five of the six borings were submitted to McCampbell Analytical, Inc. of Pacheco, California to be analyzed for the presence of petroleum hydrocarbons, methyl tert-butyl ether (MTBE), and semi-volatile compounds (SVOCs)(no samples from boring SB-C were submitted for analysis). Petroleum hydrocarbons were quantified as total petroleum hydrocarbons in all three ranges (TPHg, TPHd and TPHmo) and as benzene, toluene, ethylbenzene, and xylenes (BTEX). Standard field procedures for sample collection are presented in Appendix C.

Soil analytical results from the August 30, 2001 investigation are summarized in Tables 1 and 2. Likewise, grab groundwater analytical results from this same sampling program are summarized in Tables 5 and 6. Laboratory sheets for the soil and groundwater testing are presented in Appendix D. For completeness and to ease review of all analytical data for the Site, soil and groundwater data from prior investigations at the Site have been reproduced in Tables 1, 2, 5 and 6.

3.1.1 Field Activities

Field activities completed during the August 30, 2001 sampling program are presented in Appendix E. The results of the field investigation are presented below.

3.1.2 Soil Analytical Results

One soil sample was collected from above the water table in each of the following borings; SB-A. SB-B, SB-D, SB-E, and SB-F. These soil samples were analyzed for TPHg, TPHd, TPHmo, BTEX, MTBE, and SVOCs. Chemical concentrations were low in all of these soil samples tested in this phase of the program. The maximum chemical detections were found in the soil sample from boring SB-F in the northern portion of the Site, but even in that sample, concentrations were low (i.e. below RBSLs as discussed in Section 4, below). For example, benzene in the sample from SB-F was detected at 0.021 milligrams per kilogram (mg/kg) and the maximum total petroleum hydrocarbon detection was in the motor oil range at a concentration of 16 mg/kg TPHmo. No MTBE or SVOCs were detected in any of the soil samples collected from SB-A, SB-B, SB-D, SB-E, and SB-F.

3.1.3 Grab Groundwater Analytical Results

Analytical results for grab groundwater samples from borings SB-A, SB-B, SB-D, SB-E, and SB-F are summarized in Tables 5 and 6. Hydrocarbons were detected in all three ranges of TPH including TPHg, TPHd, and TPHmo. BTEX constituents were only detected in the groundwater sample from SB-E. Figure 4 illustrates the distribution of TPHg, TPHd, and TPHmo, and benzene in grab groundwater samples from the August 30, 2001 investigation as well as earlier grab groundwater samples collected by Baseline in their borings SB-1, SB-1A, SB-2 and SB-3. Chemical concentrations in the grab groundwater samples are highest in the northeast portion of the Site. The locations where higher concentrations were found were north of the location of the former UST identified as subsurface anomaly 1 on Figure 4.

Chemical concentrations in grab groundwater samples generally decrease away from the source area toward the Bay. As illustrated on Figure 4, this trend is shown moving from SB-1 to SB-A to SB-2.

MTBE was not detected in any grab groundwater sample from SB-A, SB-B, SB-D, SB-E, or SB-F. SVOCs were detected infrequently. Their only detection in water samples from SB-A, SB-B. B-E. SUC'S from desil or wrapping for 4575 SB-D, SB-E, and SB-F were fluoranthene and pyrene at SB-D and naphthalene at SB-E.

3.2 Foundation Investigation

Subsurface investigations completed by Baseline (Baseline, 2001) and those completed by Iris-Cambria during the drilling program of August 30, 2001, suggest that there were remnants of an old foundation at some 2 to 3 feet below the current ground surface in portions of the Site.

Additional investigations of the old foundation's extent were undertaken to assist with future property redevelopment activities and to assess the foundation's potential influence on the distribution of chemicals at the Site.

To further investigate the general extent of the remnants of the old buried foundation, Iris-Cambria contracted for magnetometer and Ground Penetrating Radar (GPR) surveys to be completed at the Site. These surveys were completed by Norcal Geophysical Consultants, Inc. (Norcal) of Petaluma, California on September 5, 2001. The geophysical surveys were ineffective in delineating the extent of the subsurface foundation at the Site. GPR transects and test pit locations are shown on Figure 5 and Norcal's entire report is reproduced in Appendix F.

Because geophysical surveys proved ineffective in determining the extent of the old foundation remnants, existing boring data were used to map its extent. Boring locations were used to map the old foundation remnants; the approximate extent of the remnants are shown on Figure 6. Figure 6 illustrates that the foundation underlies approximately two thirds of the eastern portion of the Site. The extent to the west is unclear but the foundation remnants were not encountered in either RN-A1 or RN-A2 in the western portion of the Site.

The remnants of the old foundation found between depths of 2 to 3 ft bgs are believed to be too shallow and insufficiently widespread to have a material effect on the chemical distributions at the Site. Site redevelopment activities especially grading activities should plan to encounter the remnants in the eastern portion of the Site and possibly to the west, although the occurrence of the foundation remnants to the west is less certain.

3.3 Test Pits

During geophysical surveying to investigate the old buried foundation remnants and to clear locations for drilling, the geophysical subcontractors identified four subsurface magnetic anomalies at the Site. These subsurface anomalies are illustrated on Figure 5 and the entire geophysical survey report is presented in Appendix F.

Geophysical signatures from anomalies 1 and 2 were similar to those typically seen from USTs. Anomalies 3 and 4 appeared to be buried debris. To investigate the geophysical anomalies, Iris-Cambria field staff oversaw exploration activities of Foss Environmental Services of Alameda, California, using a backhoe on September 17, 2001.

At anomaly 1, soil was excavated to approximately 6 ft bgs. Rebar and wires were encountered in the excavation between 2 and 4 ft bgs. A concrete foundation was partially detected at 2 ft bgs at this location. The foundation was cut or broken on the eastern side of the excavation, and missing entirely in the rest of the excavation. Anomaly 1 appears to be the likely location of the former UST that was reportedly removed from the Site in 1970 (Port of Oakland, 1994). Consistent with records showing the UST removal in 1970, the backhoe excavations completed by Iris-Cambria at anomaly 1 show that the former UST has indeed been removed from the Site.

At anomaly 2, rebar was unearthed prior to encountering a concrete foundation at 2 ft bgs. Anomaly 3 was not excavated because it was nearby anomaly 1 and because excavation at anomaly 1 indicated a UST had previously been removed. In addition, the geophysical signature for anomaly 3 suggested that this anomaly consisted of buried debris. At anomaly 4, soil was excavated to approximately 2 ft bgs, where a hubcap was unearthed. This finding was consistent with the geophysical signature of buried debris.

A soil sample (M-1-5) was collected at 5 ft bgs in the test pit dug at anomaly 1, the location of the former UST. The soil sample was analyzed for TPHg, TPHd, TPHmo, BTEX and MTBE (by EPA Methods 8015 and 8020). Analytical results are presented in Tables and 2. The highest total petroleum hydrocarbon detection was in the gasoline range at 2,300 mg/kg (TPHg) and benzene was detected at 1.8 mg/kg. MTBE was detected at 31 mg/kg; however, this result is considered anomalous because the sample was analyzed by EPA Method 8020 which may yield false positives for MTBE in the presence of other petroleum hydrocarbons (American Petroleum Institute, 2000). The MTBE detection in soil sample M-1-5 being a false positive is further supported by the consistent lack of MTBE detection in any other soil or groundwater sample collected and analyzed from the Site.

3.4 Monitoring Well Installation and Sampling

Iris-Cambria installed four groundwater monitoring wells (MW-1 through MW-4) at the Site on October 9, 2001. Well locations are illustrated on Figure 2, and boring logs and well completion details are presented in Appendix A. One soil sample was analyzed from each well borehole. Well installation and sampling procedures are presented in Appendix C.

Monitoring well locations were selected based on the results of previous investigations at the Site. Monitoring well locations were proposed to the Alameda County Health Care Services Agency (ACHCSA) in the September 27, 2001, *Workplan Addendum* (Iris-Cambria, 2001). ACHCSA provided written comments on October 3, 2001 (Chan, 2001) and final monitoring well locations were adopted as recommended by the ACHCSA.

3.4.1 Field Activities

Field activities completed during the October 2001 sampling program are presented in Appendix E. Well sampling forms and survey data are presented in Appendices G and H, respectively. The results of the field investigation are presented below.

3.4.2 Soil Analytical Results

Soil samples collected during the installation of monitoring wells MW-1 through MW-4 were analyzed for TPHg, TPHd, TPHmo, BTEX, SVOCs, and VOCs. Well locations are shown on Figure 2. One soil sample from boring MW-1 was analyzed for CAM 17 Metals. Metals detections in the soil sample from MW-1 were generally low (i.e. below expected background concentrations as characterized in colluvium and fill at Lawrence Berkeley National Laboratory (LBNL, 1995). A discussion of the significance of onsite metal detections in soils is addressed in the Risk Based Screening Evaluation, Section 4.0. Soil analytical results are presented in Tables 1, 2, 3 and 4.

Chemicals detected in onsite soils included TPHg, TPHd, TPHmo, BTEX, naphthalene, and metals. No MTBE was detected in soil samples from MW-1, MW-2, MW-3, or MW-4. The maximum total petroleum hydrocarbon detection in soil samples from the four monitoring well boreholes was found was at MW-2 (300 mg/kg of TPHmo) which is south of the former UST location and the maximum benzene detection was found at MW-4 (0.7 mg/kg benzene) which is north of the former UST location.

3.4.3 Groundwater Analytical Results

Groundwater was sampled three times in each monitoring well (MW-1 through MW-4). The sampling events occurred on October 19, 2001 and at two different tidal stages on December 5, 2001. The analytical results for groundwater testing are summarized in Tables 5, 6 and 7.

Figure 7 presents groundwater concentrations of TPHg, TPHd, TPHmo and BTEX detected during the sampling event of December 5, 2001. Highest chemical concentrations are generally found in the north and northeast portion of the Site furthest away from the Bay. For the majority of compounds tested, MW-4 yielded the most elevated concentrations. Exceptions to this pattern included TPHmo, 2-methylnaphthalene, and naphthalene where MW-3 contained the highest concentrations.

Detected chemical concentrations decrease rapidly toward the Bay from the former UST location. The former UST location is shown as anomaly 1 on Figures 4 and 5. An illustration of this is provided on Figure 8 where benzene contours are plotted. Benzene concentrations decrease significantly from the former UST location toward the Bay (to the south and southwest). Chemical concentrations in groundwater at MW-2 (approximately 85 feet from the Bay) are non-detect for all compounds except TPHd and TPHmo.

Generally, detected chemical concentrations in the four monitoring wells MW-1 through MW-4 have decreased from the first sampling event of October 19, 2001. Analytical results from the December 2001 sampling events likely are most representative of actual groundwater concentrations at the Site; the October 19, 2001 samples may have been biased by effects of the well installation processes.

The lower concentrations detected in groundwater in the December 2001 sampling events do not appear to be the result of groundwater elevation fluctuations. Groundwater elevations measured in the site wells on December 5, 2001 (6:50-8:05 AM) were very similar to those measured on October 19, 2001.

3.5 Conduit Survey

To evaluate if subsurface conduits identified by the geophysical surveying had the potential to act as preferential pathways for groundwater and dissolved chemical migration, Iris-Cambria performed a conduit survey. The survey inventoried all subsurface utilities beneath the Site and in nearby areas. Subsurface utilities at the Site and beneath the Embarcadero (north of the Site) were researched by Foresite Engineering Surveys, Inc. of Pleasant Hill, California, and by Underground Service Alert (USA) prior to the August and October 2001 drilling programs. Iris-Cambria mapped the utility locations and used a measuring wheel to determine distances. Iris-Cambria also reviewed topographic and subsurface utility maps developed for the Site by Watson Engineering, Inc. of Auburn, California. The utility trench locations are illustrated on Figure 9. A City of Oakland Engineering Department (City) utility and street improvement map is included in Appendix I for completeness but Figure 9 best illustrates the utility features that could intersect groundwater beneath the Site.

The conduit survey indicated that a storm drain, a sanitary sewer, an electric line, a telephone line, and a water line run parallel to Embarcadero. A small storm drain collection grate and an associated drain pipe (about 4-inch in diameter and approximately 12-inch bgs) is located near the southeast corner of the Site and drains to the Bay. Only the storm drain and sanitary sewer lines

are shown on Figure 9 since these would be the deeper utility trenches than those for electric and telephone.

A 78-inch diameter interceptor sanitary sewer line runs roughly east to west within the Embarcadero. This sanitary sewer line is the deepest utility feature found during the conduit survey. According to communication with the City of Oakland, the 78-inch diameter interceptor is a sanitary sewer line that transports wastewater to an East Bay Municipal Utilities District facility, and the top of the interceptor line is at least approximately 3 ft bgs. Therefore, the invert (or flow line) of the 78-inch diameter interceptor line is at least 9.5 ft bgs. Street elevation near the Site is approximately 11.0 ft (Port datum) which would make the invert elevation of the interceptor line approximately 1.5 ft (Port datum). Other invert elevations for nearby utility conduits are shallower and are shown on Figure 9.

Figures 9 and 10 show the approximate locations and estimated invert elevations of key utilities near the Site that could intersect groundwater. Groundwater elevations (in Port datum) at two tides (a falling tide and a rising tide) are posted on Figure 10. Figure 10 indicates that invert elevations for conduits except the 78-inch diameter interceptor line are generally above the water table. The sanitary sewer invert is close to the groundwater level measured in MW-4 on the falling tide cycle on December 5, 2001. The invert elevation of the 78-inch diameter interceptor line is deeper than groundwater elevations measured in monitoring wells anywhere on the Site. The 78-inch diameter interceptor line and possibly the sanitary sewer could be affecting local groundwater flow patterns at the Site; a possible explanation for the observed groundwater flow opposite regional flow patterns.

3.6 Wooden Bulkhead Evaluation

The City utility map initially created July 17, 1969 (reproduced in Appendix I) and a Site location map obtained from the Port (Inner Harbor, Brooklyn Basin, Test Piles Adjacent to Harbor Line Points 57&59, September 2, 1947) both show the presence of a wooden bulkhead that traverses the Site from west to east. The approximate location of the wooden bulkhead is shown on Figure 2. An additional 1936 drawing of the wooden bulkhead was obtained from the Port and is reproduced in Appendix J. The Appendix J drawing shows the wooden bulkhead to consist of riprap and creosoted sheet piles.

The depth of the wooden bulkhead is unclear from the available information. If the scale shown on the 1936 drawing in Appendix J is roughly correct, then the wooden bulkhead might extend to about 40 feet below the 1936 Site grade; however, the scale does not appear correct. The wooden

bulkhead is located near well MW-1; however, it was not encountered during any of Iris-Cambria's excavation or drilling activities.

Of interest to the current investigation, is the question of what effect the wooden bulkhead might have on the Site hydrogeologic system and whether the wooden bulkhead divides the Site into two separate hydraulic units. If the wooden bulkhead were an effective hydraulic barrier, then the response to tidal changes should be markedly different on the Bay side of the bulkhead versus the side furthest away from the Bay. Figure 10 illustrates changes in water levels in wells closest to the Bay and in wells on the other side of the wooden bulkhead furthest away from the Bay. Water levels were measured during a rising and during a falling tide some six hours apart on December 5, 2001 as discussed more completely in the following section.

Water levels in wells on both sides of the wooden bulkhead fluctuated by at least 0.5 ft over the 6-hr period. These similar hydraulic responses in wells on either side of the wooden bulkhead suggest that the wooden bulkhead is not an effective hydraulic barrier to shallow groundwater flow at the Site and should not be a significant controlling factor in affecting local groundwater flow patterns. Figure 10 also illustrates the location of the utility trenches to the north of the Site and their relative depths with respect to groundwater elevations.

3.7 Limited Tidal Influence Study

Iris-Cambria performed a limited tidal study to evaluate tidal influence in Site monitoring wells MW-1, MW-2, MW-3, and MW-4. On December 5, 2001, groundwater levels were measured in two separate events, approximately six hours apart. The 6:10-6:20 AM measuring event occurred on a falling tide, and the 11:43-11:49 AM measuring event occurred on a rising tide. On December 19, 2001, groundwater levels were measured near the daily lower low tide. During each groundwater measuring event, water levels in all wells were measured within a 20- to 30-minute period to minimize the impact of tidal fluctuation between the network of wells. Groundwater level maps for all three tidal study measuring events are shown on Figures 11, 12, and 13. Groundwater level elevations are tabulated in Tables 5 and 6. Field forms are included in Appendix G.

During the measuring events of December 5, 2001 between the falling and rising tide gauging events, water levels in wells MW-1, MW-2, and MW-4 increased by about 0.5 ft. The water level in MW-3 increased by over 1.0 ft. Figures 10, 11, and 12 map the groundwater levels across the Site during each monitoring event. Although there appears to be tidal influence at the Site, these figures show that the observed groundwater gradients are roughly constant in direction. There is no indication of gradient reversals occurring at the Site as a result of nearby tidal variations.

3.8 Sanborn Map Study

At the request of the ACHCSA (Chan, 2001), Sanborn maps of the Site were obtained from Environmental Data Resources, Inc. (EDR) covering the years 1911, 1950, 1952, 1953, 1957, 1959, 1960, 1964, 1965, 1967, and 1969, and are included as Appendix K. The approximate Site boundaries have been placed on the Sanborn maps presented in Appendix K.. The Sanborn maps are of poor quality with respect to relative distances, feature dimensions, and relative orientation. The Site boundaries were located using the best consistent anchor points, such as Railroad tracks and older buildings present on the most recent maps.

- Buildings do not appear on the Sanborn maps covering the Site until 1950 where it appears
 that a shed exists in the northeast portion of the Site and another building, possibly an office,
 appears in the north central portion of the Site.
- In the 1952 map, the northeast shed is present but drawn slightly differently. The central portion of the Site is empty.
- In the 1953 map, the Site contains only the office structure to the northeast. The shed in the 1952 map has been removed.
- In the 1957 map, a small area appears in the eastern portion of the Site including a marked area near the assumed location of the former onsite UST. A pallet handling area now shows up in the central portion of the Site.
- In the 1959 map, a yacht club appears on the western portion of the Site and a new building appears on the eastern portion of the Site. Other structures remain the same.
- The 1960, 1964,1965,1967, and 1969 maps remain the same as the 1959 map. The building on the eastern portion of the Site appears to have been used for sawing.

The maps yielded no additional information (beyond those recognized by Henshaw) suggesting historical site uses that would have involved onsite hazardous materials use or storage. The Sanborn maps do not confirm the former location of the UST.

3.9 Site Investigation Conclusions

The conclusions that are drawn from the subsurface investigation of the Site are presented under general topics of key concern. They are as follows:

3.9.1 Source of Contamination

Investigation results suggest that an underground storage tank (UST) was previously located in the northwest portion of the Site. The approximate former UST location is illustrated as subsurface anomaly 1 on Figures 4 and 5. The former UST was removed from the Site in the 1970s as reported by Henshaw Associates (Henshaw, 2001). Test pits dug in the same area

confirmed that the UST had been removed. Elevated petroleum related chemical concentrations in soil and groundwater surrounding the former UST location suggest that historical site operation of the UST, potentially including storage and dispensing of gasoline, was the source of TPHg and BTEX detected at the Site. The exact source of TPHmo, TPHd, and SVOCs is uncertain; however it appears to have been located near the former UST.

3.9.2 Chemical Distribution in Site Soil

Chemicals concentrations in soils are generally greatest at the former UST location. The highest TPHd, TPHg, and benzene concentrations found in soils anywhere on the Site were in a soil sample from a test pit dug at the former UST location (M=1-5). This soil sample contained 2,300 mg/kg of TPHg, 850 mg/kg TPHd, and 1.8 mg/kg benzene. Other petroleum compounds were also detected in this sample. The highest TPHmo concentration of 300 mg/kg was detected approximately 40 ft south of this location in a soil sample from boring MW-2.

Soil samples from immediately surrounding sample locations were likewise elevated. With the sole exception of TPHmo, concentrations of petroleum related compounds drop in magnitude as distance away from the former UST location increases suggesting that the former UST location was the source of the chemicals being detected in soils at the Site. Detected TPHmo concentrations were higher both west (at MW-3) and south (at MW-2) of the former UST location. Accordingly, TPHmo is distributed in the vicinity of the former UST, and the data do not suggest a single point source of TPHmo.

3.9.3 Chemical Distribution in Groundwater

Groundwater chemical data show that TPHg and BTEX concentrations were highest near the former UST location and in MW-4 approximately 35 ft to the north. Benzene concentrations decrease to below detection levels closer to the Bay as shown in MW-2 (Figure 7). TPHmo concentrations were highest south of the former UST location in MW-3 at 550 μ g/L. TPHmo was detected at 270 μ g/L closer to the Bay at MW-2 (Figure 7).

3.9.4 Groundwater Flow Direction

Measured groundwater levels in the four monitoring wells at the Site suggest that the hydraulic gradient in the northeast portion of the Site is to the northeast. This finding is inconsistent with regional flow which is to the southwest toward the Bay. Given that the regional groundwater flow direction is to the southwest, the hydraulic gradient calculated for the Site may be a local variation.

Discussion of local hydraulic influences such as the presence of utility conduits was presented earlier in this report. In summary, a 78-inch diameter interceptor line north of the Site has an invert elevation lower than *all* groundwater levels measured on the Site. Similarly, a sanitary sewer trench north of the Site has an invert elevation lower than *some* of the groundwater measured on the Site. Therefore, the 78-inch diameter interceptor line and the sanitary sewer could be affecting local groundwater flow patterns by causing groundwater to flow toward them. This could explain why the hydraulic gradient at the Site appears inconsistent with the regional flow patterns.

There is an old buried wooden bulkhead that crosses the Site west to east. Water levels measured on both sides of the wooden bulkhead over various tidal cycles, suggest that the wooden bulkhead is not an effective hydraulic barrier at the Site. Therefore, the wooden bulkhead should not be a significant controlling factor in affecting local groundwater flow patterns.

3.9.5 Plume Definition

The extent of chemicals in groundwater beneath the Site appears generally well defined with concentrations declining to the south, west and east of the former UST location. Chemical concentrations detected in groundwater samples collected closest to the Bay are reduced from concentrations detected in the former UST area. The distribution of dissolved chemical concentrations suggests that migration, if occurring, may be toward the north-northeast.

4.0 RISK-BASED SCREENING LEVEL EVALUATION

The purpose of the risk-based screening level (RBSL) evaluation is to assess whether chemicals detected in soils and groundwater at the Site could pose a risk to human health or the environment. Site plans call for the development of a commercial building within the footprint of the former restaurant building, with the remaining portions of the Site to be used for parking and limited landscaping. Based on the site development plans, the human populations included in this screening-level evaluation are future commercial workers and construction workers. The environmental receptors considered in this evaluation include urban-area terrestrial receptors, and aquatic organisms that may be present in the adjacent San Francisco Bay. The approach, results, and conclusions of the RBSL evaluation are discussed in the sections below.

4.1 Approach

The RBSL evaluation consists of a direct comparison of the chemicals detected in the soil and groundwater against selected risk-based soil and groundwater screening levels that are protective

of human health and ecological receptors. All analytical results from soil and groundwater investigations conducted by Baseline in May 2001 and Cambria in August, October, and December 2001 were used in the RBSL evaluation. The applicable RBSLs correspond to those developed and assembled by the San Francisco Regional Water Quality Control Board (SFRWQCB) for the protection of future onsite commercial and construction workers, aquatic life, and urban-area terrestrial flora and fauna (SFRWQCB, 2000). Specifically, the human health and ecological RBSLs are based upon the following exposure scenarios:

Surface Soil (less than 3 meters below ground surface)

- Commercial workers exposed to soil through direct contact (i.e., soil ingestion, and dermal contact pathways), inhalation of particulates, and inhalation of vapors in indoor air;
- Construction workers exposed to soil through direct contact (i.e., soil ingestion, and dermal contact pathways), inhalation of particulates, and inhalation of ambient vapors;
- Aquatic receptors exposed to chemicals in the soil that have leached down from soil, into the underlying groundwater, and migrated to surface waters; and
- Urban-area terrestrial flora and fauna exposed to surface soils.

Groundwater (not assumed to be a drinking water source)

- Commercial workers exposed to groundwater vapors that have migrated into the indoor air;
 and
- Aquatic receptors exposed to groundwater discharging to surface water.

These risk-based soil and groundwater screening levels were selected to evaluate the potential adverse health and environmental impacts posed by chemicals detected in soils and groundwater at the Site. The RBSLs for chemicals that are carcinogens (e.g., benzene) are based on a target cancer risk of 10⁻⁶ (one-in-one million). This represents the most stringent end of the acceptable risk range of 10⁻⁴ to 10⁻⁶, used by both Federal and State regulatory agencies in environmental decision-making. The RBSLs for noncarcinogens are based on a hazard index (HI) of 0.2. The RBSLs used in this evaluation are extremely conservative. Consequently, the presence of a chemical at concentrations below the RBSLs can be assumed to pose an insignificant threat to human health and the environment. However, it is important to note that the presence of a chemical detected above the RBSL does not necessarily indicate that adverse effects to human health or the environment are occurring; rather, exceedances of the RBSLs simply indicate that additional investigation and/or evaluation of potential risks may be warranted.

The analytical results from the soil and groundwater investigations conducted at the Site and the selected risk-based soil and groundwater screening levels are summarized and presented in Tables

1 through 6. The results of the RBSL evaluation for human health and ecological receptors are discussed below.

4.2 Results of the Human Health RBSL Evaluation

Soil and groundwater analytical data for the Site was directly compared to selected human health risk-based soil and groundwater screening levels. As discussed in Sections 3, TPHg, TPHd, TPHmo, BTEX, MTBE, other VOCs, and selected SVOCs were detected in soils and/or groundwater at the Site. MTBE was detected in only one soil sample and is believed to be a false positive result as discussed in Section 3.3. Metals were also detected in soils at the Site. The results of the RBSL evaluation for future commercial workers and construction workers who may be present at the Site are discussed below.

Future Commercial Workers

As indicated in Tables 1 though 4, benzene was the only compound detected in soils at concentrations that exceed the human health risk-based screening level for the commercial worker. The RBSL for benzene is 0.39 mg/kg, and assumes exposures occur via direct contact with soil, and via the inhalation of vapors migrating up through the soil column and accumulating in the indoor air of a building. The specific locations and depths where benzene was detected in soil at levels that exceed the commercial RBSL include MW-4 (at a depth of 5.3 feet bgs), MW-1 (at a depth of 8.3 feet bgs) and M-1-5 (at a depth of 5 feet bgs). Benzene was detected in soil at MW-4, MW-1, and M-1-5 at concentrations of 0.70 mg/kg, 0.48 mg/kg and 1.8 mg/kg, respectively, concentrations that exceed the RBSL of 0.39 mg/kg by a factor of between 1.2 and 4.6. The presence of benzene at these particular locations and depths would not be expected to pose a significant health risk to future commercial workers for the following reasons: 1) commercial workers will not likely have direct exposure with any soils located at depths of between 5 and 8 feet bgs; and 2) the exceedances occur in areas that will be used for parking and/or landscaping, and therefore potential indoor air impacts resulting from the transport of benzene vapors into a building will not occur.

Similar to soils, benzene was the only compound detected in groundwater at concentrations that exceed the human health risk-based screening levels for the commercial worker. As indicated in Table 5, the specific locations where benzene was detected in groundwater at levels that exceed the commercial RBSL of 84 µg/L include SB-1, SB-1A, SB-E (all groundwater grab samples), MW-1, and MW-4. All of these locations where benzene exceeded the RBSL are located either within or directly north of the former UST location. The concentrations of benzene in MW-1 and MW-4 range from 120 µg/L to 1,900 µg/L. As with the soil, the RBSL for benzene in the

groundwater is based on the potential for the benzene to volatilize from the groundwater and migrate up through the soil column into the indoor air of a building. Consequently, the sampling locations that are most relevant for determining whether the benzene may pose a risk to future commercial workers include those groundwater samples that are closest to the building, and that would be considered most representative of the groundwater conditions that may exist under the building.

As indicated in Table 5, the concentrations of benzene in groundwater at the locations surrounding the former restaurant (i.e., SB-2, SB-A, SB-B, MW-2 and MW-3) were well below the RBSL of 84 μ g/L. Benzene was reported as non-detect in groundwater in six of the nine groundwater samples collected from these locations, and was detected at a maximum concentration of 2 μ g/L at MW-3. Accordingly, based on the proposed development plans, the levels of chemicals detected in groundwater at the Site would not be expected to pose a significant health risk to future commercial workers who will be working at the Site.

Future Construction Workers

As indicated in Tables 1 through 4, all chemicals in soil were detected at concentrations that are below the RBSLs developed for the protection of a future construction worker. Accordingly, the levels of chemicals detected in soils at the Site would not be expected to pose a significant health risk to future construction workers who may be directly exposed to chemicals present in the subsurface soils.

4.3 Results of the Ecological RBSL Evaluation

Soil and groundwater analytical data for the Site was directly compared to selected ecological risk-based soil and groundwater screening levels. As discussed above, TPHg, TPHd, TPHmo, BTEX, MTBE, other VOCs, and select SVOCs were detected in soils and/or groundwater at the Site. Metals were also detected in soils at the Site. The results of the RBSL evaluation for urbanarea terrestrial receptors and aquatic life in San Francisco Bay are presented below (saltwater criteria were given priority over freshwater criteria).

Urban-Area Terrestrial Receptors

None of the chemicals detected at the Site were present at levels that exceed urban-area terrestrial RBSLs (Tables 1 through 4). Accordingly, chemicals detected in soils would not be expected to adversely impact urban-area terrestrial receptors that could potentially be using the Site.

What are there?

Aquatic-Life

As indicated in Tables 1 through 4, TPHg, TPHd, ethylbenzene, xylenes, MTBE, and 2-methylnaphthalene were detected in soils at concentrations that exceed the RBSL developed for the protection of aquatic receptors. The locations where soil concentrations exceeded the aquatic-based RBSLs include SB-1A (located north of the former UST), M-1 and MW-1 (both within the former UST location). The soil RBSL for aquatic-life protection is based on the potential leaching of the chemical from soil down into the groundwater, and the subsequent exposure of aquatic organisms to the chemicals in the groundwater as it discharges to the surface water.

Generally speaking, concentrations measured in groundwater provide a more direct estimate of the amount of a chemical that may leach from the soil and impact the groundwater. This is particularly true for situations where the release occurred many years ago, the leaching from soil to the groundwater has already occurred, and chemical equilibrium has been established. In these situations, it is preferable to rely upon direct groundwater monitoring data to determine whether residual chemicals present in the soil may adversely impact the underlying groundwater.

Tables 5 and 6 present a comparison of the concentrations of chemicals detected directly in the groundwater to the aquatic-based RBSLs. Assindicated in the Tables, TPHg, TPHd, TPHmo, BTEX, 1-methylnaphthalene, 2-methylnaphthalene and phenanthrene were detected at concentrations in groundwater above the risk-based screening levels for aquatic life. The locations where groundwater concentrations exceeded the RBSLs include: SB-1A, SB-E, SB-F, MW-4 (all located north of the former UST location); MW-1, SB-1, SB-D (all located at or adjacent to the former UST location); and SB-A and MW-3 (both located approximately 40 feet southwest of the former UST location). All of these exceedances are located on the northern portion of the Site, more than 100 feet away from the San Francisco Bay. The monitoring well ncloset to the San Francisco Bay is MW-2, located approximately 85 feet from the shoreline. As petroleum-related compounds tend to attenuate rapidly with distance from the source, the concentrations measured in the monitoring well closest to the San Francisco Bay provide the best available representation of the concentrations of chemicals from the Site that could be impacting aquatic organisms in the Bay. As indicated on Tables 5 and 6, only trace concentrations of TPHd and TPHmo were detected in MW-2, at concentrations below the aquatic-based RBSL. Accordingly, the analytic data collected to date support that the chemicals detected at the Site do not appear to be adversely impacting aquatic life.

where for the control of the control

4.4 Conclusions of Risk Screening

Based on the results of the RBSL evaluation, Iris-Cambria has drawn the conclusions presented below.

Chemicals detected in soils and groundwater in the area surrounding the location of the proposed building are below relevant health-based RBSLs for future onsite commercial workers. Accordingly, assuming that the groundwater plume is stable, and that the higher concentrations of benzene detected in groundwater around and north of the former UST do not migrate beneath the building in the future, the levels of chemicals detected at the Site would not be expected to pose a significant health risk to future commercial workers. As the location of the benzene exceedances occur at locations that will be used for parking and/or landscaping, the levels of benzene detected at and north of the former UST location would not be expected to pose a significant health risk to future commercial workers at the Site.

Chemicals detected in soils at the Site are below health-based RBSLs for construction workers, and would therefore not be expected to pose a significant threat to the health of future onsite construction workers who may have direct contact with the soil during subsurface activities.

Chemicals detected in soils at the Site were present at levels below urban-area terrestrial RBSLs. Accordingly, detected chemicals would not be expected to adversely impact terrestrial receptors that could potentially be using the Site.

Although chemicals were detected in both soil and groundwater at concentrations that exceed the RBSLs developed for the protection of aquatic life, these exceedances occurred more than 100 feet north of the San Francisco Bay. Petroleum-related compounds tend to attenuate rapidly with distance from the source, and thus the concentrations measured in monitoring well MW-2 provide the best representation of the concentrations of chemicals from the Site that could be impacting aquatic organisms in the Bay. As chemicals detected in soils and groundwater at MW-2 were below RBSLs developed for the protection of aquatic life, the analytic data collected to date support that the chemicals detected at the Site do not appear to be adversely impacting the aquatic receptors.

5.0 CONCLUSIONS AND RECOMMENDATIONS

The investigations completed for the Site have well characterized chemical occurrence in soil and groundwater in the northeast portion of the Site, and have provided a sound basis for the

assessment of potential risks to human health and the nearby aquatic environment posed by the presence of chemicals in the subsurface. Specifically:

- Based on existing subsurface investigation data and Site use information, the Site appears to be appropriate for the planned redevelopment from a health risk perspective. With confirmation of UST removal, there is no known ongoing source of contamination. The chemicals detected in the soil and groundwater would not pose an unacceptable risk to future commercial workers, construction workers, or nearby aquatic receptors based on the current redevelopment plans. We recommend that the Port request that the ACHCSA review the screening-level risk evaluation presented in this report and issue a letter concurring with the conclusion that based on current knowledge the Site is appropriate for the planned redevelopment.
- Ancadditional groundwater monitoring well should be installed in the northeastern corner of the Site. This well would be used to further characterize the extent of chemicals in groundwater in this portion of the Site and would be used to further understand local groundwater flow patterns.
- Measuring of water levels and sampling for water quality should continue in the site monitoring wells until data are collected over one complete hydrologic cycle. Monitoring wells have been sampled in October 2001 (prior to the rainy season) and again in December 2001. Water levels and water samples should be measured and collected again in March and June 2002 to complete the sampling over an entire hydrologic cycle. Groundwater samples would be tested for total petroleum hydrocarbons in the gasoline, diesel and motor oil ranges and for SVOCs and BTEX.
- Future construction work should be undertaken in accordance with a site-specific health and safety plan. The health and safety plan should take into account all contaminants identified in the subsurface identified during the investigations.
- If the two additional rounds of sampling proposed for March and June 2002 yield data consistent with the current understanding of the Site or yield concentrations trending downward, Iris-Cambria recommends that the Port request formal site closure or a No Further Action letter from ACHCSA and the Regional Water Quality Control Board (RWQCB).

6.0 REFERENCES

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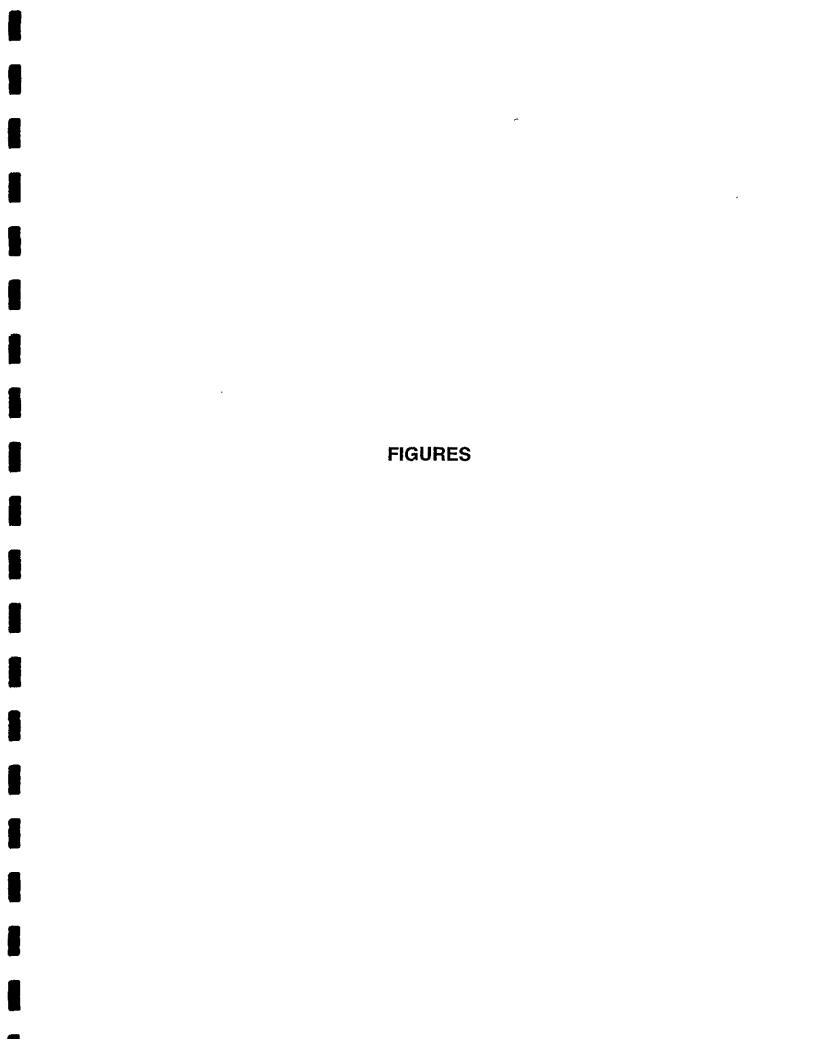
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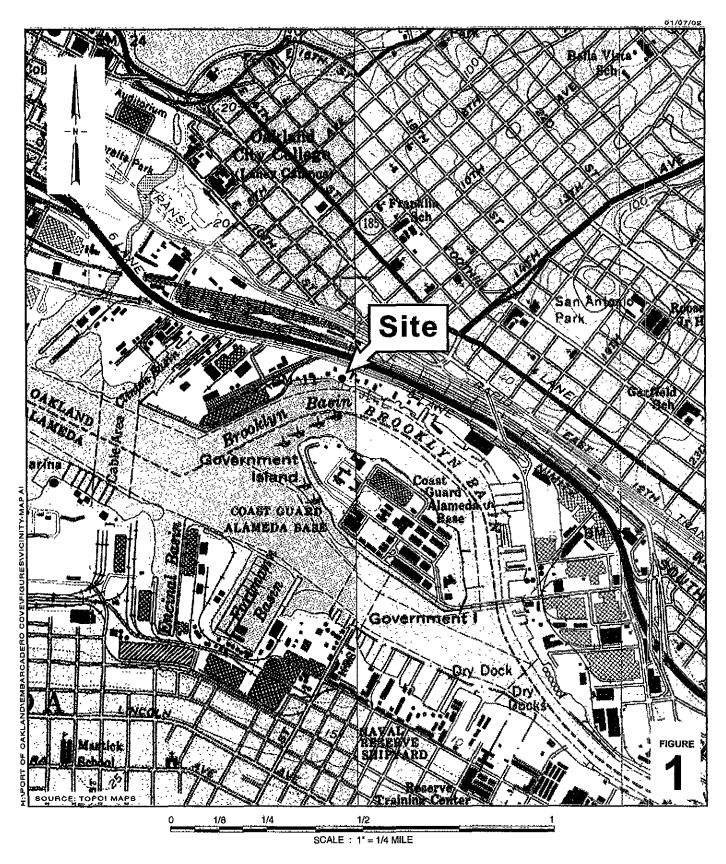
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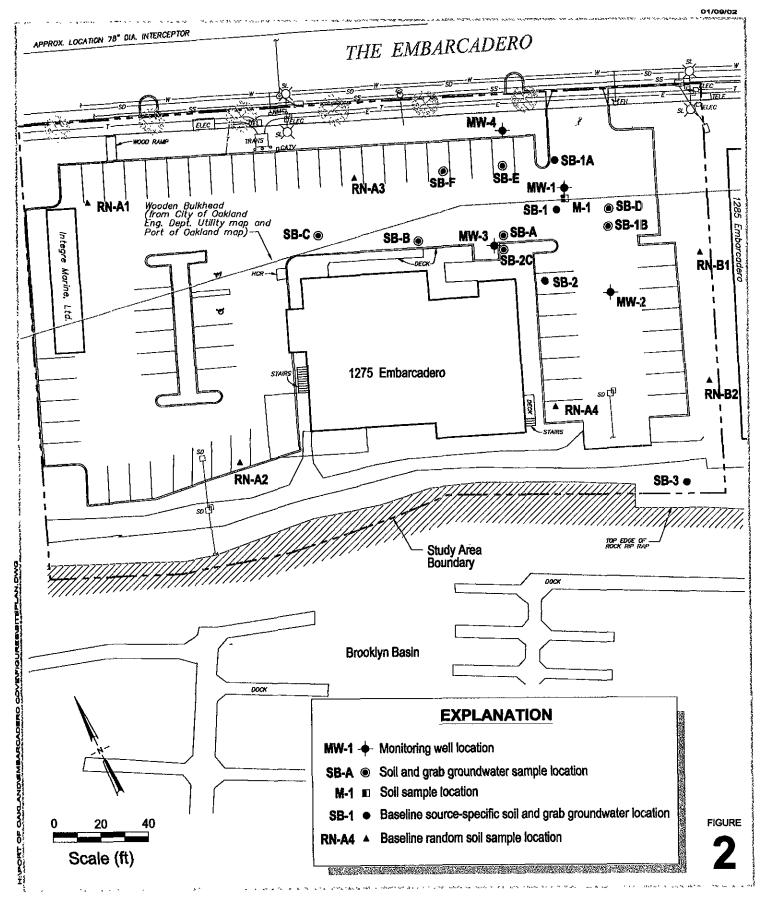


1275 Embarcadero Embarcadero Cove Project Oakland, California



Vicinity Map

CAMBRIA

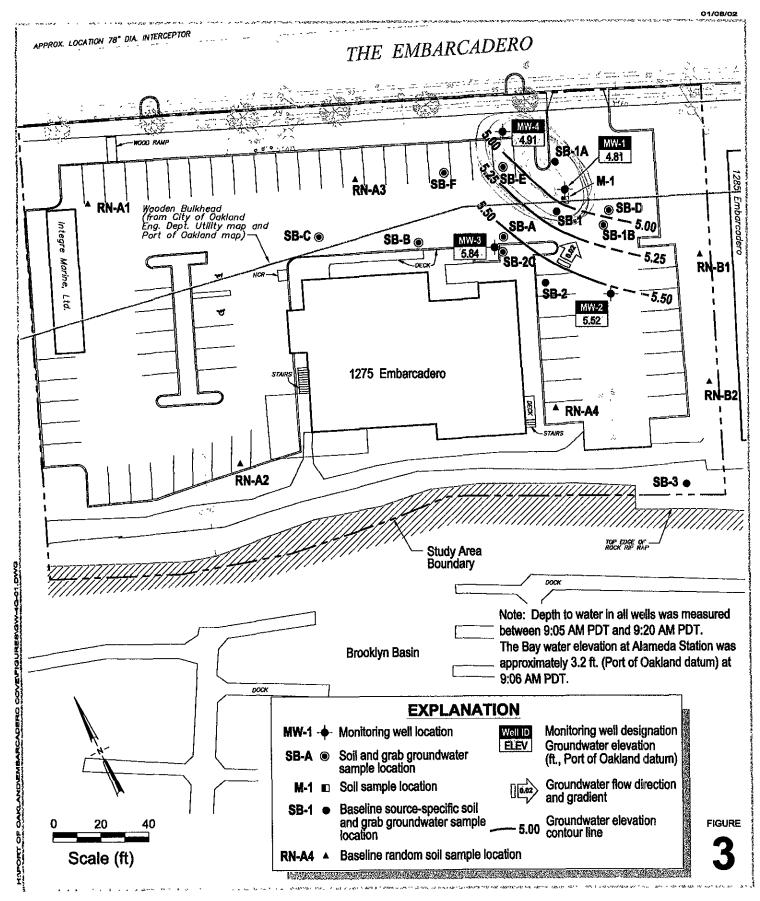


1275 Embarcadero Embarcadero Cove Project Oakland, California



Site Plan

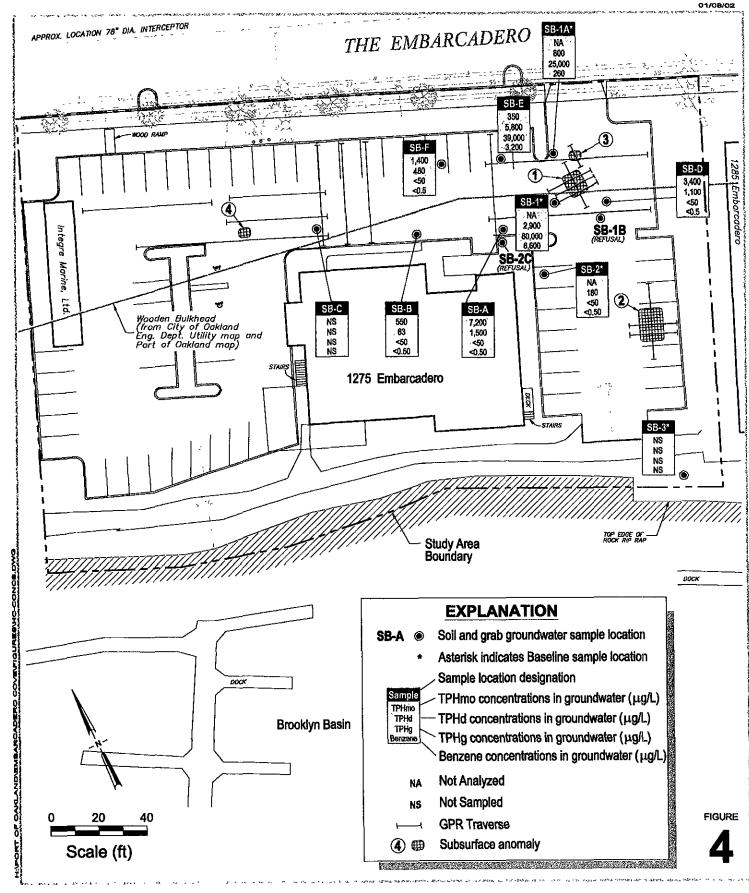
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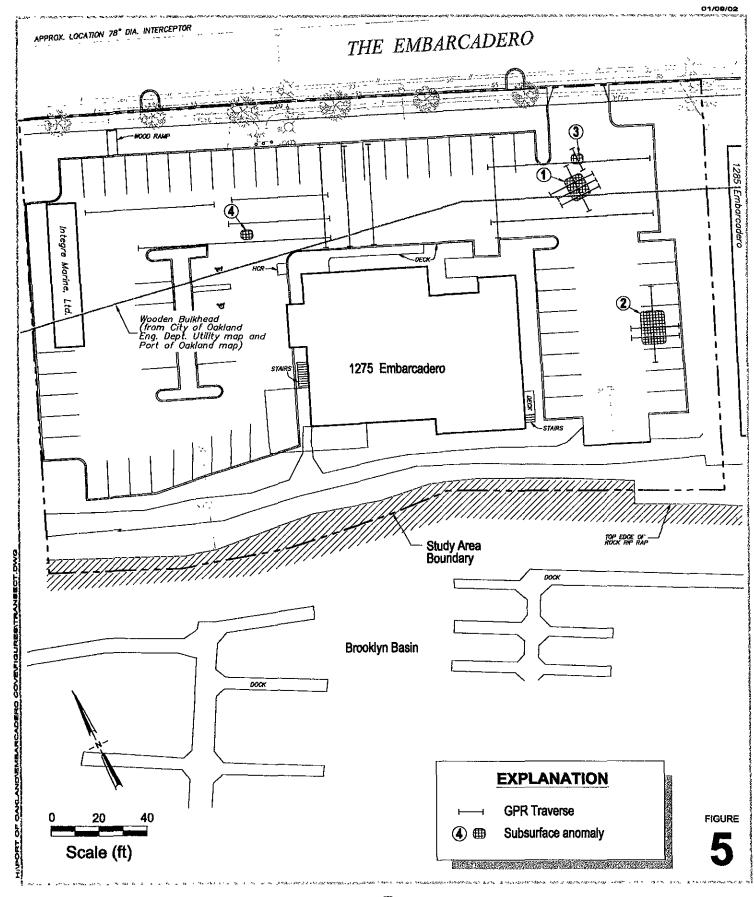
Groundwater Elevation Contours (Rising Tide)



1275 Embarcadero Embarcadero Cove Project Oakland, California



Hydrocarbon Concentrations in Grab Groundwater from Borings

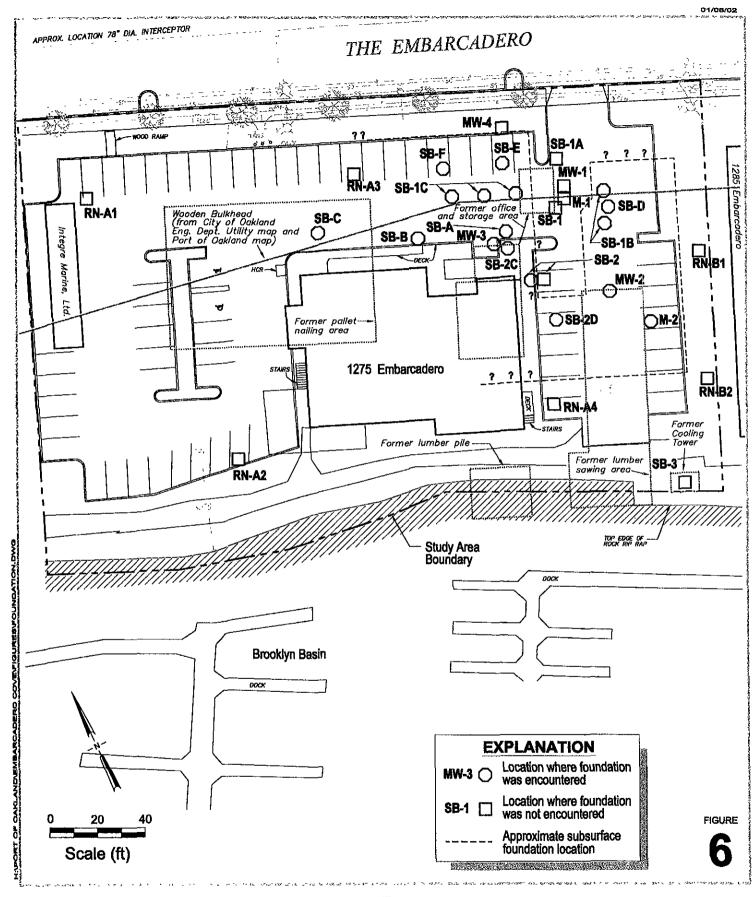


1275 Embarcadero Embarcadero Cove Project Oakland, California



Transects and Test Plt Locations

CAMBRIA

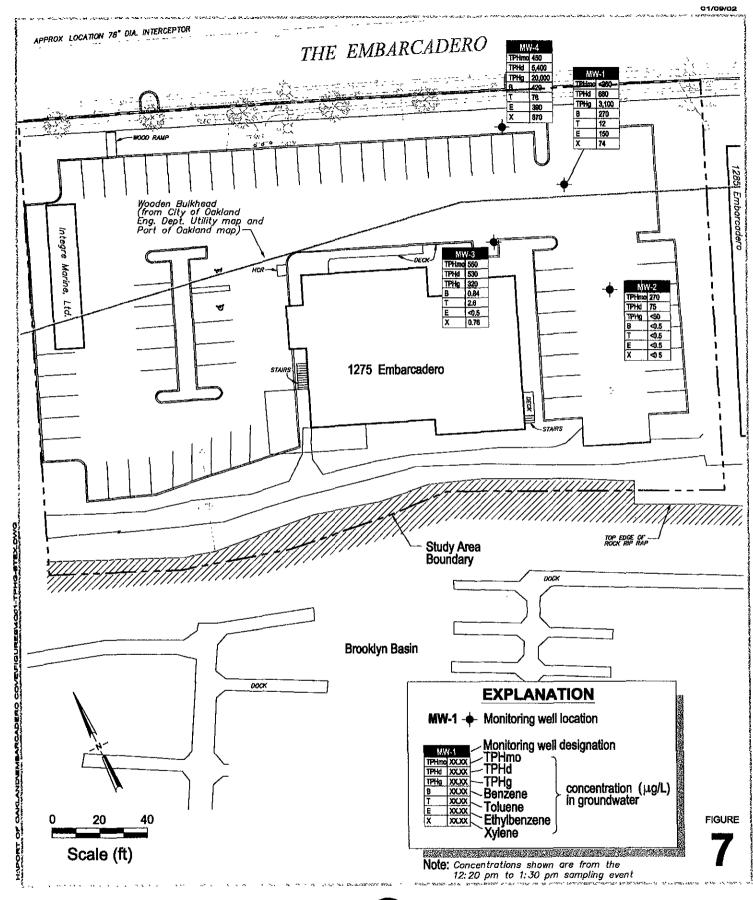


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Foundation Boundaries

CAMBRIA



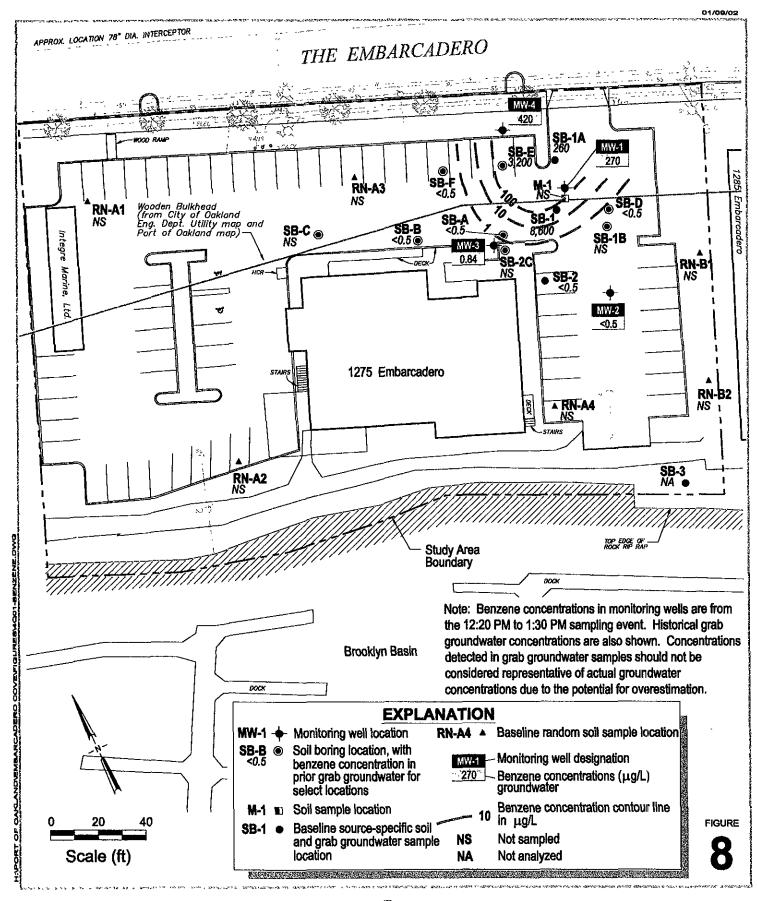
1275 Embarcadero Embarcadero Cove Project Oakland, California



Hydrocarbon Concentrations in Groundwater

CAMBRIA

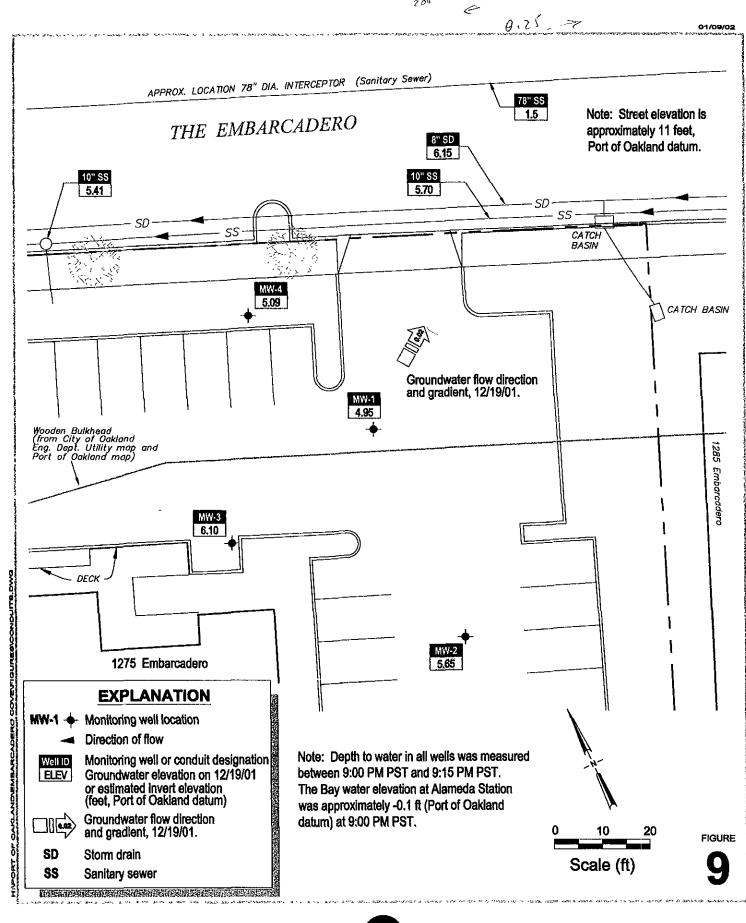
for December 5, 2001



1275 Embarcadero Embarcadero Cove Project Oakland, California



Benzene Concentrations in Groundwater



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Port of Oakland

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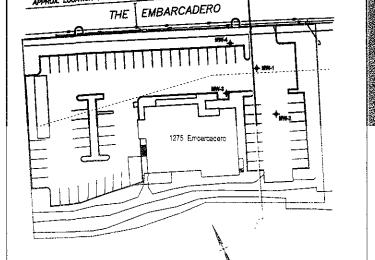


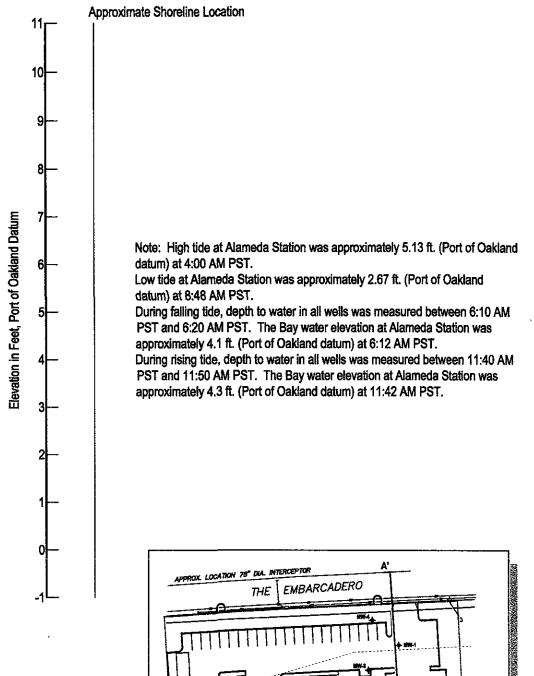
Groundwater Elevations and Deeper Subsurface Conduits



Scale (ft)

☐ Groundwater Elevation. 11:40 AM - 11:50 AM (Rising Tide)





South - Southwest

EXPLANATION

Wooden

▼ 6.69

☑ 5.54

Bulkhead MW-1

▼ 5.33

☑ 4.74

MW-4

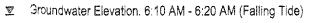
▼ 5.61

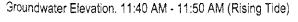
☑ 5.08

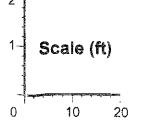
MW-2

▼ 6.11

▽ 5.66







North - Northeast

Storm
Drain Trench

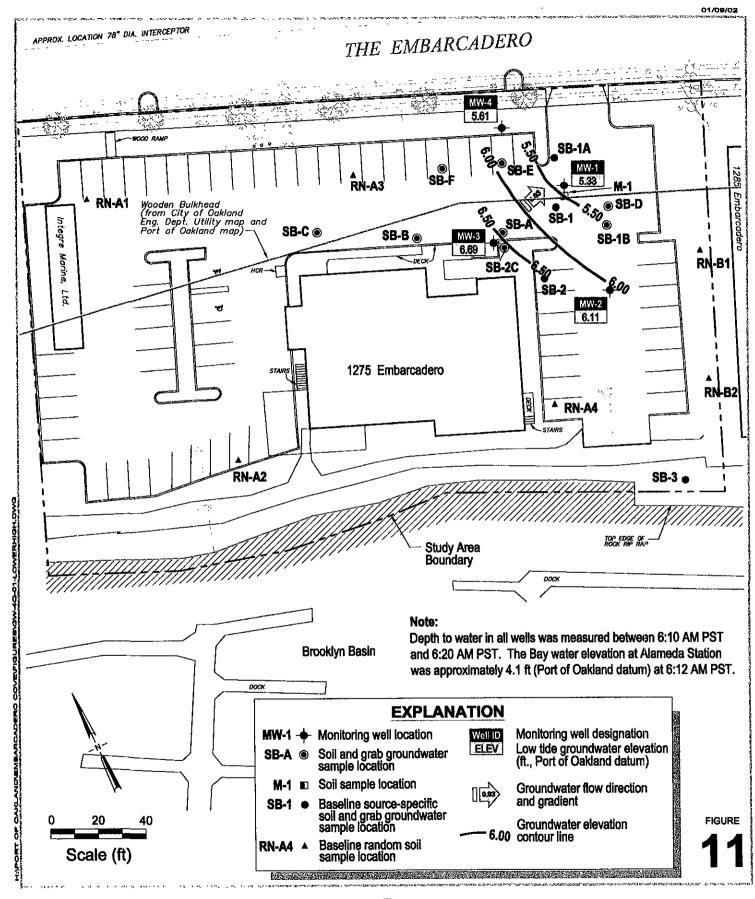
Sanitary Sewer Trench

78" interceptor Trench

10

3

FIGURE

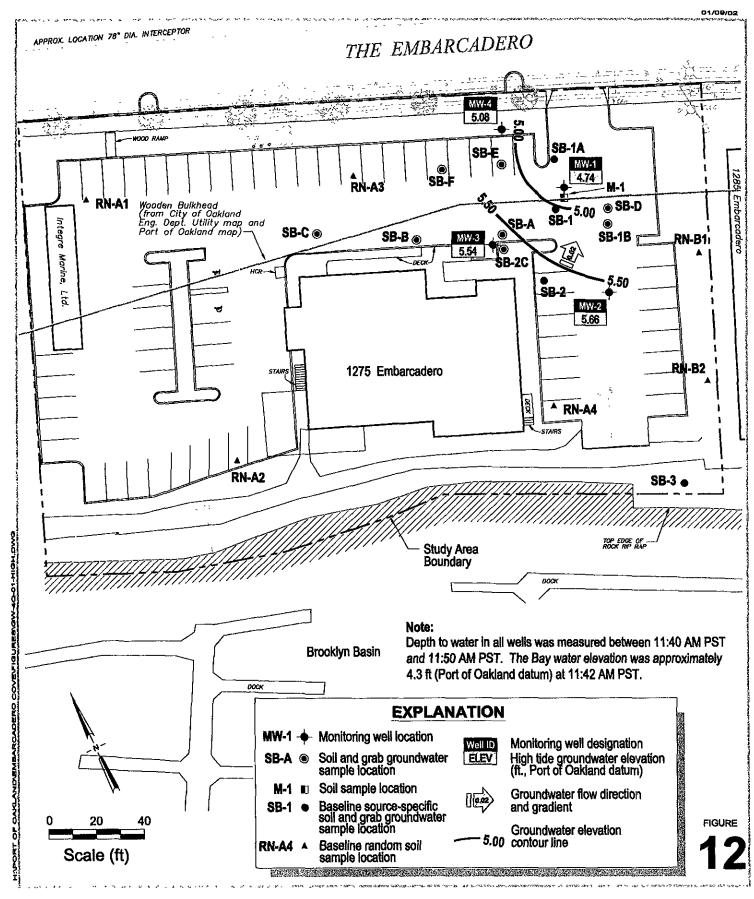


Port of Oakland

1275 Embarcadero Embarcadero Cove Project Oakland, California



Groundwater Elevation Contours (Falling Tide)

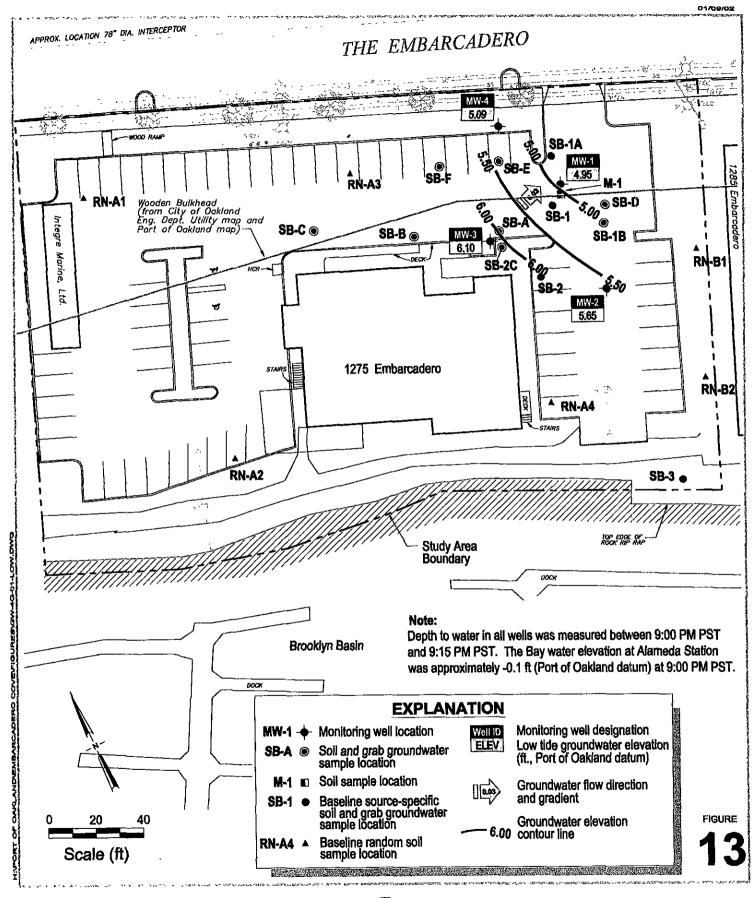


Port of Oakland

1275 Embarcadero Embarcadero Cove Project Oakland, California



Groundwater Elevation Contours (Rising Tide)



Port of Oakland

1275 Embarcadero Embarcadero Cove Project Oakland, California



Groundwater Elevation Contours (Slack Tide)

TABLES

Table 1: Soil Analytical Data - Light-Range Petroleum Hydrocarbons and MTBE - 1275 Embarcadero, Oakland, CA

Sample	Date	Sample	TPHg	Benzene	Toluene	Ethylbenzene	Xylenes	MTBE
ID	Sampled	Depth (ft)	←		m	g/kg		————
Baseline Samples			 		<u> </u>	· · · · · · · · · · · · · · · · · · ·		
RN-A1,1.5-2	05/01/01							
RN-A2;1-1.5	05/01/01	1,5		< 0.005	< 0 005	< 0.005	< 0.005	< 1.0
	05/01/01	1.0		< 0.005	< 0.005	< 0 005	< 0.005	< 1.0
RN-A3;0.5-1	05/01/01	0.5		< 0.005	< 0.005	< 0.005	< 0 005	< 1.0
RN-A4;1.0-1.5	05/01/01	1.0		< 0.005	< 0.005	< 0.005	< 0.005	< 1.0
RN-B1;1-1.5	05/01/01	1.0		< 0.005	< 0.005	< 0.005	< 0.005	< 1.0
RN-B2;1-1.5	05/01/01	1.0	₩~	< 0.005	< 0.005	< 0.005	< 0.005	< 1.0
SB-1;0.75-1.25	05/01/01	0.75	< 1.1	< 0.0056	< 0.0056	< 0.0056	< 0.0056	
SB-1;3-3.5	05/01/01	3.0	< 1.0	0.013	< 0.0052	< 0.0052	< 0.0052	
SB-1A;0-0.5	05/02/01	0.0	< 1.1	< 0.0054	< 0.0054	< 0.0054	< 0.0054	
SB-1A;5-5.5	05/02/01	5.0	500	< 0.130	F.1	5.0	16.1	
SB-1B;1-1 5	05/02/01	1.0	< 1.0	< 0.005	< 0.005	< 0.005	0.0074	
SB-2;1-1.5	05/01/01	1.0	< .98	< 0.0049	< 0.0049	< 0.0049	< 0.0049	
SB-2;4-4.5	05/01/01	4.0	< 1.1	< 0.0054	< 0.0054	< 0.0054	< 0.0054	
SB-2C;0-0.5	05/02/01	0.0	< 0.96	< 0.0048	< 0.0048	< 0.0048	< 0.0048	••
SB-2C,3-3.5	05/02/01	3.0	< 1.0	< 0.0052	< 0.0052	< 0 0052	< 0.0052	
Cambria Samples								
SB-A-3.5	08/30/01	3.5	< 1.0	< 0.005	< 0.005	< 0.005	< 0.005	< 0.05
SB-B-3.5	08/30/01	3.5	< 1.0	< 0.005	< 0.005	< 0.005	< 0.005	< 0.05
SB-D-3.5	08/30/01	3.5	< 1.0	< 0 005	< 0.005	< 0.005	< 0.005	< 0.05
SB-E-3.5	08/30/01	3.5	1.4	0.014	0.0080	< 0.005	0.003	< 0.05
SB-F-3.5	08/30/01	3.5	2.5	0.021	0.010	< 0.005	0.005	< 0.05
VETS STATE OF THE	09/17/01	5.0	2,300	=	3.7	48	7.2	
MW-1-8.3	10/09/01	8.3	30	0.48	0.067	0.70	0.52	5.1
MW-2-5.0	10/09/01	5.0	7.5	0.027	0.007	0.70	0.087	< 0.05
MW-3-5.0	10/09/01	50	1.6	< 0.005	< 0.005	_		<0.05
MW-4-5.3	10/09/01	5.3	34	0.70	0.068	< 0.005 0.41	< 0.005 0.97	<0.05 <0.05

Table 1: Soil Analytical Data - Light-Range Petroleum Hydrocarbons and MTBE - 1275 Embarcadero, Oakland, CA

Sample	Date	Sample	TPHg	Benzene	Toluene	Ethylbenzene	Xylenes	MTBE
ID	Sampled	Depth (ft)			in .	g/kg		
Soil Screening Vali								
	Commercial Worker [non-drinking water	source]					
Human Health F	Risk-Based		(11.000):	0.39	89	220	210 sat	69
Soil Leaching-B	ased for Protection of A	quatic Life	400	2.1	8.4	24	1	1
Urban Area Eco	toxicity-Based			25	150			
Construction Work	er²							
TT TT1-1 T	Risk-Based		16.000	16	520 sat	230 sat	210 sat	4900

Abbreviations and Methods:

ft = feet

mg/kg = milligrams per kilogram

-- = not available, not analyzed, or does not apply

MTBE = methyl tert-butyl ether by EPA Method 8020

Benzene, toluene, ethylbenzene, and xylenes by EPA Method 8020 or 8021B

TPHg = total petroleum hydrocarbons as gasoline by EPA Methods modified 8015, 5030, and 8020 or 602

sat = saturation limit

Notes:

Bolded values indicate exceedance of soil screening values.

¹ Soil screening values from RWQCB's (2000) Table B 2

² Soil screening values from RWQCB's (2000)

Table 2: Soil Analytical Data - Heavy-Range Petroleum Hydrocarbons and SVOCs - 1275 Embarcadero, Oakland, CA

Sample ID	Date Sampled	Sample Depth (ft)	ТРНа	TPHmo	bis(2-ethylhexyl) phthalate	Fluoranthene	2 - methyl naphthalene	Naphthalene	Pyrene
i D	Sampled	Depth (11)				mg/kg			
Baseline Samples ¹			<u> </u>						
RN-A1;1.5-2	05/01/01	1.5						< 0.0046	
RN-A2;1-1.5	05/01/01	1.0			~			< 0.005	
RN-A3;0.5-1	05/01/01	0.5					_	< 0.005	
RN-A4:1.0-1.5	05/01/01	1.0						< 0.0046	_
RN-B1;1-1.5	05/01/01	1.0	-					< 0.0049	
RN-B2;1-1,5	05/01/01	1.0	-					< 0.0047	
SB-1;0.75-1.25	05/01/01	0.75	62 ²		< 0.33	< 0.33	< 0.33	< 0.33	< 0 33
SB-1;3-3.5	05/01/01	3.0	13 ²		0.61	< 0.33	< 0.33	< 0.33	< 0.33
SB-1A;0-0.5	05/02/01	0.0	240 ²	-	< 6.60	< 6 60	< 6.60	< 6.60	< 6 60
SB-1A;5-5.5	05/02/01	5.0	40 ²		< 0.33	< 0.33	2.2	2.2	< 0.33
SB-1B;1-1.5	05/02/01	1.0	60 ²		••			***	
SB-2;1-1.5	05/01/01	1.0	43 ²		< 0.33	< 0.33	< 0.33	< 0.33	< 0.33
SB-2;4-4.5	05/01/01	4.0	43 ²	••	< 0.33	< 0.33	< 0.33	< 0.33	< 0.33
SB-2C;0-0.5	05/02/01	0.0	25 ²		***				
SB-2C;3-3.5	05/02/01	3.0	37 ²		~~				
Cambria Samples									
SB-A-3.5	08/30/01	3.5	1 4	5.2	***	< 0.25		< 0.25	< 0.25
SB-B-3.5	08/30/01	3.5	< 1.0	< 5.0		< 0.062		< 0.062	< 0.06
SB-D-3.5	08/30/01	3.5	< 1.0	< 5.0		< 0.062		< 0.062	< 0.06
SB-E-3.5	08/30/01	3.5	2.4	6.1	<u></u>	< 0.062	**	< 0.062	< 0.06
SB-F-3.5	08/30/01	3.5	4.6	16	***	< 0.25		< 0.25	< 0.25
Meron	09/17/01	50	250	97				**	
MW-1-8.3	10/09/01	8.3	5.7	< 50	< 0.33	< 0.33	< 0.33	$< 0.33 / 0.58^3$	< 0.33
MW-2-5.0	10/09/01	5.0	74	300	< 1.0	< 1.0	< 1.0	< 1.0 / < 0.005 ³	< 1.0
MW-3-5.0	10/09/01	5.0	17	160	< 0.33	< 0.33	< 0.33	< 0.33 / < 0.005 ³	< 0.33
MW-4-5.3	10/09/01	5.3	8.3	10	< 0.33	< 0.33	< 0.33	0.62 / 0.623	< 0.33
Soil Screening Values	,								
Surface Soil (<3 m) C		[non-drinking water s	sourcel 4						
Human Health Ris		,	11,000	11,000	180	6,000	280	5.7	11,000
Soil Leaching-Bas	ed for Protection of .	Aquatic Life	500	1,000	530	60	0 25	4.9	55
Urban Area Ecotor Construction Worker			-			40		40	-
Human Health Ris			16,000	16,000	1,200	12,000			

Table 2: Soil Analytical Data - Heavy-Range Petroleum Hydrocarbons and SVOCs - 1275 Embarcadero, Oakland, CA

bis(2-ethylhexyl) 2 - methyl Sample Date Sample **TPHd TPHmo** phthalate Fluoranthene naphthalene Naphthalene Pyrene ID Sampled Depth (ft) mg/kg

Abbreviations and Methods:

ft = feet

mg/kg = miligrams per kilogram

-- = not available, not analyzed, or does not apply

TPHd = total petroleum hydrocarbons as diesel by EPA method 8015

TPHd analyses with silica gel clean-up prior to extraction unless otherwise noted

TPHmo = total petroleum hydrocarbons as motor oil by EPA method 8015

SVOC = semi-volatile organic compounds by EPA Method 8270 (modified 8100) and 3550 or 625 and 3510 unless otherwise noted

Notes:

Only those compounds above laboratory reporting limits are shown

- ¹ Baseline samples analyzed for SVOCs by EPA Method 8260 or 8270
- ² No silica gel cleanup performed, prepared by shaker table.
- 3 Analyzed by EPA Method 8270 and additionally by EPA Method 8260
- ⁴ Soil screening values from RWQCB's (2000) Table B-2.
- ⁵ Soil screening values from RWQCB's (2000) Table K-3.

Bolded values indicate exceedance of soil screening values.

Table 3: Soil Analytical Data - Additional VOCs - 1275 Embarcadero, Oakland, CA

Sample ID	Date Sampled	Sample Depth (ft)	n-Butyl benzene	Isopropyl benzene	p-Isopropyl toluene	n-Propyl benzene ug/kg	1,2,4-Trimethyl benzene	1,3,5-Trimethy benzene
Cambria Samples			····					
MW-1-8.3	10/09/01	8.3	580	210	< 25	910	450	160
MW-2-5.0	10/09/01	5.0	18	< 5 0	11	17	26	11
MW-3-5.0	10/09/01	5.0	< 5.0	< 5.0	< 5.0	< 5.0	5.7	< 5.0
MW-4-5 3	10/09/01	5.3	< 25	220	< 25	870	1,000	610
Human Health	Commercial Worker Risk-Based Based for Protection of otoxicity-Based ker	_	ter source]	 	 	 	 	

Abbreviations and Methods:

µg/kg = micrograms per kilogram

-- = not available, not analyzed, or does not apply

VOC = volatile organic compounds by EPA Method 8260

Notes:

Only those compounds above laboratory reporting limits are shown

Table 4: Soil Analytical Data - Metals - 1275 Embarcadero, Oakland, CA

Sample ID	Date Sampled	Sample Depth (ft)	Barium (Ba)	Chromium (Cr) 1	Hexavalent Chromium (Cr ⁶⁺)	Cobalt (Co)	Copper (Cu) - mg/kg	Lead (Pb)	Nickel (Ni)	Vanadium (V)	Zinc (Zn)
Baseline Samples		<u></u> -									
СОМР А	04/23/01	2	41	22		5.3	17	24	25	20	40
SB-3; 0.5-1	05/01/01	0.5			<0.05		<u>-</u> -				
SB-3; 3.5-4	05/01/01	3 5			<0.05					••	
Cambria Samples											
MW-1-8 3	10/09/01	8.3	110	56	**	12	15	26	64	33	29
Soil Screening Va											****
Surface Soil (<3 n Human Health		er [non-drinking wa	•								
		n f. A. numero T. 16n	2,400	610,000	1.8	24,000	15,000	1,000	1,000	2,900	120,000
	Based for Protection of otoxicity-Based	or Aduant File	1.500	750			••				
			1,500	750	8	80	225		150	200	600
Construction Wor											
Human Health	Risk-Based		2,400	800,000	1.8	32,000	20,000	1,000	1,000	3,700	160,000

Abbreviations and Methods:

ft = feet

mg/kg = milligrams per kilogram

-- = not available, not analyzed, or does not apply

Metal analyses (CAM / CCR 17) by EPA Method 6010

Baseline SB-3 samples analyzed for hexavalent chromium by EPA Method 7196

Notes:

Only those compounds above laboratory reporting limits are shown

COMP A is four-point composite of RN-A1, RN-A2, RN-A3, and RN-A4.

¹ For purposes of this analysis, we have assumed that the total chromium detected at the Site is in the trident form. This asumption is consistent with the data collected at the Site (i.e., hexavalent chromium was not detected in samples from SB-3), and is based on our experience at similar industrial sites, where absent a source of hexavalent chromium, chromium in soils is typically found in the trivalent (reduced) form.

² Soil screening values from RWQCB's (2000) Table B-2.

³ Soil screening values from RWQCB's (2000) Table K-3.

Table 5: Groundwater Analytical and Elevation Data - Light-Range Petroleum Hydrocarbons and MTBE - 1275 Embarcadero, Oakland, CA

Sample ID	Date	Groundwater	Depth to	ТРНд	Benzene	Toluene	Ethylbenzene	Xylenes	MTBE
TOC	Sampled	Elevation	Water			μ	g/L		
(ft)		(ft 1)	(ft)						
Baseline Grab S	_								
SB-I	05/01/01		_	80,000	8,600	8,200	3,900	14,600	
SB-1A	05/02/01	-	-	25,000	260	170	760	2,290	
SB-2	05/01/01			< 50	< 0.5	< 0.5	< 0.5	< 0.5	
Cambria Grab S	amples								
SB-A	08/30/01			< 50	< 0.5	< 0.5	< 0.5	< 0.5	< 5.0
SB-B	08/30/01		-	< 50	< 0.5	< 0.5	< 0.5	< 0.5	< 5.0
SB-D	08/30/01		_	< 50	< 0.5	< 0.5	< 0.5	< 0.5	< 5.0
SB-E	08/30/01			39,000	3,200	750	1,200	3,600	< 200
SB-F	08/30/01			< 50	< 0.5	< 0.5	< 0.5	< 0.5	< 5.0
Cambria Monite	oring Well Sample	?\$							
MW-i	10/12/01	4.88	7.15						
12.03	10/19/01	4.81	7.22	11,000	900	300	470	1,000	
12,00	12/05/01 ²	5.33	6.70	13,000	1,300	180	1,200	860	< 20
	12/05/01 3	4.74	7.29	3,100	270	12	150	74	< 50
	12/19/01 4	4.95	7.08				-		
MW-2	10/12/01	5.71	5.75	••					
11.46	10/19/01	5.52	5.94	< 50	< 0.5	< 0.5	< 0.5	< 0.5	
	12/05/01 2	6.11	5.35	< 50	< 0.5	< 0.5	< 0.5	< 0.5	< 5.0
	12/05/01 3	5.66	5 80	< 50	< 0.5	< 0.5	< 0.5	< 0.5	
	12/19/01 4	5.65	5.81	-		~ 03	~ 	< 0.5 	< 5.0
MW-3	10/12/01	5.89	6.60	<u></u>					
12.49	10/19/01 5	5.84	6.65	290	2.0	6.6	0.54	1.2	***
	12/05/01 2,5	6.69	5.8	310	0.72	2.2	< 0.5	< 0.5	. # A
	12/05/01 3,5	5.54	6.95	320	0.84	2.6			< 5.0
	12/19/01 4	6.10	6.39	320			< 0.5	0.76	< 5.0
MW-4	10/12/01	4.98	8.15		 -				
13.13	10/19/01	4.91	8.22	44,000	- 1,900	 270		2 200	
-	12/05/01 2	5.61	7.52	13,000	1,900		1,500	3,300	
	12/05/01 ³	5 08	8.05			28	170	380	< 10
	12/19/01	5.09	8.03 8.04	20,000	420	78	390	870	< 20
Trip Blank	1211/101	40،۲	o U4	••	<u></u>		•		
TB	12/05/01	<u></u>		 < 50	 0.5		-0.6		
	· ~ 03/01		_	< 30	< 0.5	< 0.5	< 0.5	< 0.5	< 5.0

Table 5: Groundwater Analytical and Elevation Data - Light-Range Petroleum Hydrocarbons and MTBE - 1275 Embarcadero, Oakland, CA

Sample ID TOC (ft)	Date Sampled	Groundwater Elevation (ft 1)	Depth to Water	ТРНg	Benzene	Toluene	Ethylbenzene ug/L ————	Xylenes	мтве
Groundwater Sci Indoor Air Impac Aquatic Life Pro	ets		(4)	_ 3,700 ^a	84 700 ^b	76,000 5,000 ^b	170,000 sol 430 °	150,000 13 ^d	290000 8,000°

Abbreviations and Methods:

ft = feet

µg/L = micrograms per liter

-- = not available, not analyzed, or does not apply

msl = mean sea level

Benzene, toluene, ethylbenzene, and xylenes by EPA Method 8020

MTBE = methyl tert-butyl ether by EPA Method 8020

TPHg = total petroleum hydrocarbons as gasoline by EPA Methods modified 8015, 5030, and 8020 or 602

TOC Elev. (ft) = top of casing elevation in feet (Port of Oakland datum)

Depth to water in monitoring wells is ft below TOC.

sol = solubility threshold

Notes:

Bolded values indicate exceedance of groundwater screening values.

¹ Elevation in feet, Port of Oakland datum

² Wells gauged between 6:00 am and 6:30 am on 12/5 near lower high tide.

³ Wells gauged between 11:40 am and 12:00 pm on 12/5 near higher high tide.

⁴ Wells gauged between 9:00 pm and 9:15 pm on 12/19 at lower low tide.

⁵ Sample was collected pre-purge.

⁶ Goundwater screening values from RWQCB's (2000) Table F-2, F-4a, b, and c.

^a California Toxic Rule, Saltwater Criteria for Continuous Concentration

^b USEPA Saltwater Chronic Lowest Observable Effect Level

^c USEPA Saltwater Acute Lowest Observable Effect Level

^d USDOE Freshwater Chronic Preliminary Remedial Goal

^{*} RWQCB Saltwater Criteria for Continuous Concentration (interim)

Table 6: Groundwater Analytical and Elevation Data - Heavy-Range Petroleum Hydrocarbons and SVOCs - 1275 Embarcadero, Oakland, CA

Sample ID	Date	Groundwater	Depth to	TPHd	TPHmo	Acenaph- thalene	bis(2-ethylhexyl) phthalate	Fluoran- thene	1 - methyl- naphthalene	2 - methyl naphthalene	Naph- thalene	Phenan- threne	Pyrene
TOC	Sampled	Elevation	Water ◀					μg/L				·	
(ft)		(ft ^l)	(ft)										
Baseline Grab	•												
SB-I	05/01/01			2,900		< 94	< 94	< 94		260	610	< 94	< 94
SB-1A	05/02/01			800		<96	<96	<9.6		130	170	<96	<9.6
SB-2	05/01/01			180		< 9.7	< 97	< 9.7		< 9.7	< 9.7	< 9.7	< 9.7
Cambria Grab	Samples												
SB-A	08/30/01			1,500	7,200	< 10		< 10			< 10	< 10	. 10
SB-B	08/30/01			63	550	< 10		< 10			< 10	< 10 < 10	< 10
SB-D	08/30/01			1,100	3,400	< 10		11		 	< 10		< 10
SB-E	08/30/01			5,800	350	< 50		< 50	40		370	< 10 < 50	11
SB-F	08/30/01			480	1,400	< i0		< 10	**	**	< 10	< 50 < 10	< 50 < 10
Combain Marie	oring Well Sampl												
MW-1	oring weii Sampi 10/12/01												
12.03	10/12/01	4.88	7 15				**						••
12.03	12/05/01 2	4.81	7.22	3,300	< 250	< 10	< 10	< 10	••	54	66	< 10	< 10
	12/05/01 3.4	5.33	6.70	3,800	< 250	72		< 10	150	220	360	< 10	< 10
	12/19/01 5	4.74	7.29	680	< 250	9.6		< 1.0	18	14	22	1 3	< 1.0
MW-2	10/12/01	4.95	7 08										
11.46	10/12/01	5.71	5.75	••									
11.40	10/19/01	5.52	5.94	210	460	< 10	< 10	< 10		< 10	< 10	< 10	< 10
	12/05/01	6.11	(3 <u>3</u> 5)	150	560	< 0.5		< 0.25	< 1.0	< 1.0	< 0.25	< 0.25	< 0 25
	12/19/01	5.66	5.80	75	270	< 0.5		< 0.25	< 1.0	< 1.0	< 0.25	< 0.25	< 0.25
MW-3		5.65	5.81					••				••	
	10/12/01 10/19/01 ⁶	5.89	6.60	••									**
12.49	10/19/01 12/05/01 ^{2, 6}	5 84	6.65	1,600	1,300	< 25	< 25	< 25		670	420	< 25	< 25
	12/05/01 3,6	6.69	5.80	480	480	< 0.5	••	< 0.25	< 1.0	< 10	< 0.25	< 0.25	< 0.25
		5.54	6.95	530	550	< 0.5	••	< 0.25	< i.0	< 1.0	< 0.25	< 0.25	0.31
10014	12/19/01 5	6.10	6.39			**	**			••			
MW-4	10/12/01	4.98	8.15						••				••
13 13	10/19/01	4 91	8 22	33,000	900	< 50	< 50	< 50		< 50	< 50	< 50	< 50
	12/05/01 ²	5.61	7.52	6,400	430	24		< 10	99	190	60	18	< 10
	12/05/01 3	5.08	8.05	5,400	450	21	~~	< 10	100	180	96	12	< 10
	12/19/01 5	5.09	8.04						••			**	

Table 6: Groundwater Analytical and Elevation Data - Heavy-Range Petroleum Hydrocarbons and SVOCs - 1275 Embarcadero, Oakland, CA

Sample ID TOC	Date Sampled	Groundwater Elevation (ft ¹)	Depth to Water ◀	ТРН	TPHmo	Acenaph- thalene	bis(2-ethylhexyl) phthalate	Fluoran- thene — µg/L —	1 - methyl- naphthalene	2 - methyl naphthalene	Naph- thalene	Phenan- threne	Pyrene
	creening Values 7		<u> </u>			<u></u>						 	
Indoor Air Impa							•-	••	26,000-sol	26,000 sol	9,200		135 sol
Aquatic Life Pr	otection			640 ª	640 *	310 ^g	32 ^b	11 °	2.1 4	2.1 d	2,350 °	4.6'	300 °
Abbreviations	and Methods:						Notes:						
ft = feet							1 Elevation in feet, Po	rt of Oakland da	nun				
µg/L = microgr	ams per liter						² Wells gauged between	en 6:00 am and	6:30 am on 12/5 near	lower high tide.			
= not availab	le, not analyzed, o	does not apply					3 Wells gauged between						
msl = mean sea	level						SVOC extraction pe				36 Revision 3,	12/96.	
TOC Elev. (ft) :	= top of casing ele	vation in feet (Port of (Oakland datum)				5 Wells gauged betwe						
TPHd analyses	with silica gel clea	n-up prior to extraction	n				⁶ Sample was collecte		•				
TPHmo = total	petroleum hydroca	rbons as motor oil by	EPA method 801:	5					RWOCB's (2000) Tab	le F-2, F-4a, b, and c.			
		npound analyses perfo					Bolded values indica						
EPA Me	ethod 8270 (modif	ied 8100) and 3550 un	less otherwise no	ted			* RWQCB Saltwater			3			
		ons as diesel by EPA			0		^b USEPA Freshwater						
		ratory reporting limits					" USEPA Saltwater C						
		s is ft below TOC.					d USDOE Freshwater		•				
sol = solubility	threshold						° USEPA Saltwater A						
							USEPA Saltwater C	Inenon for Con	tinuous Concentration	<u>.</u>			
							B Ontario Ministry of						

Table 7: Goundwater Analytical and Elevation Data - Metals - 1275 Embarcadero, Oakland, CA

Sample ID	Date Sampled	Groundwater Elevation (ft 1)	Depth to Water (ft)	Antimony (Sb)	Arsenic (As)	Barium (Ba)	Beryl- lium (Be)	Cad- mium (Cd)	Chro- mium (Cr)	Cobalt (Co)	Copper (Cu)	Lead (Pb) - ug/L	Mercury (Hg)	Molyb- denum (Mo)	Nickel (Ni)	Sele- nium (Se)	Silver (Ag)	Thallium (Tl)	Vana- dium (V)	Zinc (Zn)
Cambria S	amples																			
MW-1	12/07/01 2	6.85	5.18	< 6	5.7	57	< 4	< 5	< 20	< 50	< 50	5.3	< 0.8	< 50	< 50	< 5	< 10	< 5	< 50	< 50
MW-1	12/07/01 3	7 15	<u>4</u> .88	< 6	< 5	51	< 4	< 5	< 20	< 50	< 50	< 5	< 0.8	< 50	< 50	< 5	< 10	< 5	< 50	50

Abbreviations and Methods:

ft = feet

ug/L = micrograms per liter

Metal analyses (CAM / CCR 17) by EPA Method 6010

Notes:

Only those compounds above laboratory reporting limits are shown

¹ Elevation in feet, Port of Oakland damm

² Gauged at 5 40 am on 12/7 near higher high tide

³ Gauged at 11.40 am on 12/7 near lower high tide

APPENDIX A

Soil Boring Logs



Cambria Environmental Technology, Inc 1144 - 65th St Oakland, CA 94608 Telephone (510) 420-0700 fax (510) 420-9170

SAMPLE

	JOB/S. LOCAT PROJE DRILLI DRILLI BORIN LOGGE REVIEL REMAR	ECT NUMI ER ING METH G DIAME ED BY WEO BY	HOD	ABC Hollo 8"	Drilling ow stem	auger ed to 4 l	t bgs		DRILLING STARTED DRILLING COMPLETED WELL DEVELOPMENT D GROUND SURFACE ELE TOP OF CASING ELEVA SCREENED INTERVAL DEPTH TO WATER (State	ATE (YIELD) VATION FION 4 to 19 Encountered	NA fl abov 5 fl bqs	re msl : 1 4 ft {06	
	PID (ppm)	BLOW	SAMPLE 10	EXTENT	DEPTH (ft bgs)	U.S.C.S	GRAPHIC		LOGIC DESCRIPTION		CONTACT DEPTH (ft bgs)	\ \ \ \	VELL DIAGRAM
	augeri	resenta	>					fine to coarse grained gravel, high estimated	lilf, light brown; dry; 20% s sand, 30% line to coarse of permeability tween lithologic units	ilt, 50% grained			✓ Portland Type t/il Cement ✓ Bentonite Seal ✓ Monterey Sand #3
	63	atory an	MW-4 5 3	X OI	- 5 -	MH SM ◀		20% sand, medium plant of the control of the contro	rey, moist, 20% clay, 80% estimated permeability ay wet 45% silt, 55% tine a estimated permeability. Contact between lithol	sill.	45 53 60		
ptoi	o 6	visual id	MW-4- 10 5	XXX	g of	MH I		of lithologic CLAYEY SILT(MH): Gr medium plasticity, low e SILTY SAND(SM) Gre grained sand, moderate CLAYEY SILT(MH). Gr sand, 3% gravel, mediu permeability, roots	ey, wet, 20% clay, 80% silt estimated permeability. Wet, 45% silt, 55% fine estimated permeability. ey, wet, 20% clay, 75% silt m plasticity, low estimated ey, wet, 20% clay, 80% silt.	. 2%.	103 105 118		→ 2"-diam., 0.020" Sfotted Schedule 40 PVC
- State of States in State of the State of t				w pe			alifo I	140 lbs hammer fallin rnia Modified Sample ste: All lithologic de			50		Bottom of Boring & 15 ft
) <u>;</u>				1.				•	ations unless otherwis	e noted.		ا مرد حدد	

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Cambria Environmental Technology, Inc. 1144 - 65th St. Oakland, CA 94608 Telephone: (510) 420-0700 Fax: (510) 420-9170

REMARKS	NUMBER METHOD AMETER Y BY	Em 127 458 V&\ Hol 8" K. C	75 Emba 3-1705 W Drilling Ilow-stem Cejka Schultz, I	ero Cov arcadero ng n auger RG# 70 previou	re o, Oakia r 012	it area to 6 ft bgs; located	BORING/WELL NAME DRILLING STARTED DRILLING COMPLETED WELL DEVELOPMENT DA GROUND SURFACE ELEN TOP OF CASING ELEVAT SCREENED INTERVAL DEPTH TO WATER (Station d 37.7 ft WNW of eastern pl	12- Not t above 5 ft bgs d) 8	12-Oct-01 Not Surveyed above msl 5 ft bgs I) 8.6 ft (09-Oct-01) 7.15 ft (12-Oct-01)		
39 ² 2	MW 5.	/-1-	30 5	U.S.	GRA	FILL: Gravelly sandy fine to coarse grained permeability Clayey SILT(MH): Bla	r fill; light brown; dry; 20% si I sand, 30% gravel; high est ack to grey; damp; 20% clay asticity; low estimated perme	imated	5.0		Portland Type I/II Cement Bentonite Sea Monterey Sand #2/12
0 1 2 3	MW 8.		-10	МН			nedium plasticity; shell fragn rish brown; wet; 10% clay, 8 medium plasticity	Ā	15.0		2"-diameter, 0.010" Slotted Schedule 40 PVC Bottom of Boring @ 15 ft

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CLIENT NAME	Port of Oakland	BORING/WELL NAME MW-2
JOB/SITE NAME	Embarcadero Cove	DRILLING STARTED 08-Oct-01
LOCATION	1275 Embarcadero, Oakland, CA	DRILLING COMPLETED 09-Oct-01
PROJECT NUMBER _	458-1705	WELL DEVELOPMENT DATE (YIELD) 12-Oct-01
DRILLER	V&W Drilling	GROUND SURFACE ELEVATION Not Surveyed
DRILLING METHOD	Hollow-stem auger	TOP OF CASING ELEVATION 8.26 ft above msl
BORING DIAMETER _	8"	SCREENED INTERVAL 4 to 15 ft bgs
LOGGED BY	K. Cejka	DEPTH TO WATER (First Encountered) 10.0 ft (09-Oct-01)
REVIEWED BY	R. Schultz, RG# 7012	DEPTH TO WATER (Static) 5 75 ft (12-Oct-01)
REMARKS	Hand augered to 4 ft bgs; located 28 ft WNW of ea	

CONTACT DEPTH (ft bgs) SAMPLE ID PID (ppm) BLOW COUNTS EXTENT U.S.C.S. DEPTH (ft bgs) LITHOLOGIC DESCRIPTION WELL DIAGRAM **ASPHALT** 0.5 FILL: Gravelly sandy fill; grey to brown; dry; 20% silt, 40% sand, 40% medium to coarse grained gravel; high Portland Type estimated permeability 1.5 I/II Cement CONCRETE 2.4 FILL: Gravelly sandy fill; brown; dry; 20% sitt, 50% ■ Bentonite Seal sand, 30% gravel, high estimated permeability ■ Monterey Sand #2/12 4.5 Clayey SILT(MH): Grey to brown; moist; 20% clay, 5 80% silt; low to medium plasticity; low estimated 0 MW-2permeability 5.0 Ţ МН 2"-diameter, 0.010" Slotted 10-Ā 0.6 MW-2-@ 10.0 ft: Wet; shell fragments Schedule 40 10.0 3 **PVC** 5 14.4 Silty SAND(SM): Grey to brown; wet; 15% clay, 35% SM silt, 45% fine to coarse grained sand, 5% gravel; low to moderate estimated permeability; rust-colored staining 15.0 Bottom of Boring @ 15 ft

WELL LOG (PID) H.PORTOF-11EMBARC-11GINTNECOVEMW GPJ DEFAULT GDT 1/10/02

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		FOX:	Jaic	η 420-	9170							
CLIENT	NAME		Port	of Oak	land			BORING/WELL NAME	MW-3			
JOB/SI	TE NAM	E	Emb	arcade	ro Coy				08-Oct-01			
LOCAT	ION		1275	Emba	rcader	o, Oak	land, CA		09-Oct-01			
PROJE	CT NUM		458-						ATE (YIELD)	12-	-Oct-01	
DRILLE	R		V&W	Drillin					-		t Surveyed	
DRILLII	NG MET	HOD	Hollo	w-sten					ION 9.29 f	t above	e msl	
BORING	G DIAME	TER	8"									• • • • • • • • • • • • • • • • • • • •
LOGGE	DBY		K. Ce					-				Oct-01) \(\sum_{\text{\tin}\text{\tint{\text{\ti}\text{\texi}\text{\text{\text{\text{\text{\text{\text{\text{\texi}\text{\text{\text{\text{\text{\text{\texi}\text{\text{\text{\texi}\text{\text{\text{\text{\text{\text{\texit{\texi}\text{\texit{\text{\tet{\text{\text{\texi}\text{\text{\text{\texi}\texit{\text{\t
REVIEV	VED BY		R. Sc	hultz, i				DEPTH TO WATER (Statio			6.60 ft (12	=======================================
REMAR	KS _		Hand	auger	ed to 3	3.2 ft bo		eastern planter, 44.5 ft SSV				
	1	T .	77			T				1 2	2	**************************************
PID (ppm)	BLOW	SAMPLEID	뉟	王 😧	U.S.C.S.	GRAPHIC LOG				CONTACT SEPTH (# hgs)	3"	
<u>.</u>	95	뒽	EXTENT	DEPTH (ft bgs)	S.C	A O	LITHO	LOGIC DESCRIPTION		157	W	ELL DIAGRAM
뒽	<u> </u>	₹	lω	ء ت	Þ	G. G.				ĮģĒ	-	
<u> </u>		"	╌							\ <u>\</u>	<u> </u>	0
		ĺ				· ·	CONCRETE			0.5		SI .
	ļ	}					<u>FILL</u> : Sandy fill; brov gravel; high estimate	vn; dry; 20% silt, 70% sand,	10%			1
	ļ	ļ					graver, mgn estimate	и реппеавину				Portland Type I/II Cement
ļ		1		ĺ		\bowtie						
				1						-	775 7	×
		1		ĺ			CONCRETE			2.5 2.8		 Bentonite Sea
]		- †		p 6 4	ASPHALT			3.0		
]]]	ĺ		\bowtie	CONCRETE FILL: Gravelly sandy	fill; brown; dry; 20% silt, 50	%	3.3		■ Monterey
			-			\bowtie		h estimated permeability	,,			Sand #3
]		\bowtie]
	2	l		- 5 -		\bowtie	 	 		5.0		
0	3	MW-3- 5.0					Clayey SILT (MH): Da	ark greenish grey; moist; 25° nedium plasticity; very low e	% clay,			
		0.0	X				permeability; roots	nearon pasticky, very low e	Sumateu	1	月	
	5		\bowtie	- 1					_		1 - 🗐	
6	2	MW-3-				HIII			Ä	1	· ·	1
	3	6.5	M	7								4
	4		X	- 1	İ	1111			Σ		1. 目	}
1				- 1					- <u>+</u> -	1		1
[11		j	1111	@ 8.5 ft: Dark vellow	ish brown; wet; 20% clay, 86)% silt·	1		✓ 2"-diam.,
		ı	 -	1	МН		low plasticity; roots	, , , , , , , , , , , , , , , , , ,	, , , , , , , , , , , , , , , , , , , ,			0.020" Slotted
			\mathbb{H}			Ш						PVC
j			M	-10		Ш				i		
İ	ľ			- 1	ļ	HIII						,
	l		Ц			Ш						
[l			- 1	İ	Ш	@ 11.0 ft: Greenish g	rey; wet; 20% clay, 70% silt	, 10%			
ŀ	']		\Box				tine grained sand; low	estimated permeability; roo	ts	ŀ		
- 1	1		Interest	1	- 1	{{{{}}}}}}				}		
1	1	MW-3-			- 1]
1	ŀ	12.5		╁		υц			— — —	13.0		Bottom of
	į				1	ļ						Boring @ 13 ft
]						[
	- 1				1	-						
	f			1	- 1	1	Note: Well cap dislode	ged from bottom. Surge blo	ck			ļ
- 1	- 1			l	- 1	ļ	remains in well.					
	ļ					ĺ						
}	1		[]	1	- 1	}					1]

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JOB/SIT LOCATION PROJECT DRILLER DRILLIN BORING LOGGEE	ON OT NUMI R IG METH I DIAME D BY ED BY	BER	1275 458- V&W Holld 8* K. Ce	ejka	ro Cov rcader g n auge RG# 7	ve ro, Oakl er 012		BORING/WELL NAME DRILLING STARTED DRILLING COMPLETED WELL DEVELOPMENT D GROUND SURFACE ELET TOP OF CASING ELEVAT SCREENED INTERVAL DEPTH TO WATER (First DEPTH TO WATER (Static entrance way, 3 ft NNW of r	ATE (YIELD)_ VATION	12-O Not 5 above i ft bgs) 11 8.	Oct-01 Surveyed msl 1.4 ft (09-0)ct-01) <u> </u>
PID (ppm)	BLOW	SAMPLE ID	EXTENT	DEPTH (ft bgs)	U.S.C.S.	GRAPHIC LOG		LOGIC DESCRIPTION		CONTACT DEPTH (ft bgs)	WE	LL DIAGRAM
63	2 3 4	MW-4- 5.3		- 5	мн		fine to coarse grained gravel; high estimated @ 4.5 ft: Sandy silty for 20% sand; medium placed in the company silt of the co	lill; black; moist; 5% clay, 79 asticity ey; moist; 20% clay, 80% si estimated permeability y; wet; 45% silt, 55% fine gi	grained 5% silt, ilt;	4.5 5.3 6.0		Portland Type 1/iI Cement Bentonite Sea Monterey Sand #3
0.6	1 3 5 2 2 2 3 4 4 6	MW-4- 10.5		-10-	MH SM MH		medium plasticity; low Silty SAND(SM): Grey sand; moderate estima Clayey SILT(MH): Gre	y; wet; 20% clay, 75% silt, um plasticity; low estimated	ained ∑ 2%	10.3 10.5 11.8		✓ 2"-diam., 0.020" Stotted Schedule 40 PVC Bottom of Boring @ 15 ft



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JOB/SI LOCAT PROJE DRILLE	ECT NUM ER	MBER	127 458 Viro	ibarcad 75 Emb 3-1705 onex	nbarcade 5	ove ero, Oal	BORING/WELL NAME SE DRILLING STARTED 29 Sland, CA DRILLING COMPLETED 30 WELL DEVELOPMENT DATE (GROUND SURFACE ELEVATION TOP OF CASING ELEVATION	-Aug-01 -Aug-01 YIELD) ON	NA Not	Surveyed	
BORING	G DIAM!		2"				SCREENED INTERVAL				
LOGGE				Cejka			DEPTH TO WATER (First Enco		d)8.	.0 ft (30-A	ug-01) <u> </u>
REVIEW	NED BY		R. S	<u>3chult</u> z	z, RG#	7012	DEPTH TO WATER (Static)		N	IA	Ţ
REMAR							; located 66 ft WNW of eastern planter, 40 ft SSW of north	iern pla	inter		
PID (ppm)	BLOW	SAMPLE ID	EXTENT	DEPTH (# bos)	(ft bgs) U.S.C.S.	GRAPHIC	LITHOLOGIC DESCRIPTION	The state of the s	CONTACT DEPTH (ft bgs)	WI	ELL DIAGRAM
	 	 	+	,—	+-		ASPHALT		1	W/XX	и
							FILL: Sandy fill; greyish brown; dry; 20% silt, 80% fine grained sand; very high estimated permeability CONCRETE ASPHALT CONCRETE FILL: Silty gravelly fill; light brown; 10% clay, 30% silt,		2.0 2.5 -2.7 -3.0		
3	ł	SB-A- 3.5	5	i			60% fine to coarse grained gravel; high estimated		40		Á
	Í			- 5 -	CL		permeability <u>Silty CLAY</u> (CL): Brown; dry; 60% clay, 25% silt, 15% sand; medium plasticity; very low estimated permeability <u>Sandy CLAY</u> (CH): Grey; damp; 65% clay, 30% sand 5% fine grained gravel; high plasticity; very low estimate permeability		5.0		Portland Type
4		SB-A- 7.5	5	<u></u>	СН		@ 8 ft: Brown; wet; 70% clay, 5% silt, 25% sand	Ā	9.0		
	!			ī.	CL		Silty CLAY (CL): Grey; wet; 65% clay, 35% silt;		9.0		
6		SB-A- 9.5		10	1		medium plasticity; very low estimated permeability; she fragments		10.0		Bottom of Boring @ 10 ft
	[

5.5

☑ 7.0

11.0



Cambria Environmental Technology, Inc. 1144 - 65th St. Oakland, CA 94608

				ne: (51) 10) 420-)				
CLIENT	TNAME		Por	rt of Oak	kland			BORING/WELL NAME	SB-B		
JOB/SI	TE NAME	E	Em	barcade	ero Co	ve			29-Aug-01		
LOCATI	ION		127	′5 Emba	ircade	ro, Oakl	land, CA	-			
PROJE	CT NUM	BER	458	3-1705				l			
DRILLE	.R		Virc	onex							Surveyed
DRILLIN	NG METH	HOD	Hyd	draulic pu	ush			•	_		
BORING	3 DIAME	ETER	2*						NA NA		
LOGGE	DBY		<u>к. с</u>	Cejka				_	Encountered) 7(0 ft (30-Aug-01) ∑
REVIEW	VED BY_		<u>R. S</u>	<u> Schultz, '</u>	RG# 7	012				NA	
REMAR	KS _		Han	<u>id auger</u>	ed to (3.5 ft bg	is; located 102 ft WNW c	of eatern planter, 40 ft SSW o	of northern pl	anter	
PID (ppm)	BLOW COUNTS	SAMPLEID	EXTENT	1	T	GRAPHIC LOG		DLOGIC DESCRIPTION	MANUFACTOR STORY S	CONTACT DEPTH (# bgs)	WELL DIAGRAM
	[<u> </u>			1			ASPHALT			0.5	
					:		CONCRETE ASPHALT CONCRETE FILL: Silty gravelly file	y; dry; 20% silt, 80% fine grainated permeability; cobble	.5%	2.0 2.5 2.7 3.0	
3	. J'	SB-B- 3.	5	L J	,		sand, 60% fine to coa	arse grained gravel; low plast	ticity,	1 1	

medium estimated permeability

Silty SAND (SM): Grey; damp; 40% silt, 60% fine grained sand; high estimated permeability

Silty CLAY (CH): Grey; wet; 60% clay, 40% silt; medium plasticity; low estimated permeability

3

SB-B- 6.5

SB-B-

10.5

5 -

SM

CH

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Portland Type
I/II Cement

Bottom of Boring @ 11 ft



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				,								
CLIENT	NAME		Port	of Oak	land			BORING/WELL NAME	SB-C			
JOB/SI	TE NAME		Emb	arcade	ro Cov	e		DRILLING STARTED _	29-Aug-01			
LOCAT	ION		1275	Emba	rcader	o, Oak	land, CA	DRILLING COMPLETED	29-Aug-01			
PROJE	CT NUME	BER	458-	1705				WELL DEVELOPMENT DATE (YIELD) NA				
DRILLE	R		Viron	ex				GROUND SURFACE ELEV	ATION	Not St	urveyed	
DRILLIN	NG METH	OD	Hand	lauger				TOP OF CASING ELEVATI	ON NA			
BORING	G DIAMET	rer	4"					SCREENED INTERVAL	NA			
LOGGE	DBY _		K. Ce	ejka				DEPTH TO WATER (First E	ncountered)	NA	Ϋ́	
REVIEW	VED BY_		R. Sc	chultz, l	RG# 70)12		DEPTH TO WATER (Static		NA		
REMAR	KS _		Hand	auger	ed to 4	ft bgs	; 143 ft WNW of eastern j	planter, 36 ft SSW of norther	n planter			
PID (ppm)	BLOW	SAMPLE ID	EXTENT	DEPTH (# bgs)	U.S.C.S.	GRAPHIC LOG	LITHC	LOGIC DESCRIPTION		CONTACT DEPTH (# bgs)	WELL DIAGRAM	

ASPHALT 0.5 FILL: Sandy fill; grey; dry; 10% silt, 90% fine grained sand; very high estimated permeability 2.0 CONCRETE 2.5 ASPHALT
CONCRETE
FILL: Gravel fill; Grey; 10% clay, 90% fine to coarse 2.7 3.0 grained gravel; very high estimated permeability 4.0 Bottom of Boring @ 4 ft ĝ

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WELL LOG (PID) HYPORTOF-INEMBARC-NGINTNEMBCOVE.GPJ DEFAULT.GOT 1/10/02

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BORING/WELL LOG

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JOB/SI LOCAT PROJE DRILLE DRILLE BORING LOGGE	ET NUMER NG MET G DIAME D BY VED BY	E	275 58- 'iron lydr " . Ce	Emba 1705 nex aulic pu ajka chultz, l	ro Cov rcader ush RG# 7	ve o, Oaki 012		DRILLING STARTED DRILLING COMPLETED WELL DEVELOPMENT D GROUND SURFACE ELE TOP OF CASING ELEVA SCREENED INTERVAL	VATE (YIELD) VATION NA NA Encountered	NA Not	Surveyed .5 ft (30-A	ug-01) ∑
PID (ppm)	BLOW	SAMPLE ID	EXTENT	DEPTH (# bgs)	U.S.C.S.	GRAPHIC LOG	· · · · · · · · · · · · · · · · · · ·	LOGIC DESCRIPTION		CONTACT DEPTH (ft bgs)	WE	ELL DIAGRAM
3		SB-D- 6.5 SB-D- 10.5		-10-	CL		CONCRETE FILL: Silty gravelly fill 20% silt, 75% fine to c estimated permeability Silty CLAY (CL): Dark silt; low plasticity; very	; brownish black; moist; 5% coarse grained gravel; high y	6 clay, 35% 	2.0 2.5 5.0 7.0		Portland Type I/II Cement Bottom of Boring @ 11 ft





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	E Em 12 BER 458 Viri BOD Hye TER 2" K. 6 R. 9	3-1705 onex draulic p Cejka Schultz,	ero Covarcader . ush	ve o, Oakla	and, CA	DRILLING STARTED DRILLING COMPLETED WELL DEVELOPMENT DAT GROUND SURFACE ELEVA TOP OF CASING ELEVATIO SCREENED INTERVAL	29-Aug-01 30-Aug-01 E (YIELD) TION NA NA NA	NA Not S	Surveyed 0 ft (30-At	
PID (ppm) BLOW COUNTS	SAMPLE ID	OEPTH (ft bgs)	U.S.C.S.	GRAPHIC LOG	LITHO	LOGIC DESCRIPTION		CONTACT SEPTH (ft bgs)	WE	ELL DIAGRAM
356	SB-E- 6	5	SM		CONCRETE: dyed FILL: Sandy gravelly fine to coarse gravel; v Silty SAND (SM): Bro fine gravel; medium to fragments	in; damp; 10% silt, 90% fine gh estimated permeability fill; brown; damp; 20% sand 8 very high estimated permeability; wn; wet; 30% silt, 50% sand, high estimated permeability; estimated permeability	20% shell ∑	0.5 2.0 2.5 5.0 6.5		Portland Type I/II Cement Bottom of Boring @ 8 ft



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Cambria Environmental Technology, Inc. 1144 - 65th St. Oakland, CA 94608 Telephone: (510) 420-0700 Fax: (510) 420-9170

CLIENT NAME	Port of Oakland		BORING/WELL NAME SB-	F	
JOB/SITE NAME	Embarcadero Cove		DRILLING STARTED 29-A	lug-01	
LOCATION	1275 Embarcadero	, Oakland, CA	DRILLING COMPLETED 30-A		
PROJECT NUMBER	Maria		•		
DRILLER DRILLING METHOD	Vironex				urveyed
BORING DIAMETER	2"				
LOGGED BY	K. Cejka		SCREENED INTERVAL		
REVIEWED BY	R. Schultz, RG# 70	12	•		
REMARKS			eastern planter, 12 ft SSW of north	NA	<u> </u>
			oderem planter, 12 it 35W of north		
PID (ppm) BLOW COUNTS	EXTENT DEPTH (# bgs) U.S.C.S.	COG CITHO	LOGIC DESCRIPTION	CONTACT DEPTH (# bgs)	WELL DIAGRAM
		<u>ASPHALT</u>		7 17	X/XX/X
		very high estimated p	; damp; 80% sand, 20%gravel; ermeability	2.0	
	- S	CONCRETE: dyed		2.5	
		EILL: Clayey gravelly sand, 60% gravel; high	fill; brown; wet; 30% clay, 10% h estimated permeability		
9 SB-F- 3	5 L	XXXI	•	3.5	
6 SB-F- 5.	GM 3	60% gravel; very low o	Brown; damp; 10% clay, 30% silt, estimated permeability	5.5	Portland Type I/II Cement
Sb-F- 5.	CH CH	□C	mp; 90% clay, 10% silt; high	6.0	
5 SB-F- 7.8	SM	Silty SAND (SM): Gre plasticity; high estimate	V: wet: 40% silt, 60% sand: bigh	☑ 8.0	
					Bottom of Boring @ 8 ft

APPENDIX B

Permits

PUBLIC WORKS

ALAMEDA COUNTY PUBLIC WORKS AGENCY

WATER RESOURCES SECTION
J99 ELMHURST ST. HAYWARD CA. 94544-1395
PHONE (516) 678-5554
FAX (510)781-1939

	, , , ,	
DRILLING PERMIT APPLICATION		
AND THE PROPERTY OF THE PROPER		
HRILLING PERCHANAGE CONTY		

	•
for applicant to complete	FOR OFFICE USE
•	PERMIT NUMBER WOI-941
LOCATION OF PROJECT	WELL NUMBER
PORT OF CAKLAND 1275 EMBARCADESO	ATH
OKLAND CA	
	Permut Conditions
	Circled Permit Requirement Apply
CLIENT	. crima i
Name DOAY OF OAKLAND	A. GENERAL I. A permit application should be submitted so as to
Address 530 WATER STEAT Phone STO YES BYO	arrive at the ACPWA office five days prior to
City ONKIND, CA Zip 9/104	Moposed starting date.
APPLICANT	2. Sylomit to ACPWA within 50 days after completion of
Name KRIS COSKA Ca. CAMBRIA ENVIRONMENT	permitted original Department of Water Recources.
131 510 ~ 7/0° Y1 7/0	Well Completion Report.
Address 1144 6974 St. Ste B Phone 610-420-3313	3. Permit is void if project not begun within 50 days of
CIO DANIAID, CA ZIP 14608	approval date B. WATER SUPPLY WELLS
	1. Minimum purface seal thickness is two inches of
THE AT BRAIFCT	coment grout placed by nemic.
TYPE OF PROJECT Well Construction Geolechnical Investigation	2. Minimum seal depth le 30 feet for municipal and
Carbodie Protection 5 Control 0	Industrial wells or 20 feet for domestie and irrigation
Water Supply Contamination	wells unloss a lease depth is specially approved.
Monitoring Well Oceaniction 0	C. CROUNDWATER MONITORING WELLS INCLUDING PIEZOMETERS
and a second by the second by	I. Minimum surface seal chickness is two inches of
PROPOSED WATER SUPPLY WELL USE New Domestic 0 Replacement Domestic 3	cement growt placed by treme.
Municipal 0 Intention 9	2. Minimum seal depth for manisoring wells in the
Industrial 0 Other 3	maximum depth provides die of 20 fees.
	D. GEOTECHNICAL
DRILLING METHOD:	Backfill bore hole by remie with content group or coment
Mud Rosary O Air Rosary & Auger B	grouds and mixture. Upper two-three feet replaced in kind or with compacted cumner.
Cable 0 Other 3	E. CATHODIC
DRULLER'S NAME V 4. W	Fill hale mode cone with constate placed by usinis.
	f. Well destruction
DAULER'S LICENSE NO. 720 - 904	Send a map of work site. A separate permit is required
	for wells deeper than 45 fest. G. SPECIAL CONDITIONS
WELL PROJECTS	0. Strows complitions
Drill Hale Diameter B in Management	NOTE: One application must be submitted for each well or well
Casing Diameter 2 in Depth 15 ft.	desprission. Multiple borings on one application are acceptable
Surface Small Depith . A. Owner's Well Number	for geotechnical and contamination investigations,
	•
GEDTECHNICAL PROJECTS Number of Borings Maximum	
Number of Borings Maximum Hole Diameter in Depth A.	
	1 nd 1 1/25~C
STINATED STARTING DATE 16-9- 2001	, may 7 // /
SETTIMATED COMPLETION DATE 10-10-1001	APPROVEDDATE
thereby agree to comply with all requirements of this pennu and Alameda County C	hrihanes No. 73-65./ //
1//	1 11 1
APPLICANT'S SIGNATURE DATE	10-4-2001
PLEASE PRINT NAME KROS COLKA PA	n:5:11-00
PLEASE PRINT NAME KRS COLKA RE	7070L P. 01

APPENDIX C
Standard Field Procedures for Soil Borings and Monitoring Wells

STANDARD FIELD PROCEDURES FOR SOIL BORINGS AND MONITORING WELLS

This document presents standard field methods for drilling and sampling soil borings and installing, developing and sampling ground water monitoring wells. These procedures are designed to comply with Federal, State and local regulatory guidelines. Specific field procedures are summarized below.

SOIL BORINGS

Objectives

Soil samples are collected to characterize subsurface lithology, assess whether the soils exhibit obvious hydrocarbon or other compound vapor or staining, and to collect samples for analysis at a State-certified laboratory. All borings are logged using the Unified Soil Classification System by a trained geologist working under the supervision of a California Registered Geologist (RG).

Soil Boring and Sampling

Soil borings are typically drilled using hollow-stem augers or direct-push technologies such as the Geoprobe®. Soil samples are collected at least every five ft to characterize the subsurface sediments and for possible chemical analysis. Additional soil samples are collected near the water table and at lithologic changes. Samples are collected using lined split-barrel or equivalent samplers driven into undisturbed sediments at the bottom of the borehole.

Drilling and sampling equipment is steam-cleaned prior to drilling and between borings to prevent cross-contamination. Sampling equipment is washed between samples with trisodium phosphate or an equivalent EPA-approved detergent.

Sample Analysis

Sampling tubes chosen for analysis are trimmed of excess soil and capped with Teflon tape and plastic end caps. Soil samples are labeled and stored at or below 4° C on either crushed or dry ice, depending upon local regulations. Samples are transported under chain-of-custody to a State-certified analytic laboratory.

Field Screening

One of the remaining tubes is partially emptied leaving about one-third of the soil in the tube. The tube is capped with plastic end caps and set aside to allow hydrocarbons to volatilize from the soil. After ten to fifteen minutes, a portable volatile vapor analyzer measures volatile hydrocarbon vapor concentrations in the tube headspace, extracting the vapor through a slit in the cap. Volatile vapor analyzer measurements are used along with the field observations, odors, stratigraphy and ground water depth to select soil samples for analysis.

Water Sampling

Water samples, if they are collected from the boring, are either collected using a driven Hydropunch® type sampler or are collected from the open borehole using bailers. The ground water samples are decanted into the appropriate containers supplied by the analytic laboratory. Samples are labeled, placed in protective foam sleeves, stored on crushed ice at or below 4°C, and transported under chain-of-custody to the laboratory. Laboratory-supplied trip blanks accompany the samples and are analyzed to check for cross-contamination. An equipment blank may be analyzed if non-dedicated sampling equipment is used.

Grouting

If the borings are not completed as wells, the borings are filled to the ground surface with cement grout poured or pumped through a tremie pipe.

MONITORING WELL INSTALLATION, DEVELOPMENT AND SAMPLING

Well Construction and Surveying

Ground water monitoring wells are installed to monitor ground water quality and determine the ground water elevation, flow direction and gradient. Well depths and screen lengths are based on ground water depth, occurrence of hydrocarbons or other compounds in the borehole, stratigraphy and State and local regulatory guidelines. Well screens typically extend 10 to 15 ft below and 5 ft above the static water level at the time of drilling. However, the well screen will generally not extend into or through a clay layer that is at least three ft thick.

Well casing and screen are flush-threaded, Schedule 40 PVC. Screen slot size varies according to the sediments screened, but slots are generally 0.010 or 0.020 inches wide. A rinsed and graded sand occupies the annular space between the boring and the well screen to about one to two ft above the well screen. A two ft thick hydrated bentonite seal separates the sand from the overlying sanitary surface seal composed of Portland type I,II cement.

Well-heads are secured by locking well-caps inside traffic-rated vaults finished flush with the ground surface. A stovepipe may be installed between the well-head and the vault cap for additional security.

The well top-of-casing elevation is surveyed with respect to mean sea level and the well is surveyed for horizontal location with respect to an onsite or nearby offsite landmark.

Well Development

Wells are generally developed using a combination of ground water surging and extraction. Surging agitates the ground water and dislodges fine sediments from the sand pack. After about ten minutes of surging, ground water is extracted from the well using bailing, pumping and/or reverse air-lifting through an eductor pipe to remove the sediments from the well. Surging and extraction continue until at least ten well-casing volumes of ground water are extracted and the sediment volume in the ground water is negligible. This process usually occurs prior to installing the sanitary surface seal to ensure sand pack stabilization. If development occurs after surface seal installation, then development occurs 24 to 72 hours after seal installation to ensure that the Portland cement has set up correctly.

All equipment is steam-cleaned prior to use and air used for air-lifting is filtered to prevent oil entrained in the compressed air from entering the well. Wells that are developed using air-lift evacuation are not sampled until at least 24 hours after they are developed.

Ground Water Sampling

Depending on local regulatory guidelines, three to four well-casing volumes of ground water are purged prior to sampling. Purging continues until ground water pH, conductivity, and temperature have stabilized. Ground water samples are collected using bailers or pumps and are decanted into the appropriate containers supplied by the analytic laboratory. Samples are labeled, placed in protective foam sleeves, stored on crushed ice at or below 4°C, and transported under chain-of-custody to the laboratory. Laboratory-supplied trip blanks accompany the samples and are analyzed to check for cross-contamination. An equipment blank may be analyzed if non-dedicated sampling equipment is used.

F:\TEMPLATE\SOPs\Wells-borings-gw.wpd

APPENDIX D

Laboratory Analytical Reports

110 2nd Avenue South, #D7, Pacheco, CA 94553-5560
Telephone: 925-798-1620 Fax: 925-798-1622
http://www.mccampbell.com E-mail: main@mccampbell.com

Cambria Environmental Technology	Client Project ID: #458-1705; Embarcadero Cove	Date Sampled: 12/05 & 12/07/01
1144 65 th Street, Suite C		Date Received: 12/07/01
Oakland, CA 94608	Client Contact: Bob Schultz	Date Extracted: 12/07/01
	Client P.O:	Date Analyzed: 12/07/01

12/14/01

Dear Bob:

Enclosed are:

- 1). the results of 11 samples from your #458-1705; Embarcadero Cove project,
- 2). a QC report for the above samples
- 3). a copy of the chain of custody, and
- 4). a bill for analytical services.

All analyses were completed satisfactorily and all QC samples were found to be within our control limits. If you have any questions please contact me. McCampbell Analytical Laboratories strives for excellence in quality, service and cost. Thank you for your business and I look forward to working with you again.

Edward Hamilton, Lab Director

Client Project ID: #458-1705;	Date Sampled: 12/05 & 12/07/01			
Embarcadero Cove	Date Received: 12/07/01			
Client Contact: Bob Schultz	Date Extracted: 12/10-12/12/01			
Client P.O:	Date Analyzed: 12/10-12/12/01			
	Embarcadero Cove Client Contact: Bob Schultz			

Gasoline Range (C6-C12) Volatile Hydrocarbons as Gasoline*, with Methyl tert-Butyl Ether* & BTEX*

EPA methods 5030, modified 8015, and 8020 or 602; California RWQCB (SF Bay Region) method GCFID(5030) Ethyl-% Recovery Lab ID Client ID Matrix TPH(g)+ MTBE Benzene Toluene **Xylenes** benzene Surrogate 85050 MW-1-AM W 13,000,a ND<20 1300 180 1200 860 109 85051 MW-2-AM W ND ND ND ND ND ND 111 85052 MW-3-AM W 310,a ND 0.72 2.2 ND ND 107 85053 W MW-4-AM 13,000,a,h ND<10 120 28 170 380 85054 MW-1-PM W 3100,a ND 270 12 150 74 113 85055 MW-2-PM W ND ND ND ND ND ND 107 85056 MW-3-PM W 320,a ND 0.84 2.6 ND 0.76 109 85057 W MW-4-PM 20,000,a,h ND<20 420 78 390 **_..**# 870 85058 TB W ND ND ND ND ND ND 98 Reporting Limit unless W 50 ug/L 5.0 0.5 0.5 0.5 0.5 otherwise stated; ND means not detected above S 1.0 mg/kg 0.05 0.005 0.005 0.005 0.005 the reporting limit

^{*}The following descriptions of the TPH chromatogram are cursory in nature and McCampbell Analytical is not responsible for their interpretation: a) unmodified or weakly modified gasoline is significant; b) heavier gasoline range compounds are significant(aged gasoline?); c) lighter gasoline range compounds (the most mobile fraction) are significant; d) gasoline range compounds having broad chromatographic peaks are significant; biologically altered gasoline?; e) TPH pattern that does not appear to be derived from gasoline (?); f) one to a few isolated peaks present; g) strongly aged gasoline or diesel range compounds are significant; h) lighter than water immiscible sheen is present; i) liquid sample that contains greater than ~5 vol. % sediment; j) no recognizable pattern.



^{*} water and vapor samples are reported in ug/L, wipe samples in ug/wipe, soil and sludge samples in mg/kg, and all TCLP and SPLP extracts in ug/L

[&]quot;cluttered chromatogram; sample peak coefutes with surrogate peak

Cambria Environmental Technology	Client Project ID: #458-1705;	Date Sampled: 12/05 & 12/07/01
1144 65 th Street, Suite C	Embarcadero Cove	Date Received: 12/07/01
Oakland, CA 94608	Client Contact: Bob Schultz	Date Extracted: 12/07/01
	Client P.O:	Date Analyzed: 12/07/01

Diesel Range (C10-C23) and Oil-Range (C18+) Extractable Hydrocarbons as Diesel and Motor Oil with Silica Gel Clean-Up*

EPA methods modified 8015, and 3550 or 3510; California RWQCB (SF Bay Region) method GCFID(3550) or GCFID(3510)

Lab ID	Client ID	Matrix	TPH(d) ⁺	TPH(mo)⁺	% Recovery Surrogate
85050	MW-1-AM	w	3800,d	ND	106
85051	MW-2-AM	w	150,g	560	106
85052	MW-3-AM	w	480,d,b,g	480	108
85053	MW-4-AM	w	6400,d,b,g,h	430	118
85054	MW-1-PM	w	680,d	ND	111
85055	MW-2-PM	w	75,b	270	105
85056	MW-3-PM	w	530,d,b,g	550	109
85057	MW-4-PM	w	5400,d,b,g,h	450	112
	nit unless otherwise	w	50 ug/L	250 ug/L	
	orting limit	S	1.0 mg/kg	5.0 mg/kg	

^{*}water samples are reported in ug/L, wipe samples in ug/wipe, soil and sludge samples in mg/kg, and all TCLP / STLC / SPLP extracts in ug/L

DHS Certification No. 1644

Edward Hamilton, Lab Director

^{*}cluttered chromatogram resulting in coeluted surrogate and sample peaks, or; surrogate peak is on elevated baseline, or; surrogate has been diminished by dilution of original extract.

The following descriptions of the TPH chromatogram are cursory in nature and McCampbell Analytical is not responsible for their interpretation: a) unmodified or weakly modified diesel is significant; b) diesel range compounds are significant; no recognizable pattern; c) aged diesel? is significant); d) gasoline range compounds are significant; e) medium boiling point pattern that does not match diesel (?); f) one to a few isolated peaks present; g) oil range compounds are significant; h) lighter than water immiscible sheen is present; i) liquid sample that contains greater than ~5 vol. % sediment.

Cambria Environmental	Client Project ID: #458-1705;	Date Sampled: 12/05 & 12/07/01		
Technology	Embarcadero Cove	Date Received: 12/07/01		
1144 65 th Street, Suite C	Client Contact: Bob Schultz	Date Extracted: 12/10/01		
Oakland, CA 94608	Client P.O;	Date Analyzed: 12/10-12/11/01		
Polynuclear A	romatic Hydrocarbons (PAH / PNA) by	v Liquid Chromatography		

EPA methods 8310 and 3510 Lab ID 85050 85051 85052 Reporting Limit Client ID MW-1-AM MW-2-AM MW-3-AM W, STLC S TCLP W Matrix W W Concentration* Compound ug/kg ug/L ND<10 Acenaphthene ND ND 5.0 0.5 Acenaphthylene 72 ND<0.25 ND<0.25 5.0 0.2 Anthracene ND<10 ND ND 5.0 2.0 Benzo(a)anthracene ND<10 ND<0.25 ND<0.25 5.0 0.1 Benzo(b)fluoranthene ND<10 ND ND 5.0 10 ND<10 ND Benzo(k)fluoranthene ND 5.0 2 ND<10 Benzo(g,h,i)perylene ND<0.25 ND<0.25 5.0 0.1 Benzo(a)pyrene ND<10 ND<0.25 ND<0.25 5.0 0.1 Chrysene ND<10 ND ND 5.0 5.0 ND<10 Dibenzo(a,h)anthracene ND<0.25 ND<0.25 5.0 0.1 Fluoranthene ND<10 ND<0.25 ND<0.25 5.0 0.05 Fluorene ND<10 ND<0.25 ND<0.25 5.0 0.1 Indeno(1,2,3-cd)pyrene ND<10 ND<0.25 ND<0.25 5.0 0.05 1-Methynaphthalene 150 ND ND 5.0 1.0

ND

ND<0.25

ND<0.25

ND<0.25

110

220

360

ND<10

ND<10

___#

DHS Certification No. 1644

2-Methynaphthalene

% Recovery Surrogate 1

Naphthalene

Phenanthrene

Pyrene

Edward Hamilton, Lab Director

ND

ND<0.25

ND<0.25

ND<0.25

107

5.0

5.0

5.0

5.0

1.0

0.2

0.05

0.05

___# % Recovery Surrogate 2 102 87 Comments j * water and vapor samples are reported in ug/L, soil and sludge samples in mg/kg, wipes in ug/wipe and all TCLP / STLC / SPLP extracts in ug/L. ND means not detected above the reporting limit; N/A means analyte not applicable to this analysis

[&]quot; surrogate diluted out of range or surrogate coelutes with another peak

⁽h) a lighter than water immiscible sheen is present; (i) liquid sample that contains > 1/5 vol. % sediment; (j) sample diluted due to high organic content.

Cambria Environmental	Client Project ID: #458-1705;	Date Sampled: 12/05 & 12/07/01
Technology	Embarcadero Cove	Date Received: 12/07/01
1144 65 th Street, Suite C	Client Contact: Bob Schultz	Date Extracted: 12/10/01
Oakland, CA 94608	Client P.O:	Date Analyzed: 12/10-12/11/01
Dalamatan	vomatic Undersonbane (DAII / DNA) b	T: 210)

Polynuclear Aromatic Hydrocarbons (PAH / PNA) by Liquid Chromatography						
Polynuclear Ard EPA methods 8310 and 3510	omatic Hydrocarbor	ns (PAH / PNA) by Liqu	uid Chromatograp	hy		
Lab ID	85053	85054	85055	Reporting Limit		
Client ID	MW-4-AM	MW-1-PM	MW-2-PM	s	W, STLC	
Matrix	W	W	W	3	TCLP	
Compound		Concentration*		ug/kg	ug/L	
Acenaphthene	ND<10	ND<1.0	ND	5.0	0.5	
Acenaphthylene	24	9.6	ND<0.25	50	0.2	
Anthracene	ND<10	ND	ND	5.0	2.0	
Benzo(a)anthracene	ND<10	ND<1.0	ND<0.25	5.0	0.1	
Benzo(b)fluoranthene	ND<10	ND	ND	5.0	10	
Benzo(k)fluoranthene	ND<10	ND	ND	5.0	.2	
Benzo(g,h,ı)perylene	ND<10	ND<1.0	ND<0.25	5.0	1.0	
Benzo(a)pyrene	ND<10	ND<1.0	ND<0.25	5.0	0.1	
Chrysene	ND<10	ND	ND	5.0	5.0	
Dibenzo(a,h)anthracene	ND<10	ND<1.0	ND<0.25	50	0.1	
Fluoranthene	ND<10	ND<1.0	ND<0.25	` 5.0	0.05	
Fluorene	ND<10	ND<1.0	ND<0.25	5.0	0.1	
Indeno(1,2,3-cd)pyrene	ND<10	ND<1.0	ND<0.25	5.0	0.05	
1-Methynaphthalene	99	18	ND	5.0	1.0	
2-Methynaphthalene	190	14	ND	5.0	1.0	
Naphthalene	60	22	ND<0.25	5.0	0.2	
Phenanthrene	18	1.3	ND<0.25	5.0	0.05	
Ругепе	ND<10	ND<1.0	ND<0.25	5.0	0.05	
% Recovery Surrogate 1	#	111	105			
% Recovery Surrogate 2	#	107	115			
Comments	h		j			

^{*} water and vapor samples are reported in ug/L, soil and sludge samples in mg/kg, wipes in ug/wipe and all TCLP / STLC / SPLP extracts in ug/L.

DHS Certification No. 1644

Edward Hamilton, Lab Director

ND means not detected above the reporting limit; N/A means analyte not applicable to this analysis

^{*} surrogate diluted out of range or surrogate coelutes with another peak

⁽h) a lighter than water immiscible sheen is present; (i) liquid sample that contains > 5 vol. % sediment; (j) sample diluted due to high organic content.

Cambria Environmental	Client Project ID: #458-1705;	Date Sampled: 12/05 & 12/07/01
Technology	Embarcadero Cove	Date Received: 12/07/01
1144 65 th Street, Suite C	Client Contact: Bob Schultz	Date Extracted: 12/10/01
Oakland, CA 94608	Client P.O:	Date Analyzed: 12/10-12/11/01
Polynuclear A	romatic Hydrocarbons (PAH / PNA) by	v Liquid Chromatography

Lab ID	85056	85057	Repor	ting Limit	
Client ID	MW-3-PM	NW-4-PM		W, STL	
Matrix	w	W	S	TCLP	
Compound	Concentration*			ug/L	
Acenaphthene	ND	ND<10	5.0	0.5	
Acenaphthylene	ND<0.25	21	5.0	0.2	
Anthracene	ND	ND<10	5.0	2.0	
Benzo(a)anthracene	ND<0.25	ND<10	5.0	0.1	
Benzo(b)fluoranthene	ND	ND<10	5.0	10	
Benzo(k)fluoranthene	ND	ND<10	5.0	2	
Benzo(g,h,i)perylene	ND<0.25	ND<10	5.0	0.1	
Benzo(a)pyrene	ND<0.25	ND<10	5.0	0.1	
Chrysene	ND	ND<10	5.0	5.0	
Dibenzo(a,h)anthracene	ND<0.25	ND<10	5.0	0.1	
Fluoranthene	ND<0.25	ND<10	50	0.05	
Fluorene	ND<0.25	ND<10	5.0	0.1	
Indeno(1,2,3-cd)pyrene	ND<0.25	ND<10	5.0	0.05	
1-Methynaphthalene	ND	100	5.0	1.0	
2-Methynaphthalene	ND	180	5.0	1.0	
Naphthalene	ND<0.25	96	5.0	0.2	
Phenanthrene	ND<0.25	12	5.0	0.05	
Pyrene	0.31	ND<10	5.0	0.05	
% Recovery Surrogate 1	104	#			
% Recovery Surrogate 2	105	#			
Comments	j	h			

^{*} water and vapor samples are reported in ug/L, soil and sludge samples in mg/kg, wipes in ug/wipe and all TCLP / STLC / SPLP extracts in ug/L.

DHS Certification No. 1644

Edward Hamilton, Lab Director

ND means not detected above the reporting limit; N/A means analyte not applicable to this analysis

[&]quot;surrogate diluted out of range or surrogate coelutes with another peak

⁽h) a lighter than water immiscible sheen is present; (i) liquid sample that contains >-5 vol. % sediment; (j) sample diluted due to high organic content.

Cambria Environmental Technology	Client Project ID: #458-1705;	Date Sampled: 12/05 & 12/07/01		
1144 65 th Street, Suite C	Embarcadero Cove	Date Received: 12/07/01		
Oakland, CA 94608	Client Contact: Bob Schultz	Date Extracted: 12/07/01		
	Client P.O:	Date Analyzed: 12/07-12/12/01		
	CAM / CCR 17 Metals*			

EPA methods 6010/200.7; 7470/7471/245.1/245.5 (Hg); 7060/206.2 (As); 7740/270.2 (Se); 7841/279.2 (TI); 239.2 (Pb, w); 204.2 (Sb,w)

Lab ID	85059	85060			
Client ID	MW-1-AM	MW-1-PM	I	Reporting Lim	it
Matrix	w	W	S	W	STLC,
Extraction ^o	TTLC	TTLC	TTLC	TTLC	TCLP
Compound	Conce	ntration*	mg/kg	mg/L	mg/L
Antimony (Sb)	ND	ND	2.5	0.006	0.05
Arsenic (As)	0.0057	ND	2.5	0.005	0.25
Barium (Ba)	0 057	0.051	2.5	0.05	0.05
Beryllium (Be)	ND	ND	0.5	0.004	0.01
Cadmium (Cd)	ND	ND	0.5	0.005	0.01
Chromium (Cr)	ND	ND	0.5	0.02	0.05
Cobalt (Co)	ND	ND	2.0	0.05	0 05
Copper (Cu)	ND	ND	20	0.05	0.05
Lead (Pb)	0.0053	ND	3.0	0.005	0.2
Mercury (Hg)	ND	ND	0.06	0 0008	0.005
Molybdenum (Mo)	ND	ND	2.5	0.05	0 05
Nickel (Ni)	ND	ND	2.0	0.05	0.05
Selenium (Se)	ND	ND	2.5	0.005	0.25
Silver (Ag)	ND	ND	1.0	0.01	0.05
Thallium (Tl)	ND	ND	2.5	0.005	0.5
Vanadium (V)	ND	ND	2.0	0.05	0.05
Zinc (Zn)	ND	0.050	1.0	0.05	0 05
% Recovery Surrogate	80	88		R	L
Comments			7		

^{*} water samples are reported in mg/L, soil and sludge samples in mg/kg, wipes in ug/wipe and all TCLP / STLC / SPLP extracts in mg/L

ND means not detected above the reporting limit; N/A means surrogate not applicable to this analysis

⁹ EPA extraction methods 1311(TCLP), 3010/3020(water, TTLC), 3040(organic matrices, TTLC), 3050(solids, TTLC); STLC - CA Title 22

[®] DISTLC extractions are performed using STLC methodology except that deionized water is substituted for citric acid buffer as the extraction fluid. DISTLC results are not applicable to STLC regulatory limits.

[&]quot;surrogate diluted out of range

^{*} reporting limit raised due to matrix interference

i) liquid sample that contains greater than ~2 vol % sediment; this sediment is extracted with the liquid, in accordance with EPA methodologies and can significantly effect reported metal concentrations.

QC REPORT

EPA 8015m + 8020

Date: 12/10/01	Extraction	n: TTTL	c _		Matrix:	Water	
	Concentration: ug/L			%Recovery			
Compound	Sample	MS	MSD	Amount Spiked	MS	MSD	RPD
SampleID: 120601					Instrumer	nt: Go	C-3
Surrogate1	ND	103.0	102.0	100.00	103	102	1.0
Xylenes	ND	33.7	34.0	30.00	112	113	0.9
Ethylbenzene	ND	11.2	11.2	10.00	112	112	0.0
Toluene	ND	10.9	10.9	10.00	109	109	0.0
Benzene	ND	10.5	10.4	10.00	105	104	1.0
MTBE	ND	11.0	10.4	10.00	110	104	5.6
TPH (gas)	ND	83.5	82.6	100.00	83	83	1.0

% Re covery =
$$\frac{(MS-Sample)}{AmountSpiked} \cdot 100$$

$$RPD = \frac{(MS \sim MSD)}{(MS + MSD)} \cdot 2.100$$

QC REPORT

EPA 8015m + 8020

Date: 12/07/01	Extractio	n: TTLC	: 		Matrix:	Water	
	Concentration: ug/L			%Recovery			
Compound	Sample	MS	MSD	Amount Spiked	MS	MSD	RPD
SampleID: 120501					Instrume	nt: GC-	6 A
Surrogate1	ND	109.0	104.0	100.00	109	104	4.7
TPH (diesel)	ND	8875.0	8450.0	7500.00	118	113	4.9

% Re covery =
$$\frac{(MS-Sample)}{AmountSpiked} \cdot 100$$

RPD= $\frac{(MS-MSD)}{(MS+MSD)} 2\cdot 100$

QC REPORT

SVOCs (EPA 8270/625/525)

Date: 12/10/01	Extraction	n: N/A			Matrix:	Water			
		Concen	tration:	ug/L	%Rec	overy			
Compound	Sample	MS	MS	MSD	RPD				
SampleID: 121101					Instrument: GC-8				
Surrogate1	ND	690.0	700.0	1000.00	69	70	1.4		
Pyrene	ND	610.0	620.0	1000.00	61	62	1.6		
Pentachlorophenol	ND	870.0	870.0	2000.00	44	44	0.0		
2,4-Dinitrotoluene	ND	510.0	540.0	1000.00	51	54	5.7		
4-Nitrophenol	ND	740.0	680.0	2000.00	37	34	8.5		
Acenaphtene	ND	520.0	540.0	1000.00	52	54	3.8		
4-Chloro-3-metylphenol	ND	710.0	740.0	2000.00	36	37	41		
1,2,4-trichlorobenzene	ND	520.0	550.0	1000.00	52	55	5.6		
N-nitroso-di-n-propyl	ND	490.0	470.0	1000.00	49	47	4.2		
1,4-Dichlorobenzene	ND	520.0	540.0	1000.00	52	54	3.8		
2-Chlorophenol	ND	870,0	890.0	2000.00	44	45	2.3		
Phenol	ND	830.0	800.0	2000.00	42	40	3.7		

% Re covery =
$$\frac{(MS - Sample)}{AmountSpiked} \cdot 100$$

RPD= $\frac{(MS - MSD)}{(MS + MSD)} \cdot 2000$

QC REPORT

CAM 17

Date: 12/07/01 Extraction: TTLC Matrix: Water

Date. 12/07/01						Marci			
		Concen	tration: 1	mg/L	%Rec	overy			
Compound	Sample	MS	MSD	Amount Spiked	MS	MSD	RPD		
SampleID: 120601				Instrument: P-1 AA					
Beryllium	ND	5.1	5.6	5.00	102	111	8.5		
Selenium	ND	0.0100	0.0100	0.0100	100	100	0.0		
Molybdenum	ND	4.5	4.6	5.00	90	93	2.9		
Silver	ND	0.4	0.4	0.50	87	86	1.9		
Thallium	ND	0.0110	0.0100	0.0100	110	100	9.5		
Barium	ND	4.6	4.6	5.00	93	93	0.0		
Nickel	ND	4.7	4.6	5.00	94	92	2.2		
Arsenic	ND	0.0099	0.0110	0.0100	99	110	10.5		
Vanadium	ND	4.7	4.5	5.00	93	91	2.7		
Surrogate1	ND	80.0	83.8	100.00	80	84	4.6		
Zinc	ND	5.1	5.2	5.00	102	104	1.8		
Copper	ND	4.8	4.7	5.00	97	94	3.5		
Antimony	ND	0.0100	0.0100	0.0100	100	100	0.0		
Lead	ND	0.0098	0.0100	0.0100	98	100	2.0		
Cadmium	ND	5.7	5.5	5.00	114	109	4.1		
Cobalt	ND	4.5	4.7	5.00	89	94	5.6		
Mercury	ND	0.00095	0.00093	0.00100	95	93	3.0		
Chromium	ND	4.9	4.9	5.00	97	98	1.0		

% Re covery =
$$\frac{(MS-Sample)}{AmountSpiked} \cdot 100$$

$$RPD = \frac{(MS - MSD)}{(MS + MSD)} \cdot 2 \cdot 100$$

File : D:\HPCHEM\1\DATA\12120107.D

Operator

Acquired: 12 Dec 2001 6:40 pm using AcqMethod GC3D.M Instrument: GC-3

Sample Name: GC-3
Simple Name: 5000ngGAS
Misc Info

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File : D:\HPCHEM\1\DATA\12120103.D

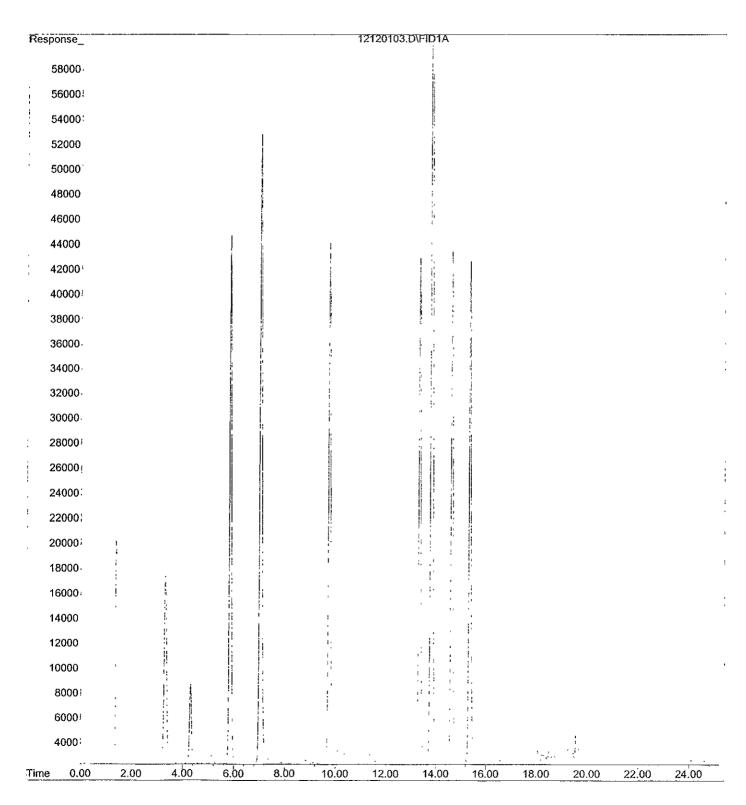
Operator

Acquired 12 Dec 2001 4:28 pm using AcgMethod GC3D.M

Instrument : GC = 3

Sample Name: (100ngBTEX)

Misc Info



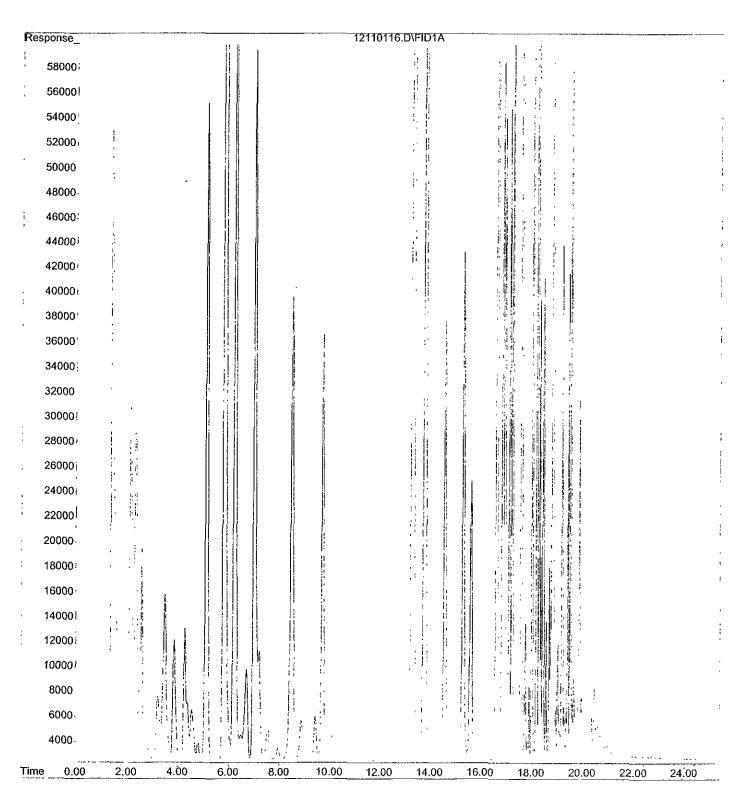
File : D:\HPCHEM\1\DATA\12110116.D

Operator

Acquired: 11 Dec 2001 8:29 pm using AcqMethod GC3D.M

Instrument : GC-3
Sample Name 85050rrW

Misc Info



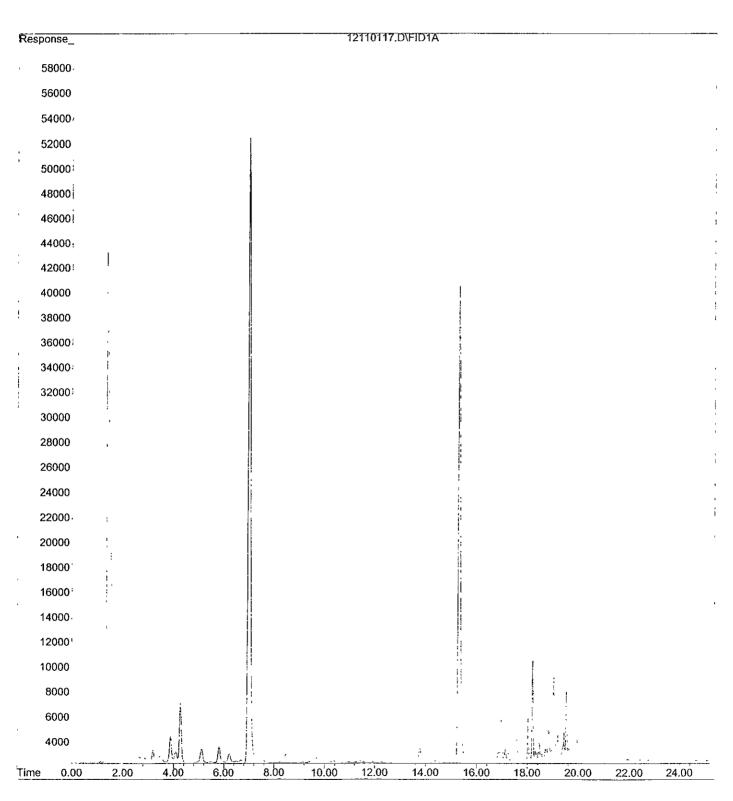
File : D:\HPCHEM\1\DATA\12110117.D

Operator

Acquired: 11 Dec 2001 9:01 pm using AcqMethod GC3D.M

Instrument : GC-3 Sample Name: 85051rrW

Misc Info



File : D:\HPCHEM\1\DATA\12120126.D

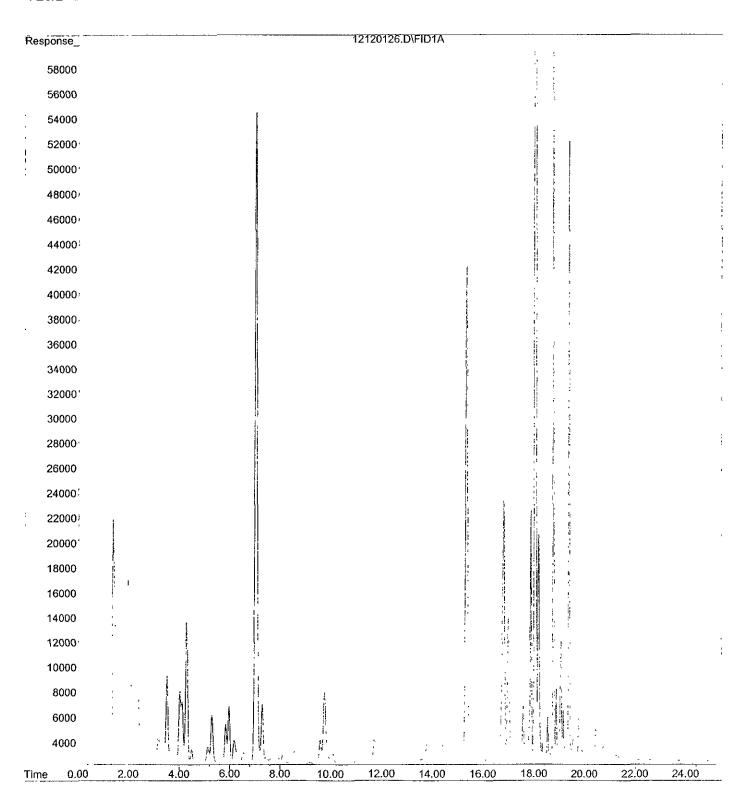
Operator

Acquired: 13 Dec 2001 4:53 am using AcqMethod GC3D.M

Instrument : GC-3

Sample Name: (85052rrrW) AF

Misc Info : — Vial Number: 26



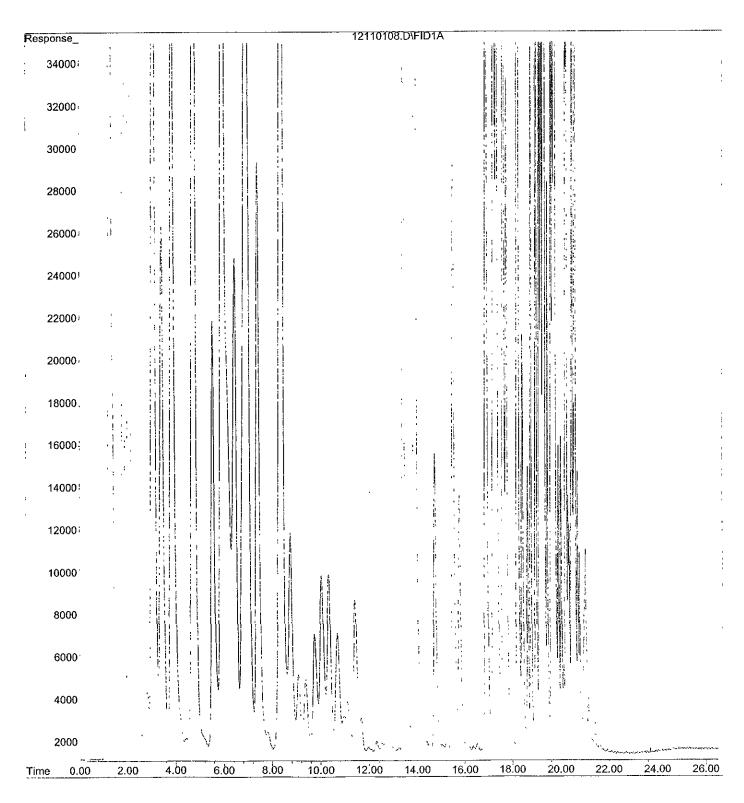
File : D:\HPCHEM\2\DATA\12110108.D

Operator

Acquired: 11 Dec 2001 3:25 pm using AcqMethod GC12A.M

Instrument : GC-12 Sample Name: 85053rrW

Misc Info : Vial Number: 8



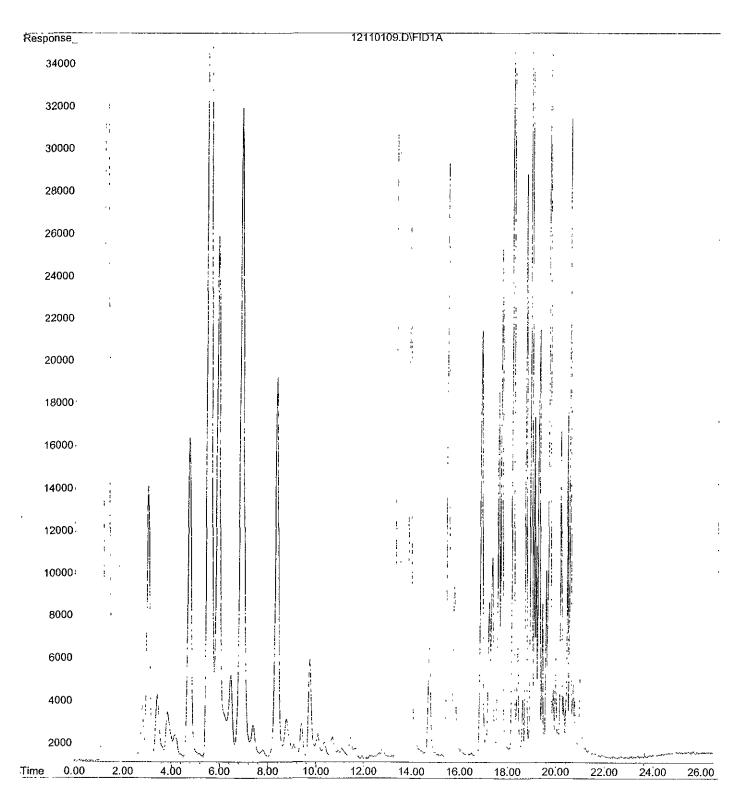
File : D:\HPCHEM\2\DATA\12110109.D

Operator

Acquired: 11 Dec 2001 3:59 pm using AcqMethod GC12A.M

Instrument: GC-12 Sample Name: 85054rrW

Misc Info : Vial Number: 9



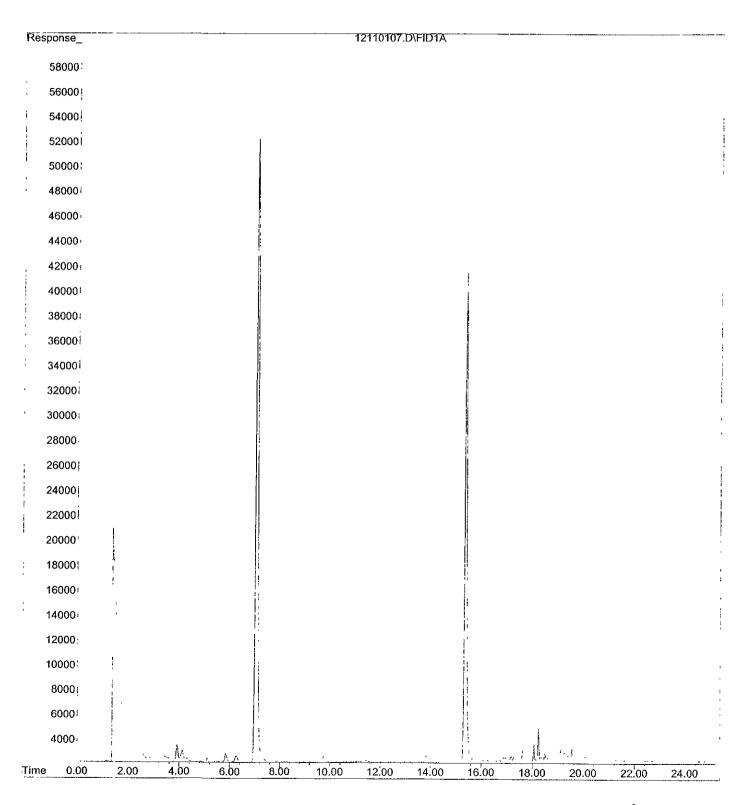
File : D:\HPCHEM\1\DATA\12110107.D

Operator

Acquired : 11 Dec 2001 3:35 pm using AcqMethod GC3D.M

Instrument: GC-3
Sample Name: 85055rrW

Misc Info



File : D:\HPCHEM\1\DATA\12120127.D

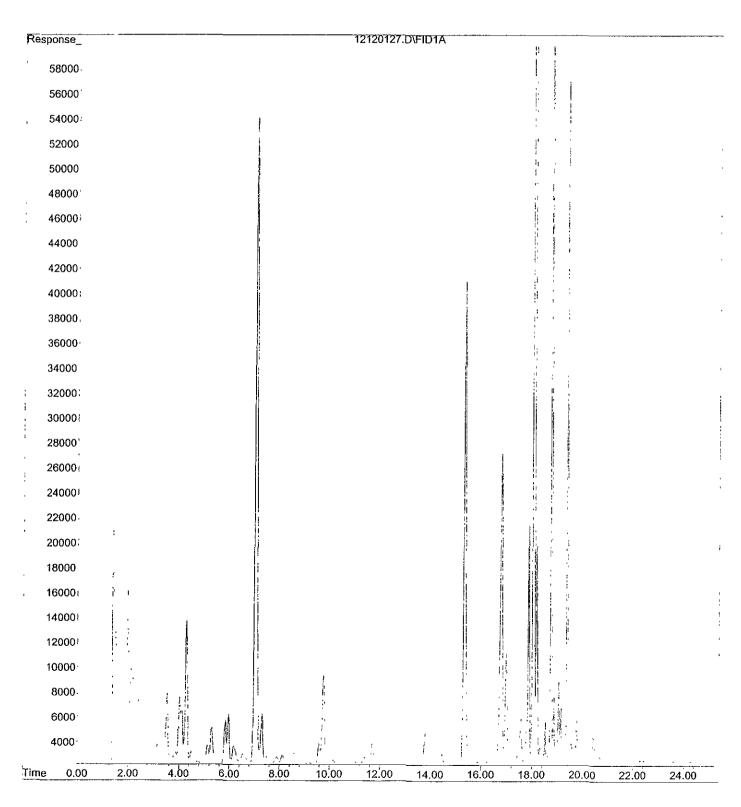
Operator

Acquired : 13 Dec 2001 5:25 am using AcqMethod GC3D.M

Instrument :

GC-3Sample Name: (85056rrrW+AF

Misc Info



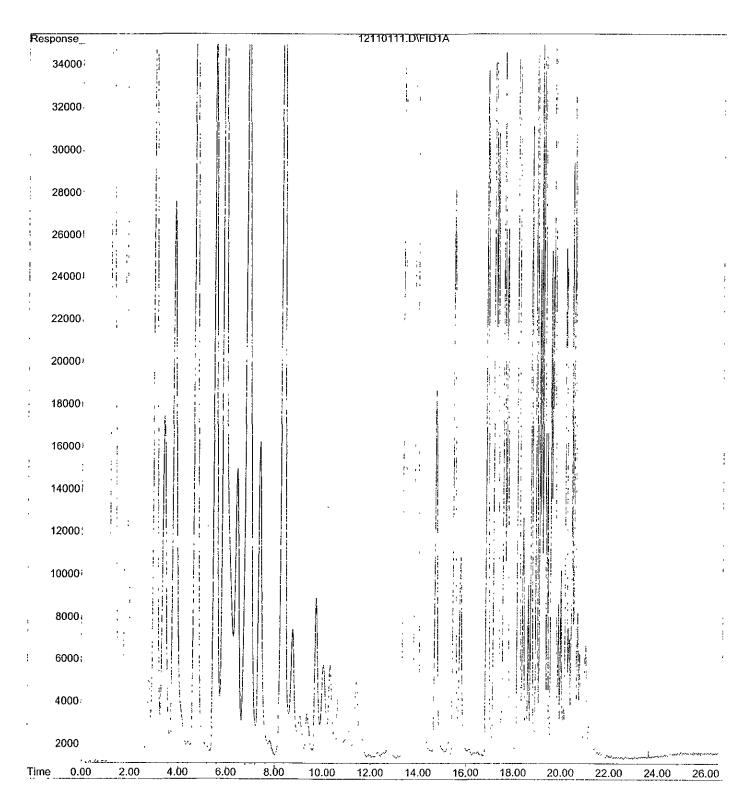
File : D:\HPCHEM\2\DATA\12110111.D

Operator

Acquired: 11 Dec 2001 5:06 pm using AcqMethod GC12A.M

Instrument: GC-12 Sample Name: 85057rrW

Misc Info : Vial Number: 11



File : D:\HPCHEM\2\DATA\12100110.D

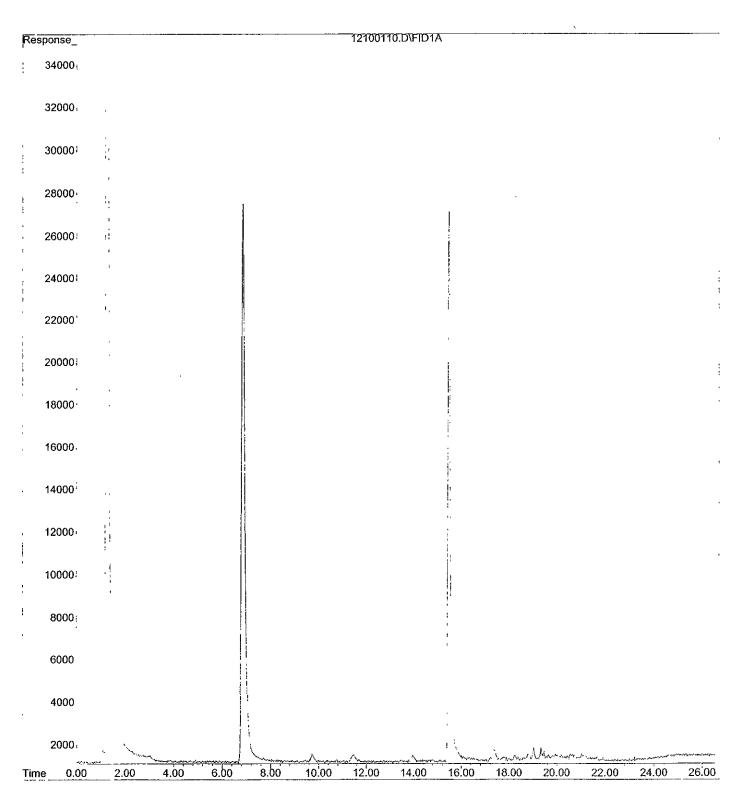
Operator

Acquired : 10 Dec 2001

8:24 pm using AcqMethod GC12A.M

Instrument: GC-12 Sample Name 85058W

Misc Info



File : E:\HPCHEM\3\DATAB\12070131.D

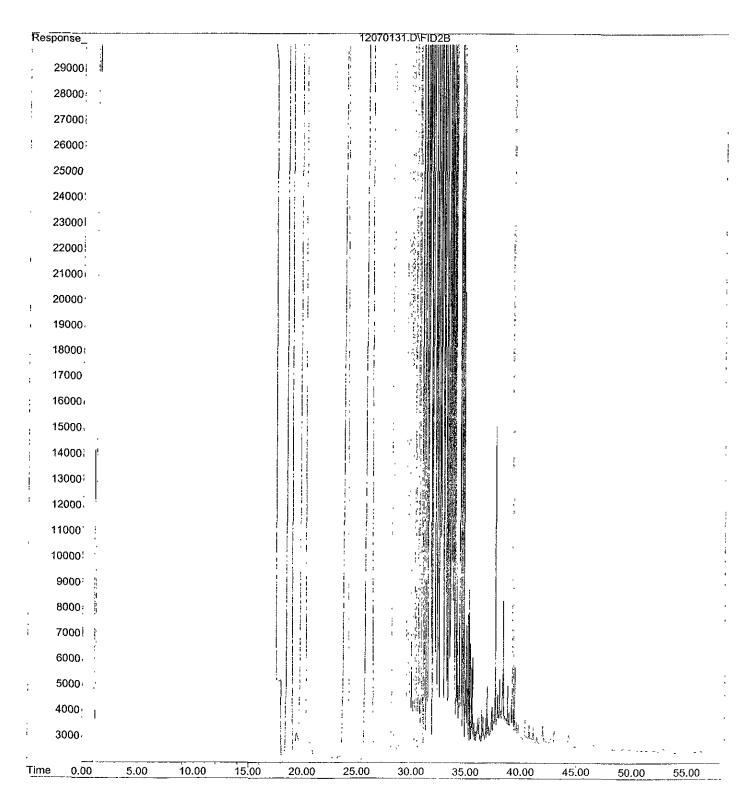
Operator : Thu

Acquired : 8 Dec 2001 11:02 am using AcqMethod GC2ANEW.M

Instrument: GC-2

Sample Name: 8 85050 W

Misc Info : Vial Number: 66



File : E:\HPCHEM\3\DATAB\12070133.D

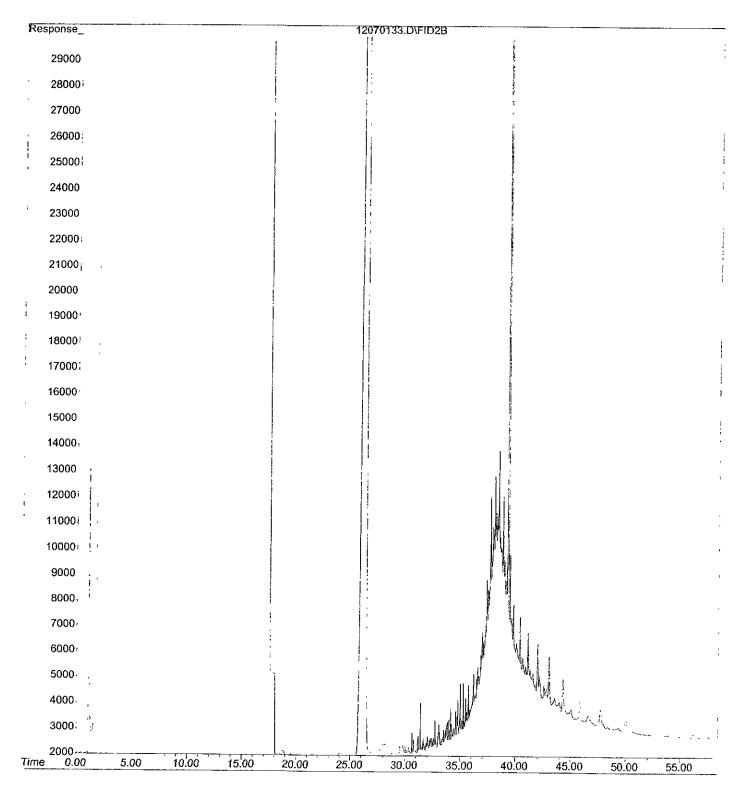
Operator : Thu

Acquired 8 Dec 2001 12:33 pm using AcqMethod GC2ANEW.M :

Instrument : GC-2

Sample Name: 8 85051 W

Misc Info Vial Number: 67



File : E:\HPCHEM\3\DATAB\12070135.D

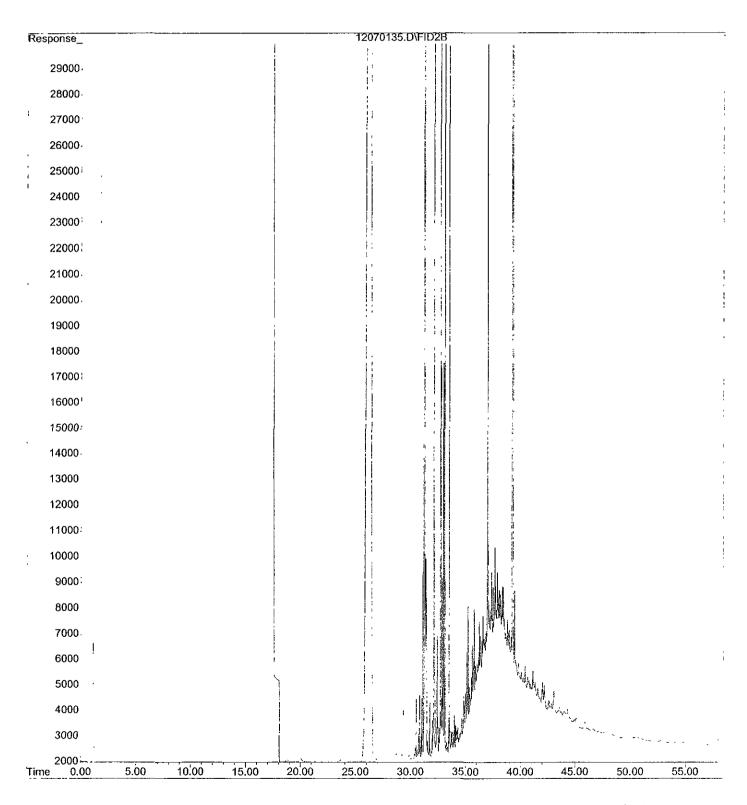
Operator : Thu

Acquired : 8 Dec 2001 2:07 pm using AcqMethod GC2ANEW.M

Instrument : GC-2

Sample Name: 8 85052 W

Misc Info :



File : E:\HPCHEM\3\DATAB\12070137.D

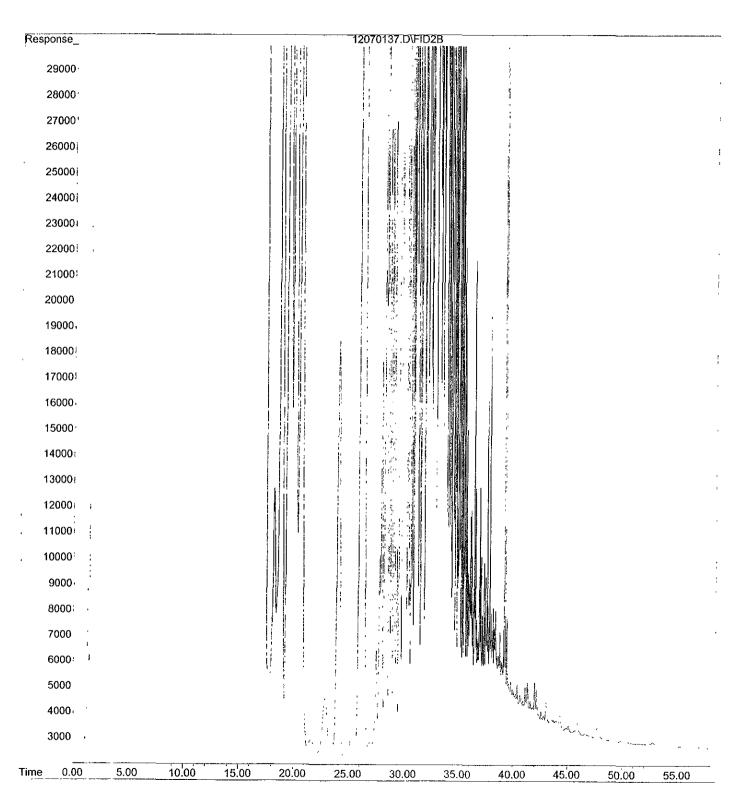
Operator : Thu

Acquired: 8 Dec 2001 3:40 pm using AcqMethod GC2ANEW.M

Instrument: GC-2

Sample Name: B 85053 W

Misc Info : Vial Number: 69



File : E:\HPCHEM\3\DATAB\12070141.D

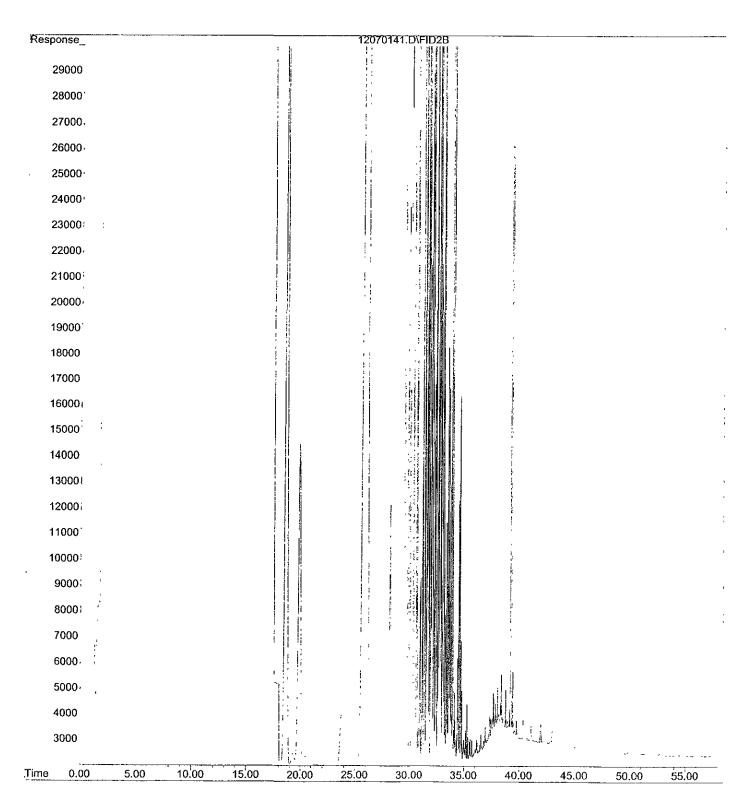
Operator : Thu

Acquired : 8 Dec 2001 6:41 pm using AcqMethod GC2ANEW.M

Instrument : GC-2

Sample Name: B 85054 W

Misc Info :



File : E:\HPCHEM\3\DATAB\12070143.D

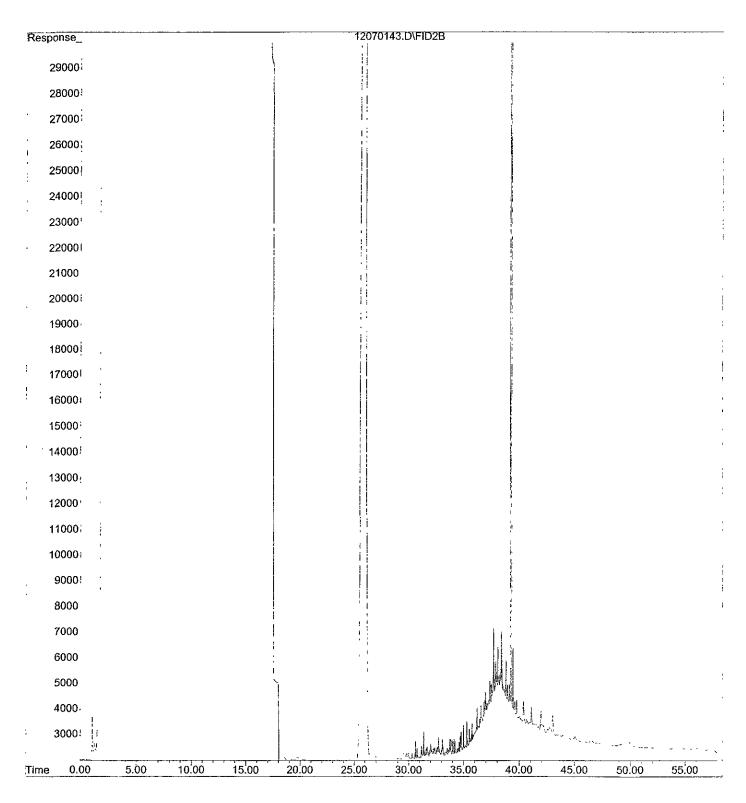
Operator : Thu

Acquired: 8 Dec 2001 8:10 pm using AcqMethod GC2ANEW.M

Instrument : GC-2

Sample Name: B 85055 W

Misc Info : Vial Number: 72



File : E:\HPCHEM\3\DATAB\12070145.D

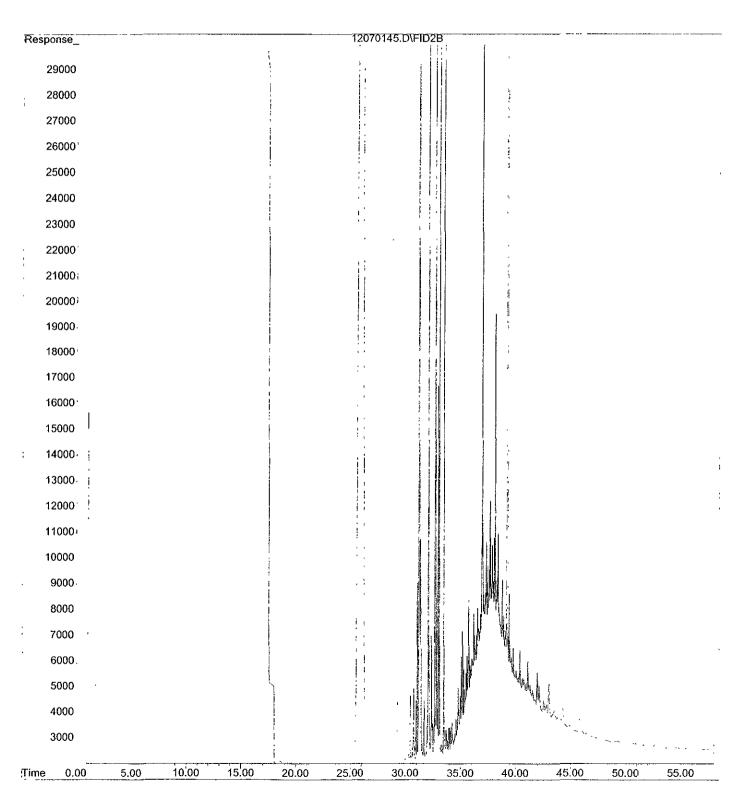
Operator : Thu

Acquired: 8 Dec 2001 9:37 pm using AcqMethod GC2ANEW.M

Instrument: GC-2

Sample Name: # 85056 W

Misc Info



File : E:\HPCHEM\3\DATAB\12070147.D

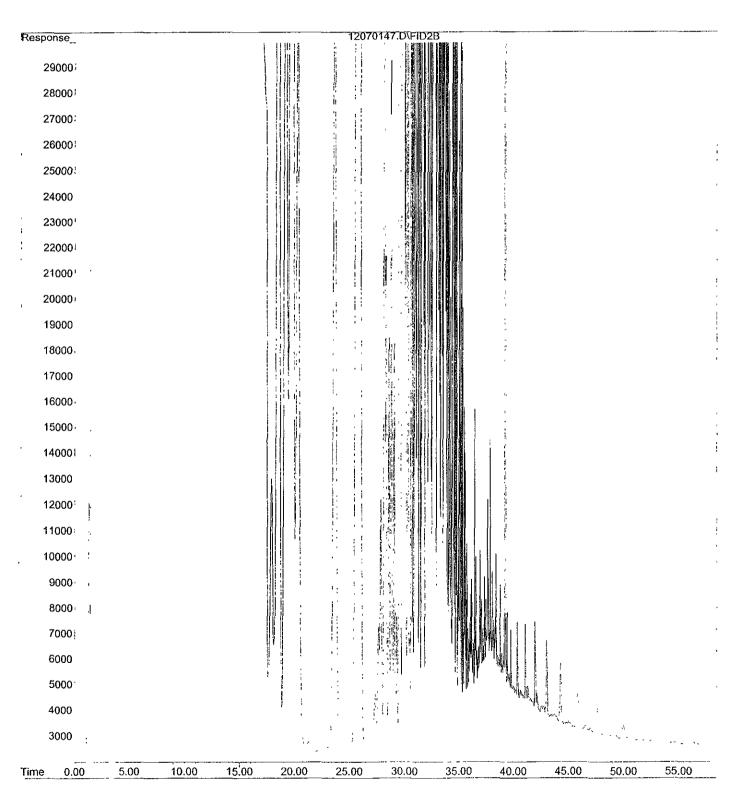
Operator : Thu

Acquired: 8 Dec 2001 11:08 pm using AcqMethod GC2ANEW.M

Instrument: GC-2

Sample Name: # 85057W

Misc Info : Vial Number: 74



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Telephon	ie: (925) 798-160	n		e.	יאי לטט	5) 798-	1622				TUI	(IV }	'KC	UN.	υH	ME								(2)	:
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Project #: 458.	-1705		Projec	t Man		<u> </u>	. 10	Con		1	520	~					8	ļ	1				/-3	Her	\setminus
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Cambria Environmental Technology	Client Project ID: #458-1705-4;	Date Sampled: 10/09/01
1144 65 th Street, Suite C	Embarcadero Ave	Date Received: 10/11/01
Oakland, CA 94608	Client Contact: Bob Schultz	Date Extracted: 10/11-11/07/01
	Client P.O:	Date Analyzed: 10/11-11/07/01

11/13/01

Dear Bob:

Enclosed are:

- 1). the results of 4 samples from your #458-1705-4; Embarcadero Ave project,
- 2). a QC report for the above samples
- 3). a copy of the chain of custody, and
- 4). a bill for analytical services.

All analyses were completed satisfactorily and all QC samples were found to be within our control limits. If you have any questions please contact me. McCampbell Analytical Laboratories strives for excellence in quality, service and cost. Thank you for your business and I look forward to working with you again.

Edward Hamilton, Lab Director

Yours truly

Cambria Environmental Technology	Client Project ID: #458-1705-4;	Date Sampled: 10/09/01
1144 65th Street, Suite C	Embarcadero Ave	Date Received: 10/11/01
Oakland, CA 94608	Client Contact: Bob Schultz	Date Extracted: 10/11/01
	Client P.O:	Date Analyzed: 10/11-10/16/01

Gasoline Range (C6-C12) Volatile Hydrocarbons as Gasoline*, with Methyl tert-Butyl Ether* & BTEX*

EPA methods 5030, modified 8015, and 8020 or 602; California RWOCR (SE Ray Region) method GCEID(5030)

Lab ID	Client ID	Matrix	TPH(g)⁺	МТВЕ	Benzene	Toluene	Ethyl- benzene	Xylenes	% Recovery Surrogate
80573	MW-1-8.3	S	30,a	ND	0.48	0.067	0.70	0.52	#
80574	MW-2-5.0	s	7.5,a	ND	0.027	0.051	0.041	0.087	#
80576	MW-3-5.0	S	1.6,g,j	ND	ND	ND	ND	ND	109
80579	MW-4-5.3	S	34, a	ND	0.70	0.068	0.41	0.97	#
									
			·						
	·								
otherwis	g Limit unless e stated; ND	W	50 ug/L	5.0	0.5	0.5	0.5	0.5	
	detected above orting limit	S	1.0 mg/kg	0.05	0.005	0.005	0.005	0.005	

^{*} water and vapor samples are reported in ug/L, wipe samples in ug/wipe, soil and sludge samples in mg/kg, and all TCLP and SPLP extracts in ug/L

^{*}The following descriptions of the TPH chromatogram are cursory in nature and McCampbell Analytical is not responsible for their interpretation: a) unmodified or weakly modified gasoline is significant; b) heavier gasoline range compounds are significant(aged gasoline?); c) lighter gasoline range compounds (the most mobile fraction) are significant; d) gasoline range compounds having broad chromatographic peaks are significant; biologically altered gasoline?; e) TPH pattern that does not appear to be derived from gasoline (?); f) one to a few isolated peaks present; g) strongly aged gasoline or diesel range compounds are significant; h) lighter than water immiscible sheen is present; i) liquid sample that contains greater than ~5 vol. % sediment; j) no recognizable pattern.



cluttered chromatogram; sample peak coelutes with surrogate peak

Cambria Environmental Technology
1144 65th Street, Suite C

Oakland, CA 94608

Client Project ID: #458-1705-4;
Embarcadero Ave

Date Sampled: 10/09/01

Date Received: 10/11/01

Client Contact: Bob Schultz

Date Extracted: 10/11/01

Client P.O:

Date Analyzed: 10/11-10/12/01

Diesel Range (C10-C23) and Oil-Range (C18+) Extractable Hydrocarbons as Diesel and Motor Oil with Silica Gel Clean-up*

EPA methods modified 8015, and 3550 or 3510; California RWQCB (SF Bay Region) method GCFID(3550) or GCFID(3510)

Lab ID	Client ID	Matrix	TPH(d) [†]	TPH(mo) ⁺	% Recovery Surrogate
80573	MW-1-8.3	S	5.7 , d	ND	100
80574	MW-2-5.0	S	74,g	300	100
80576	MW-3-5.0	s	17,g	160	102
80579	MW-4-5.3	s	8.3,d,g	10	98
Reporting Limit unle stated; ND means not of	detected above	w	50 ug/L	250 ug/L	
the reporting	limit	s	1.0 mg/kg	5.0 mg/kg	

^{*}water samples are reported in ug/L, wipe samples in ug/wipe, soil and sludge samples in mg/kg, and all TCLP / STLC / SPLP extracts in ug/L

The following descriptions of the TPH chromatogram are cursory in nature and McCampbell Analytical is not responsible for their interpretation: a) unmodified or weakly modified diesel is significant; b) diesel range compounds are significant; no recognizable pattern; c) aged diesel? is significant); d) gasoline range compounds are significant; e) medium boiling point pattern that does not match diesel (?); f) one to a few isolated peaks present; g) oil range compounds are significant; h) lighter than water immiscible sheen is present; i) liquid sample that contains greater than ~5 vol. % sediment.



cluttered chromatogram resulting in coeluted surrogate and sample peaks, or; surrogate peak is on elevated baseline, or; surrogate has been diminished by dilution of original extract.

Cambria Environmental T	echnology	Cli	ient Pr	oject II	D: #458-1705-4;	Date Sampled: 10/09/01				
1144 65 th Street, Suite C	23	En	barca	dero A	ve	Date I	Received: 10/11	ceived: 10/11/01 tracted: 10/11/01 alyzed: 10/12-10/22		
Oakland, CA 94608	į	Cli	ent Co	ntact:	Bob Schultz I	Date Extracted: 10/11/01				
		Cli	ent P.0):	Г	Date /	Analyzed: 10/12	-10/22	/01	
EPA method 8260		V	olatile	Orga	nies By GC/MS				-	
Lab ID					80573					
Client	ID				MW-1-8.3					
Matrix					S					
Compound	Concentratio	ր*	Reportin		Compound		Concentration*	Reportin	g Limit	
			W	S	•			W	S	
Acetone (b)	ND<130		5.0	25	trans-1,3-Dichloropropene		ND<25	1.0	5.0	
Benzene		260	1.0	5.0	Ethylene dibromide		ND<25	1.0	5.0	
Bromobenzene	ND<25		1.0	5.0	Ethylbenzene		800	1.0	5.0	
Bromochloromethane	ND<25		1.0	5.0	Hexachlorobutadiene		ND<130	5.0	25	
Bromodichloromethane	ND<25		1.0	5 0	lodomethane		ND<25	1.0	5.0	
Bromoform	ND<25		1.0	5.0	Isopropylbenzene		210	1.0	5.0	
Bromomethane	ND<25		1.0	5.0	p-Isopropyl toluene		ND<25	1.0	5.0	
n-Butyl benzene		580	1.0	5.0	Methyl butyl ketone (d)		ND<25	1.0	5.0	
sec-Butyl benzene	ND<25		1.0	5.0	Methylene Chloride ^(e)		ND<80	1.0	5.0	
tert-Butyl benzene	ND<25		1.0	5.0	Methyl ethyl ketone (f)		ND<50	2.0	10	
Carbon Disulfide	ND<25		1.0	5.0	Methyl isobutyl ketone (g)		ND<25	1.0	5.0	
Carbon Tetrachloride	ND<25		1.0	5.0	Methyl tert-Butyl Ether (MT	BE)		1.0	5.0	
Chlorobenzene	ND<25		1.0	5 0	Naphthalene		580	5.0	5.0	
Chloroethane	ND<25		1.0	5.0	n-Propyl benzene		910	1.0	5.0	
2-Chlorocthyl Vinyl Ether(c)	ND<25		1.0	5.0	Styrene (k)		ND<25	1.0	5.0	

Chloroform ND<25 1.0 5.0 1,1,1,2-Tetrachloroethane ND<25 1.0 5.0 Chloromethane ND<25 1.0 50 1,1,2,2-Tetrachloroethane ND<25 1.0 5.0 5.0 2-Chlorotoluene ND<25 1.0 Tetrachloroethene ND<25 1.0 5.0 ND<25 1.0 Toluene (1) 4-Chlorotoluene 5.0 1.0 5.0 Dibromochloromethane ND<25 1.0 5.0 1,2,3-Trichlorobenzene ND<130 25 1,2-Dibromo-3-chloropropane ND<50 2.0 10 1,2,4-Trichlorobenzene ND<130 25 Dibromomethane 1.0 ND<25 5.0 1,1,1-Trichloroethane ND<25 1.0 5.0 1,2-Dichlorobenzene ND<25 1.0 5.0 1,1,2-Trichloroethane ND<25 1.0 5.0 1,3-Dichlorobenzene ND<25 1.0 5.0 Trichloroethene ND<25 1.0 5.0 1,4-Dichlorobenzene 1.0 5.0 ND<25 Trichlorofluoromethane ND<25 1.0 5.0 Dichlorodifluoromethane 0.1 ND<25 5.0 1,2,3-Trichloropropane ND<25 1.0 5.0 I, I-Dichloroethane ND<25 1.0 5.0 1,2,4-Trimethylbenzene 1.0 5.0 1,2-Dichloroethane 1.0 5.0 ND<25 1,3,5-Trimethylbenzene 160 1.0 5.0 1.1-Dichloroethene 1.0 5.0 Vinyl Acetate (m) ND<25 ND<130 5.0 25 5.0 Vinyl Chloride (n) cis-1,2-Dichloroethene ND<25 1.0 ND<25 1.0 5.0 trans-1,2-Dichloroethene ND<25 1.0 5.0 Xylenes, total (6) 490 1.0 5.0 1,2-Dichloropropane ND<25 10 5.0 Comments: 1,3-Dichloropropane 5.0 ND<25 1.0 Surrogate Recoveries (%) 2,2-Dichloropropane 5.0 ND<25 1.0 Dibromofluoromethane 90 ND<25 1.0 5.0 1,1-Dichloropropene Toluene-d8 110 ND<25 50 cis-1,3-Dichloropropene 1.0 4-Bromofluorobenzene 102

water and vapor samples are reported in ug/L, soil and sludge samples in ug/kg, wipes in ug/wipe and all TCLP / SPLP extracts in ug/L

ND means not detected above the reporting limit; N/A means analyte not applicable to this analysis

(b) 2-propanone or dimethyl ketone; (c) (2-chloroethoxy) ethene; (d) 2-hexanone; (e) dichloromethane; (f) 2-butanone; (g) 4-methyl-2pentanone or isopropylacetone; (h) lighter than water immiscible sheen is present; (i) liquid sample that contains greater than ~5 vol. % sediment, (i) sample diluted due to high organic content; (k) ethenylbenzene; (l) methylbenzene; (m) acetic acid ethenyl ester; (n) chloroethene; (o) dimethylbenzenes.

DHS Certification No. 1644

Edward Hamilton, Lab Director

Cambria Environmental To	echnology			_	D: #458-1705-4;	Date S	Sampled: 10/09/	01	······
1144 65 th Street, Suite C		En	nbarcac	iero A	ve	Date F	Received: 10/11	/01	
Oakland, CA 94608		Cli	ent Co	ntact: 1	Bob Schultz	Date F	Extracted: 10/11	/01	•
Oakland, CA 34006									
		Cli	ient P.C):		Date A	Analyzed: 10/12	-10/22	/01
		<u> </u>	/olatile	Orga	nics By GC/MS	***************************************			
EPA method 8260		•	viatin	Olga	ines by Gennis				
Lab ID					80574				
Client									
Matrix		S S							
Mantx	<u> </u>		Reportin					Reportin	a 1
Compound	Concentration	*ac	W	S	Compound		Concentration*	W	S
Acetone (6)	ND<85		5.0	25	trans-1,3-Dichloroproper	ıe	ND	1.0	5.0
Benzene	ND		10	5.0	Ethylene dibromide		ND	1.0	5.0
Bromobenzene	ND		1,0	5.0	Ethylbenzene		5.3	1.0	5.0
Bromochloromethane	ND		1.0	5.0	Hexachlorobutadiene		ND	5.0	25
Bromodichloromethane	ND		1.0	5.0	lodomethane		ND	1.0	5.0
Bromoform	ND		1.0	50	Isopropylbenzene		ND	1.0	5.0
Bromomethane	ND		1.0	5.0	p-Isopropyl toluene		11	1.0	5.0
n-Butyl benzene	1	18	1.0	5.0	Methyl butyl ketone (d)		ND	0.1	5.0
sec-Butyl benzene	ND		1.0	5.0	Methylene Chloride(c)		ND<10	1.0	5.0
tert-Butyl benzene	ND		1.0	5.0	Methyl ethyl ketone (f)		ND	20	10
Carbon Disulfide	ND		1.0	5.0	Methyl isobutyl ketone (g	3	ND	1.0	5.0
Carbon Tetrachloride	ND		1.0	5.0	Methyl tert-Butyl Ether (MTBE)		1.0	5.0
Chlorobenzene	ND		1.0	5.0	Naphthalene		ND	5.0	5.0
Chloroethane	ND		1.0	5.0	n-Propyl benzene		17	1.0	5.0
2-Chloroethyl Vinyl Ether(c)	ND		1.0	50	Styrene (k)		ND	10	5.0
Chloroform	ND		1.0	50	1,1,1,2-Tetrachloroethan	d	ND	10	5.0
Chloromethane	ND		1.0	5.0	1,1,2,2-Tetrachloroethan	e	ND	1.0	5.0
2-Chlorotolucne	ND		1.0	5.0	Tetrachloroethene		ND	1.0	5.0
4-Chlorotoluene	ND		1.0	5.0	Toluene (1)		ND	1.0	5.0
Dibromochloromethane	ND		1.0	5.0	1,2,3-Trichlorobenzene		ND	5.0	25
1,2-Dibromo-3-chloropropane	ND		2.0	10	1,2,4-Trichlorobenzene		ND	5.0	25
Dibromomethane	ND		0.1	5.0	1,1,1-Trichloroethane		ND	1.0	5.0
1,2-Dichlorobenzene	ND		1.0	5.0	1,1,2-Trichloroethane		ND	1.0	5.0
1,3-Dichlorobenzene	ND		1.0	5.0	Trichloroethene		ND	1.0	5.0
1,4-Dichlorobenzene	ND		1.0	5.0	Trichlorofluoromethane		ND	1.0	50
Dichlorodifluoromethane	ND		1.0	5.0	1,2,3-Trichloropropane		ND	1.0	5.0
1,1-Dichloroethane	ND		0.1	5.0	1,2,4-Trimethylbenzene		26	1.0	5.0
1,2-Dichloroethane	ND		1.0	5.0	1,3,5-Trimethylbenzene		11	1.0	5.0
1,1-Dichloroethene	ND		1.0	5.0	Vinyl Acetate (m)		ND	5.0	25
cis-1,2-Dichloroethene	ND		1.0	5.0	Vinyl Chloride (n)		ND 21	1.0	5.0
trans-1,2-Dichloroethene	ND		1.0	5.0	Xylenes, total (0)		7.1	1.0	5.0
1,2-Dichleropropane	ND		1.0	50	Comments:				
1,3-Dichloropropane	ND		1.0	5.0		rogate Re	ecoveries (%)		
2,2-Dichloropropane	ND		10	5.0	Dibromofluoromethane				03
1.1 Dioblevensenene	MD		10	5.0	Toluene-d8			· · · · · ·	44

"water and vapor samples are reported in ug/L, soil and sludge samples in ug/kg, wipes in ug/wipe and all TCLP / SPLP extracts in ug/L

5.0

5.0

Toluene-d8

ND means not detected above the reporting limit; N/A means analyte not applicable to this analysis

ND

ND

10

1.0

(b) 2-propanone or dimethyl ketone; (c) (2-chloroethoxy) ethene; (d) 2-hexanone; (e) dichloromethane; (f) 2-butanone; (g) 4-methyl-2-pentanone or isopropylacetone; (h) lighter than water immiscible sheen is present; (i) liquid sample that contains greater than ~5 vol. % sediment; (j) sample diluted due to high organic content; (k) ethenylbenzene; (l) methylbenzene; (m) acetic acid ethenyl ester; (n) chloroethene; (o) dimethylbenzenes.

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1,1-Dichloropropene

cis-1,3-Dichloropropene



4-Bromofluorobenzene

Edward Hamilton, Lab Director

94

107

Cambria Environmental T	Technology	Clien	t Pro	oject II	D: #458-1705-4;	Pate Sa	ampled: 10/09	/01			
1144 65th Street, Suite C		Emba	тсас	dero A	ve D	ate R	eceived: 10/11	/01			
Oakland, CA 94608		Client	t Co	ntact:	Bob Schultz D	ate Ex	ktracted: 10/11	1/01			
		Client	t P.C);	D	ate A	nalyzed: 10/12	2-10/22	2/01		
EPA method 8260		Vol	atile	Orga	nics By GC/MS	 ,					
Lab II)				80576						
Client					MW-3-5.0			·			
Matrix					S S		·				
Compound	Concentratio	n* L		g Limit	Compound		Concentration* Report		Concentration*		ng Limit
	315 - 25		W	S		_		W	S		
Acetone (6)	ND<85		5.0	25	trans-1,3-Dichloropropene		ND	1.0	5.0		
Benzene	ND		.0	5.0	Ethylene dibromide		ND	1.0	5.0		
Bromobenzene	ND		.0	5.0	Ethylbenzene		ND	1.0	5.0		
Bromochloromethane	ND		.0	5.0	Hexachlorobutadiene		ND	5.0	25		
Bromodichloromethane	ND		.0	5.0	Iodomethane		ND	1.0	5.0		
Bromoform	ND		.0	5.0	Isopropylbenzene		ND	1.0	5.0		
Bromomethane	ND	1	.0	5.0	p-Isopropyl toluene		ND	1.0	5.0		
n-Butyl benzene	ND	_ 1	.0	50	Methyl butyl ketone (d)		ND	1.0	5.0		
sec-Butyl benzene	ND	1	.0	50	Methylene Chloride(e)		ND<10	1.0	5.0		
tert-Butyl benzene	ND		.0	5.0	Methyl ethyl ketone (1)		ND	2.0	10		
Carbon Disulfide	ND]	.0	5.0	Methyl isobutyl ketone (g)		ND	1.0	5.0		
Carbon Tetrachloride	ND	1.	0	5.0	Methyl tert-Butyl Ether (MTE	3E)		1.0	5.0		
Chlorobenzene	ND	1.	.0	50	Naphthalene		ND	5.0	5.0		
Chloroethane	ND	1	.0	5.0	n-Propyl benzene		ND	1.0	5.0		
2-Chloroethyl Vinyl Ether(c)	ND	1.	.0	5.0	Styrene (k)		ND	1.0	5.0		
Chloroform	ND	1.	.0	5.0	1,1,1,2-Tetrachloroethane		ND	1.0	5.0		
Older and	3.150										

water and vapor samples are reported in ug/L, soil and sludge samples in ug/kg, wipes in ug/wipe and all TCLP / SPLP extracts in ug/L

ND means not detected above the reporting limit; N/A means analyte not applicable to this analysis

ND

ND

ND

ND

ND

ND

ND

ND

ND

ND

ND

ND

ND

ND

ND

ND

ND

ИD

ND

ND

1.0

1.0

1.0

1.0

2.0

1.0

10

1.0

1.0

1.0

1.0

1.0

1.0

1.0

1.0

1.0

1.0

1.0

1.0

1.0

5.0

5.0

5.0

5.0

10

5.0

5.0

5.0

5.0

5.0

5.0

5.0

5.0

5.0

5.0

50

5.0

5.0

5.0

50

1,1,2,2-Tetrachloroethane

1,2,3-Trichlorobenzene

1,2,4-Trichlorobenzene

1,1,1-Trichloroethane

1,1,2-Trichloroethane

Trichlorofluoromethane

1,2,3-Trichloropropane

1,2,4-Trimethylbenzene

1,3,5-Trimethylbenzene

Dibromofluoromethane

4-Bromofluorobenzene

Trichloroethene

Vinyl Acetate (0)

Vinyl Chloride (11)

Xylenes, total (6)

Comments:

Toluene-d8

Tetrachloroethene

Toluene (1)

(b) 2-propanone or dimethyl ketone; (c) (2-chloroethoxy) ethene; (d) 2-hexanone; (e) dichloromethane; (f) 2-butanone; (g) 4-methyl-2-pentanone or isopropylacetone; (h) lighter than water immiscible sheen is present; (i) liquid sample that contains greater than ~5 vol. % sediment; (j) sample diluted due to high organic content; (k) ethenylbenzene; (l) methylbenzene; (m) acetic acid ethenyl ester; (n) chloroethene; (o) dimethylbenzenes.

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Chloromethane

2-Chlorotoluene

4-Chlorotoluene

Dibromomethane

1,2-Dichlorobenzene

1,3-Dichlorobenzene

1,4-Dichlorobenzene

1,1-Dichloroethane

1,2-Dichloroethane

1,1-Dichloroethene

cis-1,2-Dichloroethene

1,2-Dichloropropane

1,3-Dichloropropane

2,2-Dichloropropane

1,1-Dichloropropene

cis-1,3-Dichloropropene

trans-1,2-Dichloroethene

Dichlorodifluoromethane

Dibromochloromethane

1,2-Dibromo-3-chloropropane

__Edward Hamilton, Lab Director

Surrogate Recoveries (%)

ND

ND

ND

ND

ND

ND

ND

ND

ND

ND

ND

ND

ND

ND

5.7

1.0

1.0

1.0

5.0

5.0

1.0

1.0

1.0

1.0

10

1.0

1.0

5.0

1.0

1.0

103

95

107

5.0

5.0

5.0

25

25

5.0

5.0

5.0

5.0

5.0

5.0

5.0

25

5.0

5.0

110 2nd Avenue South, #D7, Pacheco, CA 94553-5560 Telephone: 925-798-1620 Fax: 925-798-1622

http://www.mccampbell.com E-mail: main@mccampbell.com

	Volatile Organics By GC/MS	Date Finally 2cd. 10/12-10/22/01		
	Client P.O:	Date Analyzed: 10/12-10/22/01		
Oakland, CA 94608	Client Contact: Bob Schultz	Date Extracted: 10/11/01		
1144 65 th Street, Suite C	Embarcadero Ave	Date Received: 10/11/01		
Cambria Environmental Technology	Client Project ID: #458-1705-4;	Date Sampled: 10/09/01		

2DA	method	9260	
CrA.	memoa	820U	

51 11 11 10 10 10 10 10 10 10 10 10 10 10	
Lab ID	80579
Client ID	MW-4-5.3
Matrix	S

Math	<u> </u>	S					
Compound	Concentration*	Reportin	ng Limit	Compound	Concentration*	Reporti	ng Limit
•	001100111111011	W	S	Compound	Concentiation	W	S
Acetone ^(b)	ND<130	5.0	25	trans-1,3-Dichloropropene	ND<25	10	5.0
Benzene	700	1.0	5.0	Ethylene dibromide	ND<25	1.0	5.0
Bromobenzene	ND<25	1.0	5.0	Ethylbenzene	460	1.0	5.0
Bromochloromethane	ND<25	1.0	5.0	Hexachlorobutadiene	ND<130	5.0	25
Bromodichloromethane	ND<25	10	5.0	lodomethane	ND<25	1.0	5.0
Bromoform	ND<25	1.0	5.0	Isopropylbenzene	220	1.0	5.0
Bromomethane	ND<25	1.0	5.0	p-Isopropyl toluene	ND<25	1.0	5.0
n-Butyl benzene	ND<25	1.0	5.0	Methyl butyl ketone (d)	ND<25	1.0	5.0
sec-Butyl benzene	ND<25	1.0	5.0	Methylene Chloride ^(e)	ND<80	1.0	5.0
tert-Butyl benzene	ND<25	1.0	5.0	Methyl ethyl ketone (1)	ND<50	2.0	10
Carbon Disulfide	ND<25	1.0	50	Methyl isobutyl ketone (g)	ND<25	1.0	5.0
Carbon Tetrachloride	ND<25	1.0	5.0	Methyl tert-Butyl Ether (MTBE)		1.0	5.0
Chlorobenzene	ND<25	1.0	5.0	Naphthalene	620	5.0	5.0
Chloroethane	ND<25	1.0	5.0	n-Propyl benzene	870	1.0	5.0
2-Chloroethyl Vinyl Ether(c)	ND<25	1.0	50	Styrene (k)	ND<25	1.0	5.0
Chloroform	ND<25	1.0	5.0	1,1,1,2-Tetrachloroethane	ND<25	1.0	5.0
Chloromethane	ND<25	1.0	5.0	1,1,2,2-Tetrachloroethane	ND<25	1.0	5.0
2-Chlorotoluene	ND<25	1.0	5.0	Tetrachloroethene	ND<25	1.0	5.0
4-Chlorotoluene	ND<25	1.0	5.0	Toluene (1)	41	1.0	5.0
Dibromochloromethane	ND<25	1.0	5.0	1,2,3-Trichlorobenzene	ND<130	5.0	25
1,2-Dibromo-3-chloropropane	ND<50	2.0	10	1,2,4-Trichlorobenzene	ND<130	5.0	25
Dibromomethane	ND<25	1.0	5.0	1,1,1-Trichloroethane	ND<25	1.0	5.0
1,2-Dichlorobenzene	ND<25	1.0	50	1,1,2-Trichloroethane	ND<25	1.0	5.0
1,3-Dichlorobenzene	ND<25	1.0	5.0	Trichloroethene	ND<25	1.0	5.0
1,4-Dichlorobenzene	ND<25	1.0	5.0	Trichlorofluoromethane	ND<25	1.0	5.0
Dichlorodifluoromethane	ND<25	1.0	5.0	1,2,3-Trichloropropane	ND<25	1.0	5.0
1,1-Dichloroethane	ND<25	10	5.0	1,2,4-Trimethylbenzene	1000	1.0	5.0
1,2-Dichloroethane	ND<25	1.0	5.0	1,3,5-Trimethylbenzene	610	1.0	5.0
1,1-Dichloroethene	ND<25	1.0	5.0	Vinyl Acetate (m)	ND<130	5.0	25
cis-1,2-Dichloroethene	ND<25	1.0	5.0	Vinyl Chloride (n)	ND<25	1.0	5.0
trans-1,2-Dichloroethene	ND<25	1.0	5.0	Xylenes, total (0)	990	1.0	5.0
1,2-Dichloropropane	ND<25	10	5.0	Comments:	<u></u> !		
1,3-Dichloropropane	ND<25	10	5.0	Surrogate Re	coveries (%)		
2,2-Dichloropropane	ND<25	10	5.0	Dibromofluoromethane		10)7
1,1-Dichloropropene	ND<25	1.0	5.0	Toluene-d8		9:	
cis-1,3-Dichloropropene	ND<25	1.0	5.0	4-Bromofluorobenzene		10	

water and vapor samples are reported in ug/L, soil and sludge samples in ug/kg, wipes in ug/wipe and all TCLP / SPLP extracts in ug/L

DHS Certification No. 1644

Edward Hamilton, Lab Director

ND means not detected above the reporting limit; N/A means analyte not applicable to this analysis

⁽b) 2-propanone or dimethyl ketone; (c) (2-chloroethoxy) ethene; (d) 2-hexanone; (e) dichloromethane; (f) 2-butanone; (g) 4-methyl-2-pentanone or isopropylacetone; (h) lighter than water immiscible sheen is present; (i) liquid sample that contains greater than ~5 vol. % sediment; (j) sample diluted due to high organic content; (k) ethenylbenzene, (l) methylbenzene; (m) acetic acid ethenyl ester; (n) chloroethene; (o) dimethylbenzenes.

110 2nd Avenue South, #D7, Pacheco, CA 94553-5560 Telephone: 925-798-1620 Fax: 925-798-1622 http://www.mccampbell.com E-mail: main@mccampbell.com

Cambria Environmental Technology
1144 65th Street, Suite C

Oakland, CA 94608

Client Project ID: #458-1705-4;
Embarcadero Ave

Date Sampled: 10/09/01

Date Received: 10/11/01

Client Contact: Kris Cejka

Date Extracted: 11/06/01

Client P.O:

Date Analyzed: 11/07/01

Semi-Volatile Organics By GC/MS

EPA method 625 and 3510 or 8270 and 3550

Lab ID		80573					
Client ID				MW-1-8.3			
Matrix				S			
Compand	Concentration*	Reporti	ng Limit	C	1	Repor	ting Limit
Compound	Concentration*	W	S	Compound	oncentration	w	S
Acenaphthene	ND	10	0.33	Di-n-octyl Phthalate	ND	10	0 33
Acenaphthylene	ND	10	0 33	1,2-Diphenylhydrazine	ND	10	0.33
Anthracene	ND	10	0.33	Fluoranthene	ND	10	0.33
Benzidine	ND	50	16	Fluorene	ND	10	0.33
Benzoic Acid	ND	50	1.6	Hexachlorobenzene	ND	10	0.33
Benzo(a)anthracene	ND	10	0.33	Hexachlorobutadiene	ND	10	0.33
Benzo(b)fluoranthene	ND	10	0 33	Hexachlorocyclopentadiene	ND	50	1.6
Benzo(k)fluoranthene	ND	10	0.33	Hexachloroethane	ND	10	0.33
Benzo(g,h,i)perylene	ND	10	0.33	Indeno(1,2,3-cd)pyrene	ND	10	0.33
Benzo(a)pyrene	ND	10	0.33	Isophorone	ND	10	0.33
Benzyl Alcohol	ND	20	0 66	2-Methylnaphthalene	ND	10	0.33
Bis(2-chloroethoxy)methane	ND	10	0 33	2-Methylphenol (o-Cresol)	ND	10	0.33
Bis(2-chloroethyl) Ether	ND	10	0.33	3 &/or 4-Methylphenol (m &/or p-Cresol)	ND	10	0.33
Bis(2-chlorossopropyl)Ether	ND	10	0 33	Naphthalene	ND	10	0.33
Bis(2-cthylhexyl) Phthalate	ND	10	0.33	2-Nitroanilme	ND	50	1.6
4-Bromophenyl Phenyl Ether	ND	10	0.33	3-Nitroaniline	ND	50	1.6
Butylbenzyl Phthalate	ND	10	0.33	4-Nitroaniline	ND	50	1.6
4-Chloroanaline	ND	20	0.66	2-Nitrophenol	ND	50	1.6
4-Chloro-3-methylphenol	ND	10	0.33	4-Nitrophenol	ND	50	1.6
2-Chloronaphthalene	ND	10	0.33	Nitrobenzene	ND	10	0.33
2-Chlorophenol	ND	10	0.33	N-Nitrosodiphenylamine	ND	10	0.33
4-Chlorophenyl Phenyl Ether	ND	10	0.33	N-Nitrosodi-n-propylamine	ND	10	0.33
Chrysene	ND	10	0.33	Pentachlorophenol	ND	50	1.6
Dibenzo(a,h)anthracene	ND	10	0.33	Phenanthrene	ND	10	0.33
Dibenzofuran	ND	10	0.33	Phenol	ND	10	0.33
Di-n-butyl Phthalate	ND	10	0.33	Pyrene	ND	10	0 33
1,2-Dichlorobenzene	ND	10	0.33	1,2,4-Trichlorobenzene	ND	10	0.33
1,3-Dichlorobenzene	ND	10	0.33	2,4,5-Trichlorophenol	ND	10	0.33
1,4-Dichlorobenzene	ND	10	0.33	2,4,6-Trichlorophenol	ND	10	0.33
3,3-Dichlorobenzidine	ND	20	0.66				
2,4-Dichlorophenol	ND	10	0.33	Comments:			
Diethyl Phthalate	ND	10	0 33	Surrogate Recoveries	s (%)		
2,4-Dimethylphenol	ND	10	0.33	2-Fluorophenol	<u> </u>	105	
Dimethyl Phthalate	ND	10	0.33	Phenof-d5		103	
4,6-Dinitro-2-methylphenol	ND	50	1.6	Nitrobenzene-d5		109	
2,4-Dinitrophenol	ND	50	1.6	2-Fluorobiphenyl		108	
2,4-Dinitrotoluene	ND	10	0.33	2,4,6-Tribromophenol		85	
2,6-Dinitrotoluene	ND	10	0.33	p-Terphenyl-d14		94	
	l		الــــــــــــــــــــــــــــــــــــ		1		

^{*}water samples are reported in ug/L, soil and sludge samples in mg/kg, wipes in ug/wipe and all TCLP / STLC / SPLP extracts in ug/L

ND means not detected above the reporting limit; N/A means analyte not applicable to this analysis.

[&]quot;surrogate diluted out of range

h) lighter than water immiscible sheen is present; i)liquid sample that contains greater than ~5 vol. % sediment; j) sample diluted due to high organic content

Cambria Environmental	Technology	Client	t Projec	et ID: #458-1705-4;	ate Samp	oled: 10/09/0	1	
1144 65 th Street, Suite C	1 00111.01.08)		rcader	Ave	Date Received: 10/11/01			
Oakland, CA 94608		Client	t Conta	ct: Kris Cejka Da	Date Extracted: 11/06/01			
		Client	P O	Da	ite Anali	yzed: 11/07/		
<u></u>						yzed. 11/0//		
EPA method 625 and 3510 or 8		Semi-V	⁷ olatile	Organics By GC/MS				
Lab ID				80574				
Client ID	·			MW-2-5.0				
Matrix	·			S S				•
17 14417/		Υ					r	
Compound	Concentration*	W	ng Limit	Compound		oncentration	Repor	ting Limit
Acenaphthene	ND<1.0	10	0.33	Di-n-octyl Phthalate		ND<1.0	10	0.33
Acenaphthylene	ND<1 0	10	0.33	1,2-Diphenylhydrazine		ND<1.0	10	0.33
Anthracene	ND<1.0	10	0.33	Fluoranthene		ND<10	10	0.33
Benzidine	ND<5.0	50	1.6	Fluorene		ND<1.0	10	0.33
Benzoic Acid	ND<50	50	16	Hexachlorobenzene		ND<1.0	10	0.33
Benzo(a)anthracene	ND<1.0	10	0 33	Hexachlorobutadiene		ND<1.0	10	0.33
Benzo(b)fluoranthene	ND<1.0	10	0.33	Hexachlorocyclopentadiene		ND<50	50	1.6
Benzo(k)fluoranthene	ND<1.0	10	0.33	Hexachloroethane		ND<1.0	10	0.33
Benzo(g,h,i)perylene	ND<1.0	10	0.33	Indeno(1,2,3-cd)pyrene		ND<10	10	0.33
Benzo(a)pyrene	ND<1.0	10	0 33	Isophorone		ND<1.0	10	0.33
Benzyl Alcohol	ND<2 0	20	0.66	2-Methylnaphthalene		ND<1.0	10	0.33
Bis(2-chloroethoxy)methane	ND<1.0	10	0.33	2-Methylphenol (o-Cresol)		ND<1.0	10	0.33
Bis(2-chloroethyl) Ether	ND<1.0	10	0.33	3 &/or 4-Methylphenol (m &/or p-	Cresol)	ND<1.0	10	0.33
Bis(2-chloroisopropyl)Ether	ND<1 0	10	0 33	Naphthalene	010301)	ND<1.0	10	0.33
Bis(2-ethylhexyl) Phthalate	ND<1 0	10	0.33	2-Nitroaniline		ND<50	50	1.6
4-Bromophenyl Phenyl Ether	ND<1.0	10	0.33	3-Nitroaniline		ND<5.0	50	16
Butylbenzyl Phthalate	ND<1.0	10	0 33	4-Nitroaniline		ND<5.0	50	1.6
4-Chloroanaline	ND<2.0	20	0.66	2-Nitrophenol		ND<5.0	50	1.6
4-Chloro-3-methylphenol	ND<1.0	10	0.33	4-Nitrophenol		ND<5.0	50	1.6
2-Chloronaphthalene	ND<1.0	10	0.33	Nitrobenzene		ND<1.0	10	0.33
2-Chlorophenol	ND<10	10	0 33	N-Nitrosodiphenylamine		ND<1.0	10	0.33
4-Chlorophenyl Phenyl Ether	ND<1.0	10	0.33	N-Nitrosodi-n-propylamine		ND<1.0	10	0.33
Chrysene	ND<1.0	10	0.33	Pentachlorophenol		ND<5.0	50	1.6
Dibenzo(a,h)anthracene	ND<1.0	10	0.33	Phenanthrene		ND<1.0	10	0.33
Dibenzofuran	ND<1.0	10	0 33	Phenol		ND<1.0	10	0.33
Di-n-butyl Phthalate	ND<1.0	10	0.33	Pyrene		ND<1.0	10	0.33
1,2-Dichlorobenzene	ND<1.0	10	0.33	1,2,4-Trichlorobenzene		ND<1.0	10	0.33
1,3-Dichlorobenzene	ND<1.0	10	0.33	2,4,5-Trichlorophenol		ND<1.0	10	0.33
1,4-Dichlorobenzene	ND<1.0	10	0.33	2,4,6-Trichlorophenol		ND<1.0	10	0.33
3,3-Dichlorobenzidine	ND<2.0	20	0.66					
2,4-Dichlorophenol	ND<1.0	10	0.33	Comments: j				
Diethyl Phthalate	ND<1.0	10	0.33	Surrogate I	Recoverie	s (%)		
2,4-Dimethylphenol	ND<1.0	10	0.33	2-Fluorophenol		<u> </u>	90	·
Dimethyl Phthalate	ND<1.0	10	0.33	Phenol-d5				
4,6-Dinitro-2-methylphenol	ND<5.0	50	1.6	Nitrobenzene-d5	·. -		119	
2,4-Dinitrophenol	ND<5.0	50	1.6	2-Fluorobiphenyl		··	119	
2,4-Dinitrotoluene	ND<1.0	10	0.33	2,4,6-Tribromophenol			92	
2,6-Dinitrotoluene	ND<1.0	10	0.33	p-Terphenyl-d14			114	

^{*}water samples are reported in ug/L, soil and sludge samples in mg/kg, wipes in ug/wipe and all TCLP / STLC / SPLP extracts in ug/L

ND means not detected above the reporting limit; N/A means analyte not applicable to this analysis.

[#] surrogate diluted out of range

h) lighter than water immiscible sheen is present; i)liquid sample that contains greater than ~5 vol. % sediment; j) sample diluted due to high organic content

Cambria Environmental	Technology	Client Project ID: #458-1705-4; Embarcadero Ave			Date Sampled: 10/09/01			
1144 65th Street, Suite C		Emba	ircader	o Ave	Date Received: 10/11/01			
Oakland, CA 94608		Clien	t Conta	ct: Kris Cejka	Date Extracted: 11/06/01			
		Clien	P.O:		Date Anal	yzed: 11/07/	01	
		Semi-V	olatile	Organics By GC/MS		<u></u>		
EPA method 625 and 3510 or 8								
Lab ID	<u> </u>			80576				
Client ID				MW-3-5.0				
Matrix				S				
Compound	Concentration*	Report	ng Limit	C1		1	Repor	ting Limit
Compound	Concentration	W	S	Compound		oncentration	W	S
Acenaphthene	ND	10	0.33	Di-n-octyl Phthalate		ND	10	0.33
Acenaphthylene	ND	10	0.33	1,2-Diphenylhydrazine		ND	10	0.33
Anthracene	ND	10	0.33	Fluoranthene		ND	10	0.33
Benzidine	ND	50	1.6	Fluorene		ND	10	0.33
Benzoic Acid	ND	50	1.6	Hexachlorobenzene		ND	10	0.33
Benzo(a)anthracene	ND	10	0.33	Hexachlorobutadiene		ND	10	0.33
Benzo(b)fluoranthene	ND	10	0.33	Hexachlorocyclopentadiene		ND	50	16
Benzo(k)fluoranthene	ND	10	0.33	Hexachloroethane		ND	10	0 33
Benzo(g,h,i)perylene	ND	10	0.33	Indeno(1,2,3-cd)pyrene		ND	10	033
Benzo(a)pyrene	ND	10	0.33	Isophorone		ND	10	0.33
Benzyl Alcohol	ND	20	0.66	2-Methylnaphthalene		ND	10	0.33
Bis(2-chloroethoxy)methane	ND	10	0.33	2-Methylphenol (o-Cresol)		ND	10	0.33
Bis(2-chloroethyl) Ether	ND	10	0.33	3 &/or 4-Methylphenol (m &/or	p-Cresol)	ND	10	0.33
Bis(2-chloroisopropyl)Ether	ND	10	0.33	Naphthalene	•	ND	10	0.33
Bis(2-ethylhexyl) Phthalate	ND	10	0.33	2-Nitroaniline		ND	50	1.6
4-Bromophenyl Phenyl Ether	ND	10	0.33	3-Nitroaniline		ND	50	1.6
Butylbenzyl Phthalate	ND	10	0.33	4-Nitroaniline		ND	50	1.6
4-Chloroanaline	ND	20	0.66	2-Nitrophenol		ND	50	1.6
4-Chloro-3-methylphenol	ND	10	0.33	4-Nitrophenol		ND	50	1.6
2-Chloronaphthalene	ND	10	0.33	Nitrobenzene		ND	10	0.33
2-Chlorophenol	ND	10	0.33	N-Nitrosodiphenylamine		ND	10	0.33
4-Chlorophenyl Phenyl Ether	ND	10	0.33	N-Nitrosodi-n-propylamine		ND	10	0.33
Chrysene	ND	10	0.33	Pentachlorophenol		ND	50	1.6
Dibenzo(a,h)anthracene	ND	10	0.33	Phenanthrene		ND	10	0.33
Dibenzofuran	ND	10	0.33	PhenoI		ND	10	0.33
Di-n-butyl Phthalate	ND	10	0 33	Pyrene		ND	10	0.33
1,2-Dichlorobenzene	ND	10	0.33	1,2,4-Trichlorobenzene		ND	10	0.33
1,3-Dichlorobenzene	ND	10	0.33	2,4,5-Trichlorophenol		ND	10	0.33
1,4-Dichlorobenzene	ND	10	0.33	2,4,6-Trichlorophenol		ND	10	0.33
3,3-Dichlorobenzidine	ND	20	0.66					
2,4-Dichlorophenol	ND	10	0.33	Comments:				
Diethyl Phthalate	ND	10	0.33	Surrogat	e Recoveries	s (%)	······································	
2,4-Dimethylphenol	ND	10	0.33	2-Fluorophenol		~~	102	
Dimethyl Phthalate	ND	10	0.33	Phenol-d5	·		81	
4,6-Dinitro-2-methylphenol	ND	50	1.6	Nitrobenzene-d5			105	
2,4-Dinitrophenol	ND	50	1.6	2-Fluorobiphenyl	····		105	
2,4-Dinitrotoluene	ND	10	0.33		· ·			
2,6-Dinitrotoluene	ND			2,4,6-Tribromophenol 88 p-Terphenyl-dl 4 96				

^{*}water samples are reported in ug/L, soil and sludge samples in mg/kg, wipes in ug/wipe and all TCLP / STLC / SPLP extracts in ug/L

ND means not detected above the reporting limit; N/A means analyte not applicable to this analysis.

[&]quot; surrogate diluted out of range

h) lighter than water immiscible sheen is present; i)liquid sample that contains greater than ~5 vol % sediment, j) sample diluted due to high organic content

Cambria Environmental	Fechnology			7 ID: #436-1703-4;	Date Sampled: 10/09/01				
1144 65 th Street, Suite C	:	Embarcadero Ave Date R				Received: 10/11/01			
Oakland, CA 94608		Client	Conta	ct: Kris Cejka D	ate Extra	cted: 11/06/	01		
		Client	P.O:	D	ate Anal	yzed: 11/07/	zed: 11/07/01		
EPA method 625 and 3510 or 8		Semi-V	olatile	Organics By GC/MS	 				
St. Committee Co	270 and 3550			00.500					
Lab ID				80579					
Client ID				MW-4-5.3					
Matrix			·	S					
Compound	Concentration*	Reporti	ng Limit	Compound		oncentration	Repor	ting Limit	
Compound	Concontration	W	S	Compound		oncentiation	Ŵ	S	
Acenaphthene	ND	10	0.33	Di-n-octyl Phthalate		ND	10	0.33	
Acenaphthylene	ND	10	0.33	1,2-Diphenylhydrazine		ND	10	0.33	
Anthracene	ND	10	0 33	Fluoranthene		ND	10	0.33	
Benzidine	ND	50	1.6	Fluorene		ND	10	0.33	
Benzoic Acid	ND	50	1.6	Hexachlorobenzene		ND	10	0.33	
Benzo(a)anthracene	ND	10	0.33	Hexachlorobutadiene		ND	10	0 33	
Benzo(b)fluoranthene	ND	10	0.33	Hexachlorocyclopentadiene		ND	50	16	
Benzo(k)fluoranthene	ND	10	0.33	Hexachloroethane		ND	10	0.33	
Benzo(g,h,i)perylene	ND	10	0.33	Indeno(1,2,3-cd)pyrene		ND	10	0.33	
Benzo(a)pyrene	ND	10	0.33	Isophorone		ND	10	0 33	
Benzyl Alcohol	ND	20	0.66	2-Methylnaphthalene		ND	10	0.33	
Bis(2-chloroethoxy)methane	ND	10	0.33	2-Methylphenol (o-Cresol)		ND	10	0.33	
Bis(2-chloroethyl) Ether	ND	10	0.33	3 &/or 4-Methylphenol (m &/or p	-Cresol)	ND	10	0.33	
Bis(2-chloroisopropyl)Ether	ND	10	0.33	Naphthalene		0 62	10	0.33	
Bis(2-ethylhexyl) Phthalate	ND	10	0.33	2-Nitroaniline		ND	50	16	
4-Bromophenyl Phenyl Ether	ND	10	0.33	3-Nitroaniline		ND	50	1.6	
Butylbenzyl Phthalate	ND	10	0.33	4-Nitroaniline		ND	50	1.6	
4-Chloroanaline	ND	20	0.66	2-Nitrophenol		ND	50	1.6	
4-Chloro-3-methylphenol	ND	10	0 33	4-Nitrophenol		ND	50	1.6	
2-Chloronaphthalene	ND	10	0.33	Nitrobenzene		ND	10	0.33	
2-Chlorophenol	ND	10	0.33	N-Nitrosodiphenylamine		ND	10	0.33	
4-Chlorophenyl Phenyl Ether	ND	10	0 33	N-Nitrosodi-n-propylamine		ND	10	0 33	
Chrysene	ND	10	0.33	Pentachlorophenol		ND	50	1.6	
Dibenzo(a,h)anthracene	ND	10	0.33	Phenanthrene		ND	10	0.33	
Dibenzofuran	ND	10	0.33	Phenol		ND	10	0.33	
Di-n-butyl Phthalate	ND	10	0.33	Pyrene		ND	10	0.33	
1,2-Dichlorobenzene	ND	10	0.33	1,2,4-Trichlorobenzene		ND	10	0.33	

10

10

20

10

10

10

10

50

50

10

10

0.33

0.33

0.66

0.33

0.33

0.33

0.33

1.6

1.6

0.33

0.33

2,4,5-Trichlorophenol

2,4,6-Trichlorophenol

Comments:

Phenol-d5

2-Fluorophenol

Nitrobenzene-d5

2-Fluorobiphenyl

p-Terphenyl-d14

2,4,6-Tribromophenol

ND

ND

ND

ND

ND

ND

ND

ND

ND

ND

ND

1,3-Dichlorobenzene

1,4-Dichlorobenzene

2,4-Dichlorophenol

2,4-Dimethylphenol

Dimethyl Phthalate

2,4-Dinitrophenol

2,4-Dinitrotoluene

2,6-Dinitrotoluene

4,6-Dinitro-2-methylphenol

Diethyl Phthalate

3,3-Dichlorobenzidine

Surrogate Recoveries (%)

ND

ND

10

10

101

82

106

108

86

99

0.33

0.33

^{*}water samples are reported in ug/L, soil and sludge samples in mg/kg, wipes in ug/wipe and all TCLP / STLC / SPLP extracts in ug/L

ND means not detected above the reporting limit; N/A means analyte not applicable to this analysis.

^{*} surrogate diluted out of range

h) lighter than water immiscible sheen is present; i)liquid sample that contains greater than ~5 vol. % sediment; j) sample diluted due to high organic content

Cambria Environmental Technology	Client Project ID: #458-1705-4;	Date Sampled: 10/09/01		
1144 65th Street, Suite C	Embarcadero Ave	Date Received: 10/11/01		
Oakland, CA 94608	Client Contact: Bob Schultz	Date Extracted: 10/11/01		
	Client P.O:	Date Analyzed: 10/11-10/12/01		
EPA methods 6010/200.7; 7470/7471/245.1/24	CAM / CCR 17 Metals* 5.5 (Hg); 7060/206.2 (As); 7740/270 2 (Se); 7	841/279 2 (T1); 239.2 (Pb, w); 204.2 (Sb,w)		
Lab ID 805		Paparting Limit		

	Lab ID	80573					
	Client ID	MW-1-8.3			I	Reporting Lim	it
	Matrix	S			S	w	STLC,
	Extraction®	TTLC			TTLC	TTLC	TCLP
Compound			Concen	tration*	mg/kg	mg/L	mg/L
Antimony (Sb)		ND			2.5	0.006	0.05
Arsenic (As)		ND			2.5	0.005	0.25
Barium (Ba)		110			2.5	0.05	0.05
Beryllium (Be)		ND			0.5	0.004	0.01
Cadmium (Cd)		ND			0.5	0.005	0.01
Chromium (Cr)		56			0.5	0 02	0.05
Cobalt (Co)		12			2.0	0.05	0.05
Copper (Cu)		15			2.0	0.05	0.05
Lead (Pb)		2.6	1"		 3.0	0.005	0.2
Mercury (Hg)		ND			0 06	0.0008	0.005
Molybdenum (Mo)		ND			2.5	0.05	0.05
Nickel (Ni)		64			 2.0	0.05	0.05
Selenium (Se)		ND		·····	 2.5	0.005	0.25
Silver (Ag)		ND			 1.0	0.01	0.05
Thallium (Tl)		ND			 2.5	0.005	0.5
Vanadium (V)		33			 2.0	0.05	0.05
Zinc (Zn)		29			1.0	0.05	0.05
% Recovery Surrogat	e	90					***
Comments							

^{*} water samples are reported in mg/L, soil and sludge samples in mg/kg, wipes in ug/wipe and all TCLP / STLC / SPLP extracts in mg/L ND means not detected above the reporting limit; N/A means surrogate not applicable to this analysis

[°] EPA extraction methods 1311(TCLP), 3010/3020(water, TTLC), 3040(organic matrices, TTLC), 3050(solids, TTLC); STLC - CA Title 22

[®] DISTLC extractions are performed using STLC methodology except that deionized water is substituted for citric acid buffer as the extraction fluid. DISTLC results are not applicable to STLC regulatory limits.

surrogate diluted out of range

[&]amp; reporting limit raised due to matrix interference

i) liquid sample that contains greater than ~2 vol. % sediment; this sediment is extracted with the liquid, in accordance with EPA methodologies and can significantly effect reported metal concentrations.

110 2nd Ave. South, #D7, Pacheco, CA 94553-5560 Telephone: 925-798-1620 Fax: 925-798-1622 http://www.mccampbell.com E-mail: main@mccampbell.com

QC REPORT

EPA 8015m + 8020

Date: 10/11/01					Matrix:	Soil	
_		Concen	%Rec				
Compound	Sample	MS	MSD	Amount Spiked	MS	MSD	RPD
SampleID: 100901	Extraction	: EPA 5	030		Instrumen	it: G	C-3
Surrogate1	ND	99.000	103.000	100.00	99	103	4.0
Xylenes	ND	0.332	0.354	0.30	111	118	6.4
Ethylbenzene	ND	0.109	0.111	0.10	109	111	1.8
Toluene	ND	0.106	0.115	0.10	106	115	8.1
Benzene	ND	0.101	0.109	0.10	101	109	7.6
MTBE	ND	0.112	0.111	0.10	112	111	0.9
TPH (gas)	ND	0 836	0.841	1.00	84	84	0.6
SampleID: 101501	Extraction: EPA 3550 Instrument GC-11 A						
Surrogate1	ND	103.000	102.000	100.00	103	102	1.0
TPH (diesel)	ND	140.000	152.000	150.00	93	101	8.2

% Re covery =
$$\frac{(MS-Sample)}{AmountSpiked} \cdot 100$$

RPD = $\frac{(MS-MSD)}{(MS+MSD)} \cdot 2100$

QC REPORT

VOCs (EPA 8240/8260)

Date: 10/11/01	Extraction	n: N/A			Matrix:	Soil		
		Concen	tration:	ug/kg	%Rec			
Compound	Sample	MS	MSD	Amount Spiked	MS	MSD	RPD	
SampleID: 101101					Instrumer	nt: GC	-10	
Surrogate	ND	108.0	108.0	100.00	108	108	0.0	
tert-Amyl Methyl Ether	ND	48.5	48.5	50.00	97	97	0.0	
Methyl tert-Butyl Ether	ND	50.0	50.0	50.00	100	100	0.0	
Ethyl tert-Butyl Ether	ND	52.0	53.5	50.00	104	107	2.8	
Di-isopropyl Ether	ND	49.5	51.0	50.00	99	102	3.0	
Toluene	ND	45.5	45.5	50.00	91	91	0.0	
Benzene	ND	45.0	46.0	50.00	90	92	2.2	
Chlorobenzene	ND	48.0	49.0	50.00	96	98	2.1	
Trichloroethene	ND	40.5	42.0	50.00	81	84	3.6	
1,1-Dichloroethene	ND	56.5	58.5	50.00	113	117	3.5	

% Re covery =
$$\frac{(MS-Somple)}{AmountSpiked} \cdot 100$$

RPD= $\frac{(MS-MSD)}{(MS+MSD)} \cdot 2100$

110 2nd Ave. South, #D7, Pacheco, CA 94553-5560
Telephone: 925-798-1620 Fax: 925-798-1622
http://www.mccampbell.com E-mail: main@mccampbell.com

QC REPORT

SVOCs (EPA 8270/625/525)

Date: 11/07/01	Extractio	n: N/A			Matrix:	Soil	
		Concen	tration:	mg/kg	%Rec	overy	
Compound	Sample	MS	MSD	Amount Spiked	MS	MSD	RPD
SampleID: 100701					Instrumer	nt: G	C-8
Surrogate1	ND	1000.0	1050.0	1000.00	100	105	4.9
Pyrene	ND	870.0	910.0	1000.00	87	91	4.5
Pentachlorophenol	ND	1210.0	1210.0	2000.00	61	61	0.0
2,4-Dinitrotoluene	ND	1290.0	1290.0	1000.00	129	129	0.0
4-Nitrophenol	ND	1540.0	1530.0	2000.00	77	77	0.7
Acenaphtene	ND	900.0	940.0	1000.00	90	94	4.3
4-Chloro-3-metylphenol	ND	1860.0	1890.0	2000.00	93	95	1.6
1,2,4-trichlorobenzene	ND	960.0	990.0	1000.00	96	99	3.1
N-nitroso-di-n-propyl	ND	1020.0	1040.0	1000.00	102	104	1.9
1,4-Dichlorobenzene	ND	900.0	930.0	1000.00	90	93	3.3
2-Chlorophenol	ND	1750.0	1740.0	2000.00	88	87	0.6
Phenol	ND	1580.0	1590.0	2000.00	79	80	0.6

 $\% \text{ Re covery} = \frac{\text{(}MS - Sample\text{)}}{AmountSpiked} \cdot 100$

 $RPD = \frac{(MS - MSD)}{(MS + MSD)} \cdot 2400$

RPD means Relative Percent Deviation

QC REPORT

CAM 17

Date: 10/11/01	Extractio	n: TTLC	;		Matrix:	Soil	
_		Concen	tration:	mg/kg	%Rec	overy	
Compound	/Sampia Nis Nish i			Amount Spiked	MS	MSD	RPD
SampleID: 101501					Instrumer	<u>nt:</u> P-1	AA
Beryllium	ND	5.3	5.3	5.00	106	107	0.8
Selenium	ND	9.8	9.7	10.00	98	97	1.0
Molybdenum	ND	5.4	5.5	5.00	109	109	0.2
Silver	ND	0.51	0.51	0.50	102	103	0.8
Thallium	ND	9.9	11.0	10.00	99	110	10.5
Barium	ND	5.0	4.9	5.00	101	99	2.0
Nickel	ND	5.3	5.2	5.00	106	104	1.4
Arsenic	ND	9.0	9.3	10.00	90	93	3.3
Vanadium	ND	5.1	5.2	5.00	102	104	1.0
Surrogate1	ND	90.8	89.8	100.00	91	90	1.1
Zinc	ND	5.4	5.4	5.00	107	108	0.4
Copper	ND	4.8	4.8	5.00	96	96	0.3
Antimony	ND	5.3	5.3	5.00	107	105	1.1
Lead	ND	5.3	5.2	5.00	106	104	2.1
Cadmium	ND	5.5	5.4	5.00	110	108	1.1
Cobalt	ND	5.2	5.2	5.00	103	104	0.9
Mercury	ND	0.24	0.26	0.25	96	105	8.5
Chromium	ND	5.6	5.5	5.00	111	110	1.3

% Re covery = $\frac{(MS-Sample)}{AmountSpiked} \cdot 100$

 $RPD = \frac{(MS - MSD)}{(MS + MSD)} \cdot 2 \cdot 100$

File : D:\HPCHEM\2\DATA\10160110.D

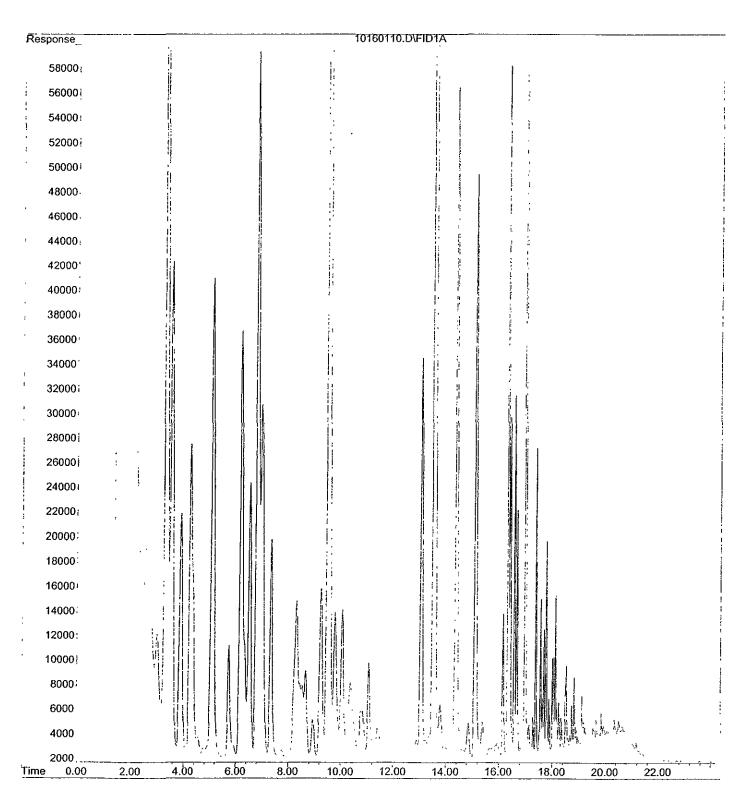
Operator

Acquired : 16 Oct 2001 3:22 pm using AcqMethod GC7B.M

Instrument : GC-7 Sample Name: 5000ngGAS

Misc Info

Vial Number: 10



File : D:\HPCHEM\2\DATA\10160108.D

Operator

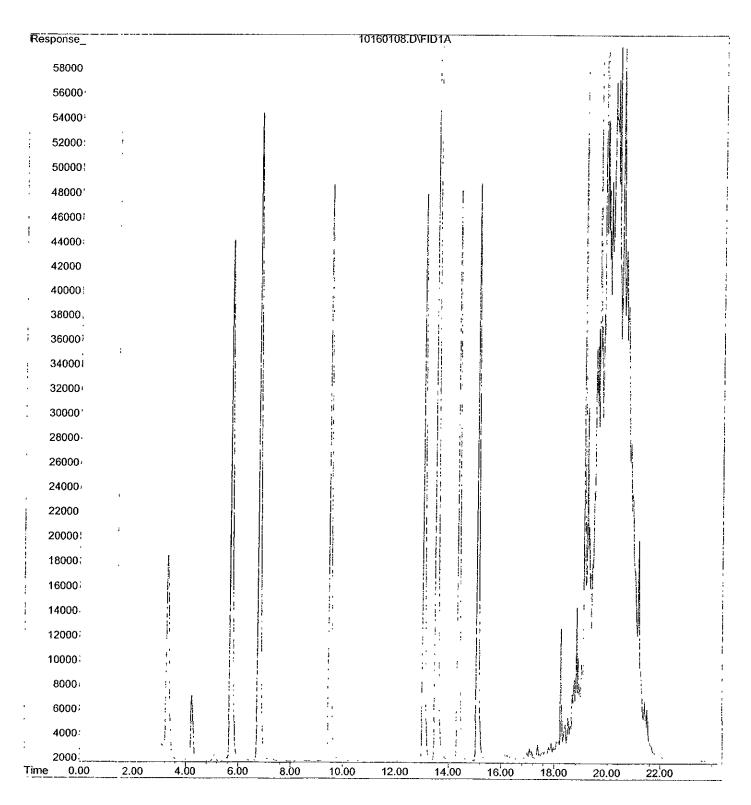
Acquired : 16 Oct 2001 2:22 pm using AcqMethod GC7B.M

Instrument : GC-7

Sample Name: (100ngBTEX

Misc Info

Vial Number: 8



IntFile : EVENTS.E

Quant Time: Nov 20 16:21 2001 Quant Results File: GC2BNEW.RES

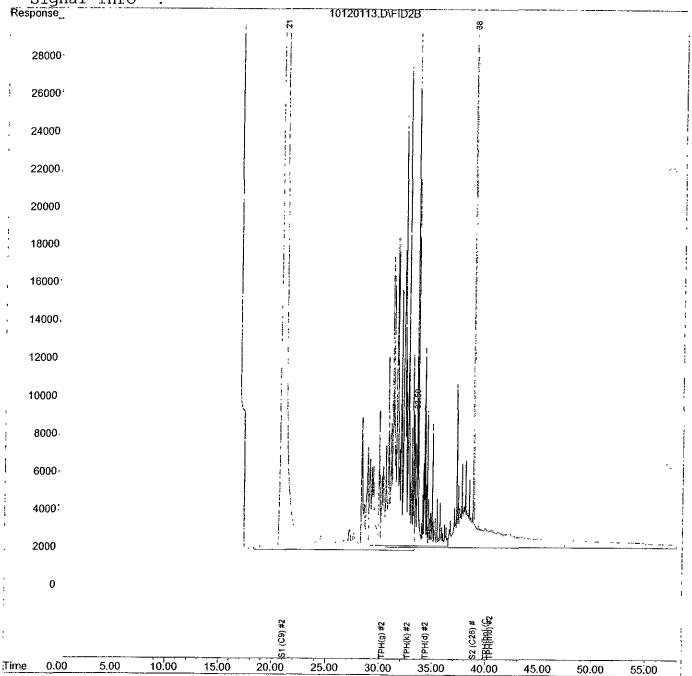
Quant Method : C:\HPCHEM\3\METHODS\GC2BNEW.M (Chemstation Integrator)

Title : DIESEL 1

Last Update : Tue Mar 27 12:53:30 2001 Response via : Multiple Level Calibration

DataAcq Meth : GC2ANEW.M

Volume Inj. : Signal Phase : Signal Info :



10120113.D GC2BNEW.M

Tue Nov 20 16:21:02 2001

BTEX3_DIESELBage 2

"MO/DIESEL C-TANDARI

Data File : D:\HPCHEM\1\DATAA\10110114.D

Acq On : 11 Oct 2001 10:14 pm

Vial: 7
Operator: Thu
Inst : GC-11
Multiplr: 2.0

Sample : 80574S Misc : D,MO,WSG IntFile : EVENTS.E

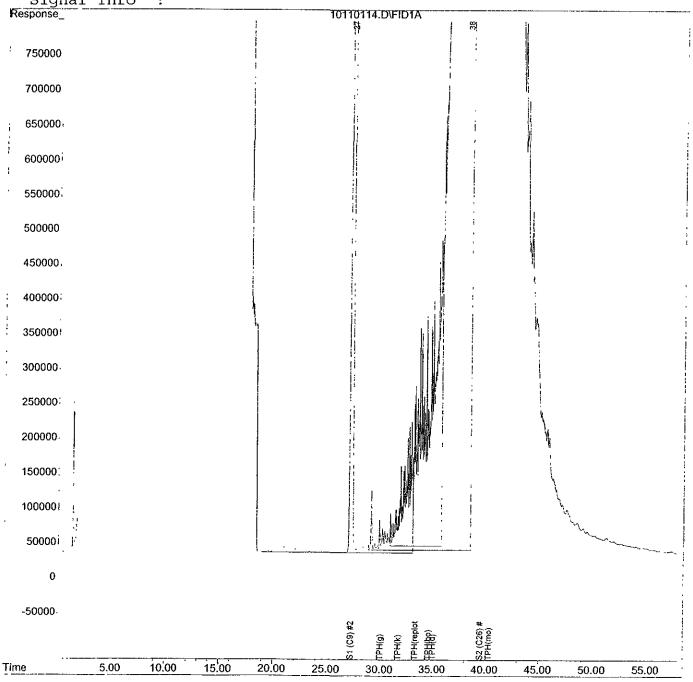
Quant Time: Nov 20 15:49 2001 Quant Results File: GC11A.RES

Quant Method : C:\HPCHEM\1\METHODS\GC11A.M (Chemstation Integrator)

Title : GC-11A

Last Update : Mon Nov 19 11:45:59 2001 Response via : Multiple Level Calibration

DataAcq Meth : GC11A.M



IntFile : EVENTS.E

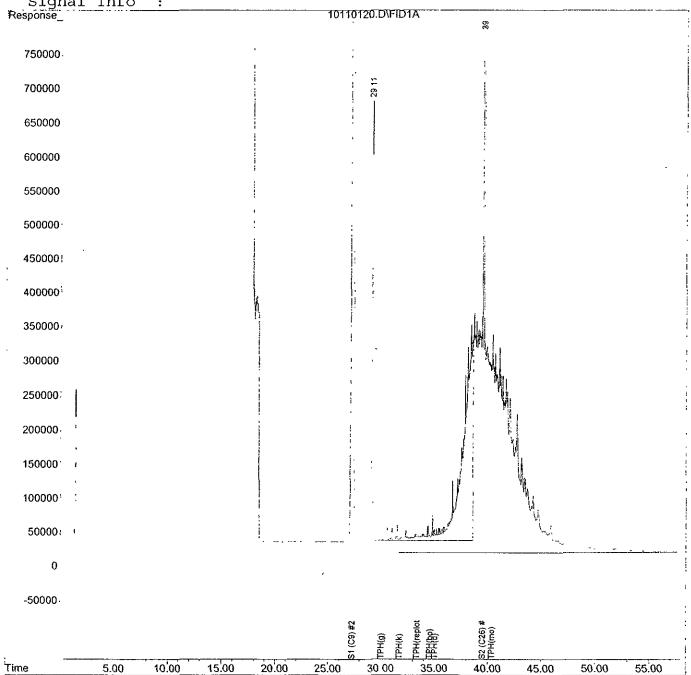
Quant Time: Nov 20 15:49 2001 Quant Results File: GC11A.RES

Quant Method: C:\HPCHEM\1\METHODS\GC11A.M (Chemstation Integrator)

Title : GC-11A

Last Update : Mon Nov 19 11:45:59 2001 Response via : Multiple Level Calibration

DataAcq Meth : GC11A.M



IntFile : EVENTS.E

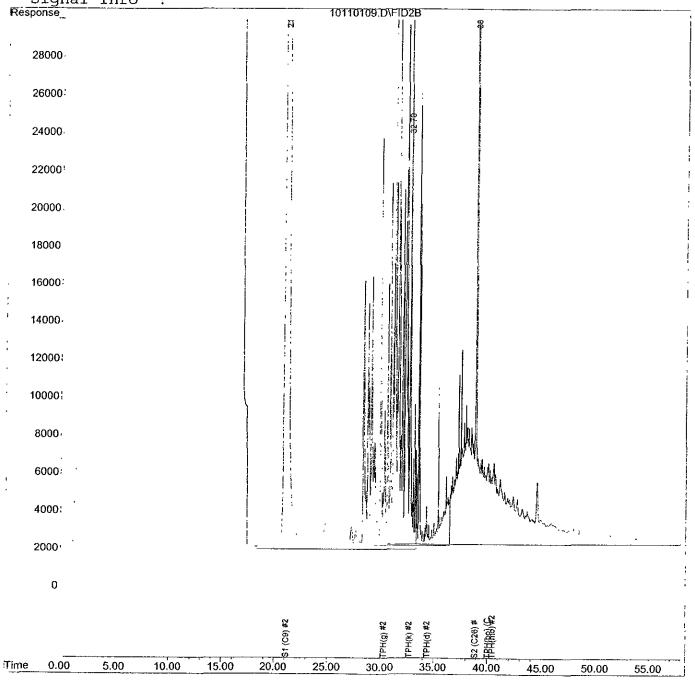
Quant Time: Nov 20 16:20 2001 Quant Results File: GC2BNEW.RES

Quant Method : C:\HPCHEM\3\METHODS\GC2BNEW.M (Chemstation Integrator)

Title : DIESEL 1

Last Update : Tue Mar 27 12:53:30 2001 Response via : Multiple Level Calibration

DataAcq Meth : GC2ANEW.M



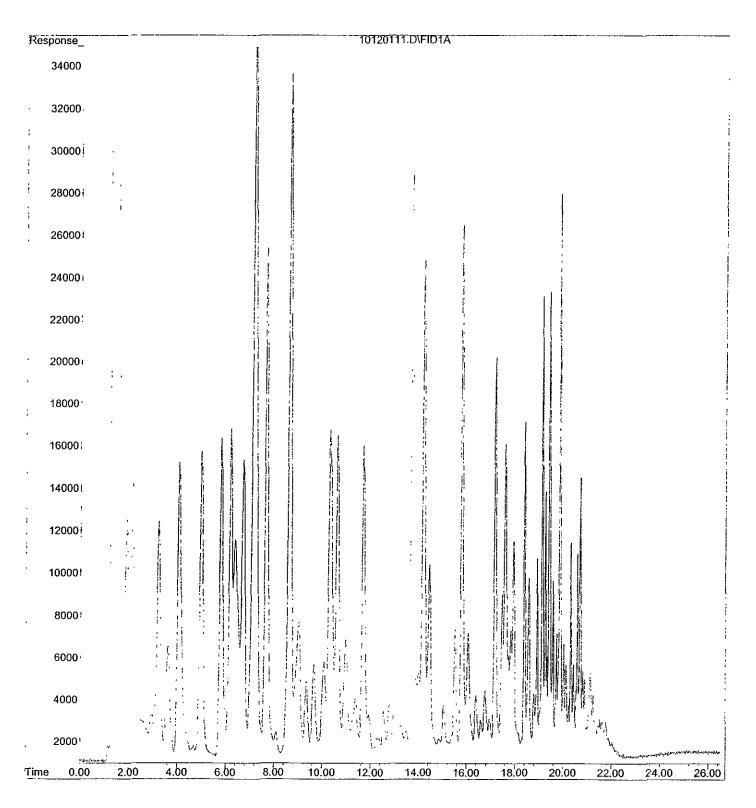
File : D:\HPCHEM\2\DATA\10120111.D

Operator

Acquired : 12 Oct 2001 9:22 pm using AcqMethod GC12A.M

Instrument : GC-12 Sample Name: 80573rrS+AF

Misc Info : \(\bar{}\)
Vial Number: 11



File : D:\HPCHEM\2\DATA\10120107.D

Operator

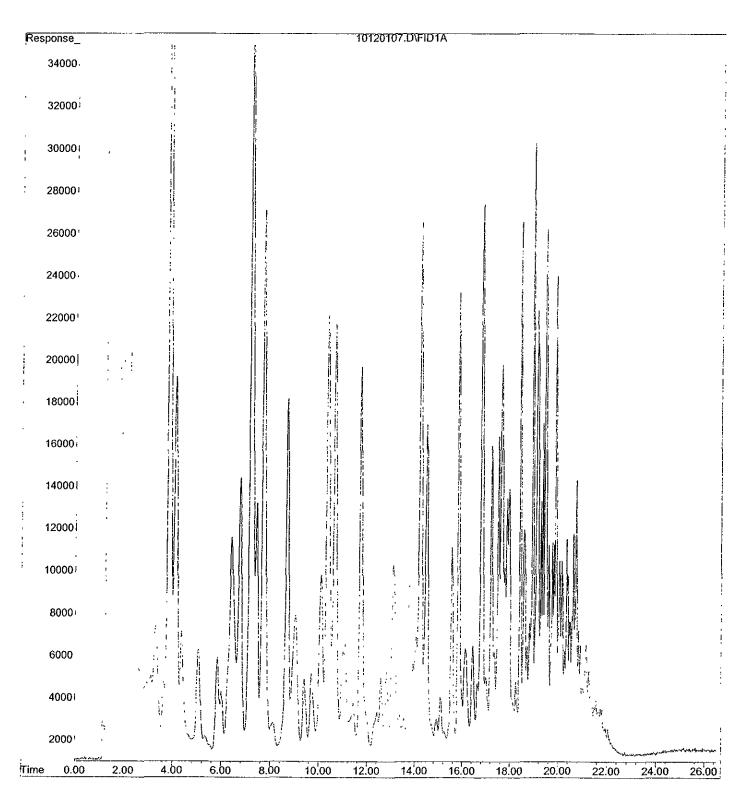
7:08 pm using AcqMethod GC12A.M Acquired : 12 Oct 2001

Instrument :

GC-12

Sample Name (80574rrS+AF)

Misc Info Vial Number: 7



: D:\HPCHEM\2\DATA\10160122.D File

Operator

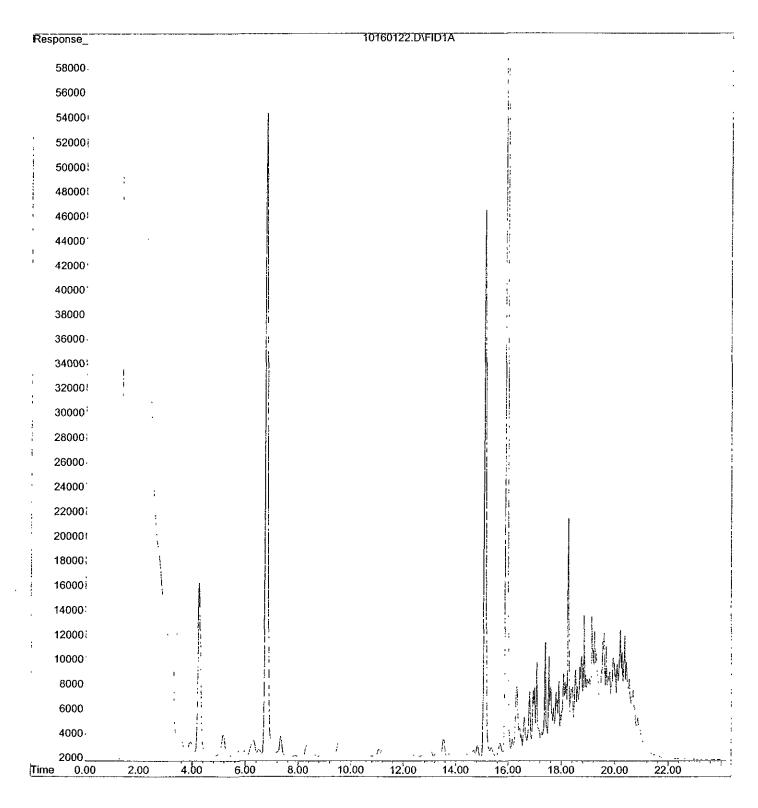
9:21 pm using AcqMethod GC7B.M Acquired : 16 Oct 2001

Instrument :

GC-7 Sample Name: (80576reSrr

Misc Info

Vial Number: 22



File : D:\HPCHEM\2\DATA\10110116.D

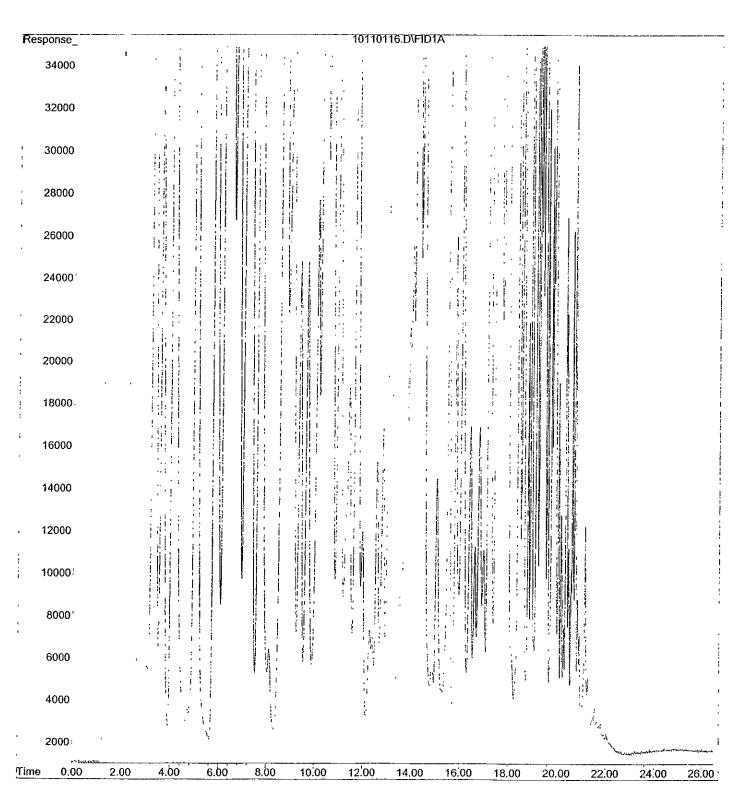
Operator

Acquired : 12 Oct 2001 1:20 am using AcqMethod GC12A.M

Instrument: GC-12 Sample Name: 80579S

Misc Info

Vial Number: 16



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	-1705 -	4	P	roject	t Nam	e: <u>{</u>	MBA	ZLADE	20	<u>ئ</u>	نك	8015)/		(552	£	6			510CS	228				7	METALS	-	馬		
Project Location:	1275	EMBAR	ecy Dar	٥								ļ t		ease	Pons	802		<u> </u>	N S	25/1		l	9	31.	34/5	#			
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				ers	Type Containers		T				T	as S	TPH as Diesel (8015)	Total Petroleum Oil & Grease (5520 E&F/B&F)	Total Petroleum Hydrocarbons (418.1)	BTEX ONLY (EPA 602 / 8020)	080	EPA 608 / 8080 PCB's ONLY	EPA 624 / 8240 / 8260 EPA 626 / 8270 Ad	PAH's / PNA's by EPA 625 / 8270 / 8310	tais	als	Lead (7240/7421/239.2/6010)	FOC, Bulk DensHy/Moistor	7				
SAMPLE ID	LOCATION		20,	Containers) tio	1 (\cdot					втех & трн	Dies	trole	Total Petroleum	Ž	EPA 608 / 8080	3 / 8	80 6	Z	CAM-17 Metals	LUFT 5 Metals	407	Ŋ	4	\$	37		
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MW-2-5.0		10-7-01	12:35	1	35	-	χ		×			X				1,			VI	7			_	\mathbf{X}'	7	X	1-		
MW-2-18.0		10-7-01	12:40	1	55		<		χ			1			· ·	-	1.7.			`\				<u>د -</u> ۱-		7 -		8057	4
MW-3-5.0		10-7-4	9:25	1	55	,			X			Ī	_						\propto	7				X		X	-		
MW-3-6.5		10-9-01	9:40	1	55				X						_											_	-	8057	Ç
MW-3-12.5		10-9-01	10:45		55				X]												_		7	f	80576	R
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MW-4-10-5		10-9-01	11:45		55)	4		X									-				_		_		_		80577	7
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Cambria Environmental Technology	Client Project ID: #458-1705;	Date Sampled: 10/19/01
1144 65 th Street, Suite C	Embarcadero Cove	Date Received: 10/23/01
Oakland, CA 94608	Client Contact: Bob Schultz	Date Extracted: 10/23/01
	Client P.O:	Date Analyzed: 10/23/01

10/30/01

Dear Bob:

Enclosed are:

- 1). the results of 4 samples from your #458-1705; Embarcadero Cove project,
- 2). a QC report for the above samples
- 3). a copy of the chain of custody, and
- 4). a bill for analytical services.

All analyses were completed satisfactorily and all QC samples were found to be within our control limits. If you have any questions please contact me. McCampbell Analytical Laboratories strives for excellence in quality, service and cost. Thank you for your business and I look forward to working with you again.

Edward Hamilton, Lab Director

110 2nd Avenue South, #D7, Pacheco, CA 94553-5560 Telephone: 925-798-1620 Fax: 925-798-1622 http://www.mccampbell.com E-mail: main@mccampbell.com

Cambria Environmental Technology	Client Project ID: #458-1705;	Date Sampled: 10/19/01
1144 65th Street, Suite C	Embarcadero Cove	Date Received: 10/23/01
Oakland, CA 94608	Client Contact: Bob Schultz	Date Extracted: 10/26-10/29/01
	Client P.O:	Date Analyzed: 10/26-10/29/01

Gasoline Range (C6-C12) Volatile Hydrocarbons as Gasoline*, with Methyl tert-Butyl Ether* & BTEX*

EPA methods 5030, modified 8015, and 8020 or 602; California RWQCB (SF Bay Region) method GCFID(5030) Ethyl-% Recovery Lab ID Client 1D Matrix TPH(g)+ **MTBE** Benzene Toluene **Xylenes** benzene Surrogate 81774 MW-1 W 11,000,a 900 300 470 1000 103 81775 MW-2 W ND ND ND ND ND 107 81776 MW-3 W 290,a,i 2.0 6.6 0.54 1.2 102 44,000,a,h 81777 MW-4 W 1900 270 1500 3300 114 Reporting Limit unless W 50 ug/L 5.0 0.5 0.5 0.5 0.5 otherwise stated; ND means not detected above S 1.0 mg/kg 0.05 0.005 0.005 0.005 0.005 the reporting limit

^{*}The following descriptions of the TPH chromatogram are cursory in nature and McCampbell Analytical is not responsible for their interpretation: a) unmodified or weakly modified gasoline is significant; b) heavier gasoline range compounds are significant(aged gasoline?); c) lighter gasoline range compounds (the most mobile fraction) are significant; d) gasoline range compounds having broad chromatographic peaks are significant; biologically altered gasoline?; e) TPH pattern that does not appear to be derived from gasoline (?); f) one to a few isolated peaks present; g) strongly aged gasoline or diesel range compounds are significant; h) lighter than water immiscible sheen is present; i) liquid sample that contains greater than ~5 vol. % sediment; j) no recognizable pattern.



^{*} water and vapor samples are reported in ug/L, wipe samples in ug/wipe, soil and sludge samples in mg/kg, and all TCLP and SPLP extracts in ug/L

^{*} cluttered chromatogram; sample peak coelutes with surrogate peak

Cambria Environmental Technology	Client Project ID: #458-1705;	Date Sampled: 10/19/01
1144 65 th Street, Suite C	Embarcadeто Cove	Date Received: 10/23/01
Oakland, CA 94608	Client Contact: Bob Schultz	Date Extracted: 10/23/01
	Client P.O:	Date Analyzed: 10/24/01

Diesel Range (C10-C23) and Oil-Range (C18+) Extractable Hydrocarbons as Diesel and Motor Oil with Silica Gel Clean-up*

EPA methods modified 8015, and 3550 or 3510; California RWQCB (SF Bay Region) method GCFID(3550) or GCFID(3510)

Lab ID	Client ID	Matrix	TPH(d)⁺	TPH(mo)⁺	% Recovery Surrogate
81774	MW-1	w	3300,d	ND	113
81775	MW-2	w	210,b,g	460	88
81776	MW-3	w	1600,b,g,i	1300	93
81777	MW-4	w	33,000,d,b,g,h	900	93
<u> </u>					
Reporting Limit stated; ND means	unless otherwise not detected above	w	50 ug/L	250 ug/L	
	rting limit	s	1.0 mg/kg	5.0 mg/kg	

^{*}water samples are reported in ug/L, wipe samples in ug/wipe, soil and sludge samples in mg/kg, and all TCLP / STLC / SPLP extracts in ug/L.

^{*}The following descriptions of the TPH chromatogram are cursory in nature and McCampbell Analytical is not responsible for their interpretation: a) unmodified or weakly modified diesel is significant; b) diesel range compounds are significant; no recognizable pattern; c) aged diesel? is significant); d) gasoline range compounds are significant; e) medium boiling point pattern that does not match diesel (?); f) one to a few isolated peaks present; g) oil range compounds are significant; h) lighter than water immiscible sheen is present; i) liquid sample that contains greater than ~5 vol. % sediment.



[&]quot;cluttered chromatogram resulting in coeluted surrogate and sample peaks, or, surrogate peak is on elevated baseline, or; surrogate has been diminished by dilution of original extract.

Cambria Environmental	Technology			(LLD: #430-1703;	ate Samp	oled: 10/19/01			
1144 65 th Street, Suite C		Emba	ircadero	D Cove	ate Rece	ived: 10/23/0	01		
Oakland, CA 94608		Clien	t Conta	cted: 10/23/01					
		Clien	t P.O:	Da	ate Analy	yzed: 10/23-10/24/01			
		Semi-V	olatile	Organics By GC/MS					
EPA method 625 and 3510 or 8	270 and 3550							~~~	
Lab ID				81774					
Client ID				MW-1 W					
Matrix									
Compound	Concentration*	Report	ing Limit S		oncentration	Repor	ting Limit		
Acenaphthene	ND	10	0.33	Di-n-octyl Phthalate		ND	10	0.33	
Acenaphthylene	ND	10	0.33	1,2-Diphenylhydrazine		ND ND	10	0.33	
Anthracene	ND	10	0.33	Fluoranthene		ND ND	10	0.33	
Benzidine	ND	50	1.6	Fluorene		ND	10	0.33	
Benzoic Acid	ND	50	1.6	Hexachlorobenzene		ND	10	0.33	
Benzo(a)anthracene	ND	10	0.33	Hexachlorobutadiene		ND	10	0.33	
Benzo(b)fluoranthene	ND	10	0.33	Hexachlorocyclopentadiene		ND	50	1.6	
Benzo(k)fluoranthene	ND	10	0.33	Hexachloroethane	ND	10	0.33		
Benzo(g,h,i)perylene	ND	10	0.33	Indeno(1,2,3-cd)pyrene	ND	10	0.33		
Benzo(a)pyrene	ND	10	0.33	Isophorone		ND	10	0.33	
Benzyl Alcohol	ND	20	0.66	2-Methylnaphthalene	54	10	0.33		
Bis(2-chloroethoxy)methane	ND	10	0.33	2-Methylphenol (o-Cresol)	ND ND	10	0.33		
Bis(2-chloroethyl) Ether	ND	10	0.33	3 &/or 4-Methylphenol (m &/or p	Cresol	ND	10	0.33	
Bis(2-chloroisopropyl)Ether	ND	10	0.33	Naphthalene	Cicson	66	10	0.33	
Bis(2-ethylhexyl) Phthalate	ND	10	0.33	2-Nitroaniline		ND	50	1.6	
4-Bromophenyl Phenyl Ether	ND	10	0.33	3-Nitroaniline		ND	50	1.6	
Butylbenzyl Phthalate	ND	10	0.33	4-Nitroaniline		ND	50	1.6	
4-Chloroanaline	ND	20	0.66	2-Nitrophenol	···	ND	50	1.6	
4-Chloro-3-methylphenol	ND	10	0.33	4-Nitrophenol		ND	50	1.6	
2-Chloronaphthalene	ND	10	0.33	Nitrobenzene		ND ND	10	0.33	
2-Chlorophenol	ND	10	0.33	N-Nitrosodimethylamine		ND	10	0.33	
4-Chlorophenyl Phenyl Ether	ND	10	0.33	N-Nitrosodiphenylamine		ND	10	0.33	
Chrysene	ND	10	0.33	N-Nitrosodi-n-propylamine	——— !	ND	10	0.33	
Dibenzo(a,h)anthracene	ND	10	0.33	Pentachlorophenol		ND	50	1.6	
Dibenzofuran	ND	10	0.33	Phenanthrene		ND	10	0.33	
Di-n-butyl Phthalate	ND	10	0.33	Phenol		ND	10	0.33	
1,2-Dichlorobenzene	ND	10	0.33	Pyrene		ND	10	0.33	
1,3-Dichlorobenzene	ND	10	0.33	1,2,4-Trichlorobenzene		ND	10	0.33	
1,4-Dichlorobenzene	ND .	10	0.33	2,4,5-Trichlorophenol		ND	10	0.33	
3,3-Dichlorobenzidine	ND	20	0.66	2,4,6-Trichlorophenol		ND	10	0.33	
2,4-Dichlorophenol	ND	10	0.33	Comments.		<u></u>		-	
Diethyl Phthalate	ND	10	0.33	Surrogate .	Recoverie	(%)			
2,4-Dimethylphenol	ND	10	0.33	2-Fluorophenol		1	43		
Dimethyl Phthalate	ND ND	10	0.33	Phenol-d5			43		
		50					66		
	ND 1								
4,6-Dinitro-2-methylphenol	ND ND	L							
	ND ND ND	50 10	1.6	2-Fluorobiphenyl 2,4,6-Tribromophenol			66		

^{*}water samples are reported in ug/L, soil and sludge samples in mg/kg, wipes in ug/wipe and all TCLP / STLC / SPLP extracts in ug/L

ND means not detected above the reporting limit; N/A means analyte not applicable to this analysis.

[&]quot; surrogate diluted out of range

h) lighter than water immiscible sheen is present; i)liquid sample that contains greater than ~5 vol. % sediment; j) sample diluted due to high organic content

ND

ND

ND

ND

ND

ND

ND

ND

ND

ND

ND

ND

ND

ND

ND

ND

ND

ND

ND

ND

ND

20

10

10

10

10

10

10

10

10

10

10

10

20

10

10

10

01

50

50

10

10

0.66

0.33

0.33

0 33

0.33

0.33

0.33

0.33

0.33

0.33

0.33

0.33

0.66

0.33

0.33

0.33

0.33

1.6

1.6

0.33

033

2-Nitrophenol

4-Nitrophenol

Nitrobenzene

Phenanthrene

Phenol

Pyrene

Comments

Phenol-d5

2-Fluorophenol

Nitrobenzene-d5

2-Fluorobiphenyl

p-Terphenyl-d14

2,4,6-Tribromophenol

N-Nitrosodimethylamine

N-Nitrosodiphenylamine

1,2,4-Trichlorobenzene

2,4,5-Trichlorophenol

2,4,6-Trichlorophenol

Pentachlorophenol

N-Nitrosodi-n-propylamine

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Cambria Environmental	Technology	Clien	t Projec	et ID: #458-1705; Date San	npled: 10/19/0	pled: 10/19/01			
1144 65 th Street, Suite C		Emba	rcadero	Date Re	ceived: 10/23/	01			
Oakland, CA 94608		Client	t Conta	ct: Bob Schultz Date Ex	racted: 10/23/	acted: 10/23/01			
		Client	P.O:	Date An	alyzed: 10/23-	10/24	/01		
EPA method 625 and 3510 or 8		Semi-V	⁷ olatile	Organics By GC/MS		 :			
Lab ID				81775					
Client ID				MW-2					
Matrix				W	· · · · · · · · · · · · · · · · · · ·				
Compound	Concentration*	Report	ng Limit	Compound	oncentration	Repor W	ting Limit		
Acenaphthene	ND	10	0.33	Di-n-octyl Phthalate	ND	10	0.33		
Acenaphthylene	ND	10	0.33	1,2-Diphenylhydrazine	ND	10	0.33		
Anthracene	ND	10	0.33	Fluoranthene	ND	10	0.33		
Benzidine	ND	50	1.6	Fluorene	ND	10	0.33		
Benzoic Acid	ND	50	1.6	Hexachlorobenzene	ND	10	0.33		
Benzo(a)anthracene	ND	10	0.33	Hexachlorobutadiene	ND	10	0.33		
Benzo(b)fluoranthene	ND	10	0 33	Hexachlorocyclopentadiene	ND	50	1.6		
Benzo(k)fluoranthene	ND	10	0.33	Hexachloroethane	ND	10	0.33		
Benzo(g,h,i)perylene	ND	10	0.33	Indeno(1,2,3-cd)pyrene	ND	10	0.33		
Benzo(a)pyrene	ND	10	0 33	Isophorone	ND	10	0.33		
Benzyl Alcohol	ND	20	0.66	2-Methylnaphthalene	ND	10	0.33		
Bis(2-chloroethoxy)methane	ND	10	0.33	2-Methylphenol (o-Cresol)	ND	10	0.33		
Bis(2-chloroethyl) Ether	NĐ	10	0.33	3 &/or 4-Methylphenol (m &/or p-Cresol)		10	0.33		
Bis(2-chloroisopropyl)Ether	ND	10	0.33	Naphthalene	ND	10	0.33		
Bis(2-ethylhexyl) Phthalate	ND	10	0.33	2-Nitroaniline	ND	50	1.6		
4-Bromophenyl Phenyl Ether	ND	10	0.33	3-Nitroaniline	ND	50	1.6		
Butylbenzyl Phthalate	ND	10	0.33	4-Nitroaniline	ND	50	1.6		

4-Chloroanaline

2-Chlorophenol

Chrysene

Dibenzofuran

4-Chloro-3-methylphenol

4-Chlorophenyl Phenyl Ether

Dibenzo(a,h)anthracene

Di-n-butyl Phthalate

1,2-Dichlorobenzene

1,3-Dichlorobenzene

1,4-Dichlorobenzene

2,4-Dichlorophenol

2,4-Dimethylphenol

Dimethyl Phthalate

2,4-Dinitrophenol

2.4-Dinitrotoluene

2,6-Dinitrotolucne

4,6-Dinitro-2-methylphenol

Diethyl Phthalate

3,3-Dichlorobenzidine

2-Chloronaphthalene

h) lighter than water immiscible sheen is present; i)liquid sample that contains greater than ~5 vol. % sediment; j) sample diluted due to high organic content

DHS Certification No. 1644

Edward Hamilton, Lab Director

Surrogate Recoveries (%)

ND

ND

ND

ND

ND

ND

ND

ND

ND

ND

ND

 $\overline{\mathsf{ND}}$

ND

50

50

10

10

10

10

50

10

10

10

10

10

10

39

67

66

74

1.6

1.6

0.33

0.33

0.33

0.33

1.6

0.33

0.33

0.33

0.33

0.33

0.33

^{*}water samples are reported in ug/L, soil and sludge samples in mg/kg, wipes in ug/wipe and all TCLP / STLC / SPLP extracts in ug/L ND means not detected above the reporting limit; N/A means analyte not applicable to this analysis.

^{*} surrogate diluted out of range

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Cambria Environmental	Technology			R ID: #458-1705;	ate Samp	oled: 10/19/0	1		
1144 65 th Street, Suite C		Emba	rcadero) Cove	ate Rece	ived: 10/23/6	01		
Oakland, CA 94608		Client	Conta	ct: Bob Schultz D	ate Extra	cted: 10/23/01			
		Client	P.O:	D	ate Analy	yzed: 10/23-10/24/01			
		Semi-V	olatile	Organics By GC/MS					
EPA method 625 and 3510 or 8	3270 and 3550								
Lab ID				81776					
Client ID	l			MW-3		·			
Matrix				W					
		Reporti	ng Limit	_			Report	ting Limit	
Compound	Concentration*	w	S	Compound		oncentration	W	S	
Acenaphthene	ND<25	10	0.33	Di-n-octyl Phthalate		ND<25	10	0.33	
Acenaphthylene	ND<25	10	0.33	1,2-Diphenylhydrazine		ND<25	10	0.33	
Anthracene	ND<25	10	0.33	Fluoranthene		ND<25	10	0.33	
Benzidine	ND<125	50	1.6	Fluorene		ND<25	10	0.33	
Benzoic Acid	ND<125	50	16	Hexachlorobenzene		ND<25	10	0.33	
Benzo(a)anthracene	ND<25	10	0.33	Hexachlorobutadiene		ND<25	10	0.33	
Benzo(b)fluoranthene	ND<25	10	0.33	Hexachlorocyclopentadiene	ND<125	50	1.6		
Benzo(k)fluoranthene	ND<25	10	0 33	Hexachloroethane	ND<25	10	0.33		
Benzo(g,h,i)perylene	ND<25	10	0.33	Indeno(1,2,3-cd)pyrene		ND<25	10	0.33	
Benzo(a)pyrene	ND<25	10	0.33	Isophorone	ND<25	10	0.33		
Benzyl Alcohol	ND<50	20	0.66	2-Methylnaphthalene	670	10	0.33		
Bis(2-chloroethoxy)methane	ND<25	10	0.33	2-Methylphenol (o-Cresol)		ND<25	10	0.33	
Bis(2-chloroethyl) Ether	ND<25	10	0.33	3 &/or 4-Methylphenol (m &/or p	-Cresol)	ND<25	10	0.33	
Bis(2-chloroisopropyl)Ether	ND<25	10	0.33	Naphthalene	/ (10301)	420	10	0.33	
Bis(2-ethylhexyl) Phthalate	ND<25	10	0.33	2-Nitroaniline		ND<125	50	1.6	
4-Bromophenyl Phenyl Ether	ND<25	10	0.33	3-Nitroaniline		ND<125	50	1.6	
Butylbenzyl Phthalate	ND<25	10	0 33	4-Nitroaniline		ND<125	50	1.6	
4-Chloroanaline	ND<50	20	0.66	2-Nitrophenol		ND<125	50	16	
4-Chloro-3-methylphenol	ND<25	10	0.33	4-Nitrophenol		ND<125	50	1.6	
2-Chloronaphthalene	ND<25	10	0.33	Nitrobenzene		ND<25	10	0.33	
2-Chlorophenol	ND<25	10	0.33	N-Nitrosodimethylamine		ND<25	10	0.33	
4-Chlorophenyl Phenyl Ether	ND<25	10	0.33	N-Nitrosodiphenylamine		ND<25	10	0.33	
Chrysene	ND<25	10	0 33	N-Nitrosodi-n-propylamine		ND<25	10	0.33	
Dibenzo(a,h)anthracene	ND<25	10	0.33	Pentachlorophenol		ND<125	50	1.6	
Dibenzofuran	ND<25	10	0.33	Phenanthrene		ND<25	10	0.33	
Di-n-butyl Phthalate	ND<25	10	0.33	Phenol		ND<25	10	0.33	
1,2-Dichlorobenzene	ND<25	10	0.33	Pyrene		ND<25	10	0.33	
1,3-Dichlorobenzene	ND<25	10	0.33	1,2,4-Trichlorobenzene		ND<25	10	0.33	
1,4-Dichlorobenzene	ND<25	10	0.33	2,4,5-Trichlorophenol		ND<25	10	0.33	
3,3-Dichlorobenzidine	ND<50	20	0.66	2,4,6-Trichlorophenol		ND<25	10	0.33	
2,4-Dichlorophenol	ND<25	10	0.33	Comments: i	^		لـــــــــــــــــــــــــــــــــــــ		
Diethyl Phthalate	ND<25	10	0.33	Surrogate	Recoverie	s (%)			
2,4-Dimethylphenol	ND<25	10	0.33	2-Fluorophenol		- (/-/	32		
Dimethyl Phthalate	ND<25	10	0.33	Phenol-d5		 	#		
4,6-Dinitro-2-methylphenol	ND<125	50	1.6	Nitrobenzene-d5	······································		60		
2,4-Dinitrophenol	ND<125	50	1.6	2-Fluorobiphenyl			59		
2,4-Dinitrotoluene	ND<25	10	0.33	2,4,6-Tribromophenol			67		
2,6-Dinitrotoluene	ND<25	10	0.33	p-Terphenyl-d14		 	49		
-11-1111000100110		لــنــا		h 101kmon), 011			77		

^{*}water samples are reported in ug/L, soil and sludge samples in mg/kg, wipes in ug/wipe and all TCLP / STLC / SPLP extracts in ug/L ND means not detected above the reporting limit; N/A means analyte not applicable to this analysis.

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___Edward Hamilton, Lab Director

surrogate diluted out of range

h) lighter than water immiscible sheen is present; i)liquid sample that contains greater than ~5 vol. % sediment; j) sample diluted due to high organic content

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Cambria Environmental	Technology			ELD: #438-1703;	ate Samp	oled: 10/19/0)1				
1144 65 th Street, Suite C		Emba	rcadere) Cove	ate Rece	ived: 10/23/	01				
Oakland, CA 94608		Client	Conta	ct: Bob Schultz Da	ate Extra	cted: 10/23/	cted: 10/23/01				
		Client	P.O:	Da	ate Analy	llyzed: 10/23-10/24/01					
		Semi-V	olatile	Organics By GC/MS							
EPA method 625 and 3510 or 8	270 and 3550				· ;•						
Lab ID	<u> </u>			81777							
Chent ID				MW-4							
Matrix				W							
		Reporti	ng Limit				Repor	ung Limit			
Compound	Concentration*	W	S	Compound		oncentration	w	S			
Acenaphthene	ND<50	10	0.33	Di-n-octyl Phthalate		ND<50	10	0.33			
Acenaphthylene	ND<50	10	0.33	1,2-Diphenylhydrazine		ND<50	10	0.33			
Anthracene	ND<50	10	0.33	Fluoranthene		ND<50	10	0.33			
Benzidine	ND<250	50	16	Fluorene		ND<50	10	033			
Benzoic Acid	ND<250	50	1.6	Hexachlorobenzene		ND<50	10	0.33			
Benzo(a)anthracene	ND<50	10	0.33	Hexachlorobutadiene		ND<50	10	0.33			
Benzo(b)fluoranthene	ND<50	10	0.33	Hexachlorocyclopentadiene		ND<250	50	<u> </u>			
Benzo(k)fluoranthene	ND<50	10	0.33	Hexachloroethane		ND<250		1.6			
Benzo(g,h,ı)perylene	ND<50	10	0.33	Indeno(1,2,3-cd)pyrene		ND<50	10 10	0.33			
	ND<50	10	0.33	Isophorone				i			
Benzo(a)pyrene Benzyl Alcohol	ND<100	20	0.55	2-Methylnaphthalene		ND<50 ND<50	10	0 33			
		10	I——				10	0.33			
Bis(2-chloroethoxy)methane	ND<50 ND<50	10	0.33	2-Methylphenol (o-Cresol)	Coursell)	ND<50	10	0.33			
Bis(2-chloroethyl) Ether	ND<50	10	0.33	3 &/or 4-Methylphenol (m &/or p- Naphthalene	-Cresoi)	ND<50	10	0.33			
Bis(2-chloroisopropyl)Ether Bis(2-ethylhexyl) Phthalate	ND<50	10	0.33	2-Nitroaniline		ND<50	10	0.33			
4-Bromophenyl Phenyl Ether	ND<50	10	0.33	3-Nitroaniline		ND<250	50	1.6			
Butylbenzyl Phthalate	ND<50	10	0.33	4-Nitroaniline		ND<250	50	1.6			
4-Chloroanaline	ND<100	20	0.33	2-Nitrophenol		ND<250 ND<250	50 50	1.6			
	ND<50	10	0 33	4-Nitrophenol				16			
4-Chloro-3-methylphenol	ND<50	10	0.33	Nitrobenzene		ND<250	50	16			
2-Chloronaphthalene 2-Chlorophenol		10	0.33	N-Nitrosodimethylamine		ND<50	10	0.33			
4-Chlorophenyl Phenyl Ether	ND<50	10				ND<50	10	0.33			
	ND<50 ND<50		0.33	N-Nitrosodiphenylamine		ND<50	10	0.33			
Chrysene		10	0.33	N-Nitrosodi-n-propylamine Pentachlorophenol		ND<50	10	0.33			
Dibenzo(a,h)anthracene	ND<50	10				ND<250	50	1.6			
Dibenzofuran	ND<50	10	0.33	Phenanthrene		ND<50	10	0.33			
Di-n-butyl Phthalate	ND<50	10	0.33	Phenol		ND<50	10	0.33			
1,2-Dichlorobenzene	ND<50	10	0.33	Pyrene		ND<50	10	0.33			
1,3-Dichlorobenzene	ND<50	10	0.33	1,2,4-Trichlorobenzene		ND<50	10	0.33			
1,4-Dichlorobenzene	ND<50	10	0 33	2,4,5-Trichlorophenol		ND<50	10	0.33			
3,3-Dichlorobenzidine	ND<100	20	0.66	2,4,6-Trichlorophenol		ND<50	10	0.33			
2,4-Dichlorophenol	ND<50	10	0 33	Comments: j,h							
Diethyl Phthalate	ND<50	10	0.33	Surrogate .	Recoverie	s (%)					
2,4-Dimethylphenol	ND<50	10	0.33	2-Fluorophenol		1	32				
Dimethyl Phthalate	ND<50	10	0.33	Phenol-d5			#				
4,6-Dinitro-2-methylphenol	ND<250	50	1.6	Nitrobenzene-d5							
2,4-Dinitrophenol	ND<250	50	16	2-Fluorobiphenyl			52				
2,4-Dinitrotoluene	ND<50	10	0.33	2,4,6-Tribromophenol			#				
2,6-Dinitrotoluene	ND<50	10	0.33	p-Terphenyl-d14			38				

^{*}water samples are reported in ug/L, soil and sludge samples in mg/kg, wipes in ug/wipe and all TCLP / STLC / SPLP extracts in ug/L ND means not detected above the reporting limit; N/A means analyte not applicable to this analysis.

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Edward Hamilton, Lab Director

^{*} surrogate diluted out of range

h) lighter than water immiscible sheen is present; i)liquid sample that contains greater than ~5 vol % sediment; j) sample diluted due to high organic content

QC REPORT

EPA 8015m + 8020

Date: 10/26/01	Extraction	: EPA 8	5030		Matrix:	Water	
		%Rec					
Compound	Sample	MS	MSD	Amount Spiked	MS	MSD	RPD
SampleID: 102601					Instrumer	nt: GO	C-3
Surrogate1	ND	105.0	101.0	100.00	105	101	3.9
Xylenes	ND	34.5	33.2	30.00	115	111	3.8
Ethylbenzene	ND	11.5	10.9	10.00	115	109	5.4
Toluene	ND	11.2	10.5	10.00	112	105	6.5
Benzene	ND	10.4	10.1	10.00	104	101	2.9
MTBE	ND	9.6	9.6	10.00	96	96	0.0
TPH (gas)	ND	84.8	84.1	100.00	85	84	0.8

% Re covery =
$$\frac{(MS-Sample)}{AmountSpiked} \cdot 100$$

$$RPD = \frac{(MS - MSD)}{(MS + MSD)} 2.100$$

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QC REPORT

EPA 8015m + 8020

Date: 10/24/01	Extractio	traction: EPA 5030				Matrix: Water			
Compound		%Recovery							
	Sample	MS	MSD	Amount Spiked	MS	MSD	RPD		
<u>SampleID:</u> 102401					Instrumer	nt: GC-	2 B		
Surrogate1	ND	115.0	108.0	100.00	115	108	6.3		
TPH (diesel)	ND	8750.0	8625.0	7500.00	117	115	1.4		

% Re covery =
$$\frac{(MS-Sample)}{AmountSpiked} \cdot 100$$

$$RPD = \frac{(MS - MSD)}{(MS + MSD)} \cdot 2.100$$

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QC REPORT

SVOCs (EPA 8270/625/525)

Date: 10/23/01-10/24/01	Extraction: N/A				Matrix: Water		
Compound	Concentration: ug/L				%Recovery		
	Sample	MS	MSD	Amount Spiked	MS	MSD	RPD
SampleID: 102301					Instrumer	nt: G	C-8
Surrogate1	ND	710.0	700.0	1000.00	71	70	1.4
Pyrene	ND	440.0	440.0	1000.00	44	44	0.0
Pentachlorophenol	ND	990.0	970.0	2000.00	50	49	2.0
2,4-Dinitrotoluene	ND	530.0	540.0	1000.00	53	54	1.9
4-Nitrophenol	ND	910.0	820.0	2000.00	46	41	10.4
Acenaphtene	ND	490.0	480.0	1000.00	49	48	2.1
4-Chloro-3-metylphenol	ND	720.0	700.0	2000.00	36	35	2.8
1,2,4-trichlorobenzene	ND	510.0	500.0	1000.00	51	50	2.0
N-nitroso-di-n-propyl	ND	320.0	320.0	1000.00	32	32	0.0
1,4-Dichlorobenzene	ND	460.0	460.0	1000.00	46	46	0.0
2-Chiorophenol	ND	720.0	710.0	2000.00	36	36	1.4
Phenol	ND	620.0	640.0	2000.00	31	32	3.2

% Re covery =
$$\frac{(MS-Sample)}{AmountSpiked} \cdot 100$$

RPD= $\frac{(MS-MSD)}{(MS+MSD)} \cdot 2\cdot 100$

Quantitation Report

(Not Reviewed)

IntFile : EVENTS.E

Quant Time: Nov 20 15:50 2001 Quant Results File: GC11A.RES

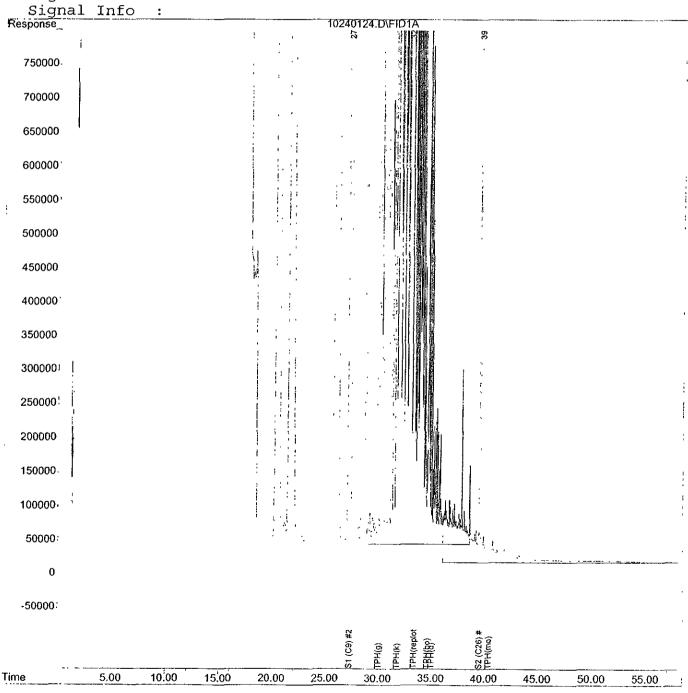
Quant Method : C:\HPCHEM\1\METHODS\GC11A.M (Chemstation Integrator)

Title : GC-11A

Last Update : Mon Nov 19 11:45:59 2001 Response via : Multiple Level Calibration

DataAcq Meth : GC11A.M

Volume Inj. : Signal Phase :



ent By: McCampbell Analytical, Inc.; 1 925 798 4612; Jan-9-02 1:23PM; Page 2 File : D:\HPCHEM\1\DATAA\10240126.D Operator : Thu Acquired : 25 Oct 2001 5:16 am using AcqMethod GC11A.M Instrument: GC-11 Sample Name: 81775W Misc Info : D,MO,WSG Vial Number: 13 Response_ 10240126.D\FID1A 750000 700000 ; 650000 600000 550000 500000 450000: 400000 350000 300000 250000 200000 150000

100000

50000

5.00

10.00

15.00

20.00

25.00

Time

55.00

50 00

45.00

40.00

30.00

35.00

(Not Reviewed)

Data File : D:\HPCHEM\1\DATAA\10240128.D

: 25 Oct 2001 6:24 am Acq On

Vial: 14 Operator: Thu : GC-11 Inst

Misc : D, MO, WSG

: 81776W

Multiplr: 1.0

IntFile : EVENTS.E

Sample

Ouant Results File: GC11A.RES Quant Time: Nov 20 15:50 2001

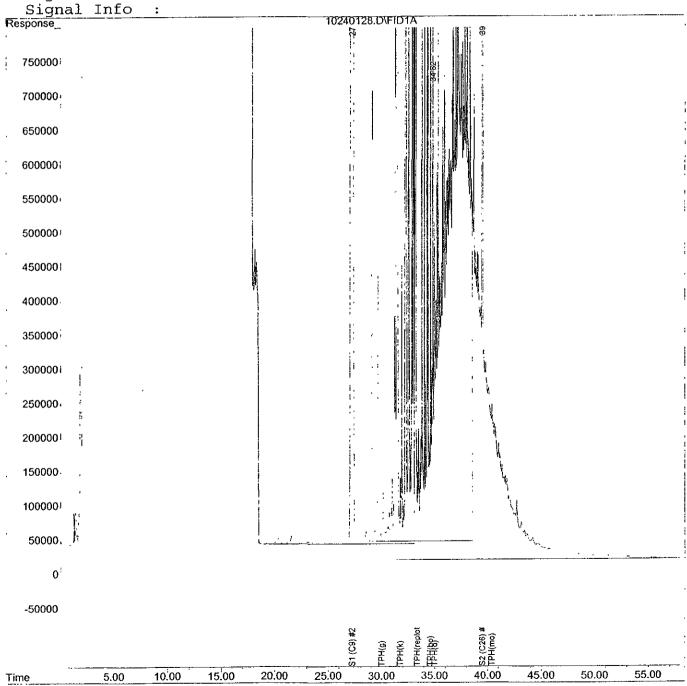
Quant Method : C:\HPCHEM\1\METHODS\GC11A.M (Chemstation Integrator)

: GC-11A Title

Last Update : Mon Nov 19 11:45:59 2001 Response via: Multiple Level Calibration

DataAcq Meth : GC11A.M

Volume Inj. Signal Phase :



IntFile : EVENTS.E

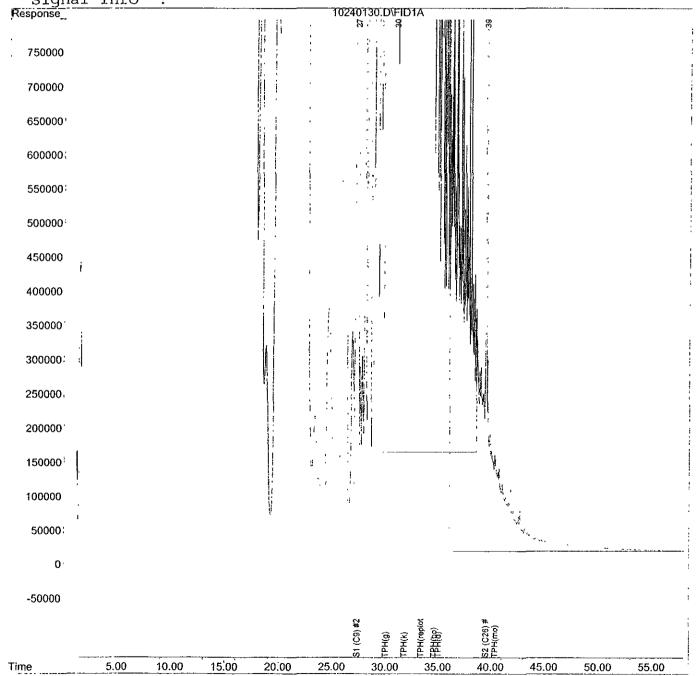
Quant Time: Nov 20 15:51 2001 Quant Results File: GC11A.RES

Quant Method : C:\HPCHEM\1\METHODS\GC11A.M (Chemstation Integrator)

Title : GC-11A

Last Update : Mon Nov 19 11:45:59 2001 Response via : Multiple Level Calibration

DataAcq Meth : GC11A.M



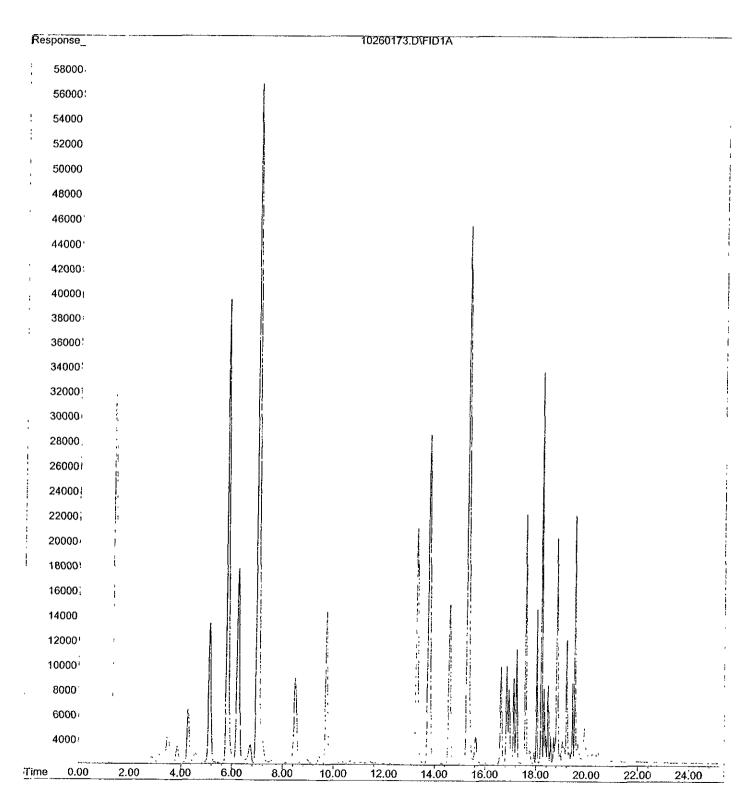
File : D:\HPCHEM\1\DATA\10260173.D

Operator

Acquired : 28 Oct 2001 4:34 am using AcqMethod GC3D.M

Instrument : GC-3 Sample Name: 81774 w

Misc Info : Vial Number: 73

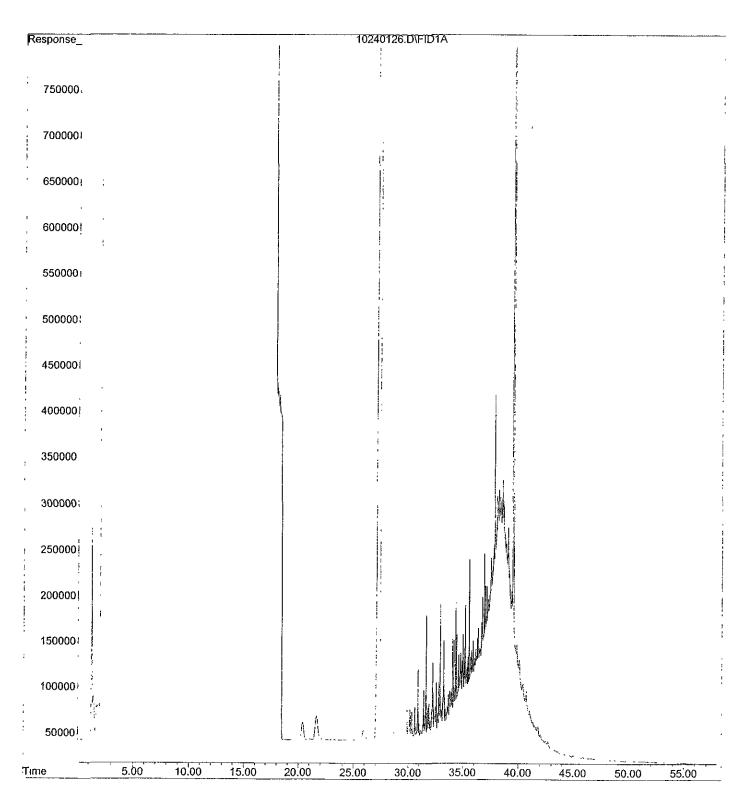


File : D:\HPCHEM\1\DATAA\10240126.D

Operator : Thu

Acquired : 25 Oct 2001 5:16 am using AcqMethod GC11A.M

Instrument: GC=11 Sample Name: 81775W Misc Info : D,MO,WSG



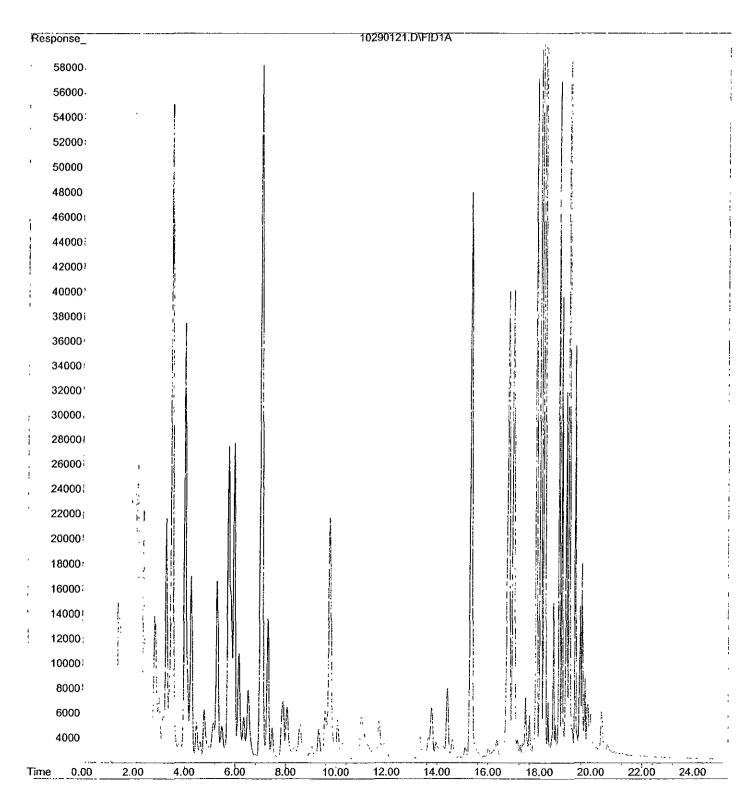
File : D:\HPCHEM\1\DATA\10290121.D

Operator

Acquired: 29 Oct 2001 10:58 pm using AcqMethod GC3D.M

Instrument : GC-3
Sample Name: (81776rrW+af)

Misc Info : Vial Number: 21



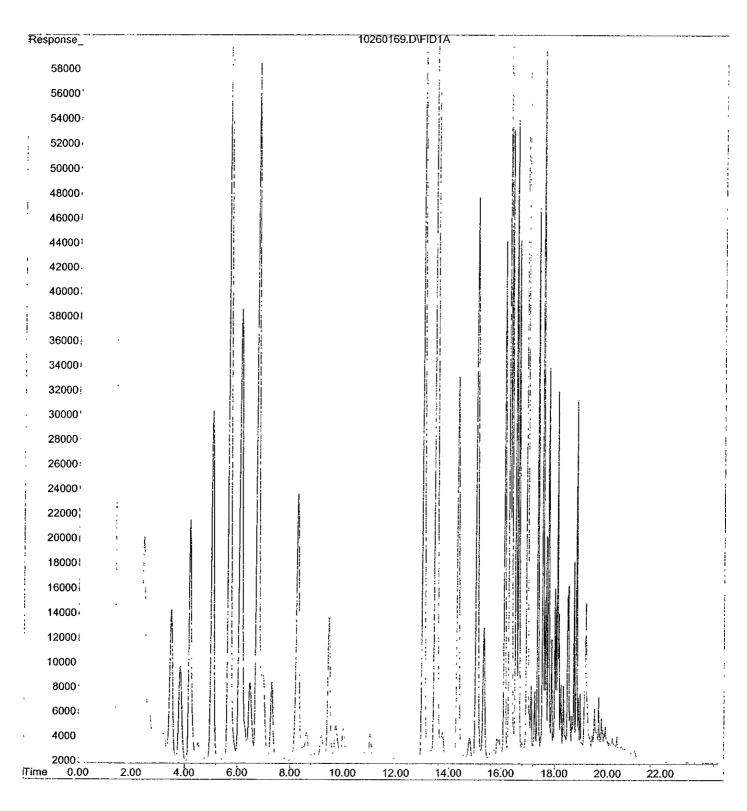
File : D:\HPCHEM\2\DATA\10260169.D

Operator

Acquired : 27 Oct 2001 9:19 pm using AcqMethod GC7B.M

Instrument : GC-7 Sample Name: 81777 w

Misc Info :



ent By: McCampbell Analytical, Inc.; 1 925 798 4612; Jan-10-02 12:17PM; Page 4/5

File : D:\HPCHEM\2\DATA\09190112.D

Operator

Operator : Acquired : 19 Sep 2001 4:32 pm using AcqMethod GC7B.M

Instrument : GC-7

Sample Name: 100ngBTEX

Misc Info : Vial Number: 12

Standarda from 78801

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ent By: McCampbell Analytical, Inc.; 1 925 798 4612; Jan-10-02 12:17PM; Page 5/5

File : D:\HPCHEM\2\DATA\09190125.D

Operator

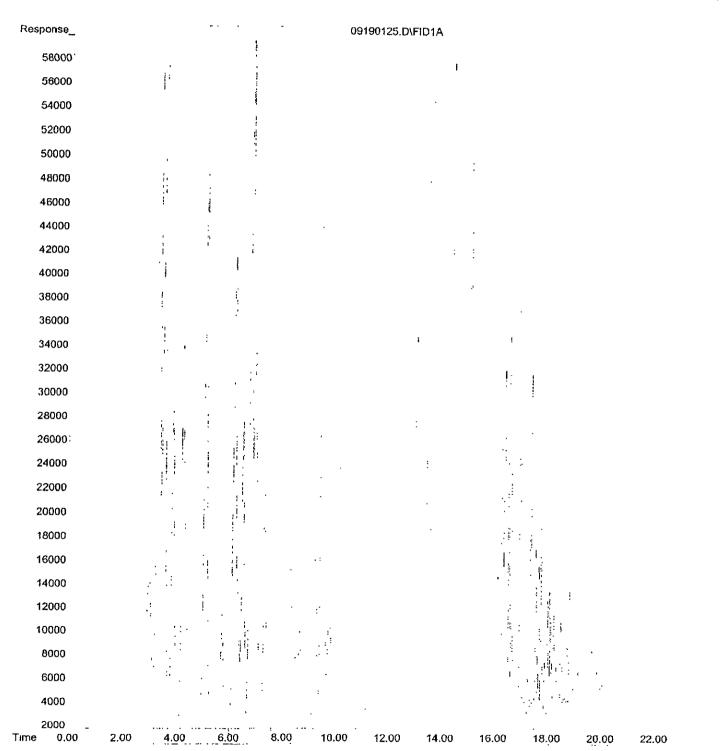
Acquired : 19 Sep 2001 11:01 pm using AcqMethod GC7B.M

Instrument : GC-7

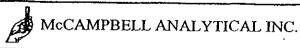
Sample Name: 5000ngGAS

Misc Info : Vial Number: 25

Stendard From 78801



McCAMPBELL ANALYTICAL INC. CHAIN OF CUSTODY RECORD 110 2" AVENUE SOUTH, #D7 TURN AROUND TIME PACHECO, CA 94553 Telephone: (925) 798-1620 Fax: (925) 798-1622 RUSH 24 HOUR 48 HOUR 5 DAY Report To: Rob Schultz
Company: Cambria Environmental Technology Bill To: Cambria Analysis Request Other Conuncuts 1144 65th Street, Suite C TPH as Diesel (8015) / TPH m.C. Total Petroleum Oil & Grease (3520 EMF/B&F) Oakland, CA 94608 W. 4h EPA 625 / 8270 / 8310 Tele: (510) 420-0700 Fax. (510) 420-9170 Project #: 453- 1705 Project Name: Embarcadero Cove Project Location: 1275 Emerates Cove Oakland, Co. Sampler Signature: BTEX ONLY (EP.A 602 / 8020) EP A 608 / 3080 PCB's ONLY Lead (7240/7421/239.2/6010) MEHIOD SAMPLING EPA 624 / 8240 / 8260 TPH as Diesel (3015) XUITAM Type Containers PRESERVED PAH's / PNA's by EPA 601/3010 EPA 608 / 8080 CAM-17 Meals EP.A 625 / 8270 BTEX & TPH 15 SAMPLE ID LOCATION BYEX Arr Sludge Date Time SVPCS Other HCI HNO, Other Soil <u>| 23</u> 12 LUFA Andre MN-1 10-14-01 9:55 X X × vou des MW-Z 10-19-01 10:40 እ $\overline{\mathsf{x}}$ × MU-3 10-14-01 10:10 × 9 MULH 10-19-01 9:20 X × 81774 81775 81776 81777 Relinquished By Date: Time. Received By: Remarks: Report results in EDF format. seemse location 1:30 VOAS LO&G METALS OTHER Unte: Time: //:50 ICEN _ PRESERVATION > Relinguished By: **APPROPRIATE** GOOD CONDITION 4:15 Jen Caro 10/23/01 1615 CONTAINERS HEAD SPACE ABSENT,



110 2nd Avenue South, #D7, Pacheco, CA 94553-5560
Telephone: 925-798-1620 Fax: 925-798-1622
http://www.mccampbell.com E-mail: main@mccampbell.com

Cambria Environmental Technology	Client Project ID: #458-1705;	Date Sampled: 09/17/01	
1144 65th Street, Suite C	Embarcadero Cove	Date Received: 09/18/01	
Oakland, CA 94608	Client Contact: Bob Schultz	Date Extracted: 09/18/01	
	Client P.O:	Date Analyzed: 09/18/01	

09/26/01

Dear Bob:

Enclosed are:

- 1). the results of 1 samples from your #458-1705; Embarcadero Cove project,
- 2). a QC report for the above samples
- 3). a copy of the chain of custody, and
- 4). a bill for analytical services.

All analyses were completed satisfactorily and all QC samples were found to be within our control limits. If you have any questions please contact me. McCampbell Analytical Laboratories strives for excellence in quality, service and cost. Thank you for your business and I look forward to working with you again.

Edward Hamilton, Lab Director



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Cambria Environmental Technology	Client Project ID: #458-1705;	Date Sampled: 09/17/01
1144 65 th Street, Suite C	Embarcadero Cove	Date Received: 09/18/01
Oakland, CA 94608	Client Contact: Bob Schultz	Date Extracted: 09/18/01
	Client P.O:	Date Analyzed: 09/19/01

Gasoline Range (C6-C12) Volatile Hydrocarbons as Gasoline*, with Methyl tert-Butyl Ether* & BTEX*

EPA methods 5030, modified 8015, and 8020 or 602; California RWQCB (SF Bay Region) method GCFID(5030)

Lab ID	Client ID	Matrix	TPH(g)⁺	мтве	Benzene	Toluene	Ethyl- benzene	Xylenes	% Recovery Surrogate
78801	M-1-5'	S	2300,a	5.1	1.8	3.7	48	7.2	#
			-						
					:				
otherwis	Limit unless stated; ND	w	50 ug/L	5.0	0.5	0.5	0.5	0.5	~~
	detected above orting limit	s	1.0 mg/kg	0.05	0.005	0.005	0.005	0.005	

^{*} water and vapor samples are reported in ug/L, wipe samples in ug/wipe, soil and sludge samples in mg/kg, and all TCLP and SPLP extracts in ug/L

^{*}The following descriptions of the TPH chromatogram are cursory in nature and McCampbell Analytical is not responsible for their interpretation: a) unmodified or weakly modified gasoline is significant; b) heavier gasoline range compounds are significant(aged gasoline?); c) lighter gasoline range compounds (the most mobile fraction) are significant; d) gasoline range compounds having broad chromatographic peaks are significant; biologically altered gasoline?; e) TPH pattern that does not appear to be derived from gasoline (?); f) one to a few isolated peaks present; g) strongly aged gasoline or diesel range compounds are significant; h) lighter than water immiscible sheen is present; i) liquid sample that contains greater than ~5 vol. % sediment; j) no recognizable pattern.



cluttered chromatogram; sample peak coelutes with surrogate peak



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Cambria Environmental Technology	Client Project ID: #458-1705;	Date Sampled: 09/17/01
1144 65 th Street, Suite C	Embarcadero Cove	Date Received: 09/18/01
Oakland, CA 94608	Client Contact: Bob Schultz	Date Extracted: 09/18/01
	Client P.O:	Date Analyzed: 09/19-09/20/01

Diesel Range (C10-C23) and Oil-Range (C18+) Extractable Hydrocarbons as Diesel and Motor Oil with Silica Gel Clean-up*

EPA methods modified 8015, and 3550 or 3510; California RWQCB (SF Bay Region) method GCFID(3550) or GCFID(3510)

Lab ID	Client ID	Matrix	TPH(d) ⁺	TPH(mo) [†]	% Recovery Surrogate
78801	M-1-5'	s	850,d	97	102
				· · · · · · · · · · · · · · · · · · ·	
Reporting Limit unless otherwise ted; ND means not detected above the reporting limit		w	50 ug/L	250 ug/L	
		S	1.0 mg/kg	5.0 mg/kg	

^{*}water samples are reported in ug/L, wipe samples in ug/wipe, soil and sludge samples in mg/kg, and all TCLP / STLC / SPLP extracts in ug/L

The following descriptions of the TPH chromatogram are cursory in nature and McCampbell Analytical is not responsible for their interpretation: a) unmodified or weakly modified diesel is significant; b) diesel range compounds are significant; no recognizable pattern; c) aged diesel? is significant); d) gasoline range compounds are significant; e) medium boiling point pattern that does not match diesel (?); f) one to a few isolated peaks present; g) oil range compounds are significant; h) lighter than water immiscible sheen is present; i) liquid sample that contains greater than ~5 vol. % sediment.



[&]quot;cluttered chromatogram resulting in coeluted surrogate and sample peaks, or; surrogate peak is on elevated baseline, or; surrogate has been diminished by dilution of original extract.

110 2nd Ave. South, #D7, Pacheco, CA 94553-5560
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http://www.mccampbell.com E-mail: main@mccampbell.com

QC REPORT

EPA 8015m + 8020

Date: 09/19/01					Matrix:	Soil	
C	Concentration: mg/kg					%Recovery	
Compound	Sample	MS	MSD	Amount Spiked	MS	MSD	RPD .
SampleID: 91801	Extraction	EPA 5	030		Instrumen	<u>l</u> G	C-7
Surrogate1	ND	98.000	99.000	100.00	98	99	1.0
Xylenes	ND	0.309	0.310	0.30	103	103	0.3
Ethylbenzene	ND	0.100	0.100	0.10	100	100	0.0
Toluene	ND	0.100	0.101	0.10	100	101	1.0
Benzene	ND .	0.093	0.096	0.10	93	96	3.2
MTBE	ND :	0.097	0.094	0.10	97	94	3.1
TPH (gas)	ND	0.953	1.001	1.00	95	100	5.0
SampleID: 91801	Extraction:	EPA 3	550		Instrument	: GC-1	i B
Surrogate1	ND	107.000	109.000	100.00	107	109	1.9
TPH (diesel)	ND :	144.500	152.000	150.00	96	101	5.1

% Re covery = $\frac{(MS-Sample)}{AmountSpiked} \cdot 100$ RPD= $\frac{(MS-MSD)}{(MS+MSD)} \cdot 2\cdot 100$ ent By: McCampbell Analytical, Inc.; 1 925 798 4612; Jan-10-02 12:16PM; Page 2 File : D:\HPCHEM\1\DATA\10260101.D Operator Acquired : 26 Oct 2001 12:40 pm using AcqMethod GC3D.M Instrument : <u>GC-3</u> Sample Name: 100ngBTEX Misc Info Vial Number: 1 Stemdard from 81774-777 Response_ 10260101.D\FID1A 58000 56000 54000 52000 50000 48000 460001 44000 42000 40000 38000 36000

Time

0.00

2.00

4.00

6,00

8.00

10.00

12.00

16.00

18.00

20.00

22.00

24.00

ent By: McCampbell Analytical, Inc.; 1 925 798 4612; Jan-10-02 12:17PM; Page 3/5 File : D:\HPCHEM\1\DATA\10260103.D Operator Acquired 26 Oct 2001 1:45 pm using AcqMethod GC3D.M Instrument : GC-3 Sample Name: 5000ngGAS Misc Info Vial Number: 3 Standard from 8174-7 Response_ 10260103.D\FID1A 58000 56000 54000 52000 50000 48000 46000 44000 42000 40000 38000 36000 34000 32000 30000 28000 26000 24000 22000 20000 18000

6000 4000 Time 0.00 2.00 4.00 6.00 8.00 10.00 12.00 14.00 16.00 18.00 20.00 22.00 24.00

Quantitation Report (Not Reviewed)

Data File : D:\HPCHEM\1\DATAB\09200105.D

Acq On : 20 Sep 2001 2:09 pm

Sample : 78801SRR

Vial: 53
Operator: Thu
Inst : GC-11
Multiplr: 100.0

Misc : D,MO,WSG IntFile : EVENTS.E

Quant Time: Nov 21 11:47 2001 Quant Results File: GC11B.RES

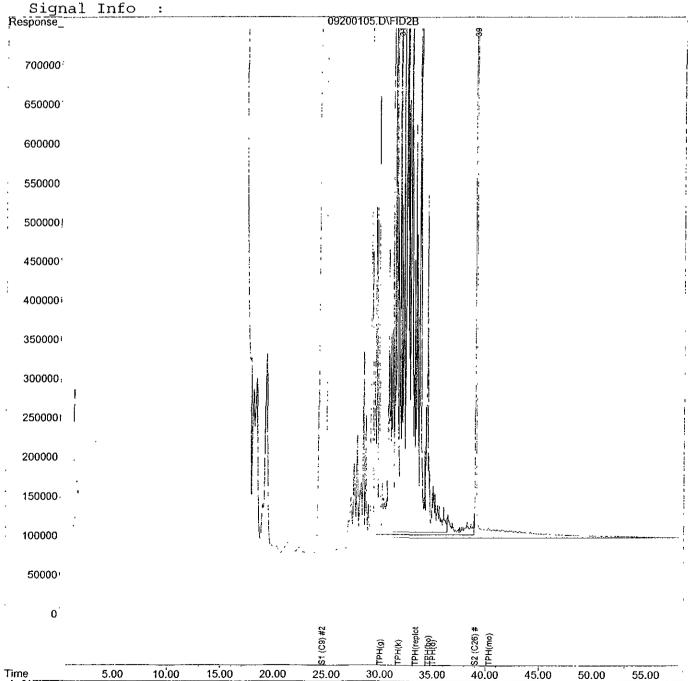
Quant Method : C:\HPCHEM\1\METHODS\GC11B.M (Chemstation Integrator)

Title : GC-11A

Last Update : Mon Nov 19 11:46:44 2001 Response via : Multiple Level Calibration

DataAcq Meth : GC11A.M

Volume Inj. : Signal Phase : Signal Info



File : D:\HPCHEM\2\DATA\09190128.D

Operator

Acquired : 20 Sep 2001 12:31 am using AcqMethod GC7B.M

Instrument : GC-7 Sample Name 78801s

Misc Info : Vial Number: 28

Res	sponse_					0919	0128.D\F	ID1A			V		
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McCAMPBELL ANALYTICAL INC. CHAIN OF CUSTODY RECORD 110 2rd AVENUE SOUTH, #D7 TURN AROUND TIME PACHECO, CA 94553 Telephone: (925) 798-1620 Fax: (925) 798-1622 RUSH 24 HOUR 48 HOUR Report To: Idin Riggi DOD SHULT 2 Bill To: Wes MAKLAND Analysis Request Other Comments Company: Cambria Environmental Technology Total Petroleum Oil & Grease (5520 E&F/B&F) 1144 65th Street, Suite C Salva, Oakland; CA 94608 PAH's / PNA's by ePA 625, 8270 / 8310 Tele: (510) 420-0700 Total Petroleum Hydrocarbons (418.1) Fax: (510) 420-9170 Project #: 458 -/705 Project Name: Emzaccaoera Con 3 BTEX ONLY (EPA 602 / 8020) Project Location: 1275 &MBARCADOU EPA 608 / 8080 PCB's ONLY TPH as Diesel (8015), MO Lead (7240/7421/239,2/6010) BTEX & TPH as Gas (602/8020 Sampler Signature: EPA 624 / 8240 / 8260 METHOD SAMPLING **MATRIX** PRESERVED Type Containers Containers CAM-17 Metals EPA 601 / 8010 EPA 608 / 8080 EPA 625 / 8270 LUFT 5 Metals SAMPLE ID LOCATION Time Sludge Date Water HNO Other Other Soil HC See Air -2' 9-17 8:45 اعتدا 78800 H. 10:00 X 9-17 × BATAL M-2-2 9-17 11:00 3146 78801 78802H Relinquished By: Date: Time: Received By: Remarks: VOAS 10&G I METALS 1 OTHER ICE/H Relinquished By Date: Time: **PRESERVATION** GOOD CONDITION **APPROPRIATE** HEAD SPACE ABSENT Réceived By:

110 2nd Avenue South, #D7, Pacheco, CA 94553-5560
Telephone: 925-798-1620 Fax: 925-798-1622
http://www.mccampbell.com E-mail: main@mccampbell.com

Cambria Environmental Technology	Client Project ID: #458-1705;	Date Sampled: 08/30/01
1144 65 th Street, Suite C	Embarcadero Cove	Date Received: 09/04/01
Oakland, CA 94608	Client Contact: Bob Schultz	Date Extracted: 09/04/01
	Client P.O:	Date Analyzed: 09/04/01

09/11/01

Dear Bob:

Enclosed are:

- 1). the results of 10 samples from your #458-1705; Embarcadero Cove project,
- 2). a QC report for the above samples
- 3). a copy of the chain of custody, and
- 4). a bill for analytical services.

All analyses were completed satisfactorily and all QC samples were found to be within our control limits. If you have any questions please contact me. McCampbell Analytical Laboratories strives for excellence in quality, service and cost. Thank you for your business and I look forward to working with you again.

Edward Hamilton, Lab Director

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Cambria Environmental Technology	Client Project ID: #458-1705;	Date Sampled: 08/30/01	
1144 65 th Street, Suite C	Embarcadero Cove	Date Received: 09/04/01	
Oakland, CA 94608	Client Contact: Bob Schultz	Date Extracted: 09/05-09/07/01	
	Client P.O:	Date Analyzed: 09/05-09/07/01	

Gasoline Range (C6-C12) Volatile Hydrocarbons as Gasoline*, with Methyl tert-Butyl Ether* & BTEX*

EPA methods 5030, modified 8015, and 8020 or 602; California RWQCB (SF Bay Region) method GCFID(5030) % Recovery Lab ID Client ID Matrix TPH(g)+ **MTBE** Benzene Toluene **Xylenes** benzene Surrogate 76862 S SB-A-3.5 ND ND ND ND ND ND 104 76865 S SB-B-3 5 ND ND ND ND ND ND 105 76868 S SB-D-3.5 ND ND ND ND ND ND 110 76871 SB-E-3.5 S 1.4,a ND 0.014 0.0080ND 0.026 ___# 76874 SB-F-3.5 S 2.5,a ND 0.021 0.010 ND 0.005 ___# 76877 SB-A W ND ND ND ND ND ND 102 76878 SB-B W ND ND ND ND ND ND 106 76879 SB-D W ND ND ND ND ND ND 105 76880 SB-E W 39,000,a ND<200 3200 750 1200 3600 104 76881 SB-F W ND ND ND ND ND ND 102

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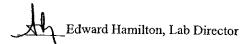
W

S

50 ug/L

1.0 mg/kg

The following descriptions of the TPH chromatogram are cursory in nature and McCampbell Analytical is not responsible for their interpretation. a) unmodified or weakly modified gasoline is significant; b) heavier gasoline range compounds are significant(aged gasoline?); c) lighter gasoline range compounds (the most mobile fraction) are significant; d) gasoline range compounds having broad chromatographic peaks are significant; biologically altered gasoline?, e) TPH pattern that does not appear to be derived from gasoline (?); f) one to a few isolated peaks present; g) strongly aged gasoline or diesel range compounds are significant; h) lighter than water immiscible sheen is present; i) liquid sample that contains greater than ~5 vol. % sediment, j) no recognizable pattern.



0.5

0.005

0.5

0.005

Reporting Limit unless

otherwise stated; ND means not detected above

the reporting limit

^{*} water and vapor samples are reported in ug/L, wipe samples in ug/wipe, soil and sludge samples in mg/kg, and all TCLP and SPLP extracts in ug/L

[&]quot; cluttered chromatogram; sample peak coelutes with surrogate peak

110 2nd Avenue South, #D7, Pacheco, CA 94553-5560
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http://www.mccampbell.com E-mail: main@mccampbell.com

Cambria Environmental Technology	Client Project ID: #458-1705;	Date Sampled: 08/30/01	
1144 65 th Street, Suite C	Embarcadero Cove	Date Received: 09/04/01	
Oakland, CA 94608	Client Contact: Bob Schultz	Date Extracted: 09/04/01	
	Client P.O:	Date Analyzed: 09/05-09/06/01	

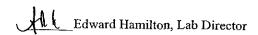
Diesel Range (C10-C23) and Oil-Range (C18+) Extractable Hydrocarbons as Diesel and Motor Oil with Silica Gel Clean-up*

EPA methods modified 8015, and 3550 or 3510; California RWQCB (SF Bay Region) method GCFID(3550) or GCFID(3510)

Lab ID	Client ID	Matrix	TPH(d) ⁺	T₽H(mo) ⁺	% Recovery Surrogate
76862	SB-A-3.5	S	1.4,g,b	5 2	106
76865	SB-B-3.5	S	ND	ND	107
76868	SB-D-3.5	s	ND	ND	105
76871	SB-E-3 5	S	2.4,g	6.1	109
76874	SB-F-3.5	S	4.6,g	16	105
76877	SB-A	w	1500,g	7200	96
76878	SB-B	w	63,g	550	104
76879	SB-D	w	1100,g	3400	104
76880	SB-E	w	5800,d	350	109
76881	SB-F	w	480,g	1400	108
	it unless otherwise	W	50 ug/L	250 ug/L	
ated, ND means not detected above the reporting limit		S	1.0 mg/kg	5.0 mg/kg	

^{*}water samples are reported in ug/L, wipe samples in ug/wipe, soil and sludge samples in mg/kg, and all TCLP / STLC / SPLP extracts in ug/L

^{*}The following descriptions of the TPH chromatogram are cursory in nature and McCampbell Analytical is not responsible for their interpretation: a) unmodified or weakly modified diesel is significant, b) diesel range compounds are significant; no recognizable pattern; c) aged diesel? is significant); d) gasoline range compounds are significant; e) medium boiling point pattern that does not match diesel (?); f) one to a few isolated peaks present; g) oil range compounds are significant; h) lighter than water immiscible sheen is present; i) liquid sample that contains greater than ~5 vol. % sediment.



^{*} cluttered chromatogram resulting in coeluted surrogate and sample peaks, or; surrogate peak is on elevated baseline, or; surrogate has been diminished by dilution of original extract.

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Cambria Environmental Technology	Client Project ID: #458-1705;	Date Sampled: 08/30/01		
1144 65th Street, Suite C	Embarcadero Cove	Date Received: 09/04/01		
Oakland, CA 94608	Client Contact: Bob Schultz	Date Extracted: 09/05/01		
	Client P.O:	Date Analyzed: 09/06-09/07/01		
7))	ATT TO TO ATT A TOWARD			

Polynuclear Aromatic Hydrocarbons (PAH / PNA) by GC-MS

EPA methods 625 (modified 610) and 3510 or 8270 (modified 8100) and 3550

EPA methods 620 (modified 610) and 3510 or 8270 (modified 8100) and 3550										
Lab ID	76862	76865	76868	76871	76874	Report	ing Limit			
Client ID	SB-A-3.5	SB-B-3.5	SB-D-3.5	SB-E-3.5	SB-F-3 5	s	W, STLC			
Matrix	S	s	s	s	S	,	TCLP			
Compound		(Concentration*			mg/kg	ug/L			
Acenaphthene	ND<0.25	ND	ND	ND	ND<0.25	0.062	10			
Acenaphthylne	ND<0.25	ND	ND	ND	ND<0.25	0 062	10			
Anthracene	ND<0.25	ND	ND	ND	ND<0.25	0.062	10			
Benzo(a)anthracene	ND<0.25	ND	ND	ND	ND<0.25	0.062	10			
Benzo(b)fluoranthene	ND<0.25	ND	ND	ND	ND<0.25	0.062	10			
Benzo(k)fluoranthene	ND<0.25	ND	ND	ND	ND<0 25	0.062	10			
Benzo(g,h,i)perylene	ND<0.25	ND	ND	ND	ND<0.25	0 062	10			
Benzo(a)pyrene	ND<0 25	ND	ND	ND	ND<0.25	0.062	10			
Chrysene	ND<0.25	ND	ND	ND	ND<0.25	0.062	10			
Dibenzo(a,h)anthracene	ND<0.25	ND	ND	ND	ND<0.25	0.062	10			
Fluoranthene	ND<0.25	ND	ND	ND	ND<0.25	0.062	10			
Fluorene	ND<0.25	ND	ŅD	ND	ND<0.25	0.062	10			
Indeno(1,2,3-cd)pyrene	ND<0.25	ND	ND	ND	ND<0.25	0.062	10			
Naphthalene	ND<0.25	ND	ND	ND	ND<0.25	0.062	10			
Phenanthrene	ND<0.25	ND	ND	ND	ND<0.25	0.062	10			
Pyrene	ND<0.25	ND	ND	ND	ND<0.25	0 062	10			
% Recovery Surrogate 1	101	96	99	95	97					
% Recovery Surrogate 2	117	106	110	105	111					
Comments	j				j					

^{*} water and vapor samples are reported in ug/L, soil and sludge samples in mg/kg, wipes in ug/wipe and all TCLP / STLC / SPLP extracts in ug/L.

ND means not detected above the reporting limit; N/A means analyte not applicable to this analysis

⁽h) a lighter than water immiscible sheen is present; (i) liquid sample that contains >~5 vol. % sediment; (j) sample diluted due to high organic content.



[&]quot;surrogate diluted out of range or surrogate coelutes with another peak

110 2nd Avenue South, #D7, Pacheco, CA 94553-5560
Telephone: 925-798-1620 Fax: 925-798-1622
http://www.mccampbell.com E-mail: main@mccampbell.com

Cambria Environmental Technology	Client Project ID: #458-1705;	Date Sampled: 08/30/01
1144 65 th Street, Suite C	Embarcadero Cove	Date Received: 09/04/01
Oakland, CA 94608	Client Contact: Bob Schultz	Date Extracted: 09/05/01
	Client P.O:	Date Analyzed: 09/06-09/07/01
		AL. CO NO

Polynuclear Aromatic Hydrocarbons (PAH / PNA) by GC-MS

EPA methods 625 (modified 610) and 3510 or 8270 (modified 8100) and 3550

Lab ID	76877	76878	76879	76880	76881	Report	ing Limit
Client ID	SB-A	SB-B	SB-D	SB-E	SB-F	C	W, STLC
Matrix	w	w	w	w	w	S	TCLP
Compound		(Concentration*			mg/kg	ug/L
Acenaphthene	ND	ND	ND	ND<50	ND	0,062	10
Acenaphthylne	ND	ND	ND	ND<50	ND	0.062	10
Anthracene	ND	ND	ND	ND<50	ND	0.062	10
Benzo(a)anthracene	ND	ND	ND	ND<50	ND	0.062	10
Benzo(b)fluoranthene	ND	ND	ND	ND<50	ND	0.062	10
Benzo(k)fluoranthene	ND	ND	ND	ND<50	ND	0.062	10
Benzo(g,h,i)perylene	ND	ND	ND	ND<50	ND	0.062	10
Benzo(a)pyrene	ND	ND	ND	ND<50	ND	0.062	10
Chrysene	ND	ND	ND	ND<50	ND	0.062	10
Dibenzo(a,h)anthracene	ND	ND	ND	ND<50	ND	0.062	10
Fluoranthene	ND	ND	11	ND<50	ND	0.062	10
Fluorene	ND	ND	ND	ND<50	ND	0.062	10
Indeno(1,2,3-cd)pyrene	ND	ND	ND	ND<50	ND	0.062	10
Naphthalene	ND	ND	ND	370	ND	0.062	10
Phenanthrene	ND	ND	ND	ND<50	ND	0.062	10
Pyrene	ND	ND	11	ND<50	ND	0.062	10
% Recovery Surrogate 1	102	111	96	105	100		
% Recovery Surrogate 2	88	99	106	98	108		
Comments							

^{*} water and vapor samples are reported in ug/L, soil and sludge samples in mg/kg, wipes in ug/wipe and all TCLP / STLC / SPLP extracts in ug/L.

⁽h) a lighter than water immiscible sheen is present; (i) liquid sample that contains >~5 vol. % sediment; (j) sample diluted due to high organic content.



ND means not detected above the reporting limit; N/A means analyte not applicable to this analysis

surrogate diluted out of range or surrogate coefutes with another peak

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http://www.mccampbell.com E-mail: main@mccampbell.com

QC REPORT

EPA 8015m + 8020

Date: 09/05/01					Matrix:	Water		
	i	Concen	%Rec	•				
Compound	Sample	MS	MSD	Amount Spiked	MS	MSD	RPD	:
SampleID: 90401	Extraction	EPA 5	030		Instrumer	<u>it:</u> G	C-3	•
Surrogate1	ND	100.0	103.0	100.00	100	103	3.0	* *
Xylenes	ND	32.4	32.7	30.00	108	109	0.9	1
Ethylbenzene	ND	10.7	10.7	10.00	107	107	0.0	1
Toluene	ND	10.7	11.0	10.00	107	110	2.8	
Benzene	ND	10.5	10.7	10.00	105	107	1.9	
МТВЕ	ND	11.0	11.1	10.00	110	111	0.9	# III
TPH (gas)	ND ·	85.3	85.2	100.00	85	85	0.1	1
SampleID: 90401	Extraction:	EPA 3	510		Instrumen	<u>t:</u> GC-11	I A	
Surrogate1	ND	108.0	106.0	100.00	108	106	1.9	;
TPH (diesel)	ND	7125.0	7400.0	7500.00	95	99	3.8	1

% Re covery =
$$\frac{(MS-Sample)}{AmountSpiked} \cdot 100$$

RPD= $\frac{(MS-MSD)}{(MS+MSD)} \cdot 2\cdot 100$

RPD means Relative Percent Deviation

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QC REPORT

SVOCs (EPA 8270/625/525)

Date: 09/07/01-09/08/01	Extractio	n: N/A			Matrix:	Soil	
1	*	Concen	%Rec	•			
Compound	Sample	MS	MSD	Amount Spiked	MS	MSD	RPD
SampleID: 91101		er erete .		!	Instrumer	<u>nt:</u> G	C-8
Surrogate1	ND	930.0	940.0	1000.00	93	94	1.1
Pyrene	ND	730.0	760.0	1000.00	73	76	4.0
Pentachlorophenol	ND	1700.0	1750 0	2000.00	85	88	2.9
2,4-Dinitrotoluene	ND	880.0	930.0	1000 00	88	93	5.5
4-Nitrophenol	ND	2170.0	2230.0	2000.00	109	112	27
Acenaphtene	ND	820.0	840.0	1000.00	82	84	2.4
4-Chloro-3-metylphenol	ND	1550.0	1590.0	2000.00	78	80	2.5
1,2,4-trichlorobenzene	ND	830.0	840.0	1000 00	83	84	1.2
N-nitroso-di-n-propyl	ND	830.0	850.0	1000.00	83	85	2.4
1,4-Dichlorobenzene	ND	0.008	830.0	1000.00	80	83	3.7
2-Chlorophenol	ND	1480.0	1530.0	2000.00	74	77	3.3
Phenol	ND	1440.0	1490.0	2000.00	72	75	3.4

% Re covery =
$$\frac{(MS-Sample)}{AmountSpiked} \cdot 100$$

$$RPD = \frac{(MS-MSD)}{(MS+MSD)} \cdot 2\cdot 100$$

RPD means Relative Percent Deviation

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QC REPORT

SVOCs (EPA 8270/625/525)

Date: 09/06/01-0/07/01	Extraction	n: N/A	Matrix:	Vater			
	!	Concen	%Reco	i			
Compound	Sample	MS	MSD	Amount Spiked	MS	MSD	RPD
SampleID: 91101			v		Instrument	<u>:</u> G	C-8
Surrogate1	ND	590.0	620.0	1000.00	59	62	5.0
Pyrene	ND	510.0	460.0	1000.00	51	46	10.3
Pentachlorophenol	ND	1320.0	1320.0	2000.00	66	66	0.0
2,4-Dinitrotoluene	ND -	510.0	500.0	1000.00	51	50	2.0
4-Nitrophenol	ND	1190.0	1190.0	2000.00	60	60	0.0
Acenaphtene	ND	490.0	440.0	1000.00	49	44	10.8
4-Chloro-3-metylphenol	ND	900.0	930.0	2000.00	45	47	3.3
1,2,4-trichlorobenzene	ND ·	500.0	460.0	1000.00	50	46	8.3
N-nitroso-di-n-propyl	ND	500.0	480.0	1000.00	50	48	4.1
1,4-Dichlorobenzene	ND	480.0	430.0	1000.00	48	43	11.0
2-Chlorophenol	ND	950.0	900.0	2000.00	48	45	5.4
Phenol	ND	830.0	850.0	2000.00	42	43	2.4

$$\% \text{ Re covery} = \frac{\left(MS - Sample \right)}{AmountSpiked} \cdot 100$$

$$RPD = \frac{(MS - MSD)}{(MS + MSD)} \cdot 2 \cdot 100$$

ent By: McCampbell Analytical, Inc.; 1 925 798 4612; Jan-10-02 4:16PM; Page 2 File : D:\HPCHEM\2\DATAB\09050115.D Operator : Thu Acquired 6 Sep 2001 2:30 am using AcqMethod GC6ANEW.M Instrument : GC-6 Sample Name: 768628 Misc Info : D,MO,WSG Vial Number: 58 Response 09050115.D\FID2B 19000 18000 17000 16000 15000 14000 13000 12000 11000 10000 9000 8000 7000 6000 5000 4000 3000 2000 1000 Time 0.00 5.00 15.00 20.00 25.00

30.00

35.00

40.00

55.00

50.00

Quantitation Report

(Not Reviewed)

IntFile : autoint1.e

Quant Time: Nov 20 16:50 2001 Quant Results File: GC6ANEW.RES

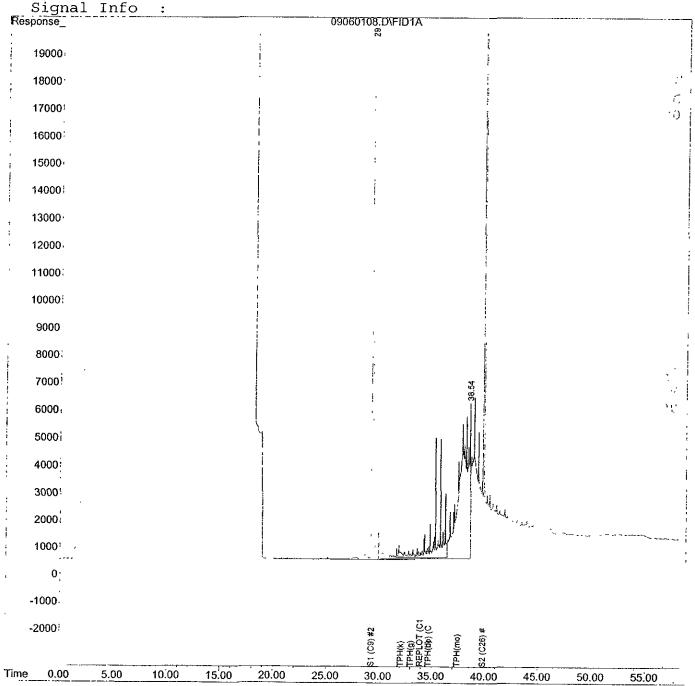
Quant Method : C:\HPCHEM\2\METHODS\GC6ANEW.M (Chemstation Integrator)

Title : GC-6A

Last Update : Mon Nov 19 11:42:54 2001 Response via : Multiple Level Calibration

DataAcq Meth : GC6ANEW.M

Volume Inj. : Signal Phase : Signal Info



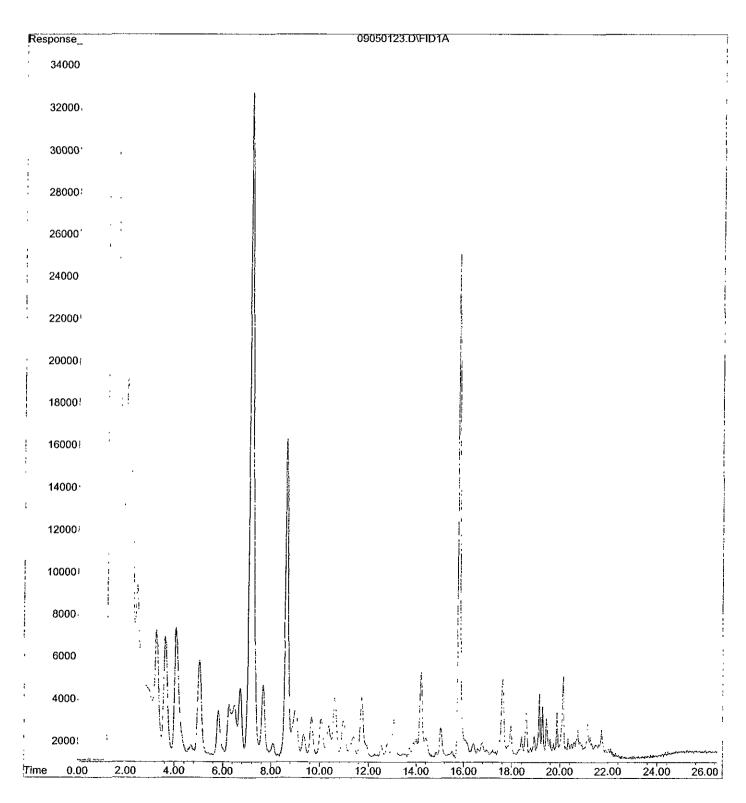
File : D:\HPCHEM\2\DATA\09050123.D

Operator

Acquired : 6 Sep 2001 5:27 am using AcqMethod GC12A.M

Instrument: GC-12 Sample Name 76871 S

Misc Info



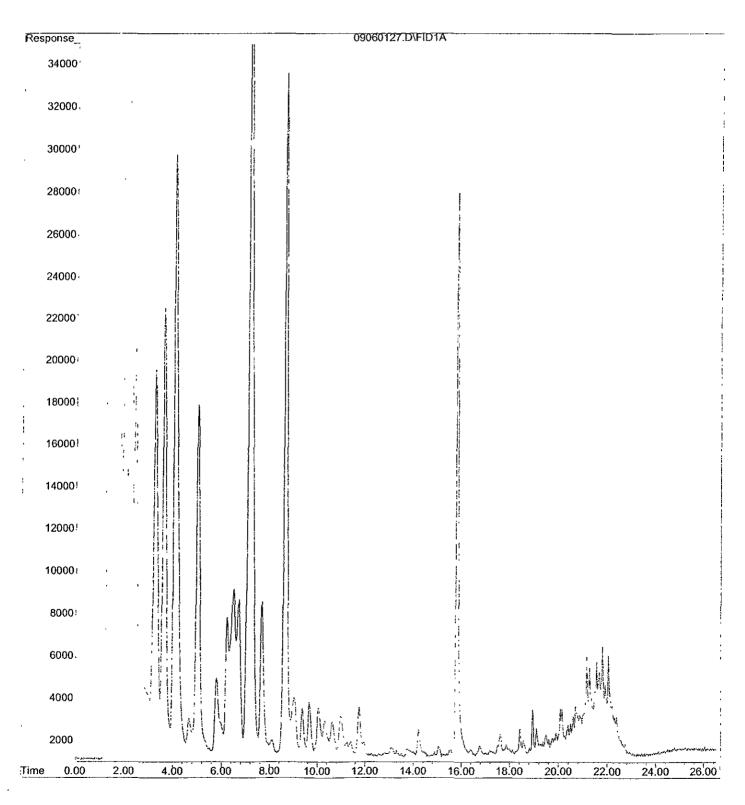
File : D:\HPCHEM\2\DATA\09060127.D

Operator

Acquired : 7 Sep 2001 6:17 am using AcqMethod GC12A.M

Instrument: GC-12 Sample Name: 76874rrS

Misc Info : Vial Number: 27



IntFile : EVENTS.E

Quant Time: Nov 20 16:54 2001 Quant Results File: GC6BNEW.RES

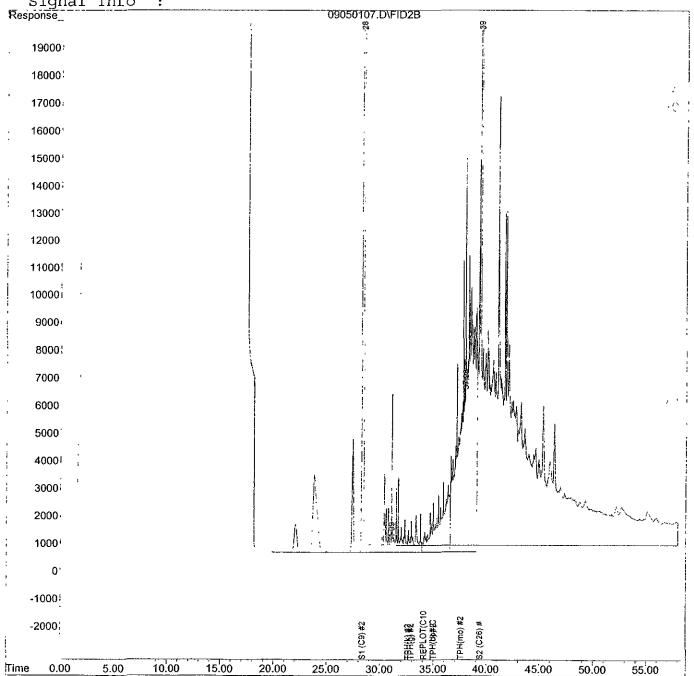
Quant Method: C:\HPCHEM\2\METHODS\GC6BNEW.M (Chemstation Integrator)

Title : GC-2B

Last Update : Fri May 05 10:34:57 2000 Response via : Multiple Level Calibration

DataAcq Meth : GC6ANEW.M

Volume Inj. : Signal Phase : Signal Info :



ent by: McCampbell Analytical, Inc.; 1 925 798 4612; Page 3/6

File : D:\HPCHEM\2\DATAA\09050120.D
Operator : Thu
Acquired : 6 Sep 2001 5:58 am using AcqMethod GC6ANEW.M

Instrument: GC-6
Sample Name: 76877W
Misc Info : D,MO,WSG

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aent By: McCampbell Analytical, Inc.; 1 925 798 4612; Jan-10-02 4:17PM; Page 4/6

: E:\HPCHEM\3\DATAB\09060115.D File

Operator : Thu

Acquired : 7 Sep 2001 4:27 am using AcqMethod GC2ANEW.M

Instrument : GC-2 Sample Name: 76878WRE Misc Info : D, MO, WSG

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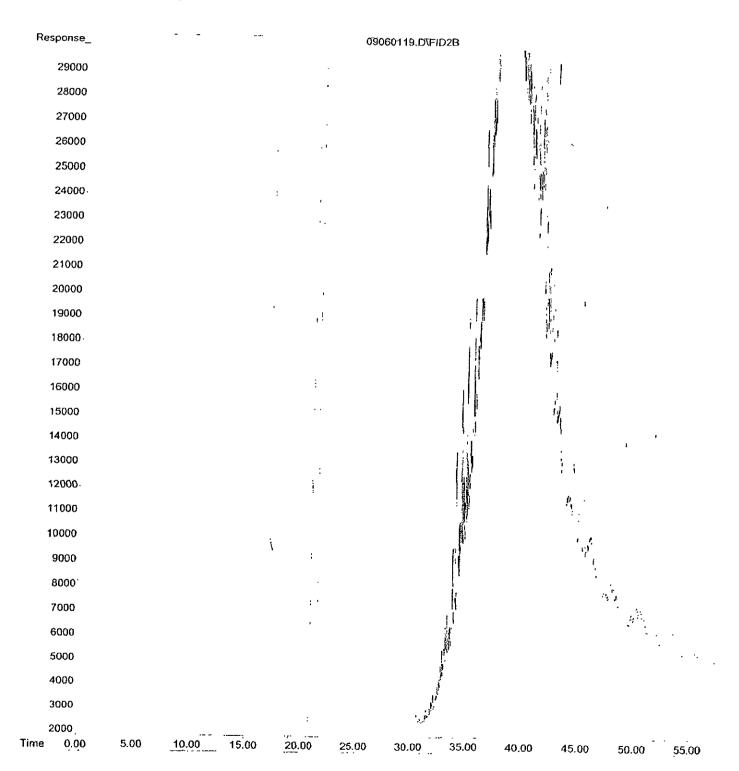
ent By: McCampbell Analytical, Inc.; 1 925 798 4612; Jan-10-02 4:17PM; Page 5/6

File : E:\HPCHEM\3\DATAB\09060119.D

Operator : Thu

Acquired : 7 Sep 2001 7:35 am using AcqMethod GC2ANEW.M

Instrument : GC-2 Sample Name: 76879WRE Misc Info : D,MO,WSG



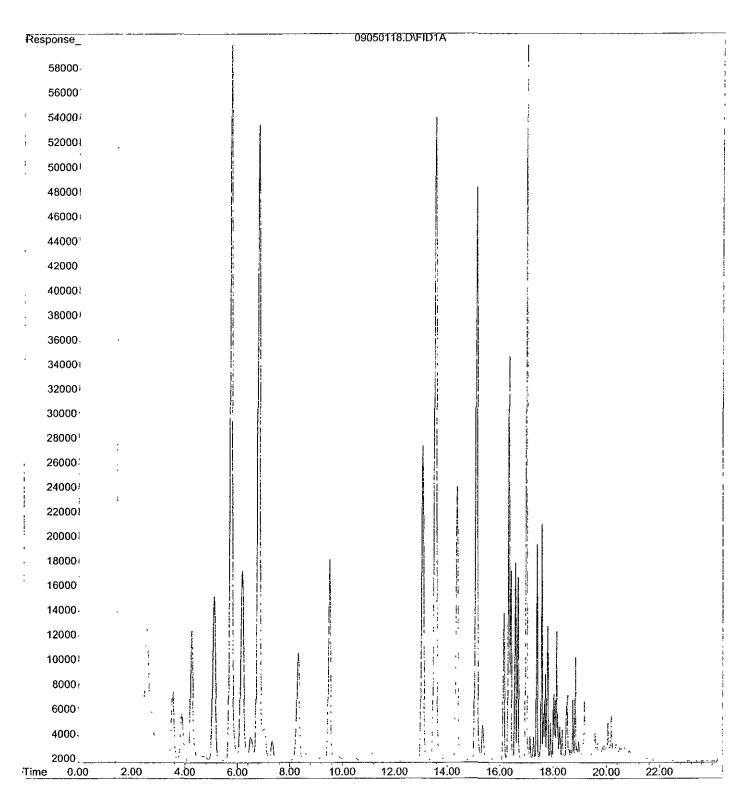
File : D:\HPCHEM\2\DATA\09050118.D

Operator

Acquired : 5 Sep 2001 9:15 pm using AcqMethod GC7B.M

Instrument: GC-7 Sample Name: 76880w

Misc Info : Vial Number: 18



Quantitation Report (Not Reviewed)

 Data File: D:\HPCHEM\2\DATAB\09050119.D
 Vial: 60

 Acq On: 6 Sep 2001 5:58 am
 Operator: Thu

 Sample: 76880W
 Inst: GC-6

 Misc: D,MO,WSG
 Multiplr: 1.00

IntFile : EVENTS.E

Quant Time: Nov 20 16:54 2001 Quant Results File: GC6BNEW.RES

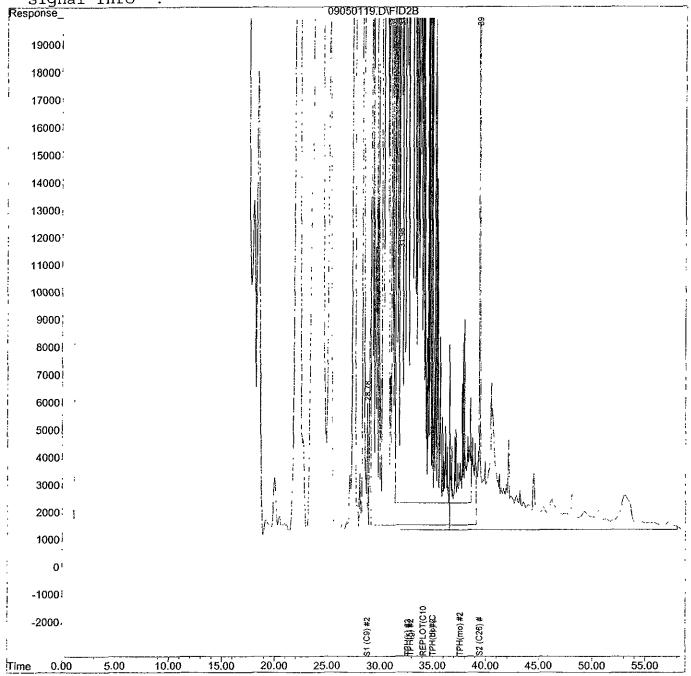
Quant Method : C:\HPCHEM\2\METHODS\GC6BNEW.M (Chemstation Integrator)

Title : GC-2B

Last Update : Fri May 05 10:34:57 2000 Response via : Multiple Level Calibration

DataAcq Meth : GC6ANEW.M

Volume Inj. : Signal Phase : Signal Info :



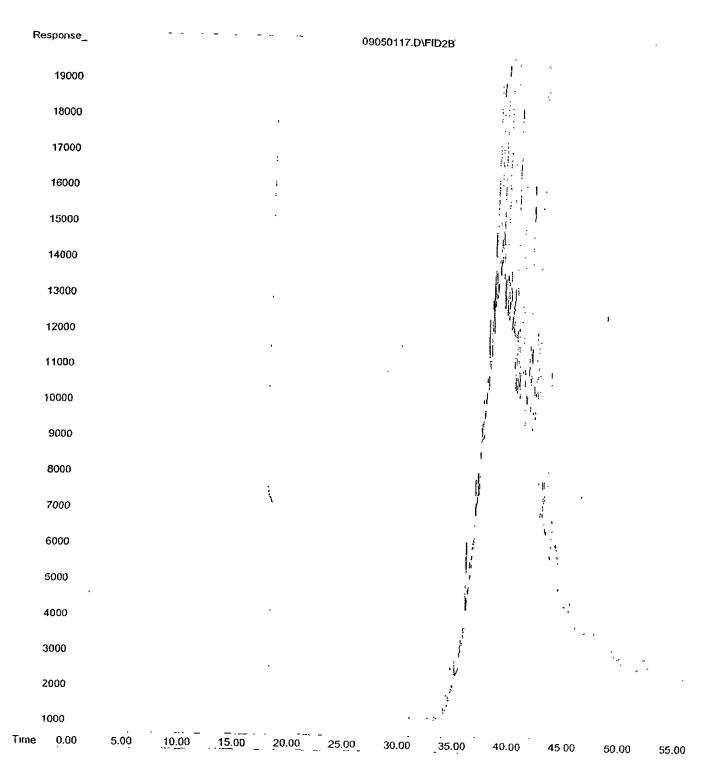
Jan-10-02 5:45PM; ent By: McCampbell Analytical, Inc.; 1 925 798 4612; Page 2/2

File : D:\HPCHEM\2\DATAB\09050117.D

Operator : Thu

Acquired 6 Sep 2001 4:15 am using AcqMethod GC6ANEW.M

Instrument : GC-6 Sample Name: 76881W Misc Info : D,MO,WSG



MCCAMPBELL AN	ALYTICAL INC.	OTTANION CONTRACTOR
110 2 [™] AVENUE	SOUTH, #137	CHAIN OF CUSTODY RECORD
PACHECO, C Telephone: (925) 798-1620	Fax: (925) 798-1622	TURN AROUND TIME
Report To: John Rissi Och Schulle	Rill To: P C Range	RUSH 24 HOUR 48 HOUR
Company: Cambria Environmental Technology	JOHN OF CHALLMON	Analysis Request Other Comments
1144 65th Street, Suite C		
Oakland, CA 94608 Tele: (510) 420-0700		76862 (0) 10 10 10 10 10 10 10 10 10 10 10 10 10
Project #: 458 - 1705	Fax: (510) 420-9170	76863 H
Project Location: OAKLAND	Project Name: EMBARCADES COL	(635)
Sampler Signature:		200 PA 625 / 8020) 2.00 PA
SAMPLING	MATRIX METHOD PRESERVED	201539.246010) 201539.246010) 201539.246010) 201539.246010) 201539.246010) 201539.246010) 201539.246010) 201539.246010) 201539.246010) 201539.246010)
SAMPLE ID LOCATION	laine laine	80 Pe 80 Pe
Date Time	Complete Com	24 / 80 ONL Y 24 / 80 ONL Y 80
	# Containers Type Containers Water Soil Air Sludge Other HCI EMO,	## 19 BTEX & TPH as Gas (602)/R010 Total Petroleum Oil & Grease Total Petroleum Hydrocarbon EPA 601 / 8010 BTEX ONLY (EPA 602 / 802 BTEX ONLY (EPA 602 / 802 EPA 608 / 8080 PCB* ONLY E
SB-A-45 3.5 85/11 930	I & V	
50-A-715 7.5 1936		DO HOU
53-A=4-5 9.5 940		76060
SB-B-4,5 3.5 11:51		76868
58-8-7-5 6.5 11:58		76869 H
58-B-42-5 O.S 1205		
SB-D-4.5 3.5 1045		768704
5B-D-75 6.5 1055	X	
SB-0 -12-5 10.5 11:00		76871
56-6-4,5-3.5 730	1 1 8	20 768724
SB-E-7.5 6.0 740		
58-E-9.5-7.5 800		76873H
58-F-4.5 3.5 635	K K	
88-F-7,0 S.S 855	I K K	76874
SB-F-4-5 7.5 Selinguished By: Date: Time:		76875H
Date: Time:	Received By:	Rentarks:
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		the Continu MIDE 8020 detections by 8260 768761
Relinquished By: Date: Time:	Received Dy:	Remarks: #= tube - Solice Gel Claim of on THATTHMI. The Confirm MIDE 8020 detections by 8200 768761 OR if detection Limit is greater than stopps.

APPENDIX E

Field Activity Descriptions

APPENDIX E

FIELD ACTIVITY DESCRIPTIONS

August 2001 Investigation

Field activities completed during the August 30, 2001 sampling program are presented below. The discussion is organized according to the nature of the individual activity.

Sampling Date: August 30, 2001.

Scope of Work: Iris-Cambria collected soil and grab groundwater samples from six

(6) onsite borings in the vicinity of a former UST.

Personnel Present: Kris Cejka, Cambria Senior Staff Geologist; John Riggi, Cambria

Project Geologist and a California Registered Geologist; Steven

Moore, Vironex Geoprobe operator.

Drilling Company: Vironex of San Leandro, California (C57# 705927).

Drilling Methods: Geoprobe[™] hydraulic push. Each boring was hand-augered to 5 ft

bgs (or less if restricted by refusal or obstacles in the borehole) as a

safeguard to prevent damage to subsurface utilities.

Number of Borings: Six (6). Boring SB-C was advanced to 4 ft bgs and could not be

completed to groundwater because of subsurface obstacles.

Boring Depths: Four to 11 ft bgs. Boring logs are included as Appendix A.

Concrete Coring: On August 29, 2001, Osborne Concrete Coring of Oakland,

California (Osborne), performed asphalt and concrete coring at the six boring locations. After coring the surface pavement, underlying gravel was excavated to expose a buried foundation(s). The buried foundation, that was encountered at all 6 locations, was then cored by Osborne using an extension device on the concrete corer. Surface

pavement was approximately 6 inches thick and the buried

foundation was approximately 6 to 12 inches thick.

Soil Sampling: Soil samples were collected continuously from all borings during

drilling and logged in accordance with the Unified Soil Classification

System.

Groundwater Sampling: Grab groundwater samples were collected through 3/8"-inner

diameter (ID) polyethylene tubing from borings SB-A, SB-B, SB-D,

SB-E, and SB-F.

Backfill Method: All borings were backfilled to the surface with neat cement and

completed with concrete patch.

Waste Disposal: All soil cuttings and rinsate were left onsite in Department of

Transportation (DOT)-approved 55-gallon drums pending disposal

by the Port.

October 2001 Investigation

Field activities completed during the October 2001 sampling program are presented below. The discussion is organized according to the nature of the individual activity.

Installation Date: October 9, 2001.

Scope of Work: Iris-Cambria advanced four (4) borings and completed all four as

monitoring wells. Soil samples were collected from all borings. The wells were developed and sampled per Tri-Regional procedural

guidelines.

Personnel Present: Kris Cejka, Cambria Senior Staff Geologist; and Barbara Jakub,

R.G., Cambria Project Geologist.

Drilling Company: V&W Drilling of Rio Vista, California (C57# 702904).

Drilling Methods: Hollow-stem auger drill rig. Each boring was hand-augered to 5 ft

bgs (or less if restricted by subsurface conditions) as a safeguard to

prevent damage to subsurface utilities.

Number of Borings/Wells: Four (4).

Boring Depths: Prior to well installation, the borings were drilled to depths of

between 13 and 15 ft bgs.

Concrete Coring: On October 8, 2001, Osborne Concrete Coring of Oakland,

California (Osborne), performed asphalt and concrete coring at the four well locations. After coring the surface pavement, underlying soil was excavated to expose a buried foundation(s) or to a maximum depth of 5 ft bgs. The buried foundation(s) (encountered in locations

MW-2 and MW-3) was then cored by Osborne using an extension device on the concrete corer. In borings MW-2 and MW-3, surface pavement was approximately 6 inches thick, and the buried foundation(s) was approximately 10 to 11 inches thick. Borings MW-1 and MW-4 were installed on dirt surfaces and no buried foundation was encountered in either boring.

Soil Sampling:

Soil samples were collected continuously from all borings during drilling and logged in accordance with the Unified Soil Classification System.

Well Development:

On October 12, 2001, Iris-Cambria developed monitoring wells MW-1 through MW-4. During development of monitoring well MW-3, the bottom cap was dislodged from the well casing and the surge block could not be retrieved from the well. Well MW-3 can be sampled, although there is an obstruction in the well at approximately 9 ft bgs. Well depth measurements are included in Appendix G.

Groundwater Gauging:

Groundwater was gauged in Site wells on October 12 (during well development), October 19 (during the first well sampling event), December 5 (near lower high tide and higher high tide, and the corresponding second and third sampling events), and December 19, 2001 (during lower low tide). Well caps were removed and water levels allowed to equilibrate for at least 15 minutes and until stable. During each gauging event, all four wells were gauged within a 30-minute time period. Field forms are included in Appendix G.

Groundwater Sampling:

All four (4) site wells were sampled three times: on October 19, 2001, on December 5, 2001, and again 6 hours later on December 5, 2001. Groundwater samples were collected from Site wells MW-1, MW-2, and MW-4 after purging three well casing volumes of water from each well and waiting for at least 80% groundwater recharge to the well. The samples for MW-3 were collected pre-purge due to an obstruction in the well. In addition, samples were collected from MW-1 only on December 7, 2001 in HNO₃-preserved polyethylene containers for metals analysis. The samples collected for volatile analyses were placed in 40-ml VOAs while samples collected for petroleum hydrocarbons and SVOCs were placed in 1 liter impreserved ambers. The sample bottles were labeled and placed in a cooled container for transport to McCampbell Analytical in Pacheco, California. Cambria's standard sampling procedures are presented in Appendix C. Well sampling forms are included in Appendix G.

Chemical Analysis:

Soil and groundwater samples were sent under chain of custody control to McCampbell Analytical, Inc., of Pacheco, California., and analyzed for: TPHg by EPA Method 8015; TPHd and TPHmo

by EPA Method 8015 with silica-gel cleanup; BTEX by EPA Method 8020; and SVOCs by EPA Method 8270. Soil samples were also analyzed for VOCs by EPA Method 8260. Groundwater samples from MW-1 on December 7, 2001 were analyzed for CAM 17 metals by EPA Method 6010. Laboratory analytical results are included in Appendix D.

Well Survey:

Virgil Chavez Land Surveying of Vallejo, California, (State of California Licensed Land Surveyor No. 6323) surveyed the well top of casing (TOC) elevations relative to Port datum TP184 on October 30, 2001. Survey data is presented in Appendix H.

Waste Disposal:

All soil cuttings and well development and purge water were left onsite in DOT-approved, 55-gallon drums pending disposal by the Port.

APPENDIX F

Geophysical Survey Report



September 27, 2001

Mr. Bob Shultz Cambria Environmental Technology, Inc 1144 65th Street, Suite A Oakland, CA 94008

Dear Mr Shultz:

This letter is submitted in reference to a ground penetrating radar (GPR) survey performed by NORCAL Geophysical Consultants, Inc. at 1275 Embarcadero Rd in Oakland, California. The GPR field survey was conducted on September 5, 2001 by NORCAL geophysicist Dan Jones. Site orientation was coordinated by Mr Kris Cejka of Cambria.

SITE DESCRIPTION AND PURPOSE

The site is a restaurant parking lot adjacent to a restaurant building that is no longer in use. Cambria advised us that a previous geophysical survey performed by others detected four isolated subsurface metal anomalies in this parking lot. The spray painted surface traces of these anomalies were visible on the site. In addition, previous environmental borings encountered concrete at a depth of approximately two feet ranging in thickness from four inches to one foot at various locations throughout the parking lot. The concrete may be that of a former building foundation at the site. The purpose of NORCAL's geophysical survey is two-fold: 1) to collect additional subsurface information at the four subsurface metal locations to determine if they may represent underground storage tanks (UST's) and 2) to delineate the boundary of the subsurface concrete (presumed foundation).

GEOPHYSICAL METHODOLOGY

Based on the anticipated required depth of investigation, the nature of the targets, and the high resolution needed to identify such objects, we chose the ground penetrating radar (GPR) method to investigate for both the UST's and the concrete foundation boundary.

GPR is a method that provides a continuous, high resolution cross-section depicting variations in the electrical properties of the shallow subsurface. The method is particularly sensitive to variations in electrical conductivity and electrical permittivity (the ability of a material to hold a charge when an electrical field is applied). The system operates by radiating electromagnetic pulses into the ground from an antenna as it is moved along the ground surface. Since most earth materials are partially transparent to electromagnetic energy, only a portion of the radar signal is reflected back to the surface from interfaces representing variations in electrical properties. When the signal encounters a metal object such as a UST, all of the incident energy is reflected. Lesser amounts of energy are reflected by non-metallic features such as as concrete blocks or backfill boundaries. The reflected signals are then received by the same antenna and are printed in cross-section form on a graphical recorder. Depending upon depth and/or thickness the resulting records can provide information regarding the location of UST's, sumps, buried debris, underground utilities, and variations in the shallow site materials.

Typically, UST's result in a high-amplitude, parabolic reflections at least as broad as the UST itself when the traverse is oriented perpendicular to the long axis of the UST. The GPR response of a concrete foundation is



Cambria Environmental Technology, Inc. September 27, 2001 Page 2

a relatively flat reflector sometimes with steeply sloping diffractions off the edge. In addition, the presence of steel reinforcing bar within the concrete can help to identify the foundation on the "cross-sections", by manifesting itself as closely spaced high-amplitude reflections.

In some cases, site-specific soil properties can limit the effectiveness of the GPR method. For instance, clayrich soil, conductive fill-material, or very wet soil can severely attenuate GPR signals, reducing the available depth of investigation and decreasing the resolution necessary to identify small subsurface variations. In addition, there may not be a significant enough contrast in electrical properties between the target of interest and the surrounding materials to identify the target

To complete the survey, we used a Geophysical Survey Systems, Inc. SIR-2000 Subsurface interface Radar System equipped with a 500 megahertz (MHz) antenna. A 900 MHz antenna was used in addition to the 500 MHz antenna for the concrete delineation portion of the survey since it provides higher resolution

GPR SURVEY AND RESULTS

UST Investigation

To begin the survey, we verified locations of the subsurface metal anomalies with a specialized metal detector (Fisher TW-6 M-Scope). We confirmed that each anomaly did provide a significant metallic response. The approximate dimensions in feet of the metal anomalies (north-south by east-west) as outlined by others and confirmed by NORCAL are as follows: A1 - 12x7, A2 - 11x6, A3 - 5x4, and A4 - 3x3.

We then established a series of multiple orthogonally oriented GPR traverses over each anomaly, A1-A4, extending the traverses at least five feet in all directions past the metal response boundaries. We did not observe any significant isolated reflectors or characteristic reflection signatures of USTs at anomalies A1 and A2. At A3, a small isolated reflector was observed at the center of the metal response. The lateral extent of this feature is approximately 2-3 feet. Neither its shape nor lateral breadth is consistent with that of a UST. At A4, an approximately 2 foot by 2 foot flat feature was observed on the crossing GPR profiles at an approximated depth of less than 2 feet. This feature also is not interpreted as representing a possible UST due also to its limited lateral breadth and non-characteristic reflection character.

As we did not observe the source of the metal anomalies at A1 and A2 on the associated GPR profiles, the sources of A1 and A2 may lie deeper than the detection limits of the GPR in these locations (approximately 3 feet). For this reason, we can not rule out the possibility of potential USTs in these locations.

Concrete Foundation Delineation

We initially established approximately 10 GPR traverses over, and adjacent to, several borings within the parking lot where concrete foundation material was encountered at a depth of 2 feet. These traverses ranged in length from 25 to 90 feet long in an attempt to locate the boundary of the concrete material. We were unable to identify a consistent reflecting horizon at the anticipated depth of about two feet (using approximate asphalt/soil/fill GPR velocities) that correlated with the top of the encountered concrete. Our analysis of the data indicated that there is no reinforcement in the concrete which contributed to the lack of detection of the



Cambria Environmental Technology, Inc. September 27, 2001 Page 3

concrete foundation. No combination of gains, filters, or alternative antennae frequencies resulted in the subsurface definition necessary to provide definitive detection of the concrete foundation boundary. We believe that there is not a significant electrical contrast between the concrete and the overlying fill material to detect this interface with confidence and subsequently map out the edge of the presumed foundation. Upon this determination, we recommended that the survey not proceed since the required information could not be obtained.

STANDARD CARE AND WARRANTY

The scope of NORCAL's services for this project consisted of using geophysical techniques to characterize the shallow subsurface. The accuracy of our findings is subject to specific site conditions and limitations inherent to the techniques used. We performed our services in a manner consistent with the level of skill ordinarily exercised by members of the profession currently employing similar techniques. No warranty, with respect to the performance of services or products delivered under this agreement, expressed or implied, is made by NORCAL.

We look forward to future investigations with Cambria Environmental Technology.

Respectfully,

NORCAL Geophysical Consultants, Inc.

Dan P. Jones

Geophysicist 9m

Kenneth G. Blom

Geophysicist GP-887

DPJ/KGB/jm

APPENDIX G

Well Sampling Forms

WELL DEPTH MEASUREMENTS

Well ID	Time	Product Depth	Water Depth	Product Thickness	Well Depth	Comments
MW-1			7.15		14.80	
MW-2			5.75		14.80	
MW-3			6.60		12.60	
MW-4			815		14.25	
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Project Name: Embricade so Case	Project Number: 458 -1705
Measured By: J. W.	Date: 10-12-01

Project Name: Embascadeso Cov	Cambria Mgr: BS	Well ID: MU-1
Project Number: 458- 170	Date: 10-12-01	Well Yield:
Site Address: 1275 Embarcades o Cove	Development Method:	Well Diameter: 2'/
Oakland, Cc		Technician(s): SC,
Initial Depth to Water: 7.15	Total Well Depth: /4-80	Water Column Height: 7.65
Volume/ft: 0.16	1 Casing Volume: 1.22	10 Casing Volumes: 12.2
Purging Device: Aco pump	Did Well Dewater?: yes	Total Gallons Purged: 5

Casing Volume = Water column height x Volume/ ft.

Well Diam	Volume/ft (gallons)
2"	0.16
4"	0.65
6"	1.47

Time	Activity	Water Depth	Gallons Purged						
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	11		3	22.5	1329	7.55	clear		
	**		4	22.4	1384	7.58			
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Project Name: Embascadeso Cov	Cambria Mgr: BS	Well ID: 9 MW-2
Project Number: 458-176	Date: 10-12-01	Well Yield:
Site Address: 1275 Embarcailero Cove	Development Method:	Well Diameter: 2'1
Oakland, Cc		Technician(s): SG
Initial Depth to Water: 5.75	Total Well Depth: 14.80	Water Column Height: 9.05
Volume/ft: 0.16	1 Casing Volume:	10 Casing Volumes: /4.4
Purging Device: Aco pump	Did Well Dewater?:	Total Gallons Purged:

Casing Volume = Water column height x Volume/ ft.

Well Diam.	Volume/ft (gallons)
2"	0.16
4"	0.65
6"	1.47

Time	Activity	Water Depth	Gallons Purged	Comments				
	no s	utge 5.7	\$ \$	temp	/ PH	Cond		
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12:13	<u> </u>		1 4.5	22.4	7.22	1321	11	
12:14			6.0	27.4	7.24	1471	7	•
12:15	~ ~		7.5	22.4	7-27	14 52	17.	
12:16	•1	i	9.0	22.7	7.25	1490		
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	ss: parcadero Cove		Devel	opment Meth	iod:		Well Dia	meter: 2'1	
	land, Cc					,	Technici	an(s): SG	
Initial Dept	h to Water:		Total	Well Depth:			Water Co	olumn Height	:•
Volume/ft:	0.,	16	1 Casi	ng Volume:			10 Casin	g Volumes:	
Purging De			Did W	ell Dewater?	:		Total Ga	llons Purged:	
Casing Volume	= Water column height x Volum	ne/ ft.				Wel	1 Diam. 2" 4" 6"	Volume/ft (gallor 0.16 0.65 1.47	<u>ns)</u>
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Project Name: Embase	adero Cov	Cambria Mgr:	SS	Well ID: MLJ-4	
Project Number: 458		Date: 10-12-0		Well Yield:	
Site Address: 1275 Embascades o Cove		Development Method:		Well Diameter: 2'/	
Oakland, Co		surge block		Technician(s): SG	
Initial Depth to Water:	8.15	Total Well Depth:	14.25	Water Column Height: 6.10	
Volume/ft:	0.16	1 Casing Volume:	0.97	10 Casing Volumes: 9.7 h	
Purging Device: Aco	nmp	Did Well Dewater?:	Λo	Total Gallons Purged:	
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Casing Volume = Water column height x Volume/ ft.

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4"	0.65
6"	1 47

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WELL DEPTH MEASUREMENTS

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MW-2			5.94		14.85	
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MW-U			8.22		14.30	
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Project Name: <u>Embascadeso Cove</u>	Project Number: 458 - 1705
Measured By: Saniv Gill	Date: 10-19-01

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WELL DEPTH MEASUREMENTS

Well ID	Time	Product Depth	Water Depth	Product Thickness	Well Depth	Comments
MW-1	6:20		6-70		14.80	
MW-2	6:10		5.35		14.80	
MW-3	6:15		5.80		obstruction	n no puise
mw-4	6:17	7	7.52		14.25	
MW-1	11.49		7.29			
MW-2	11:43		5.80			
MU-3	11:45		6.95			
MW4	11:47		8.05			
77-7-01						
MW-1-A 12-7-01	550 5:40	`	6.85			
_MW-1	11:40		7.15			

Project Name: Imbarabero Cove	Project Number:
Measured By: A. D.	Date: 12-5-01

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WELL SAMPLING FORM

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		58-1705	Date:			Weil Yiel	
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Project Name: Embarcadero C	Campas	Mgr BS		Well ID: N	·
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Project Number: 453-1705 Site Address: 1275 Embre		Method:		Well Diame	
Cakland, Ca	:	sable bailer		Techniciani	(=):
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Purgung Device: disposable bai	· 			Total Galler	
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Project Nu	mber 4 5 ź	3-1705	Date:	12-5-01		Well Yield:	
Project Na	me: Embaio	indera Co	ue : Cambria	Mga BS	5	Well ID. M	W-C)

WELL DEPTH MEASUREMENTS

Well ID	Time	Product Depth	Water Depth	Product Thickness	Well Depth	Comments
MW-1	9:00		7.08			
MW-2	9:05		5.81			
MW-3	9:10		6-39			
MW-4	9:15		8.04			
			, , , , , , , , , , , , , , , , , , ,			

Project Name:	Project Number: 458-1705
Measured By:	Date: 12-19-0

APPENDIX H

Survey Data

Virgil Chavez Land Surveying

312 Georgia Street, Sutte 225 Vallejo, California 94590-5907 (707) 553-2476 • Fax (707) 553-8698

November 6, 2001 Project No. 2003-37

Kris Cejka Cambria Environmental 1144 65th Street, Suite C Oakland, CA 94608

Subject: Monitoring Well Survey

1275 Embarcadero Cove

Oakland, CA

OF CA

Dear Kris:

This is to confirm that we have proceeded at your request to survey the ground water monitoring wells located at the above referenced location. The survey was performed on October 30, 2001. The benchmark used for the survey was a Port of Oakland benchmark "TP184". The coordinates are based on California State Coordinate System, Zone 3 (NAD 83) as per the Port of Oakland Record of Survey map recorded in Book 18 of Surveys, Page 50 . Measurements taken at approximate north side of top of box and top of casing. Benchmark Elev. = 13.42 feet (Port of Oakland datum).

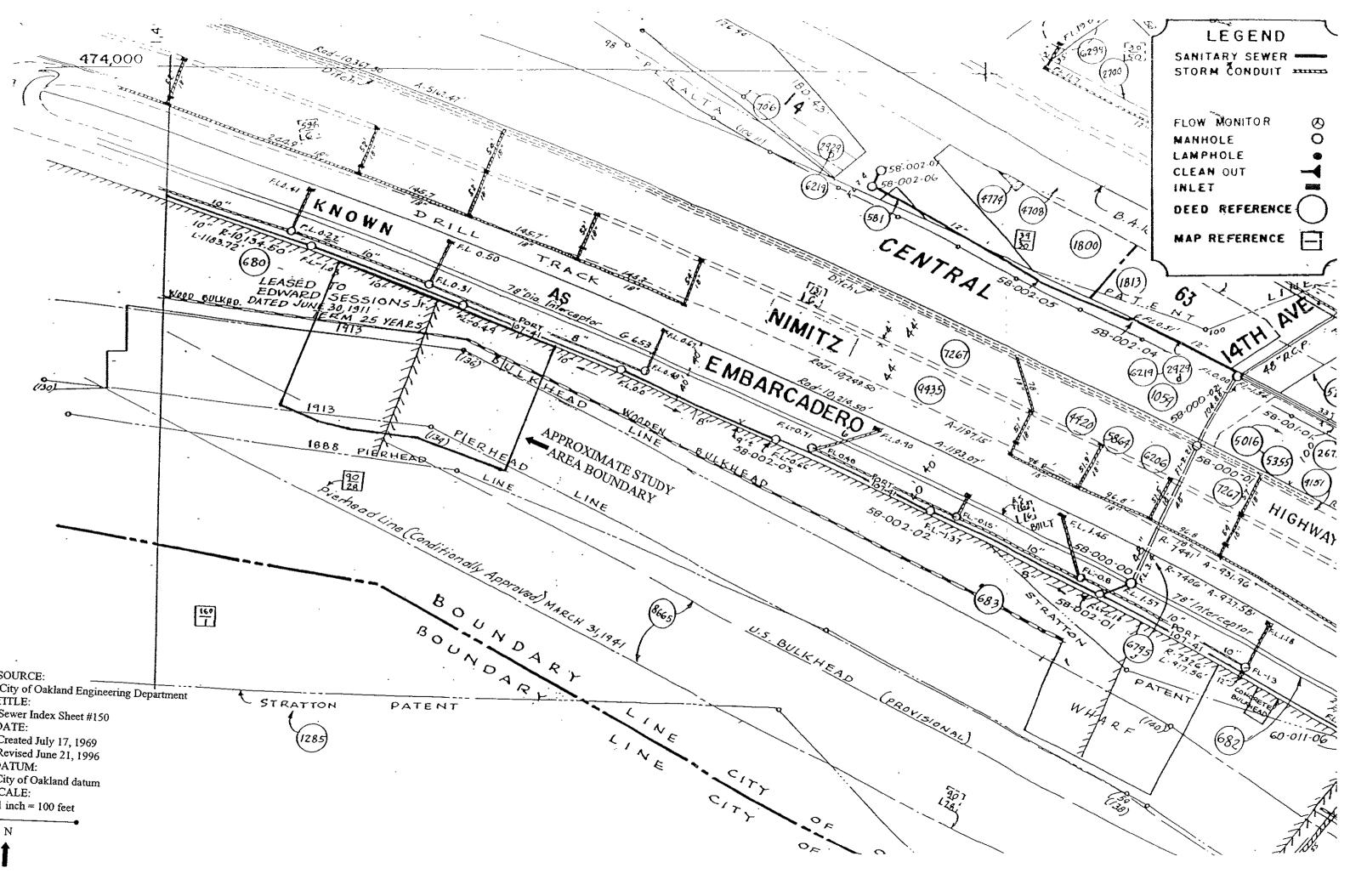
	Rim	TOC		
Well No.	Elevation	Elevation	Northing	Rasting
MW - 1	12.33'	12.03'	2114056.08	6055754.02
MW - 2	11.72'	11.46'	2114007.51	6055751.15
MW - 3	י 79.79	12.49'	2114047.29	6055715.16
MW - 4	13.28'	13.13'	2114088.29	6055740.30

Sincerely,

Virgil D. Chavez, PLS 6823

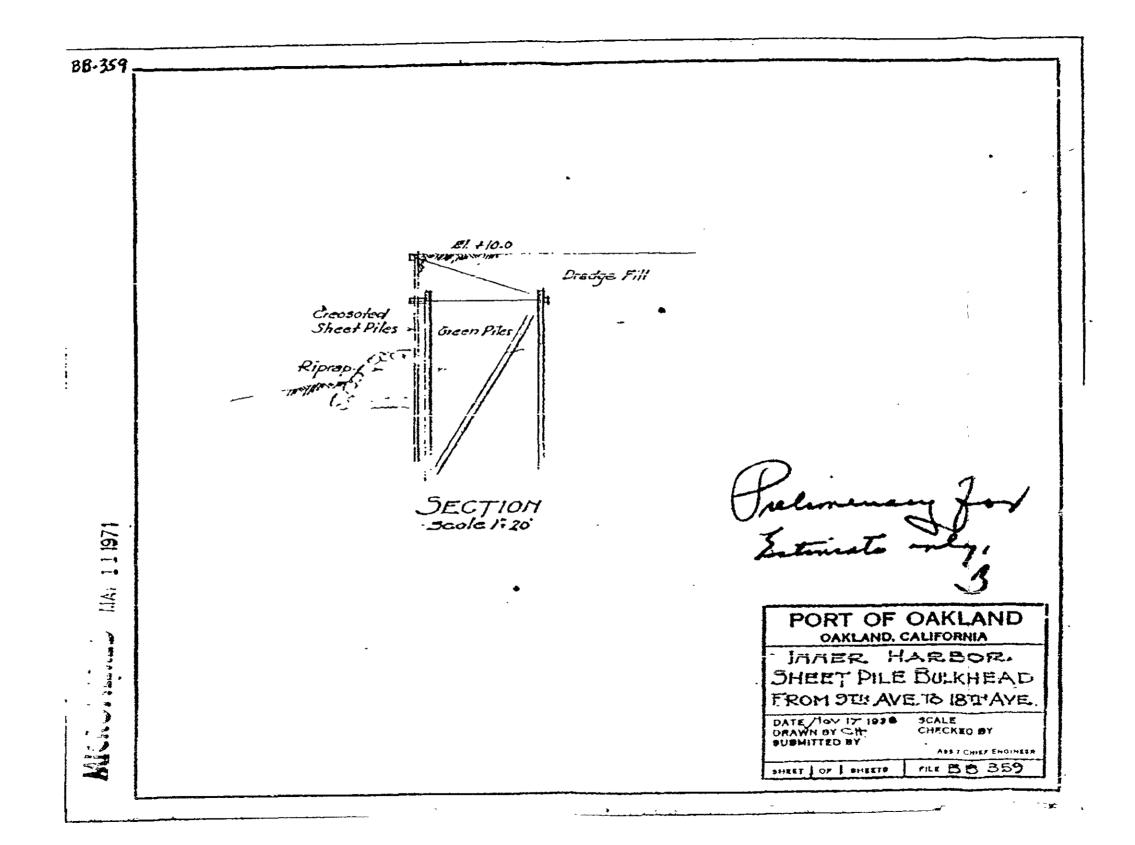
APPENDIX I

City of Oakland Utility Map



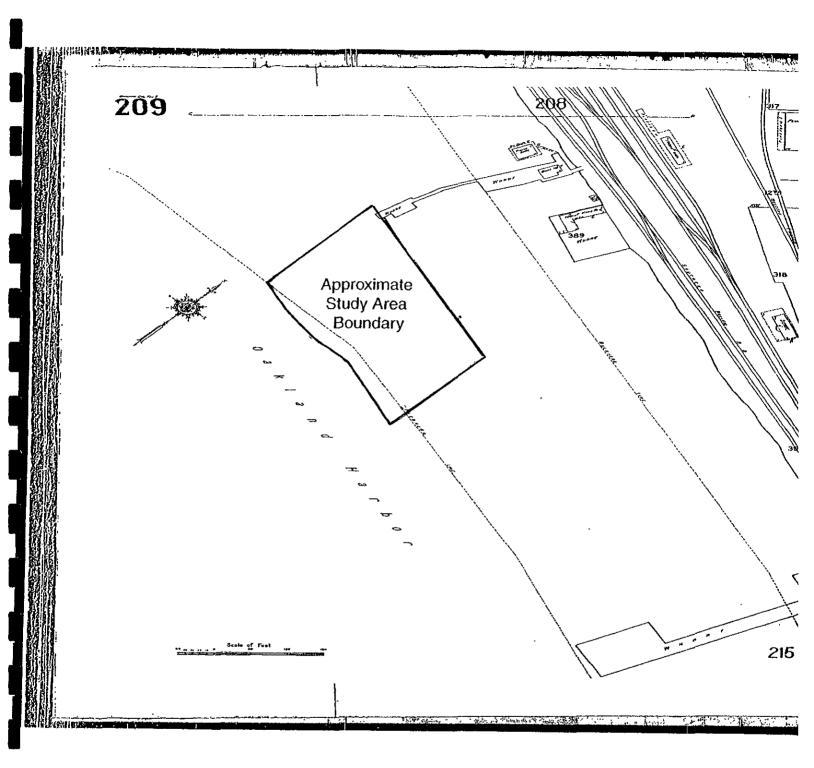
APPENDIX J

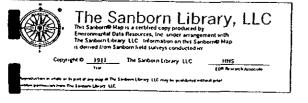
Port of Oakland Inner Harbor Sheet Pile Bulkhead Diagram

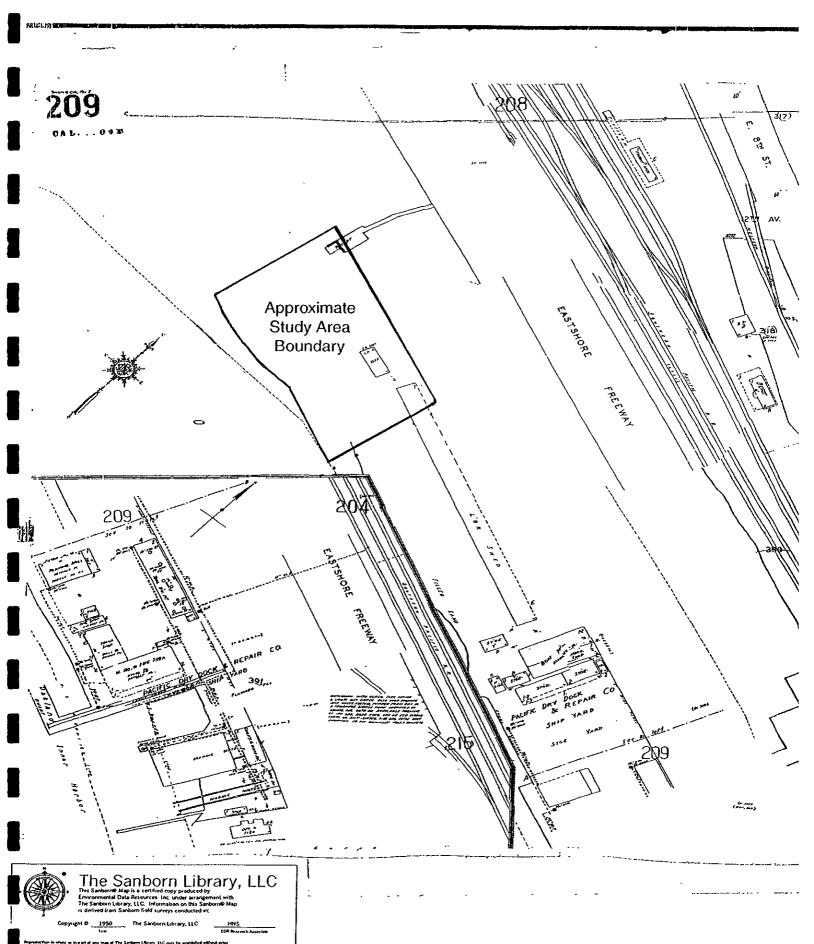


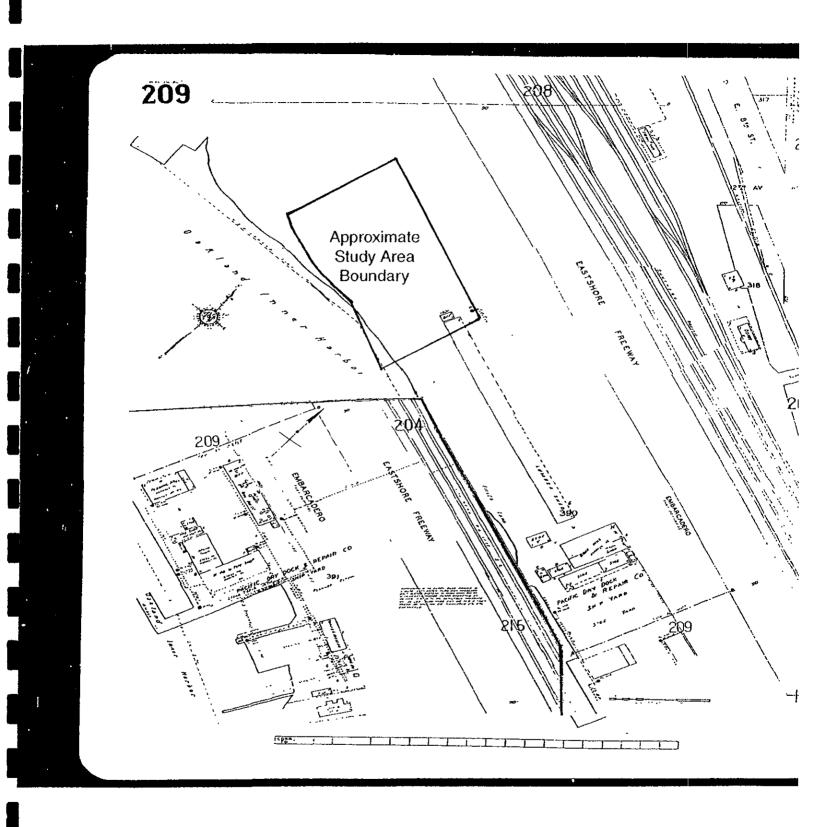
APPENDIX K

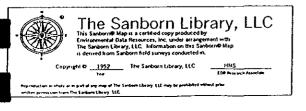
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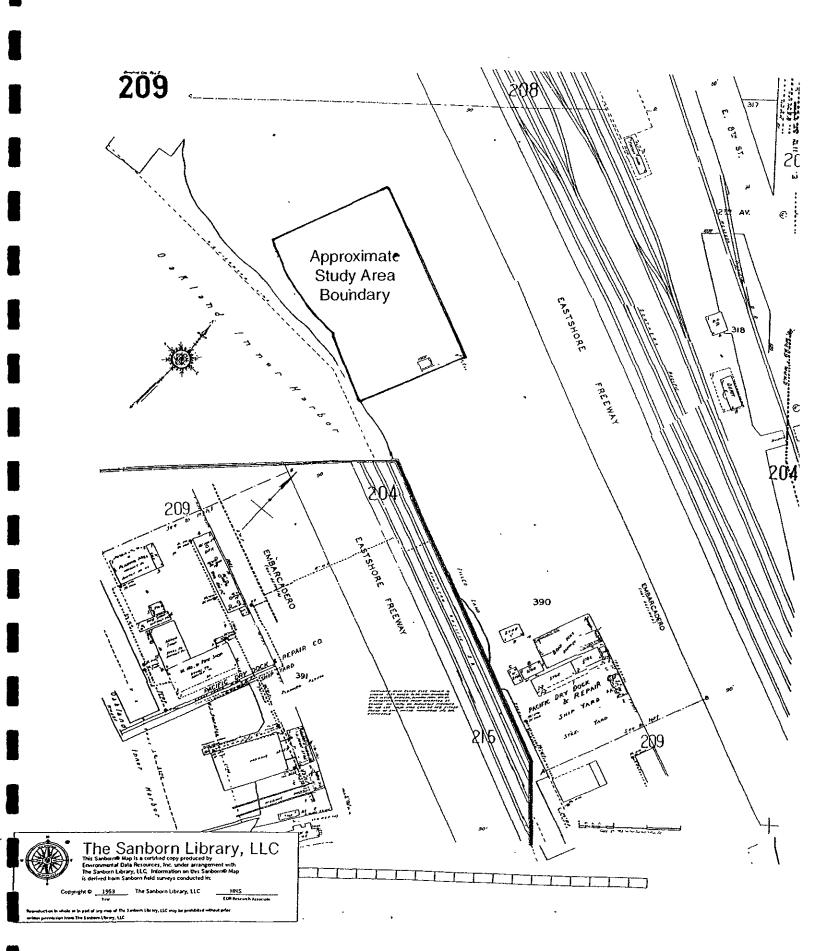


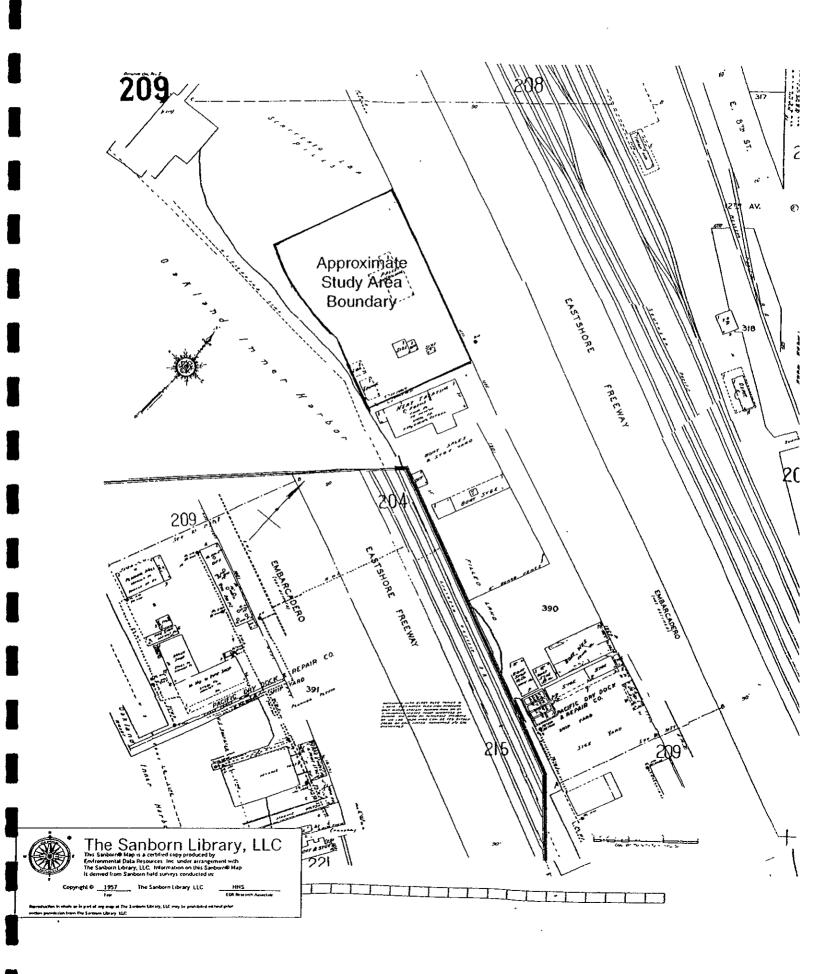


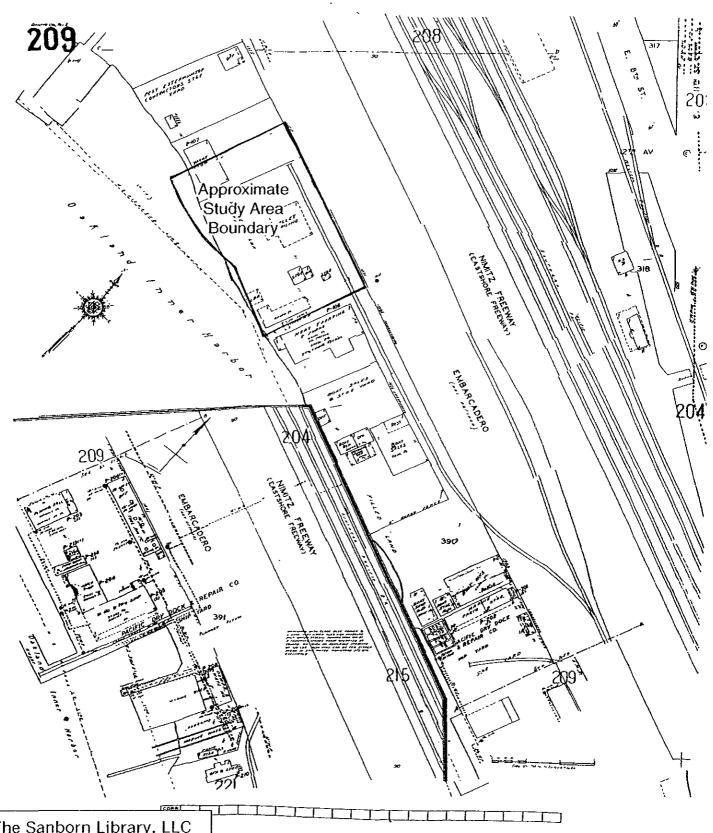












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