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SOIL AND GROUNDWATER SAMPLING REPORT AND HEALTH RISK ASSESSMENT

Rix Property 6460 Hollis Street Emeryville, California

PROTECTION

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Prepared for:

HFH Limited 1355 Ocean Avenue Emeryville, California

July 1998 Project No. 4710.01

Geomatrix Consultants

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1 July 1998 Project 4710.01

Ms. Susan Hugo Alameda County Health Agency 1131 Harbor Bay Parkway, 2nd Floor Alameda, California 94502

Dr. Ravi Arulanantham Regional Water Quality Control Board San Francisco Bay Region 2101 Webster Street Oakland, California 94612

Subject:

Soil and Groundwater Sampling Report and Health Risk Assessment

Rix Property

Emeryville, California

Dear Ms. Hugo and Dr. Arulanantham:

Geomatrix Consultants, Inc., is pleased to submit this report on behalf of HFH, Limited. This report presents the results of soil and groundwater sampling and a human health risk assessment conducted at the subject site. Please contact either of the undersigned if you have any questions or require additional information.

Sincerely,

GEOMATRIX CONSULTANTS, INC.

Jennifer L. Patterson

Project Engineer

Tom Graf, P.E. Principal Engineer

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cc:

Andy Getz; HFH, Limited

Erik Housh; MRE Commercial Real Estate



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SOIL AND GROUNDWATER SAMPLING REPORT AND HEALTH RISK ASSESSMENT RIX PROPERTY 6460 Hollis Street Emeryville, California

1.0 INTRODUCTION

This report summarizes the results of soil and groundwater sampling activities and a human health risk assessment conducted by Geomatrix Consultants, Inc. (Geomatrix), at the Rix Property located at 6460 Hollis Street in Emeryville, California (Figure 1). Work was conducted on behalf of HFH, Ltd., of Emeryville, California, in regards to their potential purchase of the property. The purpose of the work conducted at the site was to provide additional assessment of chemicals in soil and groundwater beneath the site and to address the items outlined by the Alameda County Health Care Services Agency (ACHCSA) in their 25 March 1998 letter to Mr. Frank DeWolf (property owner) that must be addressed prior to receiving site closure. This report outlines site background information, field activities, and results of the work conducted, and addresses each point outlined by the ACHCSA.

2.0 SITE BACKGROUND

The site is currently vacant and consists of one large building and a yard area. Geomatrix reviewed historical Sanborn Fire Insurance maps (Sanborn maps) of the site and reports prepared by others to obtain site background information. A description of on-site investigations and regional information are presented in this section.

The site was undeveloped on the 1911 Sanborn map. Sterling Paint Company operated on the premises from at least 1950. In 1967 and 1969, an industrial adhesives facility is shown on the Sanborn map. Sometime after 1969, Rix Industries began using the site for manufacturing of large air compressors. Rix Industries vacated the property in 1996.



Ten underground storage tanks (USTs) were located on the property. Five of these USTs were located beneath the building and five were located in the yard area east of the building. Reportedly, these USTs historically contained chlorinated solvents, methyl ethyl ketone, alcohols, and ethyl silicate. Groundwater beneath the site is located at a depth of approximately 4 feet below ground surface (bgs) and flows towards the west to southwest.

2.1 PREVIOUS ON-SITE INVESTIGATIONS

In 1992, Hageman-Aguiar, Inc. (HA), of Lafayette, California, sampled the UST contents, advanced soil borings, and installed and sampled three monitoring wells on behalf of Mr. DeWolf. Analytical results of UST contents indicated diesel fuel, secbutanol, mineral spirits, methyl isobutyl ketone (MIBK), and tetrachloroethene (PCE). Eight soil borings were advanced in the vicinity of the five USTs located beneath the building and soil samples were collected at depths of 2 or 4 feet below ground surface (bgs) depending on the compaction of the soil. Soil samples contained petroleum hydrocarbons (up to 400 milligrams per kilogram [mg/kg]) and volatile organic compounds (VOCs; up to 15 mg/kg). Monitoring wells MW-1 and MW-2 were installed inside the building and well MW-3 was installed in the yard area between the USTs and the building (Figure 2). Soil samples were collected from depths of 3 to 7.5 feet bgs during monitoring well installation. These samples contained petroleum hydrocarbons (up to 3,000 mg/kg), ketones (up to 41 mg/kg), and other VOCs (up to 31 mg/kg). These wells were sampled periodically until August 1996. Groundwater samples contained petroleum hydrocarbons (up to 21,000 micrograms per liter pg/li), alcohols (up to 17,000 μ g/l), ketones (up to 97,000 μ g/l) and other VOCs (up to 2200 μ g/l). Concentrations of detected constituents have exhibited a decreasing trend over time. Soil sample analytical results from the borings and monitoring wells are included in Appendix A. Historical analytical results for the monitoring wells are included in Appendix B.



In July 1994, the five USTs located beneath the building were closed in-place by HA. The USTs were triple-rinsed and filled with cement sand slurry. In December 1994, the five remaining USTs in the yard area and approximately 160 tons of overexcavated soil were removed from the site. Ten soil samples were collected from the sidewalls of the UST excavation at depths of 4 feet bgs (the approximate groundwater level). Soil samples contained petroleum hydrocarbons (up to 4.2 mg/kg), alcohols (up to 860 mg/kg), ketones (up to 31 mg/kg), and other VOCs (up to 0.021 mg/kg).

In February 1998, two soil borings were advanced by International Geologic (IG), of Oakland, California, on behalf of a potential buyer to collect grab groundwater samples. Groundwater samples were also collected from the monitoring wells. The borings were located approximately downgradient of the former UST locations along the western property boundary. Analytical results showed concentrations of petroleum hydrocarbons (up to 62,000 μ g/l), and VOCs (up to 86 μ g/l) in the grab groundwater samples. Groundwater samples collected from the monitoring wells contained petroleum hydrocarbons (up to 9,300 μ g/l), and VOCs (up to 173 μ g/l). These results are included in Appendix B.

2.2 REGIONAL INFORMATION

Based on information provided by the ACHCSA, environmental investigations have been conducted at several properties in the vicinity of the site. A monitoring well was installed upgradient of the site at 1372 Ocean Avenue by IG in association with a UST removal. The well is located adjacent to the eastern and upgradient boundary of the site. The analytical results from sampling conducted in March 1998 indicated concentrations of several VOCs at concentrations up to 82 µg/l and petroleum hydrocarbons at concentrations up to 120 µg/l

Environmental investigations have also been conducted at the Ryerson Steel Facility and Lowenberg Property located across Hollis Street and downgradient of the site. VOCs



were detected in groundwater samples at these properties at concentrations up to $170 \mu g/l$ in 1995.

Geomatrix conducted a regional well survey through the Alameda County Public Works Association (ACPWA) and identified one property with an irrigation well approximately 1/3-mile downgradient of the site. The total depth of this well is 470 feet. Other items identified during the well survey appear to be related to shallow groundwater investigations at other industrial properties in the area.

3.0 FIELD ACTIVITIES

In April 1998, Geomatrix performed shallow soil sampling, grab groundwater sampling, and well installation and groundwater monitoring activities. Field activities are discussed below. Prior to beginning field activities, appropriate permits were acquired (Appendix C) and Underground Service Alert (USA) was notified to clear locations for underground utilities. All activities were conducted in accordance with Geomatrix protocols (Appendix D) and samples were sent to American Environmental Network of Pleasant Hill, California, for analysis. Laboratory analyses for the samples collected were chosen based on results of previous site investigations and a review of previous site activities.

3.1 SHALLOW SOIL SAMPLING

Geomatrix collected shallow one soil sample per boring from depths of 0.5 to 2 feet bgs at ten locations (S-1 to S-10) on site. Soil samples were collected using a hand auger; locations are shown on Figure 2. Samples S-1 and S-2, S-7 and S-8, and S-9 and S-10 were composited for analyses for CAM 17 metals by U.S. Environmental Protection Agency (U.S. EPA) Methods 6000/7000 series. Sample S-6 was analyzed discretely for CAM 17 metals . Samples S-3, S-4, and S-5 were analyzed discretely for VOCs according to U.S. EPA Method 8260 and for CAM 17 metals.



3.2 GRAB GROUNDWATER SAMPLING

Four soil borings (GGW-1 through GGW-4) were advanced by Gregg Drilling, Inc., of Martinez, California, for the collection of grab groundwater and soil samples. Sampling locations are shown on Figure 2. Soil borings were advanced to depths of 6 to 12 feet bgs; lithologic logs are included in Appendix E. Soil samples were collected from depths of 1 to 2 feet bgs and analyzed discretely for CAM 17 metals. Grab groundwater samples were analyzed for VOCs according to U.S. EPA Method 8260 and CAM 17 metals. In addition, samples from borings GGW-2 and GGW-3 were analyzed for TPHd and TPHg.

3.3 MONITORING WELL INSTALLATION AND DEVELOPMENT

One monitoring well (MW-4) was installed by Gregg Drilling, Inc., adjacent to the site, downgradient of the former UST locations. The well was constructed of 2-inch-diameter polyvinyl chloride (PVC) well casing to 3 feet bgs and 10 feet of 2-inch-diameter factory-slotted PVC well screen from 3 to 13 feet bgs. The filter pack is Monterey 2/12 sand and the well is sealed with 6 inches of bentonite and 1.5 feet of cement. Well construction details are included in Appendix E.

Approximately 24-hours after installation, the well was developed by Gregg Drilling using a method of surging and then purging the well. The existing monitoring wells (MW-1 through MW-3) were also redeveloped at this time to remove any sediment that may have built up.

3.4 GROUNDWATER MONITORING

Three days after the wells were developed, Geomatrix collected groundwater samples according to Geomatrix protocols. Groundwater samples were analyzed for VOCs according to U.S. EPA Method 8260. In addition, samples from MW-2 and MW-4 were analyzed for CAM 17 metals, and an additional groundwater sample from MW-4 was



sent to Friedman & Bruya, of Seattle, Washington, for hydrocarbon fingerprinting. The groundwater samples collected for VOC and CAM 17 analyses were analyzed by the laboratory after the holding time had expired. Analytical results for MW-1 through MW-3 were similar to those from the February 1998 sampling event, so the wells were not resampled. Since no basis for comparison was available for MW-4, a second sample was collected for analysis for VOCs, which was conducted within the holding time.

4.0 ANALYTICAL RESULTS

This section presents the analytical results of soil and groundwater sampling conducted at the site by Geomatrix. Analytical results are summarized in Tables 1 through 4 and on Figures 3 through 5. Laboratory analytical data sheets are included in Appendix F.

4.1 SHALLOW SOIL SAMPLES

Thirteen metals were detected in shallow composite and discrete soil samples collected from borings S-1 through S-10 and GGW-1 through GGW-4. Analytical results for metals in soil samples are presented in Table 1 and on Figure 3. Metals concentrations are compared to concentrations considered to be indicative of regional background to assist in evaluating the data. The regional background data used for comparison were collected by the Lawrence Berkeley National Laboratory (LBNL) for the purpose of establishing background concentrations of selected metals in the vicinity of the laboratory (LBNL, 1995). A chemical was considered to be present above representative background concentrations if the maximum concentration of the site data was above the 95 percent upper tolerance limit (95% UTL) of the LBNL data. Barium, cadmium, chromium, copper, lead, mercury, and zinc were detected in one or more samples at concentrations greater than background levels. Elevated metals concentrations were detected in samples collected from both the yard area and beneath the building slab.



Based on the elevated lead concentrations in some soil samples, a waste extraction test (WET) for lead was performed on two of the composite samples (S-7,8 and S-9,10) to determine disposal options available for future construction. Lead was detected at concentrations of 15 mg/l in the samples S-7,8 and at 8.7 mg/l in sample S-9,10. The soluble threshold limit concentration (STLC) for lead is 5 mg/l. Based on these concentrations, excavated near-surface soil at the site will likely require disposal as a California hazardous waste. However, lead was not detected at concentrations above laboratory reporting limits in groundwater samples collected at the site (see Sections 4.2 and 4.3).

VOCs were detected in the three soil samples analyzed (S-3, S-4, and S-5) at concentrations ranging from 0.03 to 6.5 mg/kg. Analytical results for VOCs in soil are presented in Table 2 and on Figure 4. These samples were collected from beneath the building slab.

4.2 GRAB GROUNDWATER SAMPLES

The analytical results for the grab groundwater samples analyzed for metals and TPH are presented in Table 3. Arsenic, barium, cobalt, molybdenum, nickel, vanadium, and zinc were detected in one or more of the four grab groundwater samples analyzed at concentrations ranging from 6 to 520 μ g/l. Of the two samples analyzed for TPHg and TPHd, only TPHg was detected in the sample and duplicate sample collected from boring GGW-3, at concentrations of 220 and 240 μ g/l.

Analytical results for the four samples analyzed for VOCs are presented in Table 4 and on Figure 5. Vinyl chloride and 1,1-dichloroethane (1,1-DCA) were detected in the sample collected from GGW-2, located in Hollis Street, downgradient of the site, at concentrations of 120 and 11 µg/l, respectively. The sample and duplicate sample collected from GGW-3, located within the yard area, contained several VOCs at concentrations ranging from 5 to 120 µg/l. The samples collected from GGW-1 and



GGW-4 did not contain VOCs above laboratory reporting limits; however, the reporting limits were elevated due to matrix interferences. A sheen was noted in this well during sampling activities.

4.3 MONITORING WELL SAMPLES

Groundwater samples collected from wells MW-2 and MW-4 were analyzed for metals. Analytical results for metals in groundwater are presented in Table 3. Arsenic, barium, and nickel were detected in both samples and molybdenum was detected in the sample collected from MW-2.

Samples collected from all four monitoring wells were analyzed for VOCs. Analytical results are included in Table 4 and on Figure 5. Several VOCs were detected in the groundwater samples at concentrations ranging from 5 to 250 μ g/l. Results were generally consistent with past monitoring events in the existing wells. Two samples were analyzed from the new downgradient well, MW-4. Three constituents were detected in one of the samples; sec-butylbenzene at a concentration of 15 μ g/l, isopropylbenzene at a concentration of 16 μ g/l, and n-propylbenzene at a concentration of 5 μ g/l. The second sample did not contain any constituents above laboratory reporting limits.

The hydrocarbon fingerprinting analysis performed on the groundwater sample from MW-4 indicated the presence of diesel fuel that had undergone evaporative and biological degradation.

5.0 HEALTH RISK ASSESSMENT

This section presents an assessment of the potential risks to human health and the environment associated with the presence of chemicals detected in soil and groundwater at the site assuming continued commercial use of the property. The results of the human health risk assessment (HHRA) are discussed below. An ecological risk assessment was



not considered necessary for the following reasons. First, current and planned future use of this site is for commercial purposes, and the majority of the site is or will be covered by buildings, pavement, and landscaping. Second, the site is located within a commercial/industrial area approximately ½-half mile from the nearest surface water body, the San Francisco Bay. Therefore, no further assessment of potential risks to non-human receptors is warranted.

The HHRA is organized into sections that parallel the risk assessment process outlined by the National Academy of Science (1983) and U.S. EPA (1989), including data evaluation, exposure assessment, toxicity assessment, and risk characterization. The primary guidance used in this HHRA are California Environmental Protection Agency's (Cal-EPA's) "Supplemental Guidance for Human Health Multimedia Risk Assessments of Hazardous Waste Sites and Permitted Facilities" (1992), U.S. EPA's "Risk Assessment Guidance for Superfund, Volume 1, Human Health Evaluation Manual, Parts A and B" (1989; 1991a), and U.S. EPA's "Human Health Evaluation Manual, Supplemental Guidance: Standard Default Exposure Factors" (1991b). Cal-EPA's "Preliminary Endangerment Assessment (PEA) Guidance Manual" (1994a) was used, as appropriate, in the development of chemical-specific parameters and exposure pathway assumptions.

5.1 DATA EVALUATION

Data evaluation is the process of analyzing site characteristics and analytical data to identify chemicals of potential concern (COPCs) to be evaluated in the HHRA. As described previously, several VOCs have been detected in soil and groundwater beneath the site. For purposes of this HHRA, all of the detected organic chemicals were identified as COPCs. Several naturally occurring metals also were detected in soil and groundwater. As discussed further in Section 5.2, potential exposure to chemicals in groundwater is limited to inhalation of vapors; therefore, metals detected in groundwater are not considered further in the HHRA. For metals detected in soil, the analytical data for each chemical, including chemical name, total number of samples analyzed, total number of detections, range of detected concentrations, and range of detection limits are



summarized in Table 5. Metals in soil were identified as COPCs according to the following criteria:

Primary Criterion

If the maximum detected concentration is greater than the U.S. EPA Region 9 preliminary remediation goal (PRG) for residential soil, then that chemical was further evaluated using the Secondary Criterion described below. If the maximum detected concentration is less than the appropriate PRG, then that chemical was not considered further in the HHRA.

• Secondary Criterion

If the maximum detected concentration is greater than the concentration in soil considered indicative of regional background, then that chemical was identified as a COPC. If the maximum detected concentration is less than the background concentration, then that chemical was not considered further in the HHRA.

The representative background data used as part of the secondary criterion were collected by LBNL for the purpose of establishing background concentrations of selected metals in the vicinity of the laboratory (LBNL, 1995). The LBNL data set is discussed in more detail in Section 4.1. A chemical was considered to be present in soil at concentrations within background if the maximum detected concentration was below the 95% UTL of the LBNL data. The metals in soil identified as COPCs are presented in Table 5. As stated previously, three of the six samples analyzed for metals were each composites of two discrete samples. Even if the results from these samples were doubled (i.e., assuming that the total concentration detected was in one of the discrete samples), the same metals would be identified as COPCs in soil.

5.2 EXPOSURE ASSESSMENT

Exposure assessment is the process of measuring or estimating the intensity, frequency, and duration of human exposure. The principal elements of exposure assessment consist of:



- evaluation of the influence of fate and transport processes for the COPCs,
- identification of potential exposure scenarios (i.e., receptors) and exposure pathways,
- calculation of representative chemical concentrations, and
- estimation of potential chemical uptake.

A site conceptual model was created based on the soil and groundwater conditions, current and planned future land use, and physical and chemical characteristics of the chemicals potentially present at the site. As described in the U.S. EPA's "Guidance for Conducting Remedial Investigations and Feasibility Studies under CERCLA" (1988a), the purpose of a site conceptual model is to describe what is currently known about chemical sources, likely migration pathways, exposure routes, and possible exposure scenarios so that the data necessary to complete a risk assessment are identified. Figure 6 presents the site conceptual model for this site. Each of the components of the site conceptual model is discussed in the following sections.

The exposure assessment reflects reasonable maximum exposure (RME) conditions for each receptor. The U.S. EPA defines the RME scenario as "The highest exposure that is reasonably expected to occur at the site (U.S. EPA, 1989). In addition, the U.S. EPA states that "the intent of the RME is to estimate a conservative exposure case (i.e., well above average) that is still within the range of possible exposures" (U.S. EPA, 1989).

5.2.1 Environmental Fate and Transport of Chemicals of Potential Concern

The environmental fate of the COPCs is controlled by the cumulative interaction of transport and transformation processes. Once released into the environment, a chemical may partition among various environmental media (soil, water, air). The transport processes that define the movement of chemicals between media are highly dependent on the physicochemical properties of both the chemicals and the environmental media.

The COPCs can be divided into two categories: VOCs and metals. The following paragraphs provide a brief overview of the fate and transport characteristics of each of



these general categories. Several physical constants, including molecular weight, Henry's Law constant, and organic carbon partitioning coefficient are presented for the organic COPCs in Table 6. These physical constants were used to identify whether or not a chemical may be considered volatile. It was assumed that a COPC is volatile if its Henry's Law constant is greater than 10⁻⁵ atmospheres-cubic meters per mole (atm-m³/mole) and its molecular weight is less than 200 grams per mole (U.S. EPA, 1991a).

Volatile Organic Chemicals

VOCs have relatively high vapor pressures, and relatively high solubilities in water and organic solvents. Chemicals that are volatile and water soluble tend to have low soil adsorption and bioaccumulative potentials. Based on these characteristics, these chemicals are relatively mobile in soil and may volatilize into the air or leach to groundwater.

Metals

Lead was the only metal identified as a COPC. Lead occurs naturally in soil, is non-volatile, and adheres tightly to soil particles. In general, the mobility of lead in the environment depends on the pH of the surrounding medium. Under acidic conditions, lead may become soluble and be mobilized through soil by infiltrating water or can be dissolved directly into groundwater if present in the saturated zone. Otherwise, lead is relatively immobile in soil and generally exhibits low water solubility. Lead is not subject to biodegradation and will persist in the environment (NRC, 1994; Freeze and Cherry, 1979).

Environmental media (e.g., soil, air, groundwater) that may significantly contribute to an individual's exposure to a chemical can be determined based on this general understanding of the environmental transport mechanisms of the COPCs. As shown in Figure 6, COPCs at the site may be present in surface and subsurface soil, air, or groundwater.



5.2.2 Identification of Potential Receptors and Exposure Pathways

Based on the current and planned future use of the site for commercial purposes, potential receptors are limited to current and future on-site workers. As shown in Figure 6, current and future on-site workers may come into contact with chemicals in surface soil or air. Potential exposure to future construction workers who may intrude into subsurface soil or shallow groundwater will be addressed as part of a site-specific Health and Safety Plan at the time of construction.

Exposure pathways are the means through which an individual may contact a chemical in the environment. Exposure pathways are governed by environmental conditions (e.g., depth to groundwater or prevailing wind direction); by the potential for the chemical to move from one medium (water or air) to another; and by the general lifestyles and/or work activities of potentially exposed populations (e.g., construction work or office work). Although several potential pathways may exist, only a few may be complete. For a pathway to be complete, each of the following elements must exist:

- a source and mechanism for chemical release,
- an environmental transport medium (e.g., air, water),
- a point of potential contact with the medium, and
- an exposure route at the contact point (e.g., inhalation, dermal contact).

Current and future on-site workers may be exposed to COPCs in surface soil outside the existing building via incidental ingestion and dermal contact. In the event that the existing building is replaced, future on-site workers are not expected to come into contact with soil because, after construction, it is expected that the newly constructed building or buildings, asphalt or concrete pavement, and landscaping will cover the site. Current and future on-site workers may also be exposed to COPCs present in indoor air as vapors (originating from soil or groundwater).

Shallow groundwater beneath and in the vicinity of the site is not currently used as a drinking water source. A regional well survey identified only one irrigation well



approximately 1/3-mile downgradient of the site. The total depth of this well is 470 feet. Other wells identified during the survey appear to be related to shallow groundwater investigations at other industrial properties in the area. Given the generally poor quality of the shallow groundwater due to historical industrial practices in the City of Emeryville and the possible impacts from sewers and storm drains, it is very unlikely that shallow groundwater will be used as a municipal source of water in the foreseeable future. Therefore, potential exposure to chemicals in groundwater associated with domestic use was not evaluated in this HHRA.

5.2.3 Exposure Point Concentrations

Estimates of exposure point concentrations are necessary for evaluating chemical uptake in exposed individuals. For purposes of evaluating an RME scenario, the U.S. EPA recommends that the exposure point concentration be equal to the 95-percent upper confidence limit (95% UCL) of the arithmetic mean of a given set of chemical concentration data to account for the uncertainty in estimating the true average concentration at a site (U.S. EPA, 1992a). In this case, the COPCs in soil and groundwater were detected relatively infrequently and at relatively low concentrations. Therefore, to be conservative, the maximum detected concentrations were used (see Table 7).

The concentrations of volatile COPCs in air were estimated based on the representative concentrations in soil and groundwater. As shown in Appendix G, the emission rates of the COPCs in air above these media were estimated using the VOC emissions model (Cal-EPA, 1994a). The resulting emission rates were used in conjunction with a box model to estimate indoor air concentrations for current and future on-site workers. Several physical constants required for these models are summarized in Table 6. In some cases, physical constants were not readily available and had to be calculated; these calculations are shown in Appendix H.



5.2.4 Exposure Equations and Input Parameters

This section presents the exposure equations and input parameters used to calculate the annual average daily dose (AADD) and lifetime average daily dose (LADD) for each of the identified routes of exposure. The AADD and LADD both provide quantitative estimates of an individual's daily exposure to a chemical. The difference between the two estimates is the time over which the exposure is averaged. Noncarcinogenic health effects are assumed to occur only after a threshold dose is reached (see Section 5.3.1); therefore, the AADD represents the average daily dose during the period of exposure. Conversely, carcinogenic health risks are not considered to be threshold phenomena (see Section 5.3.2). Therefore, the LADD represents the average daily dose over a lifetime.

As discussed in Section 5.3.3, potential noncarcinogenic health risks associated with exposure to lead are evaluated using Cal-EPA's Lead Spread Model rather than calculating an AADD. Therefore, the only exposure pathway for which an AADD or LADD was calculated is inhalation of vapors. The components of this equation are presented below:

Inhalation of Vapors

$$Dose_{tv} = \frac{Cin \times IRa \times EF \times ED}{BW \times AT}$$

where:

Dose_{iv} = Dose received through inhalation of vapors originating from soil or groundwater (mg/kg-day)

or groundwater (mg/kg-day)

Cin = Chemical concentration in indoor air (mg/m³)

IRa = Inhalation rate (m^3/day)

EF = Exposure frequency (days/year)

ED = Exposure duration (years)

BW = Body weight (kg)

AT = Averaging time (days)



Values used for each input parameter, including the source and rationale, are summarized in Table 8. In all cases, the input parameters are equal to conservative default assumptions prescribed by Cal-EPA (e.g., 1992) or U.S. EPA (e.g., 1991b).

5.3 TOXICITY ASSESSMENT

The purpose of the toxicity assessment is two-fold (U.S. EPA, 1989):

- to evaluate available information regarding the potential for a chemical to cause adverse health effects in exposed individuals (hazard identification), and
- to estimate the relationship between the extent of exposure and the increased likelihood (e.g., probability or chance) and/or severity of adverse effects (dose-response assessment).

Hazard identification entails determining if a chemical can cause an increase in a particular adverse effect (e.g., cancer) and the likelihood that the adverse effect will occur in humans. The result of the hazard identification is a toxicity profile that summarizes the available toxicological information and its relevance to human exposure under conditions present at the site. Dose-response assessment entails quantifying the relationship between the dose of a contaminant and the incidence of adverse effects in the exposed population. The result of the dose-response assessment is toxicity criteria that are used in the risk characterization to estimate the likelihood of adverse effects occurring in humans at different exposure levels. The toxicity criteria used to evaluate noncarcinogenic and carcinogenic health risks are commonly referred to as reference doses (RfDs) and slope factors (SFs), respectively. The basis for these criteria is described briefly in the following sections.

5.3.1 Toxicity Criteria for Noncarcinogenic Health Risks

Observable noncancer biological effects of chemical substances occur only after a threshold dose is reached. For the purposes of establishing health criteria, this threshold dose is usually estimated from the no-observed adverse effect level (NOAEL) or the lowest-observed adverse effect level (LOAEL) determined in chronic animal exposure



studies. The NOAEL is defined as the highest dose at which no adverse effects occur, whereas the LOAEL is defined as the lowest dose at which adverse effects begin to occur. Cal-EPA, U.S. EPA, and other regulatory agencies use NOAELs and LOAELs derived from animal studies to establish RfDs to evaluate human intake of noncarcinogenic compounds. RfDs, which are expressed in terms of mg/kg-day, are criteria intended to represent the dose of a chemical that is not expected to cause adverse health effects over a lifetime of daily exposure, even in sensitive individuals, with a substantial margin of safety. The associated noncarcinogenic toxicity criteria used in the HHRA are presented in Table 9.

5.3.2 Toxicity Criteria for Carcinogenic Health Risks

Regulatory guidance assumes that chemicals that are carcinogenic should be treated as if they do not have thresholds (U.S. EPA, 1989). This approach assumes that the dose-response curve for carcinogens only allows for zero risk at zero dose (i.e., for all doses, some risk is assumed to be present). To estimate theoretically plausible responses at these low doses, various mathematical models are used. The accuracy of the projected risk depends on how well the model predicts the true relationship between dose and risk at dose levels where the relationship cannot be actually measured. The accuracy of these models is currently unknown, but they are believed not to underestimate the true risk.

Health risks for exposure to carcinogens are defined in terms of probabilities that quantify the likelihood of a carcinogenic response in an individual receiving a given dose of a particular compound. The SF, which is expressed in units of (mg/kg-day)⁻¹, is defined as the 95% UCL of the probability of a carcinogenic response per unit daily intake of a chemical over 70 years. By using the 95% UCL, the estimate of carcinogenic response will be conservative and will purposefully overestimate the actual risk posed by the chemical. The associated carcinogenic toxicity criteria used in the HHRA are presented in Table 10.



5.3.3 Toxicity Criteria for Lead

Noncarcinogenic toxicity criteria have not been established for lead. Instead, the potential for noncarcinogenic health effects associated with exposure to lead in soil were evaluated using the Cal-EPA Lead Spread Model (Cal-EPA, 1992). This mathematical model estimates blood-lead levels resulting from contact with lead in various environmental media (e.g., soil, air, food). The blood-lead level is of interest, because, unlike other chemicals, sufficient toxicology data exist to correlate the body burden of lead measured by blood-lead level with a specific adverse effect (e.g., a blood-lead level of "x" is associated with a particular adverse health effect).

5.4 RISK CHARACTERIZATION

Risk characterization represents the final step in the risk assessment process, in which the results of the exposure and toxicity assessments are integrated into quantitative or qualitative estimates of potential health risks. Noncarcinogenic and carcinogenic health risks are characterized separately, as described below.

5.4.1 Noncarcinogenic Health Risks

Potential noncarcinogenic adverse health effects were evaluated by comparing the AADD to the appropriate RfD. This comparison is expressed in terms of a "hazard quotient" and was calculated as follows:

$$Hazard\ Quotient = \frac{AADD}{RfD}$$

A hazard quotient less than or equal to 1 indicates that the predicted exposure should not pose a significant noncarcinogenic health risk (U.S. EPA, 1989). In cases where individual chemicals potentially act on the same organs or result in the same health endpoint (e.g., respiratory irritants), additive effects can be addressed by calculating a hazard index as follows:



Hazard Index =
$$\sum_{a}^{n}$$
 Hazard Quotient_a + Hazard Quotient_b +...+ Hazard Quotient_n

A hazard index of less than or equal to 1 is indicative of acceptable levels of exposure for chemicals having an additive effect. In this HHRA, a screening-level hazard index was calculated by summing the hazard quotients for all chemicals, regardless of toxic endpoint. This approach likely overestimates the potential for noncarcinogenic health effects due to simultaneous exposure to multiple chemicals; however, it can be used as a screening tool to rapidly eliminate those exposure scenarios that do not pose a noncarcinogenic health risk.

The potential noncarcinogenic health risks associated with exposure to the COPCs by current or future on-site workers are summarized in Table 11; the calculation spreadsheets are presented in Appendix I. The total hazard index is 0.009. This value is considerably less than 1, indicating that exposure to chemicals at the site should not pose a noncarcinogenic health risk to current or future on-site commercial workers under the conditions evaluated.

5.4.2 Carcinogenic Health Risks

Carcinogenic health risks are defined in terms of the probability of an individual developing cancer as the result of exposure to a given chemical at a given concentration and are dependent on the LADD and the appropriate SF. In this assessment, theoretical lifetime excess cancer risks were estimated as follows:

Theoretical Lifetime Excess Cancer Risk = LADD x SF

As with hazard indices, the estimated cancer risks for each chemical and exposure route were summed to estimate the total excess cancer risk for the exposed individual. The U.S. EPA considers estimates of lifetime excess cancer risk associated with exposure to chemicals of less than one-in-one-million $(1x10^{-6})$ to be *de minimis*. Risks within the range of one-in-one-million to one hundred-in-one-million $(1x10^{-4})$ may be acceptable



depending on other factors (e.g., site-specific considerations) (U.S. EPA, 1990a and 1990b). In addition, the California Safe Drinking Water and Toxic Enforcement Act of 1986 (Proposition 65) considers lifetime excess cancer risks less than ten-in-one-million (1x10⁻⁵) as being insignificant for purposes of the Act (i.e., no warning to the potentially exposed population is required) (22 California Code of Regulations [CCR] 12721(d)(3)).

The potential carcinogenic health risks associated with exposure to COPCs by current or future on-site workers are summarized in Table 12; the calculation spreadsheets are provided in Appendix I. The total theoretical lifetime excess cancer risk is 4×10^{-8} . This value is below the generally accepted risk range of 10^{-4} to 10^{-6} , indicating that exposure to chemicals at the site should not pose an unacceptable carcinogenic health risk to current or future on-site commercial workers under the conditions evaluated.

5.4.3 Lead Health Risks

As discussed in section 5.3.3, the Cal-EPA's Lead Spread Model was used to evaluate potential health risks associated with the presence of lead in soil. The Centers for Disease Control (CDC) have established a blood-lead level of concern at 10 micrograms per deciliter (Tg/dl) of whole blood for children. The Cal-EPA has adopted the 10 Tg/dl significance level established by the CDC for both children and adults (Cal-EPA, 1992); however, this standard may be overly conservative for evaluating potential risks to adults. A more appropriate value may be 30 Tg/dl, as recommended in the California Code of Regulations (CCR) Title 8 §1532, Lead in Construction Standard.

The output from the Lead Spread Model is presented in Appendix J. The estimated blood lead level at the 99th percentile is 4.9 Tg/dl for the current or future on-site worker. This value is well below the level of concern adopted by the Cal-EPA, indicating that exposure to lead in soil should not pose an unacceptable risk to current or future on-site workers.

5.4.4 Uncertainty Analysis

Uncertainty is inherent in many aspects of the risk assessment process, and generally arises from a lack of knowledge of (1) site conditions, (2) toxicity and dose-response of



the COPCs, and (3) the extent to which an individual will be exposed to those chemicals. The assumptions that introduce the greatest amount of uncertainty and their effect on the noncarcinogenic and carcinogenic risk estimates are discussed below. In general, assumptions were selected in a manner that purposefully biases the process toward health conservatism.

Exposure Point Concentrations

The maximum detected concentrations in soil and groundwater were used as the exposure point concentrations in this HHRA. This assumption should overestimate the concentrations to which an individual would be exposed over an extended period of time.

Exposure Assumptions and Parameters

Default Cal-EPA and U.S. EPA exposure assumptions and parameters were used throughout this HHRA. These values are established with the intent of being conservative to account for the highest exposure that could reasonably be expected to occur for a given exposure pathway at a site. For example, this HHRA assumes that a future on-site commercial worker will work at the site for 25 years; however, in 1992 the Bureau of Labor Statistics reported that the average number of years an individual works in one job is 4.5 years (Bureau of Labor Statistics, 1992). These upper-bound estimates of exposure most likely overestimate the potential health risks associated with exposure to the COPCs in soil and groundwater.

Toxicity Criteria

One of the largest sources of uncertainty in any risk assessment is associated with the toxicity criteria used to estimate noncarcinogenic and carcinogenic health risks. The noncarcinogenic toxicity criteria (i.e., oral and inhalation RfDs) incorporate multiple uncertainty factors to account for limitations in the quality of available data (e.g., animal data in lieu of human data). The carcinogenic toxicity criteria (i.e., oral and inhalation SFs) are based on a mathematical extrapolation from high to low doses. Both of these methods are conservative to ensure that the noncarcinogenic and carcinogenic health risks are far more likely to be overestimated than underestimated.



These and other assumptions contribute to the overall uncertainty in the results of the HHRA. However, given that the largest sources of uncertainty generally result in overestimates of exposure or risk, it is believed that the noncarcinogenic and carcinogenic risks presented in this HHRA represent conservative estimates of the risks, if any, posed by the site.

5.5 CONCLUSIONS OF THE RISK ASSESSMENT

The HHRA presents an evaluation of the potential human health risks associated with the presence of chemicals soil and groundwater at the site assuming continued commercial use of the property. The HHRA was completed in accordance with multiple risk assessment guidance documents provided by Cal-EPA and U.S. EPA. Potential noncarcinogenic and carcinogenic health risks were estimated for current and future on-site workers assuming conservative estimates of human exposure. Potential exposure to future on-site construction workers will be addressed as part of a site-specific Health and Safety Plan at the time of construction.

As shown in Tables 11 and 12, the estimated hazard indices and theoretical excess cancer risks for current and future on-site workers are at or below the levels generally considered acceptable by regulatory agencies (i.e., a hazard index of 1 for noncarcinogens and a cancer risk of $1x10^{-4}$ to $1x10^{-6}$ for carcinogens) under the conditions evaluated. In addition, the estimated blood-lead levels are well below a level of regulatory concern. Therefore, the presence of chemicals at the site should not pose an unacceptable noncarcinogenic or carcinogenic risk to current or future on-site workers assuming continued commercial use of the site.

6.0 CONCLUSIONS

This section will address the guidelines for evaluating sites for closure as a low risk groundwater case as identified by the ACHCSA in their 25 March 1998 letter.



The leak has been stopped and on-going sources have been removed or remediated.

The five USTs located under the building at the site were closed in-place in July 1994. The five USTs located in the yard and 161.54 tons of soil were removed in December 1994. Confirmation soil sampling from around the USTs and groundwater concentrations detected in samples collected from monitoring wells within and downgradient of the former and existing UST locations do not indicate that residual chemicals in site soil are a significant ongoing source to groundwater.

2) The site has been adequately characterized.

ACHCSA stated in their 25 March 1998 letter that "additional downgradient sampling will be required, [and] groundwater monitoring must be performed continuously to show that the plume is stable and/or shrinking." Geomatrix installed an additional monitoring well (MW-4), downgradient of the USTs and previous sampling points. As presented above, diesel and only low concentrations of isopropylbenzene and n-propylbenzene were detected in the samples collected from MW-4. Sample results from the existing monitoring wells (MW-1, MW-2, and MW-3) were consistent with the previous sampling round conducted in February 1998. Geomatrix conducted additional site sampling to assess the potential impact of previous site operations on soil and groundwater. This additional sampling is considered adequate to characterize risks posed by the site to human health and the environment. In addition, we recommend that groundwater monitoring be conducted on a semiannual basis to confirm the stability of the plume identified in the area of the underground storage tanks.

3) The dissolved hydrocarbon plume is not migrating.

A thin film of weathered diesel was encountered in MW-4, downgradient of the UST installation. Weathered diesel is not considered to be significantly soluble



and would therefore not contribute to a migrating dissolved plume. Groundwater monitoring at the site does not indicate plume migration.

4) No water wells, deeper drinking water aquifers, surface water, or other sensitive receptors are likely to be impacted.

One irrigation well is located approximately 1/3-mile downgradient of the site. However, this well is 470 feet deep. The shallow groundwater beneath Emeryville is currently not used as a drinking water source and likely will not be used in the foreseeable future based on historical practices in the area and possible effects from the sewer and storm drains. Based on results of shallow soil and groundwater sampling, it is unlikely that constituents detected at the site would affect the deeper aquifer, which is located approximately 42 feet bgs (EKI, 1995).

5) The site presents no significant risk to human health

The results of the HHRA indicate that chemicals in soil and groundwater do not present an unacceptable noncarcinogenic or carcinogenic health risk to current or future on-site workers under the conditions evaluated. Potential health risks associated with exposure to soil or groundwater by a future on-site construction worker will be addressed as part of a site-specific health and safety plan.

6) The site presents no significant risk to the environment.

No ecological risk assessment was considered warranted because the current planned future use of the site is for commercial purposes and the site is located within a commercial/industrial area approximately ½-mile from the nearest surface water body.

7.0 RECOMMENDATIONS

Based on the information presented herein, Geomatrix recommends that groundwater monitoring wells MW-3 and MW-4 continue to be sampled on a semiannual basis to



confirm stability of the existing plume. These wells are located upgradient of the USTs beneath the building (MW-3) and downgradient of all the former USTs, respectively (MW-4). We recommend that monitoring wells MW-1 and MW-2 be destroyed to allow commercial use of the building. It is our opinion that the two remaining wells will be sufficient to determine the stability of the plume beneath the site.

We also recommend that a risk management plan be developed and implemented during proposed future construction at the site. This plan will include:

- Proper handling and storage of excavated soil;
- Sampling of soil stockpiles for disposal purposes;
- Aeration of soil removed from beneath the current building slab;
- Removal of the USTs that have been closed in-place beneath the building and surrounding soil if the building is demolished;
- Development of a site-specific health and safety plan for on-site construction workers.

· Jacundwater management plan



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TABLES



TABLE 1

SOIL SAMPLE ANALYTICAL RESULTS - METALS¹

Rix Property Emeryville, California

Concentrations are in milligrams per kilogram (mg/kg)

Sample ID	Arsenic	Barium	Beryllium	Cadmium	Chromium	Cobalt	Copper	Lead	Mercury	Molybdenum	Nickel	Vanadium	Zinc
S-1,2	<20	300	<0.5	<0.5	33	8	33	84	0.07	<2	28	30	150
S-3	<20	190	<0.5	<0.5	32	7	12	7	<0.06	<2	19	35	23
S-4	<20	160	<0.5	0.6	120	8	40	60	0.51	3	37	42	140
S-5	<20	140	0.6	<0.5	68	13	20	45	0.09	<2	79	40	77
S-6	<20	1800	<0.5	1.5	37	10	53	240	0.32	<2	26	36	630
S-7,8	<11	560	<0.5	1.8	38	10	41	430	0.24	<2	41	27	720
S-9,10	<10	870	<0.5	2	71	16	120	410	0.46	<2	43	40	490
GGW1-1.0	<10	180	<0.5	<0.5	31	8	12	6	<0.06	<2	21	31	22
GGW2-2.0	10	150	<0.5	<0.5	47	6	10	7	<0.06	<2	25	36	31
GGW3-1.5	12	1600	<0.5	3	88	11	32	390	0.32	<2	29	40	900
GGW4-1.5	<10	260	0.5	0.6	41	18	24	84	0.07	<2	39	43	120
LBNL	_			i									
Background ²	19.1	323.6	1.0	2.7	99.6	22.2	69.4	16.1	0.4	7.4	119.8	74.3	106.1

Note:

- 1. Samples were collected on 22 April 1998 by Geomatrix Consultants, Inc. and analyzed for metals according to EPA Method 6000/7000 Series by American Environmental Network of Pleasant Hill, California. Samples were collected within the top 2 feet of soil.
- 2. Lawrence Berkeley National Laboratory, 1995, Protocol for Determining Background Concentrations of Metals in Soil at Lawrence Berkeley National Laboratory, August.



SOIL SAMPLE ANALYTICAL RESULTS VOLATILE ORGANIC COMPOUNDS¹

Rix Property Emeryville, California

Concentrations are in milligrams per kilogram (mg/kg)

					р-			1,2,4-	1,3,5-	
	n-	sec-		Isopropyl	Isopropylt	n-		Trimethyl-	Trimethyl-	Xylenes,
Sample ID	Butylbenzene	Butylbenzene	Ethylbenzene	benzene	oluene	Propylbenzene	Napthalene	benzene	benzene	Total
S-03	< 0.03	0.03	0.03	< 0.03	0.03	0.04	0.16	0.24	0.053	0.18
S-05	6.5	6	<3	4	<3	5.9	5.1	<3	<3	<5
S-04	<0.3	<0.3	<0.3	<0.3	0.3	0.3	<0.3	1.9	1.4	1.1

Note:

1. Soil samples were collected on 22 April 1998 by Geomatrix Consultants, Inc. and analyzed for volatile organic compounds (VOCs) according to EPA Method 8260 by American Environmental Network of Pleasant Hill, California. Only constituents detected are shown. Samples were collected from beneath the slab inside the building within the top 2 feet of soil.



GROUNDWATER ANALYTICAL RESULTS METALS AND TOTAL PETROLEUM HYDROCARBONS¹

Rix Property Emeryville, California

Concentrations reported in micrograms per liter (µg/l)

	Dated				Molyb-		Ţ <u> </u>		l	
Sample ID	Sampled	Arsenic	Barium	Cobalt	denum	Nickel	Vanadium	Zinc	TPHd	TPHg
MW-2 ²	4/27/98	7	19000	<5	<10	10	<5	<10	NA	NA
$MW-4^2$	4/27/98	18	150	<5	20	10	<5	<10	NA	NA
GGW-1	4/22/98	34	230	5	30	10	<5	110	NA	NA
GGW-2	4/22/98	<5	150	6	50	30	<5	10	<50	<50
GGW-3 ³	4/22/98	<5/<5	520/410	6/6	<10/<10	<10/<10	<5/<5	100/80	NA/<50	220/240
GGW-4	4/22/98	8	40	<5	<10	<10	32	10	NA	NA

Notes:

- Samples were collected by Geomatrix Consultants, Inc., and analyzed for CAM 17 metals according to EPA Methods 6000/7000 series and total petroleum hydrocarbons quantified as gasoline (TPHg) and as diesel (TPHd) according to EPA Method 8015 by American Environmental Network of Pleasant Hill, California. Only analytes detected are shown. TPHd samples were analyzed following silica gel cleanup procedures according to EPA Method 3630M.
- 2. Samples were analyzed outside of holding time.
- 3. A duplicate sample was collected. Both results are shown.



GROUNDWATER ANALYTICAL RESULTS¹ VOLATILE ORGANIC COMPOUNDS

Rix Property Emeryville, California

Concentrations reported in micrograms per liter (µg/l)

Sample ID	Dated Sampled	sec- Butyl- benzene	1,1 <i>-</i> DCA	•	trans-1,2 DCE	Ethyl- benzene	Iso- propyl- benzene	Naph- thalene	PCE	n-Propyl- benzene	Toluene	TCE	1,2,4- TMB	1,3,5- TMB	VC	Total Xylenes
MW-1 ²	4/27/98	8	18	5	10	<5	8	<5	<5	9	<5	13	<5	<5	<10	10
MW-2 ²	4/27/98	<5	12	61	18	7	<5	5	120	<5	<5	130	<5	<5	12	26
MW-3 ^{2,3}	4/27/98	<30/<30	<30/<30	(140/140)	<30/<30	<30/<30	<30/<30	<30/<30	40/30	<30/<30	250/220	190/190	<30/<30	<30/<30	7100/110	75/<50
MW-4 ^{2,4}	4/27/98/ 5/15/98	15/<5	<5/<5	<5/<5	<5/<5	<5/<5	16/<5	<5/<5	<5/<5	5/<5	<5/<5	<5/<5	<5/<5	<5/<5	<10/<10	<10/<10
GGW1	4/22/98	<50	<50	<50	<50	<50	<50	<50	<50	<50	<50	<50	<50	<50	<100	<100
GGW2	4/22/98	<5	(120)	<5	<5	<5	<5	<5	<5	<5	<5	<5	<5	<5	1417	<10
GGW3 ³	4/22/98	<5/<5	6/5	<5/<5	<5/<5	<5/<5	<5/<5	<5/<5	<5/<5	<5/<5	<5/<5	<5/<5	120/110	11/10	<10/<10	13/10
GGW4	4/22/98	<500	<500	<500	<500	<500	<500	<500	<500	<500	<500	<500	<500	<500 ,	<1000	<1000

Notes:

- 1. Samples were collected by Geomatrix Consultants, Inc., and analyzed for volatile organic compounds (VOCs) according to EPA Method 8260 by American Environmental Network of Pleasant Hill, California. Only constituents detected are shown.
 - 1,1-DCA = 1,1-dichloroethane; cis-1,2-DCE = cis-1,2-dichloroethene,
 - trans-1,2-DCE = trans-1,2-dichloroethene; PCE = tetrachloroethene; TCE = trichloroethylene; 1,2,4-TMB = 1,2,4-trimethylbenzene;
 - 1,3,5-TMB = 1,3,5-trimethylbenzene; VC = vinyl chloride.
- 2. Samples were analyzed after the holding time had expired.
- 3. A duplicate sample was collected. Both results are shown.



IDENTIFICATION OF CHEMICALS OF POTENTIAL CONCERN IN SOIL (METALS)

Rix Property Emeryville, California

							LBNL B	ackground)	Data ¹	
Chemical	No. of Samples	No. of Detctions	Percent (%) Detected	Range of Detections	Range of Detection Limits	Residential PRG	Minimum	Maximum	95% UTL	Selected as COPC?
Arsenic	11	2	18	10 - 12	10 - 20	0.38	0.25	63	19.1	NO ³
Barium	11	11	100	140 - 1800		5200				NO
Beryllium	11	2	18	0.5 - 0.6	0.5	150				NO
Cadmium	11	6	55	0.6 - 3	0.5	9 ²				NO
Chromium	11	11	100	31 - 120		210				NO
Cobalt	11	11	100	6 - 18		3300				NO
Copper	11	11	100	10 - 120		2800				NO
Lead	11	11	100	6 - 430		130 ²	0.5	31	16.1	YES
Mercury	11	8	73	0.07 - 0.51	0.06	22				NO
Molybdenum	11	1	9	3	2	370				NO
Nickel	11	11	100	19 - 79		150 ²				NO
Vanadium	11	11	100	27 - 43		520				NO
Zinc	11	11	100	22 - 900		22000				NO

Notes:

COPC = Chemicals of Potential Concern

UTL = Upper Tolerance Limit

PRG = U.S. EPA Region 9, Preliminary Remedation Goal (U.S. EPA, 1998)

- 1. Lawrence Berkeley National Laboratory (LBNL), 1995, Protocol for Determining Background Concentrations of metals in soil at Lawrence Berkeley National Laboratory.
- 2 California-modified PRG.
- 3. The maximum detection limit is slightly higher than the background value; however, given that the maximum detected value is 12 mg/kg and the detection limit for other samples is less than the background value, arsenic was not identified as a COPC.



PHYSICAL CONSTANTS FOR ORGANIC CHEMICALS OF POTENTIAL CONCERN

Rix Property Emeryville, California

	Molecular Weight		Henry's Law Constant at 25oC		Solubility (S) at 25oC		Diffusivity in Air (Di)	
Chemical	(g/mol)	Reference	(atm-m3/mol)	Reference	(mg/L)	Reference	(cm2/sec)	Reference
n-Butylbenzene	134.22	Montgomery, 1996	1.25E-03	Montgomery, 1996	5.00E+01	Montgomery, 1996	0.064	Calculated
sec-Butylbenzene	134.22	Montgomery, 1996	1.14E-02	Montgomery, 1996	1.71E+02	Montgomery, 1996	0.064	Calculated
1,1-Dichloroethane	98.96	Montgomery, 1996	5.62E-03	U.S. EPA, 1996	5.06E+03	U.S. EPA, 1996	0.0742	U.S. EPA, 1996
cis-1,2-Dichloroethene	96,94	trans-1,2-Dichloroethene	4.08E-03	U.S. EPA, 1996	3.50E+03	U.S. EPA, 1996	0.0736	U.S. EPA, 1996
trans-1,2-Dichloroethene	96.94	Montgomery, 1996	9.38E-03	U.S. EPA, 1996	6.30E+03	U.S. EPA, 1996	0.0707	U.S. EPA, 1996
Ethylbenzene	106.17	Montgomery, 1996	7.88E-03	U.S. EPA, 1996	1.69E+02	U.S. EPA, 1996	0.075	U.S. EPA, 1996
Isopropylbenzene	120.19	Montgomery, 1996	5.58E-03	Montgomery, 1996	7.30E+01	Howard, 1997	0.068	Calculated
p-Isopropyltoluene	134.22	Montgomery, 1996	1.10E-02	Howard, 1997	2.34E+01	Howard, 1997	0.064	Calculated
Naphthalene	128.18	Montgomery, 1996	4.83E-04	U.S. EPA, 1996	3.10E+01	U.S. EPA, 1996	0.059	U.S. EPA, 1996
				Montgomery, 1996 (average		Montgomery, 1996 (average		
n-Propylbenzene	120.19	Montgomery, 1996	1.08E-02	of three values)	8.34E+01	of three values)	0 068	Calculated
Tetrachloroethene	165.83	Montgomery, 1996	1.84E-02	U.S. EPA, 1996	2.00E+02	U.S. EPA, 1996	0.072	U.S. EPA, 1996
Toluene	92.14	Montgomery, 1996	6.64E-03	U.S. EPA, 1996	5.26E+02	U.S. EPA, 1996	0.087	U.S. EPA, 1996
Trichloroethene	131.39	Montgomery, 1996	1.03E-02	U.S. EPA, 1996	1.10E+03	U.S. EPA, 1996	0.079	U.S. EPA, 1996
1,2,4-Trimethylbenzene	12.19	Montgomery, 1996	5.70E-03	Montgomery, 1996	5.70E+01	Montgomery, 1991	0.068	U.S. EPA, 1996
1,3,5-Trimethylbenzene	120.19	Montgomery, 1996	6.73E-03	Montgomery, 1996	9.77E+01	Montgomery, 1996	0.068	Calculated
Vinyl Chloride	62.5	Montgomery, 1996	2.70E-02	U.S. EPA, 1996	2.76E+03	U.S. EPA, 1996	0 106	U.S. EPA, 1996
				U.S. EPA, 1996 (average of		U.S. EPA, 1996 (average of		U.S. EPA, 1996 (average of
Xylenes	106.17	Montgomery, 1996	6 73E-03	the three isomers)	1.75E+02	the three isomers)	0.087	the three isomers)

Notes:

Diffusívity in air coefficients were calculated using equations presented in U.S. EPA's Superfund Exposure Assessment Manual (1988b)

References:

Howard, P.H., 1997, Handbook of Environmental Fate and Exposure Data for Organic Chemicals, Volume V, Lewis Publishers, Boca Raton, New York

Montgomery, J.H., 1991, Groundwater Chemical Desk Reference, Lewis Publishers, Chelsea, Michigan

Montgomery, J.H., 1996, Groundwater Chemical Desk Reference, Second Edition, Lewis Publishers, Chelsea, Michigan

U.S. EPA, 1996, Soil Screening Guidance, Technical Background Document, EPA/540/R-95-/128.



-VRERSENTATIVE CONCENTRATIONS FOR CHEMICALS OF POTENTIAL CONCERN

Rix Property Emeryville, California

	Maximum Soil	Maximum Groundwater
Chemical	Concentration (mg/kg)	Concentration (mg/l)
n-Butylbenzene	6.5	ND
sec-Butylbenzene	6	0.015
1,1-Dichloroethane	ND	0.12
cis-1,2-Dichloroethene	ND	0.14
trans-1,2-Dichloroethene	ND	0.018
Ethylbenzene	0.03	0.007
Isopropylbenzene	4	0.016
p-Isopropyltoluene	0.3	ND
Lead	430	NA
Naphthalene	5.1	0.005
n-Propylbenzene	5.9	0.009
Tetrachloroethene	ND	0.12
Toluene	ND	0.25
Trichloroethene	ND	0.19
1,2,4-Trimethylbenzene	1.9	0.12
1,3,5-Trimethylbenzene	1.4	0.011
Vinyl Chloride	ND	(0.11)
Xylenes	1.1	0.075

Notes:

NA = not applicable



EXPOSURE PARAMETERS

Rix Property Emeryville, California

Exposure Parameters	Units	Value	Rationale
Exposure Frequency (EF)	days/year	250	Cal-EPA, 1992; U.S. EPA, 1991
Exposure Duration (ED)	years	25	Cal-EPA, 1992; U.S. EPA, 1991
Body Weight	kg	70	Cal-EPA, 1992; U.S. EPA, 1991
Averaging Time	days	25,550 (carcinogens) 9,125 (noncarciongens)	Cal-EPA, 1992; U.S. EPA, 1991
Inhalation Rate (IRa)	m³/day	20	Cal-EPA, 1992; U.S. EPA, 1991

References:

California Environmental Protection Agency (Cal-EPA), 1992, Supplemental Guidence for Human Health Multimedia Risk Assessment of Hazardous Waste Sites and Permitted Facilities.

U.S. EPA, 1991b, Human Health Evaluation Manual, Supplemental Guidence: "Standard Default Exposure Factors:" Office of Emergency and Remedial Responce, Washington, D.C.



TABLE 9

NONCARCINOGENIC TOXICITY CRITERIA FOR CHEMICALS OF POTENTIAL CONCERN
Rix Property

Rix Property
Emeryville, California

Chemical	RfDo (mg/kg-day)	Reference	RíDi (mg/kg-day)	Reference	Target Organ	Critical Effect	Reference
n-Butylbenzene	0.01	U.S. EPA, 1998	0.01	Route Extrapolation			
sec-Butylbenzene	0.01	U.S. EPA, 1998	0.01	Route Extrapolation	1		
1,1-Dichloroethane	NE	NE	0.14	U.S. EPA, 1997	Kidney	Damage	U.S. EPA, 1997
	i					Decereased Hematocrit and	
c1s-1,2-Dichloroethene	0.01	U.S. EPA, 1997	0.01	Route Extrapolation	Blood	Hemoglobin	U.S. EPA, 1997
						Increase in Serum Alkaline	1
trans-1,2-Dichloroethene	0.02	IRIS, 1998	0.02	Route Extrapolation	Liver	Phosphatase Levels	IRIS, 1998
Ethylbenzene	NE	NE	0.29	IRIS, 1998	Liver and Kidney	Toxicity	IRIS, 1998
					Kidney/ Central Nervous	Increased Weight/ Involvment/	
Isopropylbenzene	NE	NE	0.0026	U.S. EPA, 1997	System/ Nose	Irritation	U.S. EPA, 1997
p-Isopropyltoluene	NE	NE	NA	NA			
Naphthalene	NE	NE	0.00086	U.S. EPA, 1998			
n-Propylbenzene	0.01	U.S. EPA, 1998	0.01	Route Extrapolation			
n-r topytoenzene	0.01	O.B. L171, 1770	0.01	Troute Danapolation		Increased Weight/ Decereased	
Tetrachloroethene	0.01	IRIS, 1998	0.01	Route Extrapolation	Liver/ Total Body	Weight	IRIS, 1998
Toluene	NE	NE	0.11	IRIS, 1998	Liver and Kidney	Weight Changes	IRIS, 1998
Trichloroethene	0.006	U.S. EPA, 1998	0.006	Route Extrapolation	<u></u>		
1,2,4-Trimethylbenzene	NE	NE	0.0017	U.S. EPA, 1998		'	
1,3,5-Trimethylbenzene	NE	NE	0.0017	U.S. EPA, 1998			
Vinyl Chloride	NE	NE	NA	NA.	NA.	NA	NA
v my i emoride	112	1,12		- "-	Central Nervious System/	Hyperactivity /Decreased	
 Xylenes	NE	NE	0.2	U.S. EPA, 1998	Whole Body	Weight	U.S. EPA, 1997

Notes-

NA = not available

NE = not evaluated

References:

Integrated Risk Information System (IRIS), 1998, on-line search, June.

U.S. EPA, 1997, Health Effects Summary Tables, FY-1997 Annual, Office of Solid Waste and Energy Response.

U.S. EPA, 1998, U.S. EPA Region 9 Preliminary Remediation Goals (PRGs)



CARCINOGENIC TOXICITY CRITERIA FOR CHEMICALS OF POTENTIAL CONCERN

Rix Property Emeryville, California

Chemical	SFi (mg/kg-day) ⁻¹	Reference	Target Organ	Critical Effect	Reference
Tetrachloroethene	0.021	U.S. EPA, 1998	Liver	Hepatocellular adenomas/carcinomas	Cal-EPA, 1994b
[Hepatocellular	
				adenomas/carcinomas/lymphoma;	
Trichloroethene	0.01	U.S. EPA, 1998	Liver/Lung	adenocarcinomas	Cal-EPA, 1994b
			Liver/Mammary Gland/	Angiosarcomas/hepatocellular carcinomas;	
Vinyl Chloride	0.27	Cal-EPA, 1994	Lung	adenocarcimonas; carcinomas	Cal-EPA, 1994b

Reference:

Cal-EPA, 1994b, California Cancer Potency Factors: Office of Environmental Health Hazard Assessment, Sacramento, California.



HAZARD QUOTIENT SUMMARY TABLE

Rix Property Emeryville, California

Chemical	Inhalation of Vapors from Soil	Inhalation of Vapors from Groundwater	Total Hazard Index
n-Butylbenzene	3E-04	NA	3E-04
sec-Butylbenzene	2E-03	3E-06	2E-03
1,1-Dichloroethane	NA	2E-07	2E-07
cis-1,2-Dichloroethene	NA	3E-06	3E-06
trans-1,2-Dichloroethene	NA	4E-07	4E-07
Ethylbenzene	4E-07	3E-08	4E-07
Isopropylbenzene	2E-03	4E-07	2E-03
p-Isopropyltoluene	NA	NA	NA
Naphthalene	5E-05	3E-06	5E-05
n-Propylbenzene	2E-03	2E-06	2E-03
Tetrachloroethene	NA	1E-06	1E-06
Toluene	NA	2E-06	2E-06
Trichloroethene	NA	3E-05	3E-05
1,2,4-Trimethylbenzene	1E-03	2E-04	1E-03
1,3,5-Trimethylbenzene	2E-03	8E-06	2E-03
Vinyl Chloride	NA	NA	NA
Xylenes	2E-05	4E-07	2E-05
Total	9E-03	3E-04	9E-03

Notes:

NA = not applicable



CANCER RISK SUMMARY TABLE

Rix Property Emeryville, California

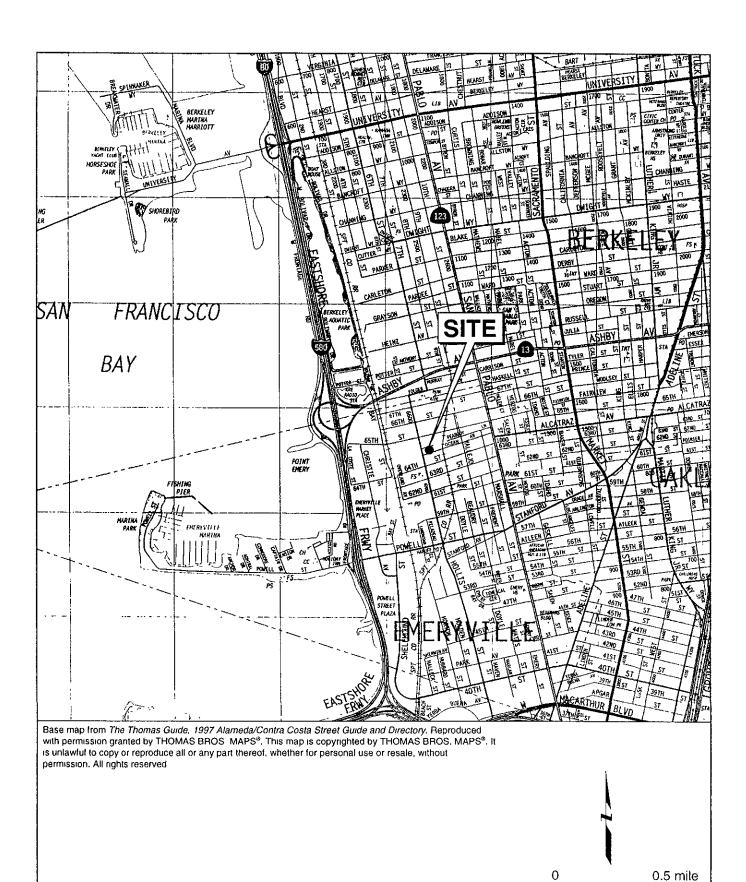
	Inhalation of Vapors
Chemical	from Groundwater
n-Butylbenzene	NA
sec-Butylbenzene	NA
1,1-Dichloroethane	NA
cis-1,2-Dichloroethene	NA
trans-1,2-Dichloroethene	NA
Ethylbenzene	NA
Isopropylbenzene	NA
p-Isopropyltoluene	NA
Naphthalene	NA
n-Propylbenzene	NA
Tetrachloroethene	5E-09
Toluene	NA
Trichloroethene	3E-09
1,2,4-Trimethylbenzene	NA
1,3,5-Trimethylbenzene	NA
Vinyl Chloride	3E-08
Xylenes	NA
Total	4E-08

Notes:

NA = not applicable

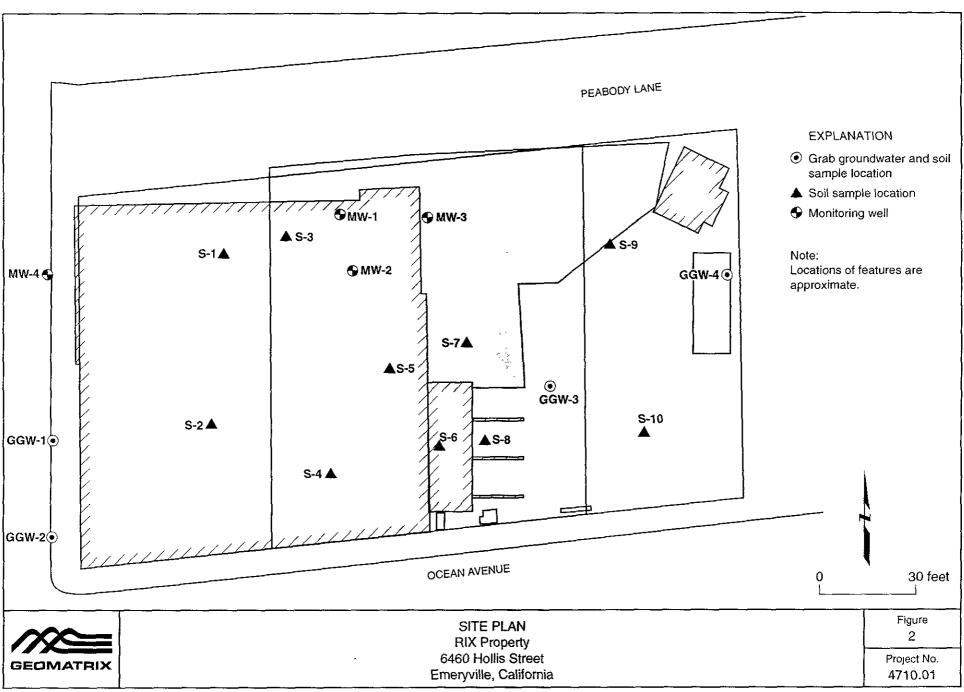


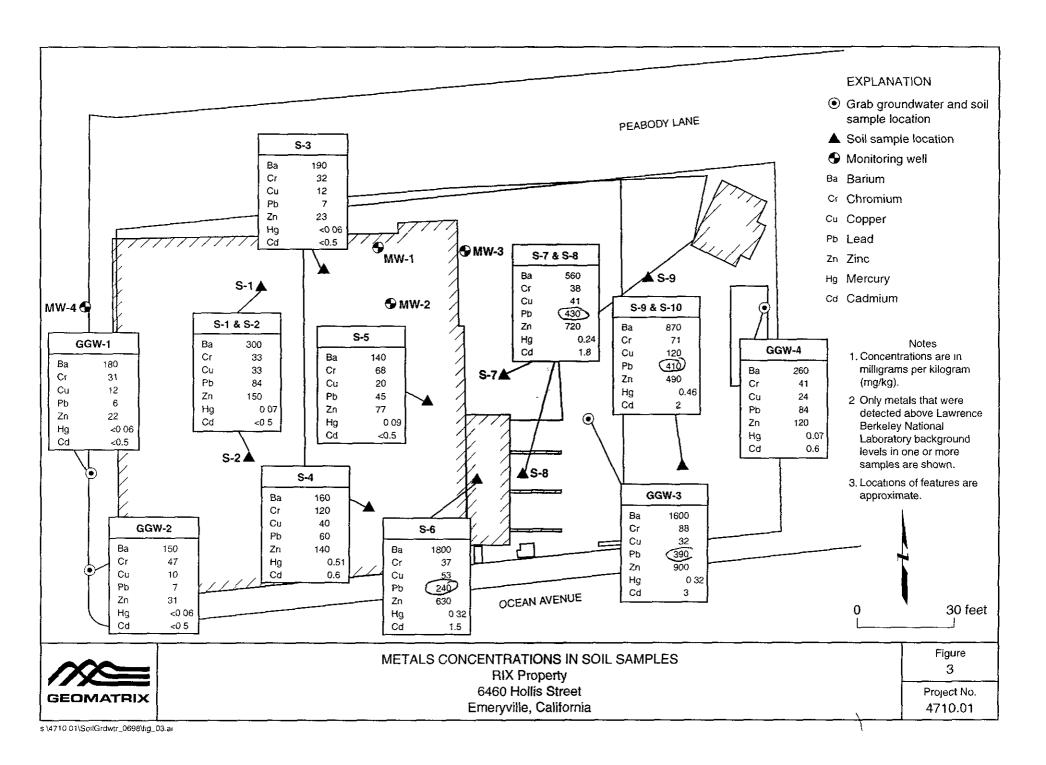
FIGURES

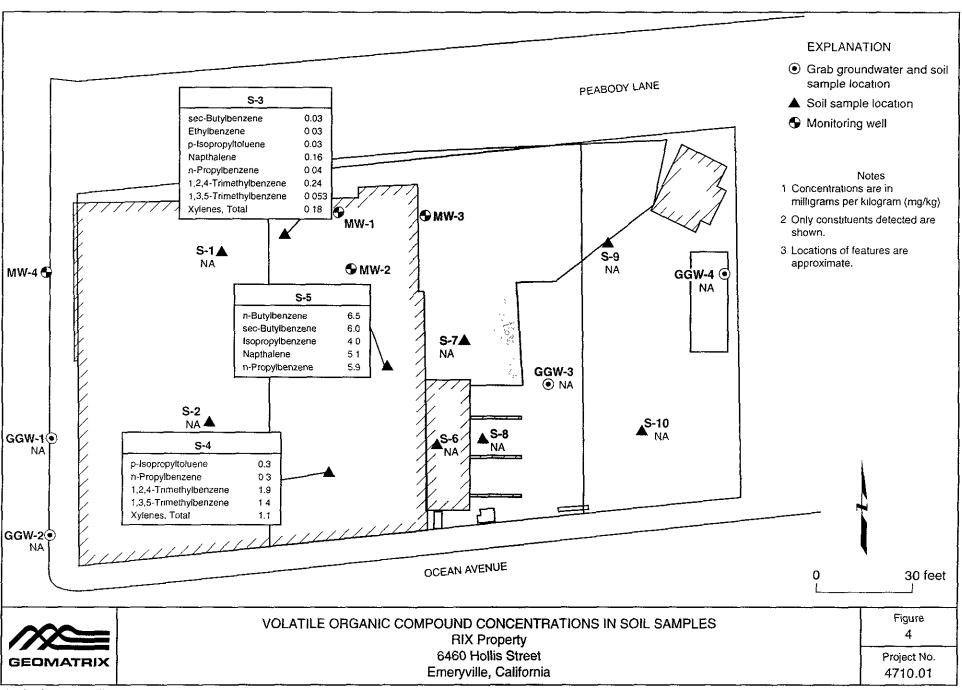


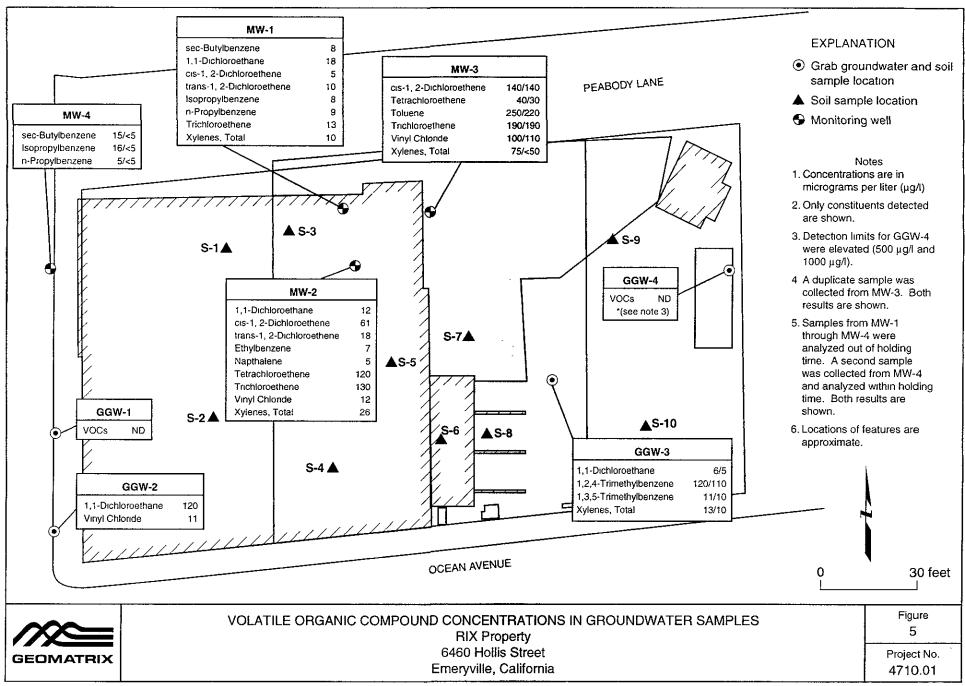


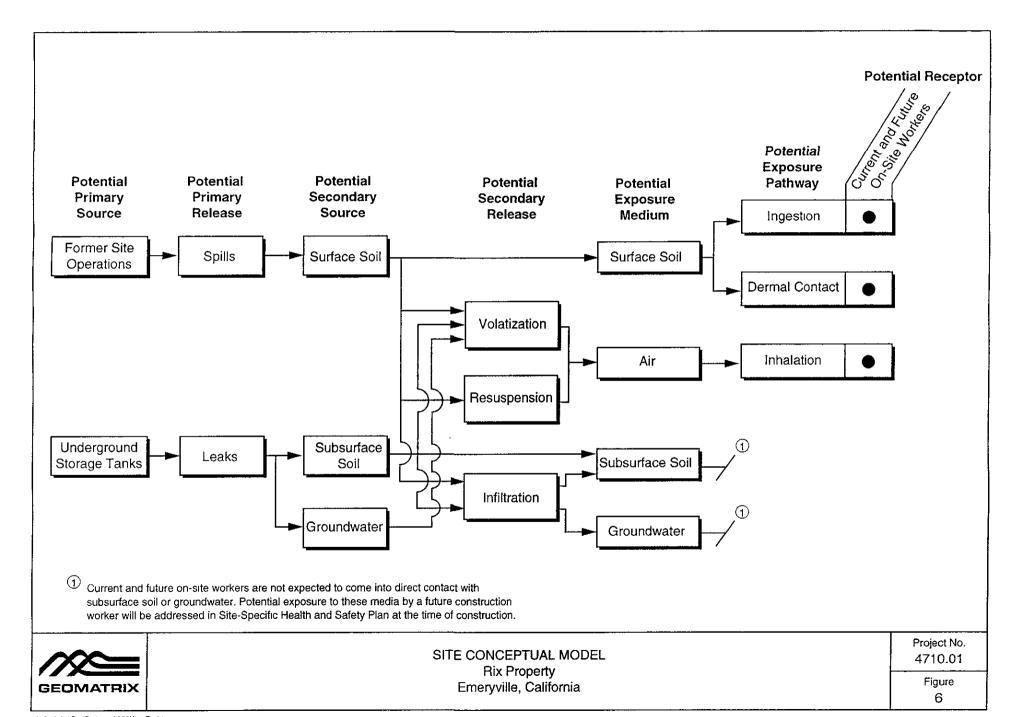
SITE LOCATION MAP RIX Property 6460 Hollis Street Emeryville, California Figure
1
Project No.
4710.01







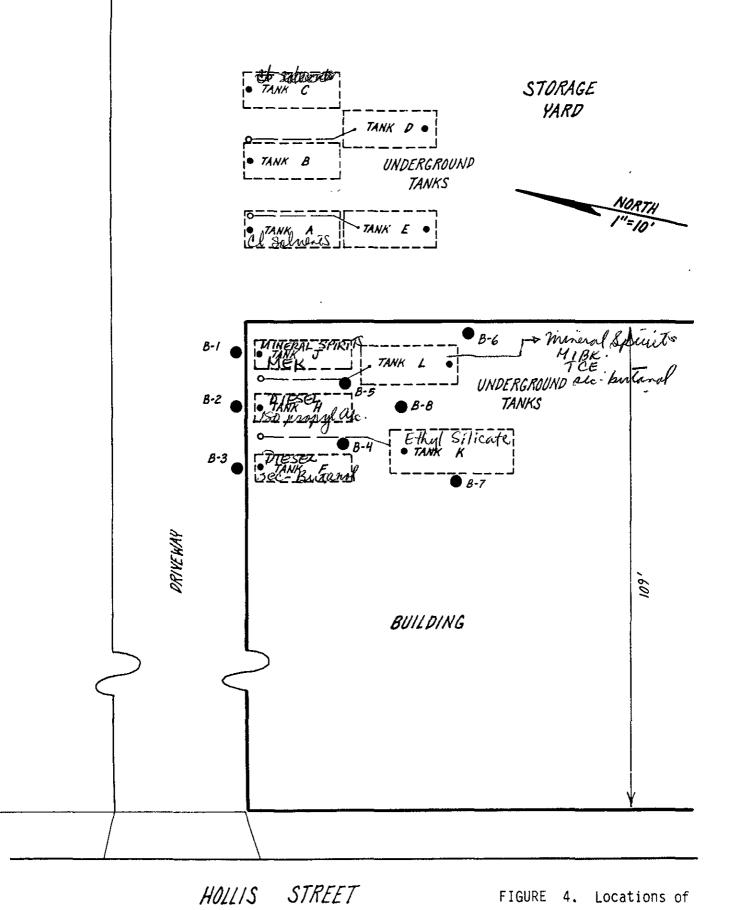






APPENDIX A

PREVIOUS SOIL SAMPLE ANALYTICAL RESULTS



Hand Soil Borings.

TABLE 3. Hand Boring Soil Sampling Results.

Chemical	· · · · · · · · · · · · · · · · · · ·	B-1	B-2	B-3	B-4	B-5	B-6	
Gasoline	(mg/kg)	2.2	12	** <u>*42</u> *	34	8.8	15	
Diesei	(mg/kg)	3.4	1.5	52	30	32	41	60
Kerosene	(mg/kg)	ND	ND	ND	ND	ND	ND	
Mineral Spirits	(mg/kg)	ND	1.8	ND	§26 P	19	20	
Methyl Ethyl Ketone (MEK)	(mg/kg)	ND	ND	ND	ND	ND	ND	
Methyl Isobutyl Ketone (MIBK)	(mg/kg)	ND	ND	ND	ND	ND	ND	
cis-1,2- Dichloroethene	(ug/kg)	ND	ND	ND	ND	2 52	ND	
Tetrachloroethene	(ug/kg)	360	340	230	160	1,5001	230	
Trichloroethene (TCE)	(ug/kg)	ND	ND	ND	ND	007 **	ND	
Benzene	(mg/kg)	ND	ND	ND	ND	@0.0073*	ND	
Toluene	(mg/kg)	0.0080	0.013	0.056	0.021	0.24	0.023	0,078
Ethyl Benzene	(mg/kg)	0.0058	ND	0.15	0.064	0.039	0.043	0.078
Total Xylenes	(mg/kg)	0.0057	0.0092	0.54	0.82	@0,91 _#	0.12	
Ethyl Silicate	(ug/kg)	ND	ND	ND	ND	ND	ND	
Oil & Grease	(mg/kg)	76	160	,100	310	310	(400)	

TABLE 3 (cont.). Hand Boring Soil Sampling Results.

Chemical		B-7	B-8
Gasoline	(mg/kg)	29	3.2
Diesel	(mg/kg)	(60 ⁰)	5.6
Kerosene	(mg/kg)	ND	ND
Mineral Spirits	(mg/kg)	ND	13
Methyl Ethyl Ketone (MEK)	(mg/kg)	ND	ND
Methyl Isobutyl Ketone (MIBK)	(mg/kg)	ND	ND
cis-1,2- Dichloroethene	(ug/kg)	ND	5.2
Tetrachloroethene	(ug/kg)	210	330
Trichloroethene (TCE)	(ug/kg)	ND	6.5
Benzene	(mg/kg)	ND	ND
Toluerie	(mg/kg)	0.020	0.078
Ethyl Benzene	(mg/kg)	0.078	0.0051
Total Xylenes	(mg/kg)	0.090	0.049
Ethyl Silicate	(ug/kg)	ND	ND
Oil & Grease	(mg/kg)	260	160

TABLE 4. Well Installation Soil Sampling Results.

Chemical		MW-1 @ 3'	MW-1 @ 5'	MW-2 @ 3'	MW-2 @ 5'	MW-3 @ 5'	MW-3 @ 7.5'
Gasoline	(mg/kg)	ND	300	140	1,800	55	85
Diesel	(mg/kg)	ND	540	500	3,000	24	6.9
Kerosene	(mg/kg)	ND	430	400	2,400	19	5.5
Mineral Spirits	(mg/kg)	ND	380	350	2,100	17	4.8
Methyl Ethyl Ketone (MEK)	(mg/kg)	ND	ND	ND	ND	41	ND
Methyl Isobutyl Ketone (MIBK)	(mg/kg)	ND	ND	ND	ND	4.9	ND
cis-1,2- Dichloroethene	(ug/kg)	ND	ND	ND	ND	720	ND
Tetrachloroethene	(ug/kg)	ND	ND	ND	ND	31,000	4,500
Trichloroethene (TCE)	(ug/kg)	ND	ND	ND	, ND	420	ND
Benzene	(mg/kg)	ND	ND	ND	ND	ND	0.096
Toluene	(mg/kg)	ND	0.12	ND	0.80	19	1.7
Ethyl Benzene	(mg/kg)	ND	ND	0.11	ND	0.78	0.96
Total Xylenes	(mg/kg)	ND	2.6	0.79	2.6	5.8	8.0
Ethyl Silicate	(ug/kg)	ND	ND	ND	ND	ND	ND
Oil & Grease	(mg/kg)	ND	ND	ND	ND	ND	ND



APPENDIX B PREVIOUS GROUNDWATER ANALYTICAL DATA

722-1.HPL SUMMARY LETTER, 6460 HOLLIS STREET, EMERYVILLE, CA

TABLE 1 CUMULATIVE RESULTS FOR WELL MW-1 LABORATORY ANALYSES OF GROUNDWATER SAMPLES FOR HYDROCARBON COMPOUNDS by EPA 5030/8015 and 8260 6460 Hollis Street

Emeryville, California

Compound Defected	7/92	8/94	11/94	2/95	6/95	10/95	4/96	8/96	2/98
TPHp	680	< 50	440	630	620	960	1,500	1,200	940 ^{abc}
ГРНа	6,100	590	1,000	1,400	1,600	2,600	760	420	9,300 ^{dec}
Kerosene	6,100	960	< 50	< 50	< 50	< 50	< 50	< 50	8,700
Stoddard Solvent	6,400	< 50	190	310	58	450	190	110	580 ^{abc}
Oil & Grease	14	NA	< 250						
Benzene	(3.8)	0.5	< 0.5	< 0.5	0.5	<0.5	< 0.5	< 0.5	< 0.5
Toluene	< 0.5	< 0.5	8.0	0.5	< 0.5	1.7	0.6	0,1	ι.ι ¹
Ethylbenzene	38	< 0.5	2.6	1.2	2.3	1.4	4.6	3.9	2,5 ⁸
Total Xylenes	3.4	< 0.5	6.2	3.6	9.6	7.9	14	16	9.8 ^h
MTBI	NA	<5							

Results in micrograms per kilograms (ug/l) = parts per million (ppb).

Data from 7/92 through 8/96 obtained from Hageman/Aguiar (see references section.)

- 1 Less than the detection limit for the analysis method.
- a. Strongly aged gasoline or diesel range compounds are significant.
- h No recognizable pattern.
- Confighter than water immiscible sheen is present.
- d Unmodified or weakly modified diesel is significant.
- c Gasoline range compounds are significant.
- 1 1 1 pph by EPA 8260.
- д 2.6 ppb by EPA 8260
- (i) 8 2 by EPA 8260
- NA Nor analyzed

722-1.HPL SUMMARY LETTER, 6460 HOLLIS STREET, EMERYVILLE, CA

TABLE 2 CUMULATIVE RESULTS FOR WELL MW-1 LABORATORY ANALYSES OF GROUNDWATER SAMPLES FOR VOCs by EPA 601 and 8260

6460 Hollis Street Emeryville, California

Compound Detected	7/92	11/94	2/95	6/95	10/95	4/96	8/96	2/98
3romotorm	< 0.5	< 0.5	< 0.5	< 0.5	< 0.5	<0.5	< 0.5	< 0.5
`hlorotorm	< 0.5	< 0.5	1.8	1.5	1.9	< 0.5	< 0.5	< 0.5
,1-Dichloroethane	36	33	32	12	< 0.5	<0.5	< 0.5	14 ⁶
,1-Dichloroethene	< 0.5	< 0.5	1.0	< 0.5	< 0.5	< 0.5	< 0.5	< 0.5
1,2-Dichtoroethane	< 0.5	< 0.5	1.1	< 0.5	< 0.5	< 0.5	< 0.5	< 0.5
1,1,1 Trichloroethane	< 0.5	< 0.5	0.7	< 0.5	< 0.5	< 0.5	< 0.5	< 0.5
Cis/Trans 1,2-Dichloroette	ne < 0.5	< 0.5	8.5	2.7	5.1	<0.5	< 0.5	3.4 ^d
Petraeliloroethene	< 0.5	< 0.5	< 0.5	< 0.5	< 0.5	< 0.5	< 0.5	t.8ª
Prichloroethene	< 0.5	< 0.5	15	4.6	<0.5	< 0.5	< 0.5	0.97 ⁴
Uncidoroflouromethane	< 0.5	NR	9.8	4.9	<0.5	<0.5	< 0.5	2.3°
Vmyl Chloride	< 0.5	< 0.5	< 0.5	< 0.5	<0.5	<0.5	< 0.5	2.3 ^e

Results in micrograms per kilograms (ug/l) = parts per willion (ppb).

Data from 7/92 through 8/96 obtained from Hageman/Aguiar (see references section,)

- ** Less than the detection limit for the analysis method.
- a < 1.0 pph by EPA 8260.
- h. It pph by EPA 8260.
- C 1 0 pph by EPA 8260
- d: 1.8 pph by EPA 8260.
- e. Not confirmed by EPA 8260; one other probable freon is present.
- NR Not reported.

722-1.HPL SUMMARY LETTER, 6460 HOLLIS STREET. EMERYVILLE, CA

TABLE 3 CUMULATIVE RESULTS FOR WELL MW-1 LABORATORY ANALYSES OF GROUNDWATER SAMPLES FOR VOCs by EPA 8260

6460 Hollis Street Emeryville, California

Compound Detected	7/92	8/94	11/94	2/95	6/95	10/95	4/96	8/96	2/98
Acetone	< 50	210	2,700	610	76	220	180	210	< 15
Isopropanol	< 500	9,100	17,000	6,400	< 500	< 500	< 500	< 500	< 200
Methyl Ethyl Ketone	< 500	230	1,500	1,300	97,000	700	1,100	1,400	< 1.0
Methyl Isobutyl Ketone	< 500	180	420	600	< 500	< 500	< 500	< 500	< 1.0
Sec-Butanol	< 500	710	< 500	< 500	< 500	< 500	< 500	< 500	<5
n-Buryl henzene	NR	NR	NR	NR	NR	NR	NR	NR	2.0
sec-Buryl henzene	NR	NR	NR	NR	NR	NR	NR	NR	6.3
Isopropythenzene	NR	NR	NR	NR	NR	NR	NR	NR	8.7
p-Isopropyl toluene	NR	NR	NR	NR	NR	NR	NR	NR	1.2
n-Propyl benzene	NR	NR	NR	NR	NR	NR	NR	NR	8.1

Results in micrograms per kilograms (ug/l) : parts per million (ppb).

Data from 7/92 through 8/96 obtained from Hageman/Aguiar (see references section.)

NR Not reported.

· _ ____

^{*} Less than the detection limit for the analysis method.

INTERNATIONAL GEOLOGIC MARCILZ, 1998

722-1.HPL SUMMARY LETTER, 6460 HOLLIS STREET, EMERYVILLE, CA

TABLE 4 CUMULATIVE RESULTS FOR WELL MW-2 LABORATORY ANALYSES OF GROUNDWATER SAMPLES FOR HYDROCARBON COMPOUNDS by EPA 5030/8015 and 8260

6460 Hollis Street Emeryville, California

Compound Detected	7/92	8/94	11/94	2/95	6/95	10/95	4/96	8/96	2/98
TPHg	1,400	4,800	810	1,000	780	1,200	1,100	890	760 ^{ahc}
грна	17,000	320	620	810	960	350	620	610	2,300 ^{dee}
Kerosene	17.000	49()	< 50	< 50	< 50	< 50	< 50	< 50	2,200
Stoddard Solvent	20,000	< 50	160	350	< 50	< 50	81	76	390 ^{ahc}
Oil & Grease	19	NA	280						
Benzenc	< 0.5	< 0.5	< 0.5	< 0.5	0.9	<0.5	< 0.5	< 0.5	< 0.5
Foluenc	12	1.2	1.2	0.9	< 0.5	0.6	< 0.5	< 0.5	0.73
Ethythenzen e	69	5 6	4.3	3.2	3.0	3.2	2.3	1.1	12 ^f
Total Xylenes	530	18	11	6.4	13	9.7	12	9.2	50 ⁸
MTBF	NA	NA	NA	NA	NA	NA	NA	NA	<5

Results in micrograms per kilograms (ug/l) - parts per million (ppb).

Data from 7/92 through 8/96 obtained from Hageman/Aguiar (see references section.)

- Less than the detection limit for the analysis method.
- a Heavier gasoline range compounds are significant (aged gasoline?).
- b. No recognizable pattern.
- e: Lighter than water immiscible sheen is present.
- d Gasoline range compounds are significant.
- e. Diesel range compounds are significant; no recognizable pattern.
- 1: II pph by EPA 8260.
- g 47 pph by EPA 8260
- NA: Not analyzed.

722-1.HPL SUMMARY LETTER, 6460 HOLLIS STREET, EMERYVILLE, CA

TABLE 5 CUMULATIVE RESULTS FOR WELL MW-2 LABORATORY ANALYSES OF GROUNDWATER SAMPLES FOR VOCs by EPA 601 and 8260

6460 Hollis Street Emeryville, California

Compound Detected	7/92	11/94	2/95	6/95	10/95	4/96	8/96	2/98
Bromoform	< 0.5	< 0.5	< 0.5	<0.5	< 0.5	< 0.5	< 0.5	<4
Chlorotom	< 0.5	< 0.5	2.7	4,9	4.4	1.8	1.6	<4
1.1-Dichtoroethane	22	17	9.6	8	< 0.5	< 0.5	< 0.5	7.7
1,1-Dicidoroethene	< 0.5	< 0.5	2	<0.5	<0.5	< 0.5	< 0.5	< 4
1,2-Dichloroethane	<0.5	< 0.5	3.2	<0.5	<0.5	<0.5	< 0.5	<4
1,1,1 Trichloroethane	< 0.5	< ().5	4.8	< 0.5	< 0.5	< 0.5	< 0.5	<4
Cis/Trans 1,2-Dichloroether	le 99	45	11	6.9	14	0.9	6.1	173 ^a
Tetrachloroethene	52	34	49	20	14	<0.5	< 0.5	110 ^b
Trichloroethene	21	20	41	33	11	36	13	170 ^b
Trichloroflouromethane	< 0.5	NR	3.6	2.7	< 0.5	< 0.5	< 0.5	<4
Vinyl Chloride	46	< 0.5	< 0.5	<0.5	<0.5	<0.5	<0.5	6.8

Results in micrograms per kilograms (ug/l) - parts per million (pph).

Data from 7/92 through 8/96 obtained from Hageman/Aguiar (see references section.)

c (sess than the detection limit for the analysis method.

a: 141 pph by EPA 8260.

b 130 pph by EPA 8260.

NR Not reported.

722-1.HPL SUMMARY LETTER, 6460 HOLLIS STREET, EMFRYVILLE, CA

TABLE 6 CUMULATIVE RESULTS FOR WELL MW-2 LABORATORY ANALYSES OF GROUNDWATER SAMPLES FOR VOCs by EPA 8260

6460 Hollis Street Emeryville, California

Compound Detected	7/92	8/ 94	11/94	2/95	6/95	10/95	4/96	8/96	2/98
Acetone	< 50	< 50	1,100	2,500	< 50	62	79	88	< 25
Isopropanol	< 500	9,100	17,000	6,400	< 500	< 500	< 500	< 500	< 2.000
Methyl Ethyl Ketone	< 500	< 500	18,000	22,000	59,000	6,100	5,200	4,900	< 10
Methyl Isobutyl Ketone	< 500	500	360	< 500	< 500	< 500	< 500	< 500	< 10
Sec-Butanol	< 500	9()	< 500	< 500	< 500	< 500	< 500	< 500	< 50
n-Bury) henzene	NR	NR	NR	NR	NR	NR	NR	NR	< 10
sec-Buryl henzene	NR	NR	NR	NR	NR	NR	NR	NR	< 10
Isopropylbenzene	NR	NR	NR	NR	NR	NR	NR	NR	< 10
p-Isopropyl toluene	NR	NR	NR	NR	NR	NR	NR	NR	< 10
n-Propyl benzene	NR	NR	NR	NR	NR	NR	NR	NR	< 10

Results in micrograms per kilograms (ug/l) = parts per million (ppb).

Data from 7/92 through 8/96 obtained from Hageman/Aguiar (see references section.)

Less than the detection limit for the analysis method.

NR: Not reported.

722-1.HPL SUMMARY LETTER, 6460 HOLLIS STREET, EMERYVILLE, CA

TABLE 7 ('UMULATIVE RESULTS FOR WELL MW-3 LABORATORY ANALYSES OF GROUNDWATER SAMPLES FOR HYDROCARBON COMPOUNDS by EPA 5030/8015 and 8260

6460 Hollis Street Emeryville, California

Compound Detected	7/92	8/94	11/94	2/95	6/95	10/95	4/96	8/96	2/98
TPHy	9,300	4,300	920	410	1,100	1,100	640	580	260 ^{µhe}
TPHd	20,000	310	< 50	900	1,200	200	420	390	710 ^{de}
Keroseik	20,000	470	< 50	< 50	< 50	< 50	< 50	< 50	630
Stoddard Solvent	21,000	< 50	< 50	370	< 50	< 50	< 50	< 50	76 ^{abc}
Oil & Greane	28	NA	NA	NA	NA	NA	NA	NA	< 2.50
Benzene	< 0.5	< 0.5	< 0.5	<0.5	0.9	< 0.5	< 0.5	<0.5	< 0.5
Foluenc	1,600	10	3.7	1.7	0.8	< 0.5	< 0.5	< 0.5	57 ^e
lithylhenzene	< 0.5	2.6	3.2	0.5	11	3.2	0.7	0.6	3.6 ^t
Total Xylenes	700	10	16	2.5	2.6	9.6	2.1	1.8	188
МТВН	NA	NΛ	NA	NA	NA	NA	NA	NA	< 5

Results in micrograms per kilograms (ug/l) = parts per million (ppb).

Data from 7/92 through 8/96 obtained from Hageman/Aguiar (see references section.)

- Less than the detection limit for the analysis method.
- a: Heavier gasoline range compounds are significant (aged gasoline?).
- b. One to a few isolated peaks present.
- er laghter than water immiscible sheen is present.
- d Unmodified or weakly modified diesel is significant.
- c. 41 pph by EPA 8260.
- 1 2.7 pph by EPA 8260.
- g 16 by EPA 8260
- NA Not analyzed

722-1 HPL SUMMARY LETTER, 6460 HOLLIS STREET, EMERYVILLE, CA

TABLE 8 CUMULATIVE RESULTS FOR WELL MW-3 LABORATORY ANALYSES OF GROUNDWATER SAMPLES FOR VOCs by EPA 601 and 8260

6460 Hollis Street Emeryville, California

Compound Detected	7/92	11/94	2/95	6/95	10/95	4/96	8/96	2/98
Bromotorn	< 0.5	< 0.5	< 0.5	< 0.5	<0.5	32	< 0.5	<3
Chloroform	< 0.5	< 0.5	4.3	3.8	17	5.2	76	< 3
1.1-Dichtoroethane	30	47	52	16	< 0.5	< 0.5	< 0.5	33 ^b
1.1-Dichloroethene	< 0.5	29	48	26	<0.5	< 0.5	< 0.5	12 ^c
1,2-Dichloroethane	< 0.5	<0.5	8.5	<0.5	< 0.5	< 0.5	< 0.5	< 3*
1,1,1 Frichloroethane	81	12	28	<0.5	<0.5	<0.5	< 0.5	8.3 ^d
1,1,2 Trichloroethane	< 0.5	< 0.5	< 0.5	< 0.5	< 0.5	<0.5	< 0.5	<3°
Cis/Trans 1,2-Dichloroeti	iene 630	327	6.6	4.9	< 0.5	3.7	0.7	105.71
Tetrachforoethene	2,200	110	54	34	< 0.5	< 0.5	< 0.5	11 ⁸
Frichloroethene	300	290	140	63	53	41	180	(140h
Trichloroflouromethane	< 0.5	NR	30	17	< 0.5	< 0.5	< 0.5	11'
Vinyl Chloride	< 0.5	67	< 0.5	< 0.5	<0.5	<0.5	<0.5	140

Results in micrograms per kilograms (ug/l) = parts per million (ppb).

Data from 7/92 through 8/96 obtained from Hageman/Aguiar (see references section,)

- < 1.ess than the detection limit for the analysis method.
- a. < 1.0 ppb by EPA 8260.
- h 29 pph hy EPA 8260.
- c: 8.2 pph by EPA 8260.
- d 6.2 pph by EPA 8260
- c. 1.6 pph by EPA 8260.
- 1: 74 pph by EPA 8260
- g 7 5 pph by EPA 8260.
- h: 100 pph by EPA 8260.
- Not confirmed by EPA §260; one other probable freon is present.
- 110 pph by EPA 8260₩

NR: Not reported

722-1.HPL SUMMARY LETTER, 6460 HOLLIS STREET, EMERYVILLE, CA

TABLE 9 CUMULATIVE RESULTS FOR WELL MW-3 LABORATORY ANALYSES OF GROUNDWATER SAMPLES FOR VOCs by EPA 8260

6460 Hollis Street Emeryville, California

Compound Detected	7/92	8/94	11/94	2/95	6/95	10/95	4/96	8/96	2/98
Acetone	< 50	< 50	810	1,300	160	340	410	420	< 15
Isopropanol	< 500	9,400	6,700	5,800	< 500	< 500	< 500	< 500	< 200
Methyl Ethyl Ketone	< 500	370	40,000	19,000	32,000	6,600	5,900	5,100	<1.0
Methyl Isobutyl Ketone	< 500	250	22,000	4,500	< 500	< 500	< 500	< 500	< 1.0
Sec-Butanot	< 500	710	< 500	< 500	< 500	<500	< 500	< 500	<5
n-Butyl henzene	NR	NR	NR	NR	NR	NR	NR	NR	< 1
sec Butyl benzene	NR	NR	NR	NR	NR	NR	NR	NR	< 1
lsopropylhenzene	NR	NR	NR	NR	NR	NR	NR	NR	<1
p-Isopropyl toluene	NR	NR	NR	NR	NR	NR	NR	NR	<1
n-Propyl benzene	NR	NR	NR	NR	NR	NR	NR	NR	< 1

Results in micrograms per kilograms (ug/l) = parts per million (ppb).

Data from 7/92 through 8/96 obtained from Hageman/Aguiar (see references section.)

Less than the detection limit for the analysis method.

NR: Not reported

722-1 HPL SUMMARY LETTER, 6460 HOLLIS STREET, EMERYVILLE, CA

TABLE 10

RESULTS FOR HYDROPUNCH LOCATIONS HP-1 and HP-2 LABORATORY ANALYSES OF GROUNDWATER SAMPLES COLLECTED FEBRUARY 21, 1998 FOR HYDROCARBON COMPOUNDS by EPA 5030/8015 and 8260

6460 Hollis Street Emeryville, California

Compound Detected	HP-1	HP-2
TPHg	80 ⁸	1.500 ^{cdc}
PHI	240 ^h	62,000 ^{fge}
Kerosene	200	61,000
Stoddard Solvent	< 50	860 ^{ede}
Oil & Grease	< 250	7,500
Benzene	<0.5	0.73
Toluene	l 6 ^h	7.3 ^k
Ethylbenzene	< 0.5	2.8
l'otal Xylenes	1.1	171
мтвп	8.6	<5

Results in micrograms per kilograms (ug/l) = parts per million (ppb).

Data from 7/92 through 8/96 obtained from Hageman/Aguiar (see references section.)

- Less than the detection limit for the analysis method.
- a: Unmodified or weakly modified gasoline is significant.
- h: Diesel range compounds are significant: no recognizable pattern.
- c: Strongly aged gasoline or diesel range compound are significant.
- d. No recognizable pattern.
- er 1 ighter than water immiscible sheen is present.
- to Unmodified or weakly modified diesel is significant.
- g: Gasoline range compounds are significant.
- le / 1.0 ppb by EPA 8260.
- 1 1.5 pph by EPA 8260.
- 1 6 4 ppb by EPA 826().
- k: 5.9 ppb by EPA 8260.
- 1 IS pph by EPA 8260.

722-1 HPL SUMMARY LETTER, 6460 HOLLIS STREFT, EMERYVILLE, CA

TABLE 11

CUMULATIVE RESULTS FOR HYDROPUNCH LOCATIONS HP-1 AND HP-2 LABORATORY ANALYSES OF GROUNDWATER SAMPLES COLLECTED FEBRUARY 21, 1998 FOR VOCs by EPA 601 and 8260

6460 Hollis Street Emeryville, California

'ompound Detected	HP-I	НР-2	
Fromoform	< 0.5	<2	
Morotorm	U 67	<2	
1.1-Dichloroediane	0.82	86 ⁸	
, (-Dichloroethene	< 0.5	18 ^b	
1,2-Dichloroethane	< 0.5	<2	
1,1,1-Trichloroethane	< 0.5	5 0 °	
1.1,2 Trichloroethane	< 0.5	<2	
Cis/Trans 1,2-Dichloroethene	<0.5	67.5 ^d	
Ferrachloroethene	< 0.5	7.0 ^e	
Trichloroethene	< 0.5	8.2 ^f	
Dichlorodiflouromethane	< 0.5	<2 ⁸	
Trichloroflourometha	n ≪ 0.5	54 ^h	
Vinyl Chloride	(1.94)	(24 ¹)	

Results in micrograms per kilograms (ug/l) = parts per million (pph).

Data from 7/92 through 8/96 obtained from Hageman/Aguiar (see references section.)

- even Less than the detection limit for the analysis method.
- a: 82 pph by EPA 8260
- h 15 pph by EPA 8260.
- 35 pph by EPA R260
- d 49.3 pph by EPA 8260.
- ct 5.5 pph by EPA 8260
- 1 5 8 ppb by EPA 8260.
- g: 42 by EPA 8260.
- h. Not confirmed by EPA 8260; one other probable freon is present.
- i. 19 pph by EPA 8260

722-1.HPL SUMMARY LETTER, 6460 HOLLIS STREET, EMERYVILLE, CA

TABLE 12

CUMULATIVE RESULTS FOR HYDROPUNCH LOCATIONS HP-1 AND HP-2 ANALYSES OF GROUNDWATER SAMPLES COLLECTED FEBRUARY 21, 1998 FOR VOCs by EPA 8260

6460 Hollis Street Emeryville, California

Compound Detected	HP-I	HP-2
Acetone	< 15	< 30
Isopropanol	< 200	< 200
Methyl Ethyl Ketone	1.1	< 2
Methyl Isobutyl Ketone	<1	< 2
Sec-Butanol	<5	~ \$
n-Buryl benzene	<1	5.3
see Butyl benzene	<1	11
Isopropylheuzeue	1.0	15
p-tsopropyl toluene	< 1	<2
n-Propyl henzene	<1	13

Results in micrograms per kilograms (ug/l) = parts per million (ppb).

Data from 7/92 through 8/96 obtained from Hageman/Aguiar (see references section.)

1.ess than the detection limit for the analysis method.

NR. Not reported



APPENDIX C

PERMITS



ALAMEDA COUNTY PUBLIC WORKS AGENCY

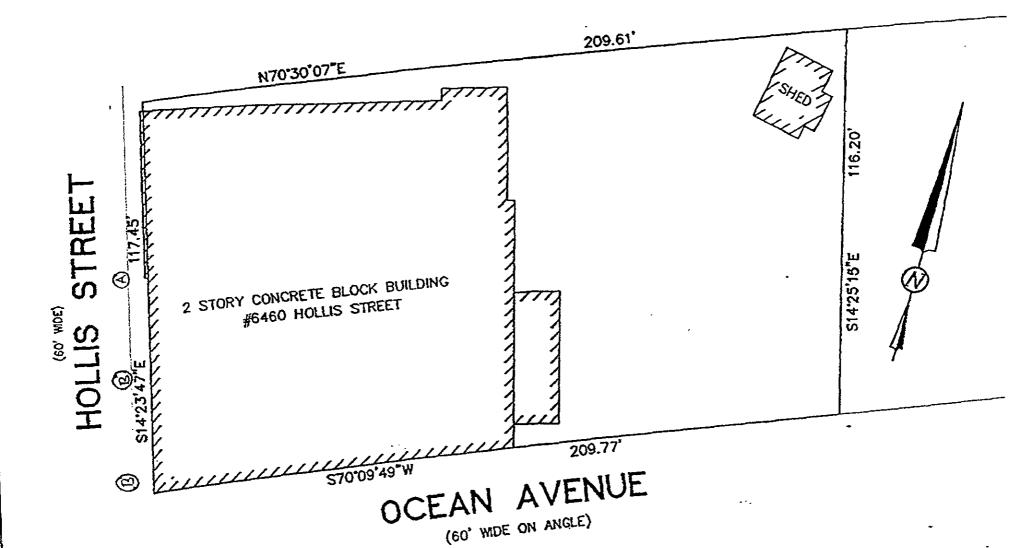
WATER RESOURCES SECTION
951 TURNER COURT, SUITE 300, HAYWARD, CA 94545-2651
PHONE (519) 670-5575 ANDREAS GODFREY FAX (510) 670-5242
(510) 670-5246 ALVIN KAN

DRILLING PERMIT APPLICATION

FOR APPLICANT TO COMPLETE	FOR OFFICE USE
LOCATION OF PROJECT CAGO HOLL'S ST	PERMIT NUMBER 98 WR163
Emaylle GA	WELL NUMBER
	APN
California Coordinates SourceR, CCER.	PERMIT CONDITIONS
CCN	Circled Permit Requirements Apply
CLIENT	(A. GENERAL
None HEN Ltd.	i. A permit application should be submitted so as to
ADDRESS 1255 December 1255 Dec	arrive at the ACPWA office five days prior to
City Emergiale, CA Zip 94608	proposed starting dieto.
	2. Submit to ACPWA within 60 days after completion
APPLICANT	permitted work the original Department of Water
Name Geomatria Consultants, Inc.	Resources Water Well Drillers Keport or equivalent
Fax(4(8) 434 - 1365	well projects, or drilling logs and location sketch for
Address 100 Pine St 10 Floor Phone (415) 434-9400	geotechnical projects.
city San Franciscu Zip 94111	3. Permit is void if project not begun within 90 days of
	approval dato.
TYPE OF PROJECT	B. WATER SUPPLY WELLS
Well Construction Geotechnical Investigation	1. Minimum surface seal thickness is two inches of
ÇEMODIE Protestion D General D	coment grout placed by tremic.
Water Supply 0 Contamination 0	2. Minimum seal depth is SO feet for municipal and
Monitoring Well Destruction	industrial wells or 20 feet for domestic and irrigance
111-1111-1111	wells unless a lesser douth is specially approved.
PROPOSED WATER SUPPLY WELL USE	(C.)GROUNDWATER MONITORING WELLS
New Domestie C. Replacement Domestic D	including piezometers
Municipal C Irrigation C	. 17 Minimum autfore seel thickness is two inches of
Industrial (7 Other O	cement growt placed by Comie.
TURRECUR. 15 Action Comments	Z. Minimum seal depth for monitoring wells is the
DRILLING METHOD:	maximum depth practicable or 20 feet.
Mud Rotary D Air Rotary D Auger X	D. GEOTECHNICAL
Cable D Other D	Backfill bore hale with compected cuttings or heavy
Capie d Galler II.	bentonite and upper two feel with compacted material.
DRULLER'S LICENSE NO. 485165	In areas of known or suspected contamination, tremise
DIGILLER'S LICENSE NO	cement grout shall be used in place of companied cutti
wet - spatcers	E. CATRODIC
WELL PROJECTS Drill Hole Diameter (a. Maximum)	Fill hole abuve anode zone with concrete placed by me
116	P. WELL DESTRUCTION
	See attached.
Surface Seal Depth 1.5 ft. Number 1	G. SPECIAL CONDITIONS
ADDRESS DE LA PROJECTO	
ODOTECHNICAL PROJECTS Number of Borings Maximum	
Hole Diameter 2 in. Depth 10 ft.	•
,	Λ Γ \to Γ
ESTIMATED STARTING DATE 4/21/98	41
ESTIMATED COMPLETION DATE ALLAIGE	APPROVEDDATE
4 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	
I hereby agree to comply with all requirements of this permit and	
Alameda County Ordinanca No. 73-68.	•
	,
	•
APPLICANT'S O D L. AUC 90	•

ENCROACHMENT PERMIT CITY OF EMERYVILLE - PUBLIC WORKS DEPARTMENT 2200 POWELL ST., 12TH FLR. EMERYVILLE, CA 94608 (510) 596 4330

DATE April 15, 1998	`
PROPERTY OWNER H. F. H., L. H.	
CONTACT PERSON Andrew Gretz	 -
ADDRESS 3 CF Occidents	
ADDRESS 1355 Ocean Avenue, Emergrille	
CONTRACTOR Gretz Construction G. LICENSE NO.382305 CLASS B	 .
CONTACT PERSON Andrew. Gietz PHONE NO. 652-4191	
ADDRESS 2915 Took CI TO 1. 1	- .
LOCATION OF WORK (INCLUDE ADDRESS AND STREET NAME AND CROSS STREETS)	_ ·
PLANNED DATE OF COMMENCEMENT April 21, 1998 COMPLETION Complete April 24	
DESCRIPTION OF WORK (INCLUDE AVERAGE DEPTH OF EXCAVATION, MAXIMUM DEPTH, AVERAGE WIDTH, LENGTH, AND ESTIMATED COST OF WORK)	л
mstall ground water maniforing well in singually	· ·
on first side of Hollis Stated, roughly allamine	*ee+
de it traject sold edge of Ocean Avenue right of ha	<u> </u>
THE THE PART THE	matk. Z
CURRENT BUSINESS LICENSE ON FILE YES? NO?	ŝ
CONTRACTOR SIGNATURE Awken 62	9 8
DO NOT WRITE BELOW THIS LINE	gue.
24 HOUR NOTICE PRIOR TO START OF WORK PLAN TO BE SUBMITTED	デボ
	ξ ξ.
	رج ع
NOTE: PROOF OF ADEQUATE INSURANCE MUST BE PRESENTED PRIOR TO START OF WORK OR THIS PERMIT IS VOID.	Oc.
BEE ATTACHED ENCROACHMENT PERMIT GENERAL PROVISIONS	7 8
FOR INSPECTION UPON COMPLETION OF WORK, PLEASE CALL JUAN ARREGUIN AT (510) 596-4333. 96-4336. PLEASE REFER TO THE PERMIT NUMBER LISTED ABOVE.	~ Ave.
ASPECTION COMPLETED ONBY	ēs Q
EFUNDABLE DEPOSIT RETURNED ON BY	2
Maure Laufu S. P. 15	d d
GNATUREY (TITLE)	مي



Amonitoring well

B one-time shallow groundwater sample

PRELIMINARY SURVEY: PLAT

#6460 HOLLIS STREET APRIL 16, 1998 SCALE 1" = 30"

CITY OF EMERYVILLE DEPARTMENT OF PUBLIC WORKS ENCROACHMENT PERMIT GENERAL PROVISIONS

- 1. Authority: Each Encroachment Permit is issued in accordance with Title 7, Chapters 2 and 3 of the Emeryville Municipal Code.
- 2. Revocation: Encroachment Permits are revocable on twenty-four (24) hours' notice.
- 3. Responsible Party: No party other than the named permittee or their agent is authorized to work under any permit.
- 4. Acceptance of Provisions: It is understood and agreed by the permittee that the doing of any work under this permit shall constitute an acceptance of the provisions of this permit and all attachments.
- Notice Prior to Starting Work: Before starting work under he Encroachment Permit, the permittee shall notify the designated City representative two (2) working ays prior to initial start of work. When work has been interrupted for more than five (5) working an additional 24-hour notification is required before estarting work unless a prerranged agreement has been made with the City's representative. inless otherwise specified, all work hall be performed on weekdays and during normal working hours f the City's representative.
- 6. Standards of Construction:
 Il work performed within the City
 right of way shall conform to
 ecognized standards of
 onstruction and the current City's

- Standard Specifications, Standard Plans and Manual on High ad Low Risk Facilities Within Highway Rights-of-Way and any Special Provisions relating thereto.
- 7. Inspection and Approval by City: All work shall be subject to monitoring inspection, and approval by the City. The permittee shall request a final inspection and acceptance of the work within three (3) working days from the completion of the work.
- 8. Permit on the Work Site: The Encroachment Permit or a copy thereof shall be kept at the site of the work and must be shown to any representative of the City or any law enforcement officer on demand. WORK MAY BE SUSPENDED IF PERMIT IS NOT AT JOB SITE AS PROVIDED.
- 9. Conflicting Permits: If a prior encroachment conflicts with the proposed work, the new permittee must arrange for any necessary removal or relocation with the prior permittee. Any such removal or relocation will be at no expense to the City.
- Agencies: The permittee shall, whenever required by law, secure the written authorization for any work that must be approved or permitted by any federal, state or local agency having jurisdiction. Failure to comply with the law, as noted above, will invalidate the City's permit.

- 11. Provisions for Pedestrians: Where facilities exist, a minimum sidewalk and bikepath width of four feet (4) shall be maintained at all times for safe passage through the At no time shall work area. pedestrians be diverted onto a portion of the street used for vehicular traffic. At locations where adjacent alternate walkways cannot be provided, appropriate signs and barricades shall be installed at the limits construction and in advance of the closure at the nearest crosswalk or intersection to divert pedestrians across the street.
- 12. Protection of Traffic: Adequate provisions shall be made for the protection for the traveling pubic. Warning signs, lights and safety devices and other measures required for the public safety, shall conform to the requirements of the Manual of traffic Controls issued by Caltrans. Traffic control for day or nighttime lane closures shall be in conformance with Caltrans Standard Plans for Traffic Control Systems.
- 13. Minimum Interference with Traffic: All work shall be planned and carried out so that there will be the least possible inconvenience to the traveling public. The permittee is authorized to place properly attired flagger(s) to stop and warn conventional traffic. Traffic shall not be unreasonably delayed. Flagging procedures shall be in conformance with the Instructions to Flaggers pamphlet and/or Manual of Traffic Controls for

construction and Maintenance Work Zones issued by Caltrans.

- 14. Care of Drainage: If the work contemplated in any Encroachment Permit shall interfere with the established drainage, ample provision shall be made by the permittee to provide for it as may be directed by the City.
- 15. Water Control: The Contractor shall use Best Management Practices established by the Alameda County Urban Runoff Clean Water Program to prevent construction water, debris, or groundwater from entering storm drains.
- 16. Restoration: In every case, the permittee shall be responsible for restoring to the specifications set forth in Emeryville Municipal Code section 7-2.08 any portion of the City facility which as been excavated or otherwise disturbed by permittee. In addition, all brush, timber, scraps, material, etc., shall be entirely removed and the rightof-way shall be left in as presentable a condition as existed before work started. The permittee shall maintain the surface over acilities placed under any permit. If the facility is not restored as herein provided for, or if the City lects to make repairs, permittee agrees by acceptance of this permit b bear the cost thereof. Permittee nderstands that under Emeryville Municipal Code section 7-2.12, it is able for any subsidence in the racilities disturbed by permittee that occurs within one (1) year from the ate of completion.
 - 7. Work Area Maintenance: ermittee is solely responsible for

maintaining any area it uses in a safe, orderly, and clean condition so as to prevent any injury to damage to property or person.

- 18. Cost of Work: Unless otherwise stated on the permit or other separate written agreement, all costs incurred for work within the City right-of-way pursuant to this Encroachment permit shall be borne by the permittee, and permittee hereby waives all claims for indemnification or contribution from the City for such work.
- 19. Inspection Billing: When the permittee is to be billed for inspections, such costs will be at the current hourly rate established by the City for Permits.
- 20. Submit Plan: For installation of all underground facilities, and all surface work or other activity of consequence, the permittee shall furnish I (one) set of plans showing location and construction or other activity with its application. Thirty (30) days after completion and acceptance of the work, one (1) set of as-built plans shall be submitted to the City.
- 21. Deposit: This permit shall not be effective for any purpose unless, and until the permittee submits to the City a deposit of one thousand dollars (\$1,000).
- 22. Release and Indemnification: Permittee agrees that the City of Emeryville and its officers, employees, agents and volunteers shall not be liable in any manner for injury to or death of any person or for damage to property from any cause. The permittee shall be responsible for any liability, cost,

expense, or claim associated with injuries to or death of any person or damage to property related directly or indirectly to any act or failure to act arising under this permit, or resulting from defects or obstructions, or from any cause whatsoever during the progress of the work, or other activity, or at any subsequent time work or other activity is being performed under the obligations provided by and contemplated by the permit.

The permittee shall indemnify and save harmless the City Emeryville and its officers. employees, agents and volunteer from all claims, suits, actions, expenses, including but not limited to attorney's and expert's fees and costs, resulting from the performance of work or other activity under the permit, or arising out of the failure on the permittee's part to perform his obligations under this permit in respect to maintenance or any obligations, or resulting from defects or obstructions, or from any cause whatsoever during the progress of the work, or other activity or at any subsequent time work or other activity is being performed under the obligations provided by and contemplated by the permit, except as otherwise provided by statute. The duty of the permittee to indemnify and save harmless includes the duties to defend. The permittee waives any and all rights to any type of expressed or implied indemnity against the City, its officers or employees. It is the intent of the parties that the permittee will indemnify and hold harmless the City, its officers, employees, agents and volunteers from any and all

claims, suits or actions as set forth above regardless of the existence or degree of fault or negligence, whether active or passive, primary or secondary, on the part of the City, the permittee, persons employed by the permittee, or persons acting in behalf of the permittee.

this permit is issued with the understanding that any particular action is not to be considered as establishing any precedent (1) on the question of the expediency of permitting any certain kind of encroachment to be erected within right-of-way of the City of Emeryville or (2) as to any utility of the acceptability of any such permits as to any other or future situation.

24. Compliance with laws: Permittee is solely responsible for ompliance with all applicable federal, state and local laws and egulations, including but not mited to all environmental laws and regulations.

Future Moving Installations: If the Encroachment ermit was issued at the request of me Permittee, it is understood that whenever City construction. construction or maintenance work on the City right-of-way requires the installation to be woved, adjusted or relocated, the permittee, at his sole expense, upon quest of the City shall comply th said request.

Attorneys fees: In the event that either party institutes any action, suit, or other dispute olution proceeding based on this

Permit against the other party, the prevailing party is entitled to receive from the losing party all costs or expenses of the proceeding, including but not limited to reasonable attorney fees and court costs.



APPENDIX D GEOMATRIX PROTOCOLS

PROTOCOL

DRILLING AND DESTRUCTION OF SOIL BORINGS

1.0 INTRODUCTION

This protocol describes the procedures to be followed during drilling and destruction of soil borings. The soil borings will provide information about geologic conditions, soil engineering properties, and/or soil quality. If the soil boring is utilized for well installation, the well will be installed in accordance with the protocol INSTALLATION OF WELLS.

The procedures presented herein are intended to be of general use and may be supplemented by a work plan and/or health and safety plan. As the work progresses and if warranted, appropriate revisions may be made by the project manager. Detailed procedures in this protocol may be superseded by applicable regulatory requirements.

If required, permits for drilling of soil borings will be acquired from the appropriate agency(s) before drilling is initiated, and an underground utility check will be conducted before drilling begins. An underground utility check will, at a minimum, consist of contacting a local utility alert service, if available.

2.0 DRILLING

A DAILY FIELD RECORD will be completed for each day of fieldwork, and the original will be kept in the project files.

The soil borings will be drilled using rotary, hollow stem auger, direct-push, or other appropriate method. In all rotary borings, compressed air will be filtered to remove oils before being circulated into the borehole. In mud rotary borings, appropriate drilling fluid additives, such as bentonite, will be used to maintain an open hole and to carry cuttings to the

surface. However, organic drilling fluid additives will only be used with prior project manager approval. The drilling mud will be circulated into a settling tank or basin located near the boring. The viscosity of the drilling fluid will be assessed periodically by the driller and will be controlled throughout the drilling operation to achieve the required results (hole stability, sample return, and mud cake thickness along borehole wall). Only potable water will be used as makeup water for drilling fluid. Exploratory borings drilled using the hollow stem auger method generally do not require the use of drilling fluid. If required, potable water from a municipal supply will be used to maintain boring stability.

The planned depth of each soil boring will be determined by the project manager before drilling. The Geomatrix field geologist/engineer will specify to the drill rig operator the depth of soil sample collection, method of sample retrieval, and other matters pertaining to the satisfactory completion of the borings. Geomatrix staff will observe the volume of drill cuttings returned to assess whether significant cavitation has occurred. Drill cuttings, unused soil samples, and drilling fluids generated during drilling of soil borings will be stored properly for future disposal by the client, unless other arrangements have been made.

The drill rods, augers, hoses, bits, and other components that fluids and cuttings contact will be steam-cleaned before drilling each boring, as well as at the beginning of each project and at the completion of field activities. Drive samplers will be cleaned with Alconox and water or steam-cleaned before each sampling event. Only potable water from a municipal supply will be used for decontamination of drilling equipment. Decontamination rinsate will be collected and stored properly for future disposal by the client, unless other arrangements have been made.

3.0 SAMPLING AND LOGGING

3.1 OBTAINING SAMPLES

Borings will be continuously cored or sampled at depth intervals specified by the project manager, based on the intended use of the boring. Continuous sampling is recommended; however, samples and/or cuttings will be obtained for logging purposes at least every 5 feet for all borings. Drive samples will be used to log hollow stem auger borings if continuous cores are not collected. The samples and/or drill cuttings will be collected and described. A lithologic log of these samples will be made. Samples for chemical analysis will be collected in accordance with the protocol SOIL SAMPLING FOR CHEMICAL ANALYSIS.

3.1.1 Discrete Sampling

For discrete sampling of mud rotary or auger borings, sampling will be accomplished by driving or pushing a split-barrel sampler or Shelby tube. The field geologist/engineer will record information on the BORING LOG pertaining to the sampling, such as rate of penetration, hydraulic ram pressure or drive-hammer blow count, coring smoothness, and sample recovery. In general, the split-barrel sampler will be opened for observation and logging of the retrieved core.

At selected depth intervals, the split-barrel sampler may be fitted with brass or stainless steel liners for collection of soil samples for possible subsequent chemical or physical testing. Samples may be retained for future review and/or preserved for chemical or physical testing, as specified by the project manager. The samples will be stored and labeled to show project number, boring number, and cored interval denoted either by depth or a sequential numbering system. Procedures for preservation and transport of soil samples retained for chemical analysis are presented in the protocol SOIL SAMPLING FOR CHEMICAL ANALYSIS.

3.1.2 Collecting Drill Cuttings

The field geologist/engineer may observe drill cuttings from the drilling fluid return for lithologic information to supplement discrete sampling. Sampling and logging cuttings will be performed as follows:

- 1. The height of the drilling table above ground surface, lengths of the drill bit, sub and drill collars, and length of drill rods should be taken into account in calculating the depth of penetration.
- 2. In mud rotary drilling, a small-diameter, fine mesh hand screen will be used to obtain a sample of the cuttings from the borings by holding the screen directly in the flow of the drill fluid return line. In air rotary drilling, cuttings will be collected after discharge from the cyclone.
- 3. In rotary drilling, a composite sample may be obtained from the return line by leaving the screen in place during the time it takes the driller to advance the boring to a preselected depth.
- 4. In rotary drilling, the travel time for cuttings to reach the surface may be estimated each time the driller adds a new length of drill rod by timing the first arrival of cuttings after circulation is resumed. This travel time can be used along with the depth of penetration to estimate the start and finish of each 5-foot sampling interval.

3.2 LOGGING OF EXPLORATORY BORINGS

The observations of the field geologist/engineer will be recorded on a BORING LOG OR WELL LOG at the time of drilling. The drill rig operator and the field geologist/engineer will discuss significant changes in material penetrated, drilling conditions, hydraulic pressure, drilling action, and drilling fluid circulation rate. The field geologist/engineer will be present during drilling of soil borings and will observe and record such changes by time and depth.

Drill cuttings and core samples will be observed in the field. A lithologic description will be recorded on the BORING LOG using the Unified Soil Classification System (USCS) as described in the American Society of Testing and Materials (ASTM) Standard D 2488-90. This description will include the USCS soil type, grain sizes and estimated percentages of

each, moisture content, color according to the Munsell color charts (Kollmorgen Instruments Corp.), plasticity for fine-grained materials, consistency, and other pertinent information, such as degree of induration, calcareous content, presence of fossils and other distinctive materials.

The original field logs will be retained by the Geomatrix office for review by the responsible professional and for storage in the project files.

4.0 GEOPHYSICAL LOGS

Following completion of drilling, downhole geophysical logs may be performed after the drilling fluid has been circulated to decrease the amount of suspended sediment in the return fluid. Geophysical methods and equipment will be selected to provide stratigraphic or hydrogeologic data appropriate for the project. Geophysical logging will be done as quickly and promply after drilling as feasible, while the boring sidewall is still in stable condition, to reduce the possibility of bridging. Instruments on the logging unit will be adjusted to try to give the maximum definition of strata boundaries. All downhole geophysical equipment will be cleaned before and after use in each borehole.

5.0 FIELD SCREENING

Soil samples collected from the borings may be screened using a portable meter such as a photoionization detector (PID), a flame ionization detector (FID), a lower explosion limit (LEL) meter or other organic vapor meter. The meter may be used to assess the presence of volatile organic compounds (VOCs) or other gases in soil samples. Additional field screening techniques for chemical characterization of soils may include x-ray fluorescence (XRF) and thin-layer chromatography (TLC). Procedures for field screening are described in the protocol SOIL SAMPLING FOR CHEMICAL ANALYSIS.

DRILLING AND DESTRUCTION OF SOIL BORINGS
REVISION DATE: MARCH 1996

Page 5 of 6 DD-5 6.0 DESTROYING SOIL BORINGS

Soil borings that are not completed as monitoring wells will be destroyed by filling the holes

with a neat cement grout, cement/sand grout, or cement/bentonite grout. A high-solids

bentonite grout may be used if appropriate. Geomatrix field staff will calculate the borehole

volume and compare it to the volume of grout used to evaluate whether bridging has occurred.

These calculations and the actual volume emplaced will be noted on the BORING LOG. The

grout will be placed in continuous lifts from the bottom of the boring to a depth of 20 feet

above the water table. The grout will be emplaced by pumping it through the hollow stem

augers, drill pipe, tremie pipe, or flexible hose initially lowered to the bottom of the borings

and raised incrementally as placement proceeds. If hollow stem augers are used, the augers

should be raised incrementally as grout emplacement proceeds. Augers will not be raised in

increments greater than 20 feet or greater than allowed by borehole stability. Borings that are

terminated above the water table and not greater than 20 feet deep may be destroyed by

continuous lifts originating at the ground surface. The grout will be pumped or poured until a

return of fresh grout is visible at the surface. Additional grout may need to be added to the

soil boring if significant settlement has occurred after the grout has set.

Attachments: Daily Field Record

Boring Log

Well Log

DRILLING AND DESTRUCTION OF SOIL BORINGS REVISION DATE: MARCH 1996

DAILY FIELD RECORD



Page 1 of _____

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DAILY FIELD RECORD (continued)



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PROTOCOL

SOIL SAMPLING FOR CHEMICAL ANALYSIS

1.0 INTRODUCTION

This protocol describes the procedures to be followed for collecting soil samples for chemical analysis and conducting soil field screening in conjunction with drilling soil borings and excavating soil. The laboratory must be certified by the appropriate regulating agency for the analyses to be performed.

If required, permits will be acquired from the appropriate agency, and an underground utility check will be performed before drilling or excavating begins. An underground utility check will, at a minimum, consist of contacting a local utility alert service, if available.

The procedures presented herein are intended to be of general use and may be supplemented by a work plan and/or health and safety plan. As the work progresses and if warranted, appropriate revisions may be made by the project manager. Detailed procedures in this protocol may be superseded by applicable regulatory requirements.

2.0 SAMPLE COLLECTION

Soil samples may be collected during drilling or excavating activities. The procedures for sample collection are discussed below.

2.1 SAMPLE COLLECTION DURING DRILLING

The drilling of soil borings will be conducted in accordance with the protocol DRILLING AND DESTRUCTION OF SOIL BORINGS. The soil sampler either will be washed with laboratory grade detergent-water solution to remove soil present and rinsed with potable water, or it may be steam-cleaned prior to and between sampling. Soil samples will be collected in clean brass or stainless steel liners that have been washed with detergent-water solution and

rinsed with potable water or steam-cleaned. The liners will generally be placed in a 2-inch- or 2.5-inch-diameter split-spoon sampler and then driven or pushed into the soil at the selected sampling depth. The sample will be parted at the joints between the liners using a clean, sharp stainless steel knife or spatula. Alternatively, a subsample for chemical analyses may also be collected by driving a smaller-diameter liner into the center of the larger core sample, taking care to reduce the potential for sample disturbance and air space within the liner. If the soil sample is collected using a hand auger, a subsample should be collected from the core of the auger, again taking care to reduce the potential for sample disturbance and air space within the liner. If the sample is to be analyzed for non-volatiles only, a loose sample may be placed in a glass jar. Samples to be analyzed for metals may be homogenized before analysis either in the field or by the laboratory to provide results more representative of average concentrations in the sampling interval.

2.2 SAMPLE COLLECTION DURING EXCAVATION

Excavated soil will be sampled as required under the appropriate agency guidelines, if applicable, or as necessary to provide the data desired. The lateral and vertical dimensions of the excavation, as well as the sample location and depth, will be mapped, and the volume estimated. If possible, samples will be collected from the backhoe or excavator bucket without entering the excavation. Samples may be collected directly from the walls or floor of the excavation, provided Occupational Safety and Health Administration (OSHA) regulations are followed before entering an excavation.

Soil stockpiles also may be sampled after completion of excavation. If they are sampled, the stockpile location, dimensions, and sample locations will be mapped, and the stockpile volume will be estimated. If compositing of soil samples containing volatile compounds is required, it should be performed by the laboratory.

The soil from excavations or stockpiles should be sampled by scraping away 3 to 6 inches of surface soil or hand augering to a known depth. A clean glass jar, brass tube, or stainless

steel tube will be forced into the soil to completely fill the container, or a clean hammer sampler may be used in conjunction with brass or stainless steel liners.

3.0 SAMPLE HANDLING AND PRESERVATION

Soil samples will be handled using the following procedures:

- 1. Clean gloves appropriate for the chemicals of concern will be worn by the sampler before touching the sample containers, and care will be taken to avoid contact with the sample.
- 2. The sample will be quickly observed for color, appearance, and composition. The ends of the liners will be immediately covered with Teflon* sheeting and/or aluminum foil, capped with plastic end caps, and sealed with tape. Glass jars will be immediately sealed with a lid.
- 3. The sample container will be labeled before or immediately after sampling with a self-adhesive label having the following information written in waterproof ink:
 - Geomatrix
 - Project number
 - Sample ID number
 - Date and time sample was collected
 - Initials of sample collector
- 4. The sample will be placed in a chest, that contains ice or blue ice if required, for transport to the laboratory. Table 1 lists common analyses performed and the appropriate storage and handling requirements.

4.0 DOCUMENTATION

4.1 FIELD DATA SHEETS

A DAILY FIELD RECORD will be completed for each day of fieldwork. Locations and unique identification of soil samples collected from soil borings will be recorded on the BORING LOG or WELL LOG. Locations and unique identification of soil samples collected from excavations or stockpiles will be recorded on a DAILY FIELD RECORD, site map, and/or other appropriate form. Samples may also be recorded on a SAMPLE CONTROL

SOIL SAMPLING FOR CHEMICAL ANALYSIS

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LOG SHEET or in the DAILY FIELD RECORD as a means of identifying and tracking the samples. Following review by the project manager, the original field records will be kept in the project file.

4.2 CHAIN-OF-CUSTODY PROCEDURES

After samples have been collected and labeled, they will be maintained under chain-of-custody procedures. These procedures document the transfer of custody of samples from the field to the laboratory. Each sample sent to the laboratory for analysis will be recorded on a CHAIN-OF-CUSTODY RECORD, which will include instructions to the laboratory on the analytical services required.

Information contained on the triplicate CHAIN-OF-CUSTODY RECORD will include:

- Project number
- Signature of sampler
- Date and time sampled
- Sample I.D.
- Number of sample containers
- Sample matrix (soil, water, or other)
- Analyses required
- Remarks, including preservatives, special conditions, or specific quality control measures
- Turnaround time and person to receive laboratory report
- Method of shipment to the laboratory
- Release signature of sampler and signatures of all people assuming custody
- Condition of samples when received by laboratory (to be completed by the laboratory)

Blank spaces on the CHAIN-OF-CUSTODY RECORD will be crossed out between the last sample listed and the signatures at the bottom of the sheet.

The field sampler will sign the CHAIN-OF-CUSTODY RECORD and will record the time and date at the time of transfer to the laboratory or an intermediate person. A set of signatures is required for each relinquished/received transfer, including transfer within Geomatrix. The

original imprint of the CHAIN-OF-CUSTODY RECORD will accompany the sample containers; a duplicate copy will be kept in the Geomatrix project file.

If the samples are to be shipped to the laboratory, the original CHAIN-OF-CUSTODY relinquishing the samples will be sealed inside a plastic bag within the ice chest, and the chest will be sealed with custody tape which has been signed and dated by the last person listed on the chain-of-custody. U.S. Department of Transportation shipping requirements will be followed and the sample shipping receipt will be retained in the project files as part of the permanent chain-of-custody document. The shipping company (e.g., Federal Express, UPS, DHL) will not sign the chain-of-custody forms as a receivor; instead the laboratory will sign as a receivor when the samples are received.

5.0 SOIL FIELD SCREENING

Soil will occasionally be screened using a field instrument or method. Readings should be recorded on the BORING LOG, WELL LOG, DAILY FIELD RECORD, or another form prepared for this purpose. Two screening methods are described below.

5.1 ORGANIC VAPOR METERS

A portable photoionization detector (PID), flame ionization detector (FID), lower explosive limit meter (LEL), or other type of organic vapor meter (OVM) may be used to screen soil. The purpose of the field screening is to assess the presence of volatile organic compounds (VOCs) in the soil. The meter measures total VOCs in the air in parts per million (ppm) by volume in reference to a selected standard. The meter will be calibrated each day prior to the soil sampling. The meter cannot specifically identify each volatile compound, but can be adjusted to be sensitive to selected volatile organics. Before choosing a meter, the response factor of the meter to the chemicals of concern at the site should be considered. Soil should be screened as soon as possible after being exposed to the atmosphere. The general procedure for screening is as follows:

- 1. Using a clean tool, dig a hole to expose fresh soil in a backhoe bucket or stockpile, or separate the brass liners from a driven sample.
- 2. Insert the probe of the OVM into the hole, taking care not to clog the probe with soil. Alternatively, headspace readings may be taken by placing soil in a covered (e.g., aluminum foil or Teflon sheet) clear glass jar or plastic resealable bag, and after several minutes have elapsed, introducing the probe into the headspace area. No soil sample used for headspace screening will be submitted to the laboratory for chemical analysis.
- 3. Record the results in ppm for PIDs and FIDs, and in percent of the lower explosive limit for LELs.
- 4. Ensure that the instrument returns to a zero measurement before the next reading. If necessary, move to an area without measurable organic vapors to zero-out the instrument.

6.0 EQUIPMENT CLEANING

The sampler, brass or stainless steel liners, spatula, and tools used in assembly and disassembly of the soil sampler will be cleaned before and after each use. All soil will be removed from the tools and parts, and the tools will be steam-cleaned or washed in laboratorygrade detergent water with a brush, followed by rinsing in potable water. Decontamination rinsate will be collected and stored properly for future disposal by the client unless other arrangements have been made.

Attachments:

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Table:

Water and Soil Analytical Methods and Sample Handling

Figures:

Daily Field Record Boring Log

Well Log

Chain-of-Custody Record Sample Control Log Sheet

SOIL-SAMPLING FOR CHEMICAL ANALYSIS **REVISION DATE: MARCH 1996**

TABLE 1

WATER AND SOIL ANALYTICAL METHODS AND SAMPLE HANDLING

Parameter	Method	Water Containers ¹	Preservation ¹	Maximum Holding Time ¹	
Total Petroleum Hydrocarbons: as diesel as gasoline	GCFID (3550) ² GCFID (5030) ²	2 - 1 liter amber glass 2 - 40 ml VOA glass	cool on ice HCL to pH 2 in water samples: cool on ice	14 days (unacidified water, 7 days) 14 days (unacidified water, 7 days)	
Benzene, Toluene, Xylene, and Ethylbenzene	EPA 8020	2 - 40 ml VOA glass	HCL to pH 2 in water samples: cool on ice	14 days (unacidified water, 7 days)	
Oil and Grease	5520 E & F (soil) ³ 5520 C & F (water) ³	2 - 1 liter amber glass	H ₂ SO ₄ to pH <2 in water samples: cool on ice	28 days	
Volatile Organics	EPA 8010 EPA 8240 ⁵	2 - 40 ml VOA glass 2 - 40 ml VOA glass	cool on ice ⁴ HCL to pH 2 in water samples: cool on ice	14 days (unacidified water, 7 days) 14 days (unacidified water, 7 days)	
Semi-volatile Organics	EPA 8270	2 - 1 liter amber glass	cool on ice	7 days for extraction, water 14 days for extraction, soil 40 days for analysis	
Polynuclear Aromatic Hydrocarbons	EPA 8310	2 - 40 ml VOA glass	cool on ice	7 days, water 14 days, soil	
Metals (dissolved)	EPA 7000 series for specific metal	1 - 500 ml plastic	Water Samples: field filtration (0.45 micron filter) and field acidify to pH 2 with HNO ₃ except: Cr ⁺⁶ - cool on ice	6 months, except: Hg - 28 days Cr ⁺⁶ - 24 hours, water; 24 hours after prep, soil	

Notes:

- All soil samples should be collected in full, clean brass liners, capped with aluminum foil or Teflon and plastic caps, and sealed with tape. If soil samples are to be analyzed for metals, they may be placed in laboratory-prepared clean glass jars. Soil should be cooled as indicated under "preservation" and maximum holding times apply to both soil and water unless otherwise noted.
- ² For analysis in California, use California DHS recommended procedure as presented in LUFT manual using gas chromatography with a flame ionization detector. In other states, local requirements should be followed.
- ³ Method to be used in California Regional Water Quality Control Board North Coast and Central Valley Regions. In other areas, local requirements should be followed.
- 4 If EPA Methods 8010 and 8020 are to be run in sequence, HCL may be added. Check with the project manager before adding acid.
- 5 Chloroethylvinylether may be detected at concentrations below 50 parts per billion due to degradation of HCL.

References

U.S. EPA, 1986, Test Methods for Evaluating Solid Waste - Physical/Chemical Methods - SW-846, Third Edition, July, and final amendments.

California State Water Resources Control Board, 1989, Leaking Underground Fuel Tank (LUFT) Field Manual, Tables 3-3 and 3-4, October.

California Regional Water Quality Control Boards, North Coast, San Francisco Bay, and Central Valley Regions, 1990, Regional Board Staff Recommendations for Initial Evaluation and Investigation of Underground Tanks, 10 August.

DAILY FIELD RECORD



Page 1 of _ Project and Task Number: Date: Field Activity: Project Name: Weather: Location: Time of OVM Calibration: Time Time PERSONNEL: Name Company ln Out PERSONAL SAFETY CHECKLIST Tyvek Coveralls Steel-toed Boots Hard Hat Safety Goggles Rubber Gloves 1/2-Face Respirator DRUM I.D. **DESCRIPTION OF CONTENTS AND QUANTITY** LOCATION TIME **DESCRIPTION OF WORK PERFORMED**

DAILY FIELD RECORD (continued)



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SAMPLE CONTROL LOG



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PROTOCOL

INSTALLATION AND DESTRUCTION OF WELLS

1.0 INTRODUCTION

This protocol describes the procedures to be followed during the installation or destruction of monitoring, groundwater extraction, and vapor extraction wells. Drilling and logging of soil borings for the well installation will be in conformance with the protocol DRILLING OF SOIL BORINGS. The procedures presented herein are intended to be of general use and may be supplemented by a work plan and/or health and safety plan. As the work progresses and if warranted, appropriate revisions may be made by the project manager. Detailed procedures in this protocol may be superseded by applicable regulatory requirements.

2.0 WELL INSTALLATION

A DAILY FIELD RECORD will be completed for each day of fieldwork, and the original will be kept in the project files. If required, permits will be acquired from the appropriate agency(s), and an underground utility check will be performed before drilling begins. An underground utility check will, at a minimum, consist of contacting a local utility alert service, if available.

After well installation, well completion report(s) will be completed and filed with the State Department of Water Resources or the appropriate agency.

Each groundwater monitoring well will be designed to enable measurement of the potentiometric surface and to permit water sampling of a specific water-bearing zone. Each vapor monitoring well will be designed to enable measurement of pressure conditions and permit sampling of a specific zone. The field geologist/engineer, in consultation with the project geologist or engineer, who will be licensed in the state in which the work is performed if required, will specify the screened interval using the lithologic log and geophysical log (if performed) and will select the well materials and techniques for well completion to be compatible with the subsurface conditions and the intended use of the well. Construction of all wells will be in conformance with the following provisions. A TYPICAL MONITORING WELL CONSTRUCTION DIAGRAM is attached.

2.1 WELL SCREEN AND CASING

The well casing will generally consist of threaded stainless steel or schedule 40 (minimum) polyvinyl chloride (PVC) casing. The inside diameter of the casing will be large enough to permit easy passage of an appropriate water-level probe and equipment for purging wells and water sample collection.

The well screen will generally consist of machine-slotted PVC or wire-wrapped stainless steel screen. The slot sizes will be compatible with the selected filter material. The screened sections will provide flow between the target zone and the well, allowing efficiency in well development and collection of representative samples.

2.2 FILTER MATERIAL

Filter material will be well-graded, clean sand (generally less than 2 percent by weight passing a No. 200 sieve and less than 5 percent by weight of calcareous material). The filter material will be either a standard sand gradation designed for a range of anticipated soil types or a sand gradation specifically designed to fit the soils collected from anticipated well completion zone.

2.3 SETTING SCREENS AND RISER CASING

Upon completion of drilling and/or geophysical logging, the boring will be sounded to verify the total depth, and the well casing will be assembled and lowered into the boring. Well casing materials will be measured to the nearest 0.1 foot and steam-cleaned before being lowered into the borehole. The well assembly will be designed so that the well screen is opposite the target zone. The bottom of the well will be fitted with a secure bottom-end cap. No PVC cement or other solvents will be used to fasten the well casing joints, well screen joints, or end caps. When installing wells in an open borehole, stainless steel centralizers will

INSTALLATION AND DESTRUCTION OF WELLS

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be used immediately above and below the well screen and approximately every 30 to 50 feet along the length of the casing. Centralizers need not be placed on well assemblies installed within augers or drill casings because the auger or drill casing will adequately center the well casing and screen in the borehole.

For borings drilled by the mud rotary method, potable water may be added to the drill mud and circulated in the borehole after completion of the boring. Circulation will continue until the suspended sediment in the return fluid has been decreased. If borehole conditions are relatively stable, the mud will be thinned before the casing assembly is lowered to the specified depth. This is preferred because it reduces the potential for clogging the well screen with thick mud. Conversely, if borehole conditions are relatively unstable, the mud will be thinned after the casing is placed at the specified depth but prior to installation of annular fill materials. After installation of the well assembly, a slurry of filter sand and potable water will then be tremied into the annular space.

For borings drilled using the hollow stem auger method, the filter sand will be placed after the well assembly has been lowered to the specific depth through the augers. The augers will be incrementally raised as the filter sand is placed by free fall through the augers. The depth to the top of the filter pack will be measured after each increment to detect possible bridging. If bridging occurs, it will be broken by washing the filter materials into proper place with potable water or by repeatedly raising and lowering the augers slightly. The amount of water, if any, added to the borehole should be noted on the BORING LOG or DAILY FIELD RECORD.

For monitoring wells, the filter sand will be placed in a calculated quantity sufficient to fill the annular space to a level of about 1 to 2 feet above the top of the well screen. For extraction wells, the level of filter sand above the well screen will be based on site conditions. The depth to the top of the filter pack will be verified by measuring, using a tremie pipe or a weighted tape. Groundwater extraction wells or monitoring wells may be surged before placement of the transition seal to promote filter material settlement, as specified by the project manager.

-INSTALLATION AND DESTRUCTION OF WELLS
REVISION DATE: MARCH 1996

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Once the depth to the top of the filter material has been verified, bentonite or fine sand may be placed in the annular space as a transition seal between the filter material and the grout. A sufficient quantity of bentonite or fine sand will be poured to fill the annular space to a level of about 2 feet above the top of the filter pack. If bentonite is to be placed below standing water, a high-solids bentonite grout will be pumped through a tremie pipe, or pellets may be poured through the annulus. If bentonite is to be placed above standing water, a high-solids bentonite grout should be used or pellets may be placed in 6-inch lifts. Unless prohibited by well conditions, each lift should be hydrated using approximately 1 gallon of potable water per lift of pellets. The completed bentonite transition seal will be allowed to hydrate for at least 30 minutes prior to placing the grout. If a layer of fine sand is placed as the transition seal, the fine sand will be mixed with potable water and placed as a slurry through the tremie pipe or poured dry through the annulus. The depth to the top of the transition seal will be verified by measuring, using the tremie pipe or a weighted tape.

A neat cement grout, cement/sand grout, cement/bentonite grout, or high-solids bentonite grout will be placed from the top of the transition seal to the ground surface. The grout seal will be placed by pumping through a tremie pipe lowered to within 5 feet of the top of the transition seal in mud rotary borings. The grout seal will be placed in hollow stem auger borings by free fall through the augers as they are incrementally raised or by pumping through flexible hose or tremie pipe lowered to near the bottom of the zone to be grouted. The grout must be tremied if there is standing water in the augers above the transition seal. Grout/additive/water mixtures will be determined on a site-specific basis. Typical specifications of grout mixtures include: (a) neat cement/bentonite grout, consisting of a mixture of one sack (94 pounds) of Portland Type I/II cement, approximately 2 to 5 percent by weight (of cement) powdered bentonite, and approximately 6 to 8 gallons of water; (b) neat cement grout, consisting of one sack of Portland cement and approximately 5 to 6 gallons of water; and (c) cement/sand grout, consisting of no more than two parts sand to one part cement and approximately 7 gallons of water. Only potable water will be used to prepare the grout. No work will be done on the monitoring well until after the grout has set approximately 24 hours.

2.4 SURFACE COMPLETION

Upon completion of the well, a suitable slip-on cap, threaded end cap, or waterproof cap will be fitted on the top of the riser casing to reduce the potential for entry of surface runoff or foreign matter. Either a steel protective well cover (e.g., stovepipe) or a vault which may have a traffic-rated cover will be completed at the ground surface. All wells will be locked for security and will be designed to limit surface water infiltration.

2.5 DEVELOPMENT OF GROUNDWATER MONITORING OR EXTRACTION WELLS

When the well installation is complete and the grout has cured a minimum of 24 hours, the well will be developed by surging, bailing, and/or pumping, or other appropriate method as specified by the project manager. The objectives of well development are to remove sediment that may have accumulated during well installation, to consolidate the filter pack around the well screen, and to enhance the hydraulic connection between the target zone and the well. In most instances, a bailer will be used to remove sediment and turbid water from the bottom of the well. A surge block may then be used within the entire screened interval to flush the filter pack of fine sediment. Surging will be conducted slowly to reduce disruption to the filter pack and screen. The well will be bailed again to remove sediment drawn in by the surging process until suspended sediment is reduced.

Following bailing and surging, the well may be further developed using air-lift or pumping methods. A bailer may be used for low-yield wells. If possible, the well will be developed at a higher pumping rate than the anticipated rate of future purging. During development, the turbidity of the water will be monitored, and the pH, specific conductance, and temperature of the return water will be measured. Drawdown and recovery will be measured during and at the end of the development process, respectively, using an electric sounder. Well development will proceed until, in the judgment of the Geomatrix field personnel, the return water is of sufficient clarity. If the screened interval is too long to be developed adequately in one stage, multiple stages will be employed, in which the end of the pump intake will be raised or lowered to various depths, as required.

2.6 DOCUMENTATION

A well construction diagram for each well will be completed in the field on the WELL LOG by the field geologist/engineer and submitted to the reviewing geologist or engineer upon completion of each well. Well installation and construction data will be summarized on the DAILY FIELD RECORD or on a specialized form produced for this purpose. Well development notes and field measurements of water quality parameters will be summarized on a WELL SAMPLING AND/OR DEVELOPMENT RECORD. Following review by the project manager, the original records will be kept in the project file.

3.0 CLEANING OF DRILLING EQUIPMENT

Cleaning of the drill rig and associated drilling equipment will follow the procedures discussed in Section 2.0 of the protocol DRILLING AND DESTRUCTION OF SOIL BORINGS.

All well casing materials will be cleaned before they are installed. Well development equipment will be cleaned before use. The following cleaning procedure has been found to be effective and will be used or adapted as appropriate for general conditions of materials or equipment to be cleaned.

- 1. Steam-rinse with potable water or rinse in deionized or organic-free water.
- 2. Cover with clean plastic to protect materials and equipment from contact with chemical products, dust, or other contaminants.

Alternatively, well casing materials that have been steam-cleaned and sealed in individual airtight plastic bags by the factory can be used.

Decontamination rinsate will be collected and stored properly for future disposal by the client, unless other arrangements have been made.

4.0 WELL DESTRUCTION

Destruction of wells will be completed in accordance with applicable state and local requirements. If required, permits for destruction will be obtained from the appropriate regulatory agency. As part of destruction design and implementation, care will be taken to seal groundwater pathways between multiple aquifers and to limit surface water infiltration through the destroyed borehole.

If practical, the well casing will be removed from the borehole. If the well casing cannot be removed, the casing should be cut and/or pressure-grouted in accordance with regulating agency requirements. For shallow wells and if the well has been completed in the uppermost aquifer, the casing may be pulled from the borehole before auger entry. Alternatively, and if the well has been completed below the uppermost aquifer, the annular fill may be drilled out with hollow stem augers and the casing removed from the borehole through the augers. If the well casing is PVC or other similar material and cannot be removed as described above, it may be removed by drilling out the casing and annular fill using a tricone or drag bit and a rotary drilling method. The borehole will be redrilled to the same or a slightly larger diameter than the original borehole. The redrilled borehole will be plumb and adequately centered, and all of the well casing will be removed.

The borehole will be filled with a neat cement, cement/sand, cement/bentonite grout, or a high-solids bentonite grout. Before its initial set, the grout will be placed in one continuous pour from the bottom of the boring to the ground surface. The grout will be emplaced by pumping through a tremie pipe or flexible hose which is initially lowered to the bottom of the borehole. The augers should be raised incrementally as emplacement proceeds, but not exceed increments of 20 feet or increments greater than allowed by borehole stability. Boreholes that are terminated above the water table and are not greater than 20 feet deep may be grouted by a continuous pour originating at the ground surface. If the aquifer is confined and the head pressure is great, the grout may need to be placed under pressure.

The volume of sealing material used will be calculated and compared to the casing or borehole volume to ensure that bridging has not occurred during well destruction. If the well is in an urban area and if the casing remains in the borehole, a hole will be excavated around the well to a depth of 5 feet, and the casing will be removed to the bottom of the excavation. The sealing material will be allowed to spill over into the excavation to form a cap. The remainder of the excavation will be backfilled with either native material, grout, or concrete.

Attachments:

Daily Field Record

Typical Monitoring Well Construction Diagram

Well Log

Well Sampling and/or Development Record

DAILY FIELD RECORD



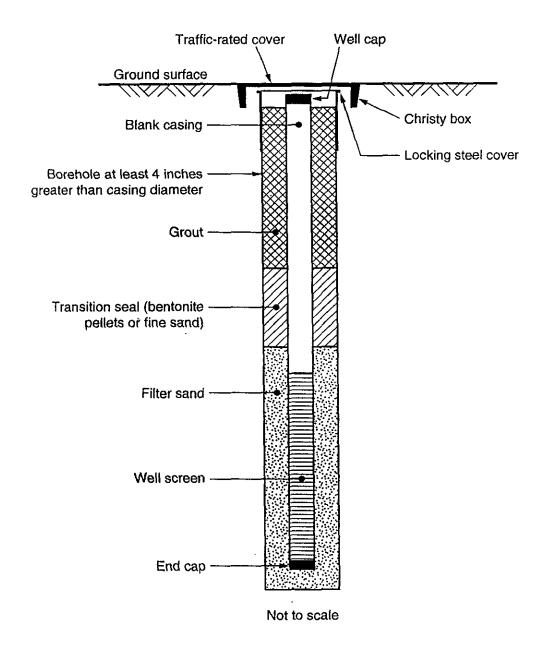
Page 1 of ___ Project and Task Number: Date: Project Name: Field Activity: Location: Weather: Time of OVM Calibration: Time Time PERSONNEL: Name Company Out In 🧬 PERSONAL SAFETY CHECKLIST Tyvek Coveralls Steel-toed Boots Hard Hat Safety Goggles 1/2-Face Respirator **Rubber Gloves** DESCRIPTION OF CONTENTS AND QUANTITY LOCATION DRUM I.D. DESCRIPTION OF WORK PERFORMED TIME

DAILY FIELD RECORD (continued)



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TYPICAL MONITORING WELL CONSTRUCTION DIAGRAM



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WELL SAMPLING AND/OR DEVELOPMENT RECORD

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PROTOCOL

SAMPLING OF GROUNDWATER MONITORING WELLS AND WATER SUPPLY WELLS

1.0 INTRODUCTION

This protocol describes the procedures to be followed during sampling of groundwater monitoring wells and water supply wells for laboratory chemical analysis. The laboratory must be certified by the appropriate regulating agency for the analyses to be performed.

The procedures presented herein are intended to be of general use and may be supplemented by a work plan and/or health and safety plan. As the work progresses and if warranted, appropriate revisions may be made by the project manager. Detailed procedures in this protocol may be superseded by applicable regulatory requirements.

2.0 SAMPLING

2.1 SAMPLE COLLECTION

A. Monitoring Wells

Methods for purging and sampling monitoring wells with dedicated and nondedicated equipment are described in this Section. When practical, the purging and sampling technique adopted for a given site will remain consistent from one sampling event to the next.

A.1 Purging Monitoring Wells

A submersible pump, diaphragm pump, positive displacement pump, which may contain a bladder, or a bailer will be used for evacuating (purging) the monitoring well casing. If the well is to be sampled using equipment that must be separately introduced into the well, the purge intake will be located near the top of the water column for removal of at least one casing volume to remove stagnant water above the screened interval in the well

casing; the pump may then be moved to the midscreen interval to complete the purging progress, if required. If a bailer is used to purge the monitoring well, it will be gently lowered into the well to reduce the potential for aeration of water. Purging will progress at a rate intended to minimize differential drawdown between the interior of the well screen and the filter material to limit cascading water along the inside of the well casing. Procedures for purging slowly recharging wells are discussed in Section A.3.

A minimum of four well casing volumes or one saturated borehole volume, whichever is greater, will be removed to purge the well prior to collection of groundwater samples if the well will be purged with non-dedicated equipment. If a low-flow capacity pump is dedicated in the well, the micropurge method described in Section A.4 may be used to reduce the purge volume. If the well goes dry before four casing volumes are removed, the procedure discussed in Section A.3 will be followed. The saturated borehole volume is the volume of water in the well casing plus the volume of water in the filter pack. For a well with a dedicated pump and packer, a casing volume is defined as the volume of water in the well casing below the inflated packer.

Periodic observations of turbidity and measurements of temperature, pH, and specific electrical conductance (SEC) will be made with field equipment during purging to evaluate whether the water samples are representative of the target zone. Samples will be collected when: (1) a minimum of four sets of parameter readings have been taken; and (2) the temperature, pH, and SEC reach relatively constant values, and the turbidity has stabilized.

A.2 Sampling Monitoring Wells

The sampler will wear clean gloves appropriate for the chemicals of concern while collecting the sample. Samples will be collected directly in laboratory-prepared bottles from the sampling device.

Each sampling episode or day should generally begin with the well having the least suspected concentrations of target compounds. Successive wells should generally be sampled in sequence of increasing suspected concentration.

A Teflon bailer, new disposable bailer, stainless steel positive displacement Teflon bladder pump with Teflon tubing, or a clean electric submersible pump with low-flow sampling capacity will be used to collect the water samples for laboratory chemical analysis.

If a bailer is being used to collect the sample, it will be gently lowered into the well below the point where the purge device was located. Samples will collected in the following order: (1) volatile organic compounds; (2) semivolatile organic compounds; (3) metals; (4) other analytes.

If a bladder pump or electric submersible pump is being used to sample the well for volatile compounds, the flow rate will be adjusted to either 1) approximately 100 milliliters per minute; 2) a rate specifically selected for the well based on groundwater flow rates and well hydraulic conditions; or 3) as low as possible. This rate will be maintained until the discharge line has been purged and the sample collected.

- Purging and Sampling Wells With Slow Recharge A.3 Wells that recharge very slowly may be purged dry once, allowed to recharge, and then sampled as soon as sufficient water is available. In this case, at least two sets of parameter readings of field water quality should be taken, one initially and one after recharge.
- Purging and Sampling Wells Using "Micropurge" Sampling Method A.4 Based on current research, a low-flow-rate, reduced purge method may be used to purge and sample a well with a dedicated pump (Barcelona et al., 1994; Kearl et al., 1994). This method may be used if acceptable to applicable agencies. This method assumes the water within the screened interval is not stagnant, and a small change to the natural flow rate in the screened interval will result in samples with particulates and colloidal material representative of groundwater. The pump should be preset in the screen interval at least 24 hours before the sampling event. A minimum of two pump plus riser pipe volumes should be purged at a flow rate of approximately 100 milliliters per minute or as low as possible based on groundwater flow and well hydraulic conditions. Purging should progress until water quality parameters (pH, SEC, temperature) have reached relatively constant values. Dissolved oxygen readings are recommended, if practical.

B. Water Supply Wells

Water supply wells will be sampled by purging the wells for a period of time adequate to purge the pump riser pipe. Alternatively, if the volume of the riser pipe is unknown, the pressure tank will be drained until the pump cycles on, or the well may be purged until three successive field measurements performed 5 to 10

minutes apart have stabilized. If the well is currently pumping, the sample can be taken without purging the well. Water samples will then be collected from the discharge point nearest the well head. Samples will be collected directly into laboratory-prepared bottles.

C. Extraction Wells

Extraction wells will be sampled while extraction is occurring. Samples will be collected from an in-line sampling port after purging the sampling line. Samples will be collected directly into laboratory-prepared bottles.

A WELL SAMPLING AND/OR DEVELOPMENT RECORD will be used to record the following information:

- Sample I.D.
- Duplicate I.D., if applicable
- Date and time sampled
- Name of sample collector
- Well designation (State well numbering system for water supply wells, and unique sequential number for other wells)
- Owner's name, or other common designation for water supply wells
- Well diameter
- Depth to water on day sampled
- Casing volume on day sampled
- Method of purging (bailing, pumping, etc.)
- Amount of water purged
- Extraordinary circumstances (if any)
- Results of instrument calibration/standardization and field measurements (temperature, pH, specific electrical conductance) and observed relative turbidity
- Depth from which sample was obtained
- Number and type of sample container(s)
- Purging pump intake depth
- Times and volumes corresponding to water quality measurement
- Purge rate

2.2 SAMPLE CONTAINERS AND PRESERVATION

Appropriate pre-cleaned sample containers and preservatives for the analyses to be performed will be obtained from the subcontracted analytical laboratory. Frequently requested analyses and sample handling requirements are listed in Table 1.

2.3 SAMPLE LABELING

Sample containers will be labeled before or immediately after sampling with self-adhesive tags having the following information written in waterproof ink:

- Geomatrix
- Project number
- Sample I.D. number
- Date and time sample was collected
- Initials of sample collector

2.4 QUALITY CONTROL SAMPLES

In order to evaluate the precision and accuracy of analytical data, quality control samples, such as duplicates and blanks, will be periodically prepared. These samples will be collected or prepared and analyzed by the laboratory, as specified in the project Quality Assurance Project Plan (QAPP) or by the project manager.

2.5 HANDLING, STORAGE, AND TRANSPORTATION

Efforts will be made to handle, store, and transport supplies and samples safely. Exposure to dust, direct sunlight, high temperature, adverse weather conditions, and possible contamination will be avoided. Immediately following collection, samples will be placed in a clean chest that contains ice or blue ice (if cooling is required), and will be transported to the subcontracted laboratory as soon as practical, or in accordance with the project QAPP.

3.0 FIELD MEASUREMENTS

Field measurements of temperature, pH, and SEC will be performed on aliquots of groundwater that will not be submitted for laboratory analysis. Field water quality measurements and instrument calibration details will be recorded on the WELL SAMPLING AND/OR DEVELOPMENT RECORD.

3.1 TEMPERATURE MEASUREMENTS

Temperature measurements will be made with a mercury-filled thermometer or an electronic thermistor, and all measurements will be recorded in degrees Celsius.

3.2 pH MEASUREMENT

The pH measurement will be made as soon as possible after collection of the sample, generally within a few minutes. The pH will be measured by immersing the pH probe into an aliquot of groundwater.

The pH meter will be calibrated at the beginning of and once during each sampling day and whenever appropriate, in accordance with the equipment manufacturer's specifications, as outlined in the instruction manual for the specific pH meter used. Two buffers (either pH-4 and pH-7, or pH-7 and pH-10, whichever most closely bracket the anticipated range of groundwater conditions) will be used for instrument calibration.

3.3 SPECIFIC ELECTRICAL CONDUCTANCE MEASUREMENT

SEC will be measured by immersing the conductivity probe into an aliquot of groundwater. The probes used should automatically compensate for the temperature of the sample. Measurements will be reported in units of micro-Siemens (μ S) per square centimeter (equivalent to micromhos or μ mhos) at 25 degrees Celsius.

The SEC meter will be calibrated at the beginning and once during each sampling day in

accordance with the equipment manufacturer's specifications, as outlined in the instruction manual for the SEC meter used. The SEC meter will be calibrated with the available standardized potassium chloride (KCI) solution that is closest to the SEC expected in groundwater below the site.

4.0 DOCUMENTATION

4.1 FIELD DATA SHEETS

A DAILY FIELD RECORD will be completed for each day of fieldwork. A WELL SAMPLING AND/OR DEVELOPMENT RECORD will be used for each well to record the information collected during water quality sampling. Samples may also be recorded on a SAMPLE CONTROL LOG SHEET or in the DAILY FIELD RECORD as a means of identifying and tracking the samples. Following review by the project manager, the original records will be kept in the project file.

4.2 CHAIN-OF-CUSTODY PROCEDURES

After samples have been collected and labeled, they will be maintained under chain-of-custody procedures. These procedures document the transfer of custody of samples from the field to the laboratory. Each sample sent to the laboratory for analysis will be recorded on a CHAIN-OF-CUSTODY RECORD, which will include instructions to the laboratory for analytical services.

Information contained on the triplicate CHAIN-OF-CUSTODY RECORD will include:

- Project number
- Signature of sampler(s)
- Date and time sampled
- Sample I.D.
- Number of sample containers
- Sample matrix (water)
- Analyses required

- Remarks, including preservatives, special conditions, or specific quality control measures
- Turnaround time and person to receive laboratory report
- Method of shipment to the laboratory
- Release signature of sampler(s), and signatures of all people assuming custody.
- Condition of samples when received by laboratory

Blank spaces on the CHAIN-OF-CUSTODY RECORD will be crossed out between the last sample listed and the signatures at the bottom of the sheet.

The field sampler will sign the CHAIN-OF-CUSTODY RECORD and will record the time and date at the time of transfer to the laboratory or to an intermediate person. A set of signatures is required for each relinquished/reserved transfer, including transfer within Geomatrix. The original imprint of the chain-of-custody record will accompany the sample containers. A duplicate copy will be placed in the project file.

If the samples are to be shipped to the laboratory, the original CHAIN-OF-CUSTODY will be sealed inside a plastic bag within the ice chest, and the chest will be sealed with custody tape which has been signed and dated by the last person listed on the chain-of-custody. U.S. Department of Transportation shipping requirements will be followed and the sample shipping receipt will be retained in the project files as part of the permanent chain-of-custody document. The shipping company (e.g., Federal Express, UPS, DHL) will not sign the chain-of-custody forms as a receivor; instead the laboratory will sign as a receivor when the samples are received.

5.0 EQUIPMENT CLEANING

Bailers, sampling pumps, purge pumps, and other non-dedicated purging or sampling apparatus will be cleaned before and after sampling each well. Factory new and sealed disposable bailers may be used for sampling, but may not be reused. Thermometers, pH

electrodes, and SEC probes that will be used repeatedly will be cleaned before and after sampling each well and at any time during sampling if the object comes in contact with foreign matter.

Purged waters and solutions resulting from cleaning of purging or sampling equipment will be collected and stored properly for future disposal by the client, unless other arrangements have been made.

Cleaning of reusable equipment that is not dedicated to a particular well will consist of the following:

- Bailers the inside and outside of bailers will be cleaned in a solution of laboratory-grade detergent and potable water, followed by a rinse with deionized (DI) water. They may also be steam-cleaned, followed by a DI water rinse. If samples are to be collected for metals analysis, the Teflon° bailer may be rinsed with a pH2 nitric acid solution followed by a double DI rinse.
- Purge Pumps All downhole, reusable portions of purge pumps will be steam-cleaned on the outside. If the pump does not have a backflow check valve, the inside of the pump and tubing also should be steam-cleaned. For a purge pump with a backflow check valve, the interior of the pump and tubing may be cleaned by pumping a laboratory-grade detergent and potable water solution through the system followed by a potable water rinse, or by steam-cleaning.
- Water Quality Meters All meters will be cleaned by rinsing the probe portions in DI water, and allowing to air dry.
- Bailer Tripod The tripod cable will be steam-cleaned or rinsed with DI water.

Sample bottles and bottle caps will be cleaned by the subcontracted laboratory using standard EPA-approved protocols. Sample bottles and bottle caps will be protected from contact with solvents, dust, or other contamination. Sample bottles will not be reused.

6.0 REFERENCES

Barcelona, M.J., et al., 1994, Reproducible Well-Purging Procedures and VOC Stabilization Criteria for Ground-Water Sampling: *Groundwater*, January-February.

Kearl, P.M., et al., 1994, Field Comparison of Micropurging vs. Traditional Ground Water Sampling: *Ground Water Monitoring Review*, Fall.

Attachments:

Water and Soil Analytical Methods and Sample Handling

Well Sampling and/or Development Record

Daily Field Record Chain-of-Custody Record Sample Control Log Sheet

TABLE 1 WATER AND SOIL ANALYTICAL METHODS AND SAMPLE HANDLING

Parameter	Method	Water Containers ¹	Preservation ¹	Maximum Holding Time ¹
Total Petroleum Hydrocarbons: • as diesel • as gasoline	GCFID (3550) ² GCFID (5030) ²	2 - 1 liter amber glass 2 - 40 ml VOA glass	cool on ice HCL to pH 2 in water samples: cool on ice	14 days (unacidified water, 7 days) 14 days (unacidified water, 7 days)
Benzene, Toluene, Xylene, and EPA 8020 Ethylbenzene		2 - 40 ml VOA glass	HCL to pH 2 in water samples: cool on ice	14 days (unacidified water, 7 days)
Oil and Grease	5520 E & F (soil) ³ 5520 C & F (water) ³	2 - 1 liter amber glass	H ₂ SO ₄ to pH <2 in water samples: cool on ice	28 days
Volatile Organics	EPA 8010 EPA 8240 ⁵	2 - 40 ml VOA glass 2 - 40 ml VOA glass	cool on ice ⁴ HCL to pH 2 in water samples: cool on ice	14 days (unacidified water, 7 days) 14 days (unacidified water, 7 days)
Semi-volatile Organics	EPA 8270	2 - 1 liter amber glass	cool on ice	7 days for extraction, water 14 days for extraction, soil 40 days for analysis
Polymelear Aromatic Hydrocarbons	EPA 8310	2 - 40 ml VOA glass	cool on ice	7 days, water 14 days, soil
Metals (dissolved)	EPA 7000 series for specific metal	1 - 500 ml plastic	Water Samples: field filtration (0.45 micron filter) and field acidify to pH 2 with HNO ₃ except: Cr ⁺⁶ - cool on ice	6 months, except: Hg - 28 days Cr ⁺⁶ - 24 hours, water; 24 hours after prep, soil

Notes:

- All soil samples should be collected in full, clean brass liners, capped with aluminum foil or Teflon and plastic caps, and sealed with tape. If soil samples are to be analyzed for metals, they may be placed in laboratory-prepared clean glass jars. Soil should be cooled as indicated under "preservation" and maximum holding times apply to both soil and water unless otherwise noted.
- For analysis in California, use California DHS recommended procedure as presented in LUFT manual using gas chromatography with a flame ionization detector. In other states, local requirements should be
- Method to be used in California Regional Water Quality Control Board North Coast and Central Valley Regions. In other areas, local requirements should be followed.
- If EPA Methods 8010 and 8020 are to be run in sequence, HCL may be added. Check with the project manager before adding acid.
- Chloroethylvinylether may be detected at concentrations below 50 parts per billion due to degradation of HCL.

U.S. EPA, 1986, Test Methods for Evaluating Solid Waste - Physical/Chemical Methods - SW-846, Third Edition, July, and final amendments.

California State Water Resources Control Board, 1989, Leaking Underground Fuel Tank (LUFT) Field Manual, Tables 3-3 and 3-4, October.

California Regional Water Quality Control Boards, North Coast, San Francisco Bay, and Central Valley Regions, 1990, Regional Board Staff Recommendations for Initial Evaluation and Investigation of Underground Tanks, 10 August.



WELL SAMPLING AND/OR DEVELOPMENT RECORD

Well ID:_							lnit	ial Dep	th to V	Vater:				
Sample I	D:	[De	pth to V	Nater	after Sampling:				
Sample [Depth:													
Project a	nd Task N	lo.:												
Project N	ame:		<u> </u>				1 Casing/Borehole Volume =							
Date:							(Circle one)							
Sampled	Ву:							casing/ rcle on		ole Volumes =				
Method o	of Purging	· 					•		•	prehole				
Method o	of Samplin	ng:	 _							ved:				
Time	Intake Depth	Rate (gpm)	Cum. Vol. (gal.)	Ten (°C		pl- (uni	ts)	Spec Electr Conduc (µS/c	ical tance	Remarks (color, turbidity, and sediment)				
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	ent Readi	ng												
SPEC	IFIC ELE	CTRICAL	CONDU	CTANC	E - C	ALIB	RAT	ION 🦠	Mod	el or Unit No.:				
KCL Solution (μS/cm = μmhos/cm)														
Temper	ature °C								1					
Instrum	ent Readi	ng			<u> </u>				<u> </u>					
Notes:				 –										
			-,					 						
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DAILY FIELD RECORD



Page 1 of _____

Project and Tas	sk Number:		Date:							
Project Name:			Field Activity:							
Location:			Weather:							
Time of OVM C	alibration:				,					
PERSONNEL:	Name		Company	elep graphic com	Time Time In Out					
				i						
	AFETY CHECKLIS	T								
Steel-toe		Hard Ha		Tyvek Co						
Rubber		Safety C	Committee of the control of the cont	CONTRACTOR AND AND ADDRESS OF THE PARTY OF T	Respirator					
DRUM I.D.	, , , , DESCRIPT	ION OF CONTEN	TS AND QUANTITY		LOCATION					
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DAILY FIELD RECORD (continued)



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Project and Tas	sk Number:	Date:		
Time	#	Description of Work Performed		
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CHAIN-OF-CUSTODY RECORD							N₂						Date:							Page	of							
Project N	o.:										ANA	ALYS	SES											REMARK	<u>s</u>			
Samplers (Signatures):			EPA Method 8010	A Method 8020	EPA Method 8020 (BTEX only)	EPA Method 8240	EPA Method 8270	TPH as gasoline	TPH as diesel													Cooled	Soil (S), Water (W), or Vapor (V)	Acidified	Number of containers	Addit	ional Comme	ents
Date	Time	Sample Number		<u> </u>	<u></u>	<u> </u>	<u> </u>	=						_		-					-	3	00	×	Z			
 			Tt	ımarı	ound t	ime:	<u> </u>	<u> </u>	<u> </u>	LI	Re	sults	to:		<u> </u>					Tota	No.	of co	ntain	ers:				
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SAMPLE CONTROL LOG Laboratory:_____ Project Name: _____ Page _____ of ____ Project and Task No.: Turnaround Time, Sample Location, Handling Notes, Chain-of-Custody Remarks, et. (Duplicate, blank info, etc.) Date Date Sample Number Sampling Analyses Requested Sent to Results C.O.C. Sampling Number Lab Due Time (ID) Date '



APPENDIX E LITHOLOGIC LOGS AND WELL CONSTRUCTION DETAILS

PROJECT. RIX PROPERT 6460 Hollis Str Emeryville, Ca	reet	Log of Borin	g No. GGW1
BORING LOCATION: Hollis	Street (Approximately 80 feet north of Ocean Ave.)	ELEVATION AND DATUM: Not surveyed	· · · · · · · · · · · · · · · · · · ·
DRILLING CONTRACTOR: (Gregg Drilling, Inc.	DATE STARTED: 4/22/98	DATE FINISHED: 4/22/98
DRILLING METHOD: Direct	t push	TOTAL DEPTH: 12.0 feet bgs	MEASURING POINT: Ground surface
DRILLING EQUIPMENT: Ge	poprobe 5400	PERTITO WATER FIRST	COMPL.
SAMPLING METHOD: Maci	ro-core system 1 3/4-inch	LOGGED BY: Preston Gaines	
HAMMER WEIGHT:	DROP:	RESPONSIBLE PROFESSION C. Crocker	DNAL: REG. NO. RG 6114
HAMMER WEIGHT: SAMPLES Sample	e. cementation, react w/HCl geo. inter nedium sand, 25% r gravel % fine sand, siley 10G 4/1), hard 50% medium e subangular 55% coarse sand, vel		
13-			
Project No. 4710.01	Geomatrix Consulta	ants	B-1 (12/95) Figure

PROJECT. RIX PROPER 6460 Hollis St Emeryville, Ca	reet	Log of Borin	g No. GGW2
BORING LOCATION: Corn	er of Hollis Street and Ocean Avenue	ELEVATION AND DATUM: Not surveyed	
DRILLING CONTRACTOR.	Gregg Drilling, Inc.	DATE STARTED: 4/22/98	DATE FINISHED: 4/22/98
DRILLING METHOD. Direc	et push	TOTAL DEPTH: 12.0 feet bgs	MEASURING POINT: Asphalt road surface
DRILLING EQUIPMENT: G	eoprobe 5400	DEBTH TO WATER FIRS	
SAMPLING METHOD: Mac	pro-core system 1 3/4-inch	LOGGED BY: Preston Gaines	100.090
HAMMER WEIGHT:	DROP:	RESPONSIBLE PROFESSIO	DNAL: REG. NO. RG 6114
DEPTH (feet) Sample No. Sample Sample Foot Foot (ppm)	DESCRIPTION NAME (USCS Symbol) color, moist, % by weight , plast , consistency, structu Surface Elevation:	<u> </u>	REMARKS
	4 inches asphalt pavement		
1 -	SILTY SAND with GRAVEL (SM) Dark brown (7.5YR 3/2), dry, 55% fine to 25% low plasticity fines, 20% fine subang subrounded gravel		
3-	LEAN CLAY (CL) Very dark gray (7.5YR 3/1), moist, 95% fi medium plasticity	ines, 5% fine sand,	
	Clayey sand (SC)		
	Color change to dark greenish gray (20	aley 10G 4/1), firm	
5-	SANDY LEAN CLAY with GRAVEL (CL) Brown (7.5YR 4/2), moist, 50% fines, 35% sand, 15% fine subrounded gravel, mediu		
6-	Color change to dark greenish gray (19	OGY 4/1) -	
7-	Hard	-	
9-	FAT CLAY (CH) Yellowish brown (10YR 5/4), moist, 100% firm	fines, high plasticity,	
	Trace fine gravel	} -	
10-	CLAYEY SAND (SC) Brown (7.5YR 4/2), wet, 60% fine sand, 4 plasticity fines	0% medium	
12-	FAT CLAY (CH) Very dark gray (Gley1 3/N), wet, 100% fir firm	nes, high plasticity,	
13-	Bottom of boring at 12.0 feet bgs.	-	
14-		-	
15			B-1 (12/95)
Project No. 4710 01	Geomatrix Consult	ants	Figure

PROJECT: RIX PROPERTY 6460 Hollis Street Emeryville, California	a	Log of Borin	g No. GGW3
BORING LOCATION: Behind buil	lding - low depression area	ELEVATION AND DATUM: Not surveyed	
DRILLING CONTRACTOR: Gregg	DATE STARTED: 4/22/98	DATE FINISHED. 4/22/98	
DRILLING METHOD: Hand auge	TOTAL DEPTH: 5.0 feet bgs	MEASURING POINT: Ground surface	
DRILLING EQUIPMENT: Hand au	ıger (only)	DEDTIL TO WATER. FIRST	
SAMPLING METHOD: Drive sam	npler (soil)	LOGGED BY: Preston Gaines	
HAMMER WEIGHT:	DROP:	RESPONSIBLE PROFESSIO C. Crocker	NAL: REG. NO. RG 6114
DEPTH (feet) Sample No. Sample Blows/ Floot COVM Reading (ppm)	DESCRIPTION (USCS Symbol) color, moist, % by weight, plast, consistency, structure Surface Elevation:	e, cementation, react w/HCl, geo inter.	REMARKS
1- 1- 1- 1- 1- 1- 1- 1- 1- 1- 1- 1- 1- 1	SILTY GRAVEL with SAND (GM) Dark brown (7.5YR 3/2), dry, 50% fine to colow plasticity fines, 20% medium to coarse LEAN CLAY with SAND (CL) Dark reddish gray (2.5YR 3/1), moist, 85% medium sand, medium plasticity, soft Wet SILTY SAND with GRAVEL (SM) Very dark gray (7.5YR 3/1), wet, 50% medians and some plasticity fines, 20% fine to coarse SILT with SAND (ML) Black (7.5YR 2.5/1), wet, 85% fines, 15% fines, 10w plasticity, very soft Bottom of boring at 5.0 feet bgs.	fines, 15% fine to	
Project No. 4710.01	Geomatrix Consulta	nts	B-1 (12/95) Figure

PROJECT: RIX PROPER 6460 Hollis St Emeryville, Ca	reet	Log of Boring No. GGW4			
 	crete slab area behind building	ELEVATION AND DATUM: Not surveyed	ELEVATION AND DATUM:		
DRILLING CONTRACTOR:	Gregg Drilling, Inc.	DATE STARTED: 4/22/98	DATE FINISHED: 4/22/98		
DRILLING METHOD: Hand	d auger	TOTAL DEPTH: 6.0 feet bgs	MEASURING POINT: Top of slab		
DRILLING EQUIPMENT: Ha	and auger	PERTUTO MATER: FIR			
SAMPLING METHOD: Driv	re sampler and large 2-in. diam. disposable bailer	LOGGED BY: Preston Gaines	o leet bys ; 3.0 leet bys		
HAMMER WEIGHT:	DROP:	RESPONSIBLE PROFESSIONAL: REG. NO. C. Crocker RG 6114			
T SAMPLES S	DESCRIPTION	O. Olockei	, AG 0114		
(feet) (feet) Sample No. Sample Blows/ Foot COVM Reading	DESCRIPTION NAME (USCS Symbol) color, moist, % by weight, plast., consistency, structure, cementation, react, w/HCl geo		inter REMARKS		
S Sa Sa V					
	8-inch concrete slab		-		
1 - GGW4	SANDY LEAN CLAY with GRAVEL (CL) Dark brown (7.5YR 5/4), moist, 50% fine medium sand, 20% fine to coarse gravel	s, 30% fine to			
2- 15 3-	LEAN CLAY with GRAVEL (CL) Yellowish brown (10YR 5/4) with very da mottling, moist, 75% fines, 15% fine grav trace wood fragments, medium plasticity				
4-	Fine to medium sand to 15%, increas coarse gravel to 5%, wet		-		
5-	Color change to bluish gray (Gley2 6/	4)]		
	Color change to black (1Gley 2.5/N)	')			
6-			-		
4	Bottom of boring at 6.0 feet bgs.				
7-			-		
-			_		
8-					
9-					
10-					
_			-		
11-			4		
-			-		
12-			+		
+			-		
13-			1		
14-					
7					
15			B-1 (12/9)		

PROJECT: RIX PROPERTY 6460 Hollis Street Emeryville, California				Log of Well No. MW-4				
BORING LOCATION: Hollis Street (north end of building)			ELEVATION AND DATUM: Ground surface of Hollis Street					
DRILLING CONTRACTOR: Gregg Drilling, Inc.			DATE STARTED: 4/22/98	DATE STARTED:				
DRILLING METHOD: 8 1/4-inch hollow stem auger			TOTAL DEPTH: 13.0 feet		4/22/98 SCREEN INTERVAL: 3.0-13.0 feet			
DRILLING EQUIPMENT: Simco limited access rig (2400)			DEPTH TO FIRS	COMPL.	CASING: 2-inch ?????			
SAMPLIN	IG M	ETH	OD:	Direc	t push, Geoprobe 54	LOGGED BY:)	2-inch fffff
HAMMER WEIGHT: DROP:			<u> </u>	C. Grocker RESPONSIBLE P C. Grocker	ROFESSIONAL:	REG. NO.		
e e d	SAM		Foot	OVM Reading (ppm)	DESCRIPTION NAME (USCS Symbol), color, moist, % I consistency, structure, cementation, react	by weight., plast ,	į.	ONSTRUCTION DETAILS R DRILLING REMARKS
Sa		S a	5 04	ð	Surface Elevation.			
1	ļ				Asphalt			Traffic-rated G-5 Christy Box
1-					WELL-GRADED SAND with SI Strong brown (7.5YR 5/6), mois coarse sand, 10% low plasticity Large cobble	st, 90% fine to		Locking water-tight cap 81/4-inch diameter borehole.
3-					LEAN CLAY with SAND (CL) Greenish gray (Gley 5/5GY), m 20% fine to medium sand, trace			2-inch diameter Sch. 40 PVC blank casing ——— Cement/bentonite
4-					plasticity, firm Increased clay content, med	lium plasticity		grout Bentonite chip seal
5-					and the state of t			Monterey 2/12 sand filter pack
6- 7-			1		Zone with up to 30% fine to	coarse sand		2-inch diameter Sch 40 PVC 0.020-inch slotted screen
8-			;		LEAN CLAY (CL) Greenish gray (Gley 5/5GY), m 5-10% fine to medium sand, me firm			
10-					Zone with up to 50% angula	ar gravel		
11-					SANDY LEAN CLAY (CL) Greenish gray (Gley 5/5GY), w 65-70% fines, 30-35% fine to c sand, trace gravel, low plasticit	oarse		
12-					Zone of clayey sand	,,, 550		Threaded PVC end
13-					LEAN CLAY (CL) Mottled greenish gray (Gley 5/5 olive brown (2.5Y 5/3), 95% fin coarse sand, trace white angul medium plasticity, hard	es, 5% fine to		cap
15					Bottom of boring at 13 feet bgs	S		
10						trix Consultants		W-1 (12/9



APPENDIX F LABORATORY ANALYTICAL DATA SHEETS

American Environmental Network

Certificate of Analysis

DOHS Certification: 1172

AIHA Accreditation: 11134

PAGE 1

GEOMATRIX CONSULTANTS 100 PINE ST., SUITE 1000 SAN FRANCISCO, CA 94111

ATTN: JENNIFER PATTERSON CLIENT PROJ. ID: 4710

C.O.C. NUMBER: 4217

REPORT DATE: 06/02/98

DATE(S) SAMPLED: 04/22/98

DATE RECEIVED: 04/22/98

AEN WORK ORDER: 9804229

PROJECT SUMMARY:

On April 22, 1998, this laboratory received 10 soil sample(s).

Client requested 6 sample(s) be composited into 3 and analyzed with 4 discreet sample(s) for chemical parameters. Results of analysis are summarized on the following page(s). Please see quality control report for a summary of QC data pertaining to this project.

Samples will be stored for 30 days after completion of analysis, then disposed of in accordance with State and Federal regulations. Samples may be archived by prior arrangement.

If you have any questions, please contact Client Services at (510) 930-9090.

Reviewed by:

William Subra

3440 Vincent Road • Pleasant Hill, CA 94523 • (510) 930-9090 • FAX (510) 930-0256

SAMPLE ID: S-03

AEN LAB NO: 9804229-05 AEN WORK ORDER: 9804229 CLIENT PROJ. ID: 4710 DATE SAMPLED: 04/22/98 DATE RECEIVED: 04/22/98

REPORT DATE: 06/02/98

ANALYTE	METHOD/ CAS#	RESULT	REPORTING LIMIT UNITS	DATE ANALYZED
#Digestion, for Furnace	EPA 3050	-	Prep Date	e 04/30/98
#Digestion for ICP/AA	EPA 3050A	-	Prep Date	04/30/98
Arsenic	EPA 6010	ND	. 10 mg/kg	05/08/98
Selenium	EPA 6010	ND	20 mg/kg	05/08/98
EPA 8260A - Soil Benzene Bromobenzene Bromochloromethane Bromodichloromethane Bromoform Bromomethane n-Butylbenzene sec-Butylbenzene tert-Butylbenzene Carbon Tetrachloride Chlorobenzene Chlorotethane Chlorotethane Chlorotoluene Dibromochloromethane 1.2-Dibromo-3-chloropropane 1.2-Dibromoethane Dibromomethane 1.2-Dichlorobenzene 1.3-Dichlorobenzene 1.4-Dichlorobenzene 1.1-Dichloroethane 1.1-Dichloroethane 1.1-Dichloroethene cis-1.2-Dichloroethene trans-1.2-Dichloroethene 1.3-Dichloropropane 1.3-Dichloropropane 1.3-Dichloropropane 1.1-Dichloropropane 1.1-Dichloropropane 1.1-Dichloropropane 1.1-Dichloropropane	EPA 8260A 71-43-2 108-86-1 74-97-5 75-27-4 75-25-2 74-83-9 104-51-8 135-98-8 98-06-6 56-23-5 108-90-7 75-00-3 67-66-3 74-87-3 95-49-8 106-43-4 124-48-1 96-12-8 106-93-4 74-95-3 95-50-1 541-73-1 106-46-7 75-71-8 75-34-3 107-06-2 75-35-4 156-59-2 156-60-5 78-87-5 142-28-9 594-20-7 563-58-6 100-41-4	ND ND D D D D D D D D D D D D D D D D D	30 ug/Kg 30 ug/Kg 30 ug/Kg 30 ug/Kg 50 ug/Kg 50 ug/Kg 30 ug/Kg	05/03/98 05/03/98

SAMPLE ID: S-03 AEN LAB NO: 9804229-05 AEN WORK ORDER: 9804229 CLIENT PROJ. ID: 4710

ANALYTE	METHOD/ CAS#	RESULT	REPORTING LIMIT UNIT	DATE S ANALYZED
Hexachlorobutadiene Isopropylbenzene p-Isopropyltoluene Methylene Chloride Naphthalene n-Propylbenzene Styrene 1.1.1.2-Tetrachloroethane 1.1.2.2-Tetrachloroethane Tetrachloroethene Toluene 1.2.3-Trichlorobenzene 1.1.1-Trichlorobenzene 1.1.2-Trichloroethane Trichloroethene Trichlorofluoromethane 1.2.3-Trichloropropane 1.2.4-Trimethylbenzene 1.3.5-Trimethylbenzene Vinyl Chloride Xylenes. Total Acetone 2-Butanone Carbon Disulfide cis-1.3-Dichloropropene trans-1,3-Dichloropropene 2-Hexanone 4-Methyl-2-pentanone Vinyl Acetate 2-Chloroethyl Vinyl Ether	87-68-3 98-82-8 99-87-6 75-09-2 91-20-3 103-65-1 100-42-5 630-20-6 79-34-5 127-18-4 108-88-3 87-61-6 120-82-1 71-55-6 79-00-5 79-01-6 75-69-4 96-18-4 95-63-6 108-67-8 75-01-4 1330-20-7 67-64-1 78-93-3 75-15-0 10061-01-5 10061-02-6 591-78-6 108-05-4 110-75-8	ND	50 ug/Kg 30 ug/Kg	05/03/98 05/03/98
CCR 17 Metals Silver Barium Beryllium Cadmium Cobalt Chromium Copper Mercury Molybdenum Nickel Lead	EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 7471 EPA 6010 EPA 6010	ND 190 * ND 7 * 32 * ND ND 19 *	0.5 mg/kg 0.5 mg/kg 1 mg/kg 2 mg/kg 2 mg/kg 0.06 mg/kg 2 mg/kg	05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/03/98 05/01/98 05/01/98

GEOMATRIX CONSULTANTS

SAMPLE ID: S-03

AEN LAB NO: 9804229-05 AEN WORK ORDER: 9804229 CLIENT PROJ. ID: 4710 **DATE SAMPLED:** 04/22/98

DATE RECEIVED: 04/22/98

REPORT DATE: 06/02/98

ANALYTE	METHOD/ CAS#	RESULT	REPORTING LIMIT	UNITS	DATE ANALYZED
Antimony Thallium Vanadium Zinc	EPA 6010 EPA 6010 EPA 6010 EPA 6010	ND ND 35 * 23 *	50 m 2 m	g/kg g/kg g/kg g/kg	06/08/98 05/01/98 05/01/98 05/01/98

Reporting limits elevated for EPA 8260 due to high levels of target compounds. Sample run at dilution.

ND = Not detected at or above the reporting limit
 * = Value at or above reporting limit

SAMPLE ID: S-05 AEN LAB NO: 9804229-06 AEN WORK ORDER: 9804229 CLIENT PROJ. ID: 4710

ANALYTE	METHOD/ CAS#	RESULT	REPORTING LIMIT	UNITS	DATE ANALYZED
#Digestion, for Furnace	EPA 3050	-		Prep Date	04/30/98
#Digestion for ICP/AA	EPA 3050A	-	•	Prep Date	04/30/98
Arsenic	EPA 6010	ND	10	mg/kg	05/08/98
Selenium	EPA 6010	ND	20	mg/kg	05/08/98
EPA 8260A - Soil Benzene Bromobenzene Bromochloromethane Bromodichloromethane Bromoform Bromomethane n-Butylbenzene sec-Butylbenzene tert-Butylbenzene Carbon Tetrachloride Chlorobenzene Chlorotenane Chloroform Chloromethane 2-Chlorotoluene Dibromochloromethane 1,2-Dibromo-3-chloropropane 1,2-Dibromoethane Dibromomethane 1,2-Dichlorobenzene 1,3-Dichlorobenzene 1,4-Dichloroethane 1,1-Dichloroethane 1,1-Dichloroethene cis-1,2-Dichloroethene trans-1,2-Dichloroethene trans-1,2-Dichloroethene 1,2-Dichloropropane 1,3-Dichloropropane 1,3-Dichloropropane 1,1-Dichloropropane 1,1-Dichloropropane	EPA 8260A 71-43-2 108-86-1 74-97-5 75-27-4 75-25-2 74-83-9 104-51-8 135-98-8 98-06-6 56-23-5 108-90-7 75-00-3 67-66-3 74-87-3 95-49-8 106-43-4 124-48-1 96-12-8 106-93-4 74-95-3 95-50-1 541-73-1 106-46-7 75-71-8 75-34-3 107-06-2 75-35-4 156-59-2 156-60-5 78-87-5 142-28-9 594-20-7 563-58-6 100-41-4	** ** ** ** ** ** ** ** ** ** ** ** **	3000 3000 5000 3000 3000 3000 3000 3000	ug/Kg ug/Kg ug/Kg	05/03/98 05/03/98

SAMPLE ID: S-05 AEN LAB NO: 9804229-06 AEN WORK ORDER: 9804229 CLIENT PROJ. ID: 4710 DATE SAMPLED: 04/22/98

DATE RECEIVED: 04/22/98 REPORT DATE: 06/02/98

ANALYTE	METHOD/ CAS#	RESULT	REPORTING LIMIT UNITS	DATE ANALYZED
Hexachlorobutadiene Isopropylbenzene p-Isopropyltoluene Methylene Chloride Naphthalene n-Propylbenzene Styrene 1.1.2-Tetrachloroethane 1.1.2-Tetrachloroethane Tetrachloroethene Toluene 1.2.3-Trichlorobenzene 1.1.1-Trichlorobenzene 1.1.2-Trichloroethane Trichloroethene Trichlorofluoromethane Trichlorofluoromethane 1.2.3-Trichloropropane 1.2.4-Trimethylbenzene 1.3,5-Trimethylbenzene Vinyl Chloride Xylenes, Total Acetone 2-Butanone Carbon Disulfide cis-1,3-Dichloropropene trans-1,3-Dichloropropene 2-Hexanone 4-Methyl-2-pentanone Vinyl Acetate 2-Chloroethyl Vinyl Ether	87-68-3 98-82-8 99-87-6 75-09-2 91-20-3 103-65-1 100-42-5 630-20-6 79-34-5 127-18-4 108-88-3 87-61-6 120-82-1 71-55-6 79-00-5 79-01-6 75-69-4 96-18-4 95-63-6 108-67-8 75-01-4 1330-20-7 67-64-1 78-93-3 75-15-0 10061-01-5 10061-02-6 591-78-6 108-05-4 110-75-8	ND * 1.000 * ND * 1.000 * 1.00	3000 ug/Kg 3000 ug/Kg 3000 ug/Kg 3000 ug/Kg 3000 ug/Kg 3000 ug/Kg 3000 ug/Kg 3000 ug/Kg 3000 ug/Kg 3000 ug/Kg 3000 ug/Kg 3000 ug/Kg 3000 ug/Kg 3000 ug/Kg 3000 ug/Kg 3000 ug/Kg 3000 ug/Kg 3000 ug/Kg 5000 ug/Kg	05/03/98 05/03/98
CCR 17 Metals Silver Barıum Beryllium Cadmium Cobalt Chromium Copper Mercury Molybdenum Nickel Lead	EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 7471 EPA 6010 EPA 6010	ND 140 * 0.6 * ND 13 * 68 * 20 * 0.09 * ND 79 * 45 *	0.5 mg/kg 5 mg/kg 0.5 mg/kg 0.5 mg/kg 1 mg/kg 2 mg/kg 2 mg/kg 0.06 mg/kg 2 mg/kg 4 mg/kg	05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/03/98 05/01/98 05/01/98

GEOMATRIX CONSULTANTS

SAMPLE ID: S-05

AEN LAB NO: 9804229-06 AEN WORK ORDER: 9804229 CLIENT PROJ. ID: 4710 DATE SAMPLED: 04/22/98 DATE RECEIVED: 04/22/98 REPORT DATE: 06/02/98

ANALYTE	METHOD/ CAS#	RESULT	EPORTING LIMIT	UNITS	DATE ANALYZED
Antimony Thallium Vanadium Zinc	EPA 6010 EPA 6010 EPA 6010 EPA 6010	ND ND 40 * 77 *	50 m 2 m	ng/kg ng/kg ng/kg ng/kg	06/08/98 05/01/98 05/01/98 05/01/98

Reporting limits elevated for EPA 8260 due to high levels of non-target compounds. Sample run at dilution.

ND = Not detected at or above the reporting limit

* = Value at or above reporting limit

GEOMATRIX CONSULTANTS

SAMPLE ID: S-04

AEN LAB NO: 9804229-07 AEN WORK ORDER: 9804229 CLIENT PROJ. ID: 4710 DATE SAMPLED: 04/22/98

DATE RECEIVED: 04/22/98

REPORT DATE: 06/02/98

ANALYTE	METHOD/ CAS#	RESULT	REPORTING LIMIT	UNITS	DATE ANALYZED
#Digestion, for Furnace	EPA 3050	-		Prep Date	04/30/98
#Digestion for ICP/AA	EPA 3050A	-		Prep Date	04/30/98
Arsenic	EPA 6010	ND	10	mg/kg	05/08/98
Selenium	EPA 6010	ND	20	mg/kg	05/08/98
EPA 8260A - Soil Benzene Bromobenzene Bromochloromethane Bromodichloromethane Bromoform Bromomethane n-Butylbenzene sec-Butylbenzene tert-Butylbenzene Carbon Tetrachloride Chlorobenzene Chlorotoluene Chlorotoluene 4-Chlorotoluene 0ibromochloromethane 1,2-Dibromo-3-chloropropane 1,2-Dibromoethane Dibromomethane 1,2-Dichlorobenzene 1,3-Dichlorobenzene 1,4-Dichlorobenzene 1,1-Dichloroethane 1,1-Dichloroethane 1,1-Dichloroethane 1,2-Dichloroethene trans-1,2-Dichloroethene trans-1,2-Dichloroethene 1,2-Dichloropropane 1,3-Dichloropropane 1,3-Dichloropropane 1,3-Dichloropropane 1,1-Dichloropropane 1,1-Dichloropropane 1,1-Dichloropropane 1,1-Dichloropropane	EPA 8260A 71-43-2 108-86-1 74-97-5 75-27-4 75-25-2 74-83-9 104-51-8 135-98-8 98-06-6 56-23-5 108-90-7 75-00-3 67-66-3 74-87-3 95-49-8 106-43-4 124-48-1 96-12-8 106-93-4 74-95-3 95-50-1 541-73-1 106-46-7 75-71-8 75-34-3 107-06-2 75-35-4 156-59-2 156-60-5 78-87-5 142-28-9 594-20-7 563-58-6 100-41-4		300 300 500 300 300 300 300 300 500 300 3	ug/Kg ug/Kg	05/03/98 05/03/98

SAMPLE ID: S-04 AEN LAB NO: 9804229-07 AEN WORK ORDER: 9804229 CLIENT PROJ. ID: 4710

ANALYTE	METHOD/ CAS#	RESULT	REPORTING LIMIT UNITS	DATE ANALYZED
Hexachlorobutadiene Isopropylbenzene p-Isopropyltoluene Methylene Chloride Naphthalene n-Propylbenzene Styrene 1.1.1.2-Tetrachloroethane 1.1.2.2-Tetrachloroethane Tetrachloroethene Toluene 1.2.3-Trichlorobenzene 1.2.4-Trichlorobenzene 1.1.1-Trichloroethane Trichloroethene Trichlorofluoromethane 1.2.3-Trichloropropane 1.2.3-Trimethylbenzene 1.3.5-Trimethylbenzene 1.3.5-Trimethylbenzene Vinyl Chloride Xylenes, Total Acetone 2-Butanone Carbon Disulfide cis-1.3-Dichloropropene trans-1.3-Dichloropropene 2-Hexanone 4-Methyl-2-pentanone Vinyl Acetate 2-Chloroethyl Vinyl Ether	87-68-3 98-82-8 99-87-6 75-09-2 91-20-3 103-65-1 100-42-5 630-20-6 79-34-5 127-18-4 108-88-3 87-61-6 120-82-1 71-55-6 79-00-5 79-01-6 75-69-4 96-18-4 95-63-6 108-67-8 75-01-4 1330-20-7 67-64-1 78-93-3 75-15-0 10061-01-5 10061-02-6 591-78-6 108-05-4 110-75-8	ND	300 ug/Kg 300 ug/Kg 300 ug/Kg 300 ug/Kg 300 ug/Kg 300 ug/Kg 300 ug/Kg 300 ug/Kg 300 ug/Kg 300 ug/Kg 300 ug/Kg 300 ug/Kg 300 ug/Kg 300 ug/Kg 300 ug/Kg 300 ug/Kg 500 ug/Kg	05/03/98 05/03/98
CCR 17 Metals Silver Barium Beryllium Cadmium Cobalt Chromium Copper Mercury Molybdenum Lead	EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 7471 EPA 6010 EPA 6010	ND 160 * ND 0.6 * 8 * 120 * 40 * 0.51 * 3 * 60 *	0.5 mg/kg 5 mg/kg 0.5 mg/kg 0.5 mg/kg 1 mg/kg 2 mg/kg 2 mg/kg 0.06 mg/kg 2 mg/kg 4 mg/kg	05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/03/98 05/01/98 05/01/98

GEOMATRIX CONSULTANTS

SAMPLE ID: S-04

AEN LAB NO: 9804229-07 AEN WORK ORDER: 9804229 CLIENT PROJ. ID: 4710

DATE SAMPLED: 04/22/98

DATE RECEIVED: 04/22/98

REPORT DATE: 06/02/98

ANALYTE	METHOD/ CAS#		PORTING LIMIT UNITS	DATE ANALYZED
Antimony	EPA 6010	ND	5 mg/kg	06/08/98
Thallium	EPA 6010	ND	50 mg/kg	05/01/98
Vanadium	EPA 6010	42 *	2 mg/kg	05/01/98
Zinc	EPA 6010	140 *	5 mg/kg	05/01/98

Reporting limits elevated for EPA 8260 due to high levels of target compounds. Sample run at dilution.

ND = Not detected at or above the reporting limit
 * = Value at or above reporting limit

GEOMATRIX CONSULTANTS

SAMPLE ID: S-06

AEN LAB NO: 9804229-08 AEN WORK ORDER: 9804229 CLIENT PROJ. ID: 4710

DATE SAMPLED: 04/22/98 DATE RECEIVED: 04/22/98 REPORT DATE: 06/02/98

ANALYTE	METHOD/ CAS#	RESULT	REPORTING LIMIT	UNITS	DATE ANALYZED
#Digestion, for Furnace	EPA 3050	-		Prep Date	04/30/98
#Digestion for ICP/AA	EPA 3050A	-		Prep Date	04/30/98
Arsenic	EPA 6010	ND	10	mg/kg	05/08/98
Selenium	EPA 6010	ND	20	mg/kg	05/08/98
CCR 17 Metals Silver Barium Beryllium Cadmium Cobalt Chromium Copper Mercury Molybdenum Nickel Lead Antimony Thallium Vanadium Zinc	EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 7471 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010	ND 1,800 * ND 1.5 * 10 * 37 * 53 * 0.32 * ND 26 * 240 * ND 36 * 630 *	0.5 0.5 1 2 0.06 2 4 5 50 2	mg/kg mg/kg mg/kg mg/kg mg/kg mg/kg mg/kg mg/kg mg/kg mg/kg mg/kg mg/kg mg/kg mg/kg mg/kg	05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98

ND = Not detected at or above the reporting limit
* = Value at or above reporting limit

SAMPLE ID: S-07,08 (COMP) **AEN LAB NO:** 9804229-11 AEN WORK ORDER: 9804229 CLIENT PROJ. ID: 4710

ANALYTE	METHOD/ CAS#	RESULT	REPORTIN LIMIT	G UNITS	DATE ANALYZED
#CA Waste Extraction	CA Title 22	-		Extrn Date	05/15/98
Lead in WET Extract	EPA 7420	15	* 0.1	mg/L	05/18/98
#Digestion, for Furnace	EPA 3050	<u></u>		Prep Date	04/30/98
#Digestion for ICP/AA	EPA 3050A	-		Prep Date	04/30/98
Arsenic	EPA 6010	11	* 10	mg/kg	05/08/98
Selenium	EPA 6010	ND	20	mg/kg	05/08/98
CCR 17 Metals Silver Barium Beryllium Cadmium Cobalt Chromium Copper Mercury Molybdenum Nickel Lead Antimony Thallium Vanadium Zinc	EPA 6010 EPA 6010	10 38 41 0.24 ND 41	* 0.5 * 0.5 * 2 * 2 * 0.06 2 * 4 50 *	mg/kg mg/kg mg/kg mg/kg mg/kg mg/kg mg/kg mg/kg mg/kg	05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/03/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98

ND = Not detected at or above the reporting limit
* = Value at or above reporting limit

SAMPLE ID: S-01,02 (COMP) AEN LAB NO: 9804229-12 AEN WORK ORDER: 9804229 CLIENT PROJ. ID: 4710

ANALYTE	METHOD/ CAS#	RESULT	REPORTING LIMIT	UNITS	DATE ANALYZED
#Digestion, for Furnace	EPA 3050	-		Prep Date	04/30/98
#Digestion for ICP/AA	EPA 3050A	-		Prep Date	04/30/98
Arsenic	EPA 6010	ND	10	mg/kg	05/08/98
Selenium	EPA 6010	ND	20	mg/kg	05/08/98
CCR 17 Metals Silver Barium Beryllium Cadmium Cobalt Chromium Copper Mercury Molybdenum Nickel Lead Antimony Thallium Vanadium Zinc	EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 7471 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010	ND 300 * ND 8 * 33 * 0.07 * ND 28 * ND ND 30 * 150 *	5 0.55 1 2 2 0.06 2 2 4 5 5 2 2 4 5 5 2 2 4 5 5 2 5 6 5 6 6 6 6 6 6 6 6 6 6 6 6 6 6	mg/kg mg/kg mg/kg mg/kg mg/kg mg/kg mg/kg mg/kg mg/kg mg/kg mg/kg	05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98

ND = Not detected at or above the reporting limit
* = Value at or above reporting limit

GEOMATRIX CONSULTANTS

SAMPLE ID: S-09,10 (COMP) AEN LAB NO: 9804229-13 AEN WORK ORDER: 9804229 CLIENT PROJ. ID: 4710

ANALYTE	METHOD/ CAS#	RESULT	REPORTING LIMIT	UNITS	DATE ANALYZED
#CA Waste Extraction	CA Title 22	-		Extrn Date	05/15/98
Lead in WET Extract	EPA 7420	8.7 %	0.1	mg/L	05/18/98
#Digestion, for Furnace	EPA 3050	-		Prep Date	04/30/98
#Digestion for ICP/AA	EPA 3050A	-		Prep Date	04/30/98
Arsenic	EPA 6010	ND	10	mg/kg	05/08/98
Selenium	EPA 6010	ND	20	mg/kg	05/08/98
CCR 17 Metals Silver Barium Beryllium Cadmium Cobalt Chromium Copper Mercury Molybdenum Nickel Lead Antimony Thallium Vanadium Zinc	EPA 6010 EPA 6010	ND 870 7 ND 2.0 7 16 7 120 7 0.46 7 ND 43 7 410 7 ND ND 40 7	5.5.5.1.2.2.6.2.2.4.5.0.2.2.2.4.5.0.2.2.2.4.5.0.2.2.2.4.5.0.2.2.2.4.5.0.2.2.2.4.5.0.2.2.2.4.5.0.2.2.2.4.5.0.2.2.2.4.5.0.2.2.2.4.5.0.2.2.2.4.5.0.2.2.2.4.5.0.2.2.2.4.2.2.2.2.2.2.2.2.2.2.2.2.2.2.2		05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98

ND = Not detected at or above the reporting limit

^{* =} Value at or above reporting limit

AEN (CALIFORNIA) QUALITY CONTROL REPORT

AEN JOB NUMBER: 9804229 CLIENT PROJECT ID: 4710

Quality Control and Project Summary

All laboratory quality control parameters were found to be within established limits.

<u>Definitions</u>

Matrix Spike(s): Aliquot of a sample (aqueous or solid) with added quantities of specific compounds and subjected to the entire analytical procedure. Matrix spike and matrix spike duplicate QC data are advisory.

Method Blank: An analytical control consisting of all reagents, internal standards, and surrogate standards carried through the entire analytical process. Used to monitor laboratory background and reagent contamination.

Not Detected (ND): Not detected at or above the reporting limit.

Relative Percent Difference (RPD): An indication of method precision based on duplicate analyses.

Reporting Limit (RL): The lowest concentration routinely determined during laboratory operations. The RL is generally 1 to 10 times the Method Detection Limit (MDL). Reporting limits are matrix, method, and analyte dependent and take into account any dilutions performed as part of the analysis.

Surrogates. Organic compounds which are similar to analytes of interest in chemical behaviour, but are not found in environmental samples. Surrogates are added to all blanks, calibration and check standards, samples, and spiked samples. Surrogate recovery is monitored as an indication of acceptable sample preparation and instrument performance.

- D: Surrogates diluted out.
- I: Interference.
- 1. Indicates result outside of established laboratory QC limits.

WORK ORDER: 9804229

QUALITY CONTROL REPORT

PAGE QR-2

ANALYSIS: Lead

MATRIX: WET Extract

METHOD BLANK SAMPLES

SAMPLE TYPE: 8lank-Method/Med INSTRUMENT: Video 12 aa sped UNITS: mg/L METHOD:		LAB ID: WET_PBW PREPARED: ANALYZED: 05/18/98			INSTR RUN: AA V12\980518101500/1/ BATCH ID: WET051598 DILUTION:			
ANALYTE Lead in Water by EPA 7420	RESULT ND	REF RESULT	REPORTING LIMIT 0.1	SPIKE VALUE	RECOVERY (%)	REC LIMITS (% LOW HIGH) RPD (%)	RPD LIMIT (%)

OTHER QC SAMPLES

INSTRUMENT:	Spike-Post-diges: Video 12 aa speci mg/L			LAB ID: PD04229-13B PREPARED: ANALYZED: 05/18/98			INSTR RUN: AA_V12\980518101500/4/2 BATCH IO: WET051598 DILUTION: 1.000000			
ANALYTE	by EPA 7420	RESULT 17.3	REF RESULT 8.70	REPORTING LIMIT	SPIKE VALUE 8.00	RECOVERY (%) 108	REC LIMITS (%) LOW HIGH		RPD LIMIT (%)	
INSTRUMENT:	Spike-Post-digesi Video 12 aa speci mg/L	tion-Dup trometer		LAB ID: PREPARED: ANALYZED:		3B		V12\98051 T051598 000000	8101500/5/3	
ANALYTE	by EPA 7420	RESULT 17.3	REF RESULT 17.0	REPORTING LIMIT	SPIKE VALUE	RECOVERY (%)	REC LIMITS (%) LOW HIGH	RPO (%) 1.75	RPO LIMIT (%)	
INSTRUMENT:	Spike-Post-digesi Video 12 aa speci mg/L	tion trometer		LAB ID: PREPARED: ANALYZED:		3B		V12\98051 T051598 000000	8101500/3/2	
ANALYTE Lead in Water	by EPA 7420	RESULT 17.0	REF RESULT 8.70	REPORTING LIMIT	SPIKE VALUE 8.00	RECOVERY (%) 104	REC LIMITS (%) Low High	RPD (%)	RPD LIMIT (%)	

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QUALITY CONTROL REPORT

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ANALYSIS: Mercury

MATRIX: Soil/Bulk

METHOD BLANK SAMPLES

SAMPLE TYPE: Blank-Method/Media blank LAB ID: HGS BLNK INSTR RUN: HG\980503123000/1/
INSTRUMENT: Coleman Hg Analyzer 50D PREPARED: BATCH ID: HGS053598
UNITS: mg/kg ANALYZED: 05/03/98 DILUTION: 1.000000

METHOD:

SPIKE RECOVERY REC LIMITS (%) RPD VALUE (%) LOW HIGH RPD (%) LIMIT (%) REF REPORTING RESULT RESULT ANALYTE

LIMIT ANALYTE RESULT
Mercury in soil EPA 7471 ND 0.06

LABORATORY CONTROL SAMPLES

SAMPLE TYPE: Spike-Method/Media blank LAB ID: HGS_LCD INSTR RUN: HG\980503123000/3/1

BATCH ID: HGS053598 DILUTION: 1.000000 INSTRUMENT: Coleman Hg Analyzer 50D PREPARED:

ANALYZED: 05/03/98 UNITS: mq/kq

METHOD:

REF REPORTING SPIKE RECOVERY REC LIMITS (%) RPD

ANALYTE RESULT RESULT LIMIT VALUE (%) LOW HIGH RPD (%) LIMIT (%)

Mercury in soil EPA 7471 0.383 ND 0.06 0.400 95.8 75 125

_______ _____ _____

SAMPLE TYPE: Spike-Method/Media blank LAB ID: HGS_LCS PREPARED: INSTR RUN: HG\980503123000/2/1 BATCH ID: HGS053598 DILUTION: 1.000000 INSTRUMENT: Coleman Hg Analyzer 50D

ANALYZED: 05/03/98 UNITS:

METHOD:

RECOVERY REC LIMITS (%)

REF REPORTING SPIKE RE ANALYTE RESULT RESULT LIMIT VALUE Mercury in soil EPA 7471 0.383 ND 0.06 0.400 (%) LOW HIGH RPD (%) LIMIT (%) 95.8 75 125

LABORATORY CONTROL DUPLICATES

INSTR RUN: HG\980503123000/4/2 BATCH ID: HG\$053598 DILUTION: 1.000000 SAMPLE TYPE: Method Spike Sample Duplicate LAB ID: HGS_LCR
INSTRUMENT: Coleman Hg Analyzer 50D PREPARED:

INSTRUMENT: Coleman Hg Analyzer 50D

UNITS: mg/kg

ANALYZED: 05/03/98 METHOD:

RECOVERY REC LIMITS (%)
(%) LOW HIGH RPD (%) LIMIT (%)
25 REF REPORTING SPIKE RECOVERY
ANALYTE RESULT RESULT LIMIT VALUE (%)
Mercury in soil EPA 7471 0.383 0.383 0.06

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QUALITY CONTROL REPORT

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ANALYSIS: Metals Scan by ICP

MATRIX: Soil/Bulk

METHOD BLANK SAMPLES

AMPLE TYPE: Blank-Method/Medi NSTRUMENT: TJA Enviro 36 WITS: mg/kg NETHOD:	ia blank		LAB ID: PREPARED: ANALYZED:	SAND_PBS 05/01/98		INSTR RUN: ICP\980430115500/1/ BATCH ID: IFS043098-Q DILUTION: 1.000000
NALYTE ty ty ty ty ty ty ty ty ty t	RESULT ND ND ND ND ND ND ND ND ND ND ND ND ND	REF RESULT	REPORTING LIMIT 0.5 5.5 0.5 2.2 2.2 2.4 50 2.5	SPIKE VALUE	RECOVERY (%)	REC LIMITS (%) RPD LOW HIGH RPD (%) LIMIT (%)
SAMPLE TYPE: Blank-Method/Medi (NSTRUMENT: TJA Enviro 36 UNITS: mg/kg METHOD:	ia blank		LAB ID: PREPARED: ANALYZED:	IFS_PBS_Z 05/08/98		INSTR RUN: ICP\980508123100/1/ BATCH ID: IFS050698-Z DILUTION: 1.000000
ANALYTE Ag Silver As Arsenic Ba Barium Be Beryllium Cd Cadmium Co Cobalt Cr Chromium Cu Copper Mo Molybdenum Ni Nickel Pb Lead Be Selenium Thallium V Vanadium Zn Zinc	RESULT ND ND ND ND ND ND ND ND ND ND ND ND ND	REF RESULT	REPORTING LIMIT 0.5 10 5 0.5 0.5 2 2 2 2 2 4 20 50 50	SPIKE VALUE	RECOVERY	REC LIMITS (%) RPD LOW HIGH RPD (%) LIMIT (%)
SAMPLE TYPE: Blank-Method/Med INSTRUMENT: TJA Enviro 36 UNITS: mg/kg METHOD:	ia blank		LAB ID: PREPARED: ANALYZED:	IFS_PBS_Z 06/08/98	40 47 TO 10 10 10 10 10 10 10 10 10 10 10 10 10	INSTR RUN: ICP\980508123100/15/ BATCH ID: IFS050698-Z DILUTION: 1.000000
NALYTE b Antimony	RESULT ND	REF RESULT	REPORTING LIMIT 5	SPIKE VALUE	RECOVERY (%)	REC LIMITS (%) RPD LOW HIGH RPD (%) LIMIT (%)

LABORATORY CONTROL SAMPLES

SAMPLE INSTRU UNITS: METHOD	mg/kg		~	LAB ID: SAND_LCD INSTR RUN: ICP\98043011550 PREPARED: BATCH ID: IFS043098-Q ANALYZED: Q5/01/98 DILUTION: 1.000000						5500/3/1
			REF	REPORTING	SPIKE	RECOVERY	REC LIM		(113	RPD
ANALYT	Ę	RESULT	RESULT	LIMIT	VALUE	(%)	LOW	HIGH	RPD (%)	LIMIT (%)
Ag	Silver	9.76	ND	0.5	10.0	97.6	75	125		
Ba	Barium	209	ИĎ	5	200	105	75	125		
Вe	8ervllium	9.41	ND	0.5	10.0	94.1	75	125		
Cd	Cadmium	9.72	NĎ	0.5	10.0	97.2	75	125		
Co	Coba I t	103	ÖN	1	100	103	75	125		
Cr	Chromium	103	ND	2	100	103	75	125		

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QUALITY CONTROL REPORT

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ANALYSIS: Metals Scan by ICP

MATRIX: Soil/Bulk

LABORATORY CONTROL SAMPLES

AMPLE TYPE: Spike-Method/ NSTRUMENT: TJA Enviro 36 NITS: mg/kg	Media blank		LAB ID: PREPARED: ANALYZED:	SAND_LCD 05/01/98			CP\98043011: S043098-Q .000000	5500/3/1
NALYTE u Copper o Molybdenum i Nickel b Lead l Thallium Vanadium n Zinc	RESULT 99.5 94.9 102 101 398 102 98.9	REF RESULT ND ND ND ND ND ND ND	REPORTING LIMIT 2 2 2 2 4 50 2	SPIKE VALUE 100 100 100 100 400 100	RECOVERY (%) 99.5 94.9 102 101 99.5 102 98.9	REC LIMITS (%) LOW HIGH 75 125 75 125 75 125 75 125 75 125 75 125 75 125	RPD (%)	RPD LIMIT (%)
AMPLE TYPE: Spike-Method/ NSTRUMENT: TJA Enviro 36	Media blank		LAB ID: PREPARED:	SAND_LCS 05/01/98		INSTR RUN: IC BATCH ID: IF	CP\980430119 S043098-Q	5500/2/1
NALYTE g Silver a Barium e Beryllium d Cadmium o Cobalt r Chromium u Copper lo Molybdenum i Nickel b Lead l Thallium Vanadium in Zinc	RESULT 9.89 210 9.37 9.89 103 104 99.7 90.5 102 103 401 103 98.5	REF RESULT ND NO ND ND NO NO ND ND ND ND ND	REPORTING LIMIT 0.5 0.5 0.5 1 2 2 2 2 2 4 50 2	SPIKE VALUE 10.0 200 10.0 10.0 100 100 100 100 100 100 100	RECOVERY (%) 98.9 105 93.7 98.9 103 104 99.7 90.5 102 103 100 103 98.5	REC LIMITS (%) LOW HIGH 75 125 75 125 75 125 75 125 75 125 75 125 75 125 75 125 75 125 75 125 75 125 75 125 75 125 75 125 75 125 75 125	RPD (%)	RPD LIMIT (%)
AMPLE TYPE: Spike-Method/ NSTRUMENT: TJA Enviro 36 NITS: mg/kg	'Media blank i		LAB ID: PREPARED: ANALYZED:	IFS_LCD_Z 05/08/98		INSTR RUN: IC BATCH ID: IF DILUTION: 1.	CP\98050812: S050698-Z 000000	3100/4/2
NALYTE g Silver is Arsenic a Barium d Cadmium d Cobalt r Chromium d Copper lo Molybdenum li Nickel b Lead e Selenium Thallium Zinc	17.7 194 49.8 46.6	ND ND ND NO NO		SPIKE VALUE 5.00 10.0 1000 5.00 5.50 50.0 50.0 50.0	RECOVERY (%) 91.2 75.1 101 93.0 92.7 98.6 98.0 96.0 91.8 96.6 98.0 88.5 92.4 99.6 93.2	75 125 75 125 75 125	RPD (%)	RPD LIMIT (%)
AMPLE TYPE: Spike-Method/ NSTRUMENT: TJA Enviro 36 NITS: mg/kg ETHOD:			LAB ID: PREPARED: ANALYZED:	06/08/98		INSTR RUN: IC BATCH ID: IF DILUTION: 1.	S050698-Z .000000	
NALYTE Sb Antimony	RESULT 45.8	REF RESULT ND	REPORTING LIMIT 5	VALUE 50.0	RECOVERY (%) 91.6	REC LIMITS (%) LOW HIGH 75 125	RPD (%)	LIMIT (%)

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QUALITY CONTROL REPORT

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ANALYSIS: Metals Scan by ICP

MATRIX: Soil/Bulk

LABORATORY CONTROL SAMPLES

SAMPLE INSTRU UNITS: METHOD	mg/kg	dia blank		LAB ID: PREPARED ANALYZED	IFS_LCS_Z : : 05/08/98		INSTR BATCH DILUTI	ID: IF	P\98050812 5050698-Z 000000	3100/3/2
PIE TITOL	' •		REF	REPORTING	SPIKE	RECOVERY	REC LIM	ITS (%)		RPD
ANALYT	Έ	RESULT	RESULT	LIMIT	VALUE	(%)	LOW	HIGH	RPO (%)	LIMIT (%)
Ag	Silver	4.42	ND.	0.5	5.00	88.4	75	125	5 (10)	-41121 (74)
As	Arsenic	7.55	ND	10	10.0	75.5	75	125		
Ba	Barium	98.7	ИD	5	100	98.7	75	125		
Be	Beryllium	4.49	ND ND	0.5	5.00	89.8	75	125		
Cd	Cadmium	5.06	ND	0.5	5.50	92.0	75	125		
Co	Cobalt	47.7	ND	0.5	50.0	95.4	75	125		
Cr	Chromium	47.7	ND	1	50.0	95.4	75 75	125		
Cu	Copper	46.3	NO.	2	50.0	92.6	75	125		
Ma	Molybdenum	43.6	ND ND	2	50.0	87.2	75 75	125		
N1	Nickel	46.8	ND ND	2	50.0	93.6	75	125		
Pb	Lead	40.0	ND ND	2	50.0	93.0	75 75	125		
	Selenium	52.7	ND	4	55.0	95.8	75	125		
Se 11	Thallium	16.4	ND ND	20 50	20.0	82.0	75 75	125		
i i		187	NU		210	89.0		125		
v Zn	Vanadium Zinc	48.2	ND	2	50.0	96.4	75	125		
711	2 Inc	44.6	ND	5	50.0	89.2	75 	125 		
INSTRU		dia blank		LAB ID: PREPARED			BATCH	ID: IF	S050698-Z	3100/17/16
UNITS:				ANALYZED	: 06/08/98		DILUTI	ON: 1.	000000	
ANALY1		RESULT 44.1	REF RESULT NO	REPORTING LIMIT 5	SPIKE VALUE 50.0	RECOVERY (%) 88.2	REC LIM LOW 75	ITS (%) HIGH 125	RPD (%)	RPD LIMIT (%)

LABORATORY CONTROL DUPLICATES

AMPLE TYPE: Method Spike Sample Duplicate NSTRUMENT: TJA Enviro 36 NITS: mg/kg ETHOD:	LAB ID: SAND_LCR PREPARED: ANALYZED: 05/01/98	INSTR RUN: ICP\980430115500/4/2 BATCH ID: IFS043098-Q DILUTION: 1.000000
g Silver 9.76	REF REPORTING SPIKE ESULT LIMIT VALUE 9.89 0.5 210 5	RECOVERY REC LIMITS (%) RPD (%) LOW HIGH RPD (%) LIMIT (%) 1.32 25 0.477 25
de Beryllium 9.41 d Cadmium 9.72 do Cobalt 103	9.37 0.5 9.89 0.5 103 1	0.426 25 1.73 25 0 25
	104 2 99.7 2 90.5 2 102 2	0.966 25 0.201 25 4.75 25 0 25
b Lead 101 1 Thallium 398 7 Vanadium 102 In Zinc 98.9	103 4 401 50 103 2 98.5 5	0.477 25 0.426 25 1.73 25 0 25 0.966 25 0.201 25 4.75 25 0 25 1.96 25 0.751 25 0.976 25 0.976 25 0.405 25

SAMPLE TYPE: Method Spike Sample Duplicate INSTRUMENT: TJA Enviro 36 UNITS: mg/kg METHOD:				PREPARED: BATCH ID: IFS				P\980508123100/5/3 \$050698-Z 000000		
ANALYTE Ag As Ba Be B Cd	Silver Arsenic Barium Geryllium Cadmium Cobalt	RESULT 4.56 7.51 101 4.65 5.10 49.3	REF RESULT 4.42 7.55 98.7 4.49 5.06 47.7	REPORTING LIMIT 0.5 10 5 0.5 0.5	SPIKE VALUE	RECOVERY (%)	REC LIMITS (%) LOW HIGH RPD (%) 3.12 0.531 2.30 3.50 0.787 3.30	RPD LIMIT (%) 25 25 25 25 25 25		

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QUALITY CONTROL REPORT

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ANALYSIS: Metals Scan by ICP

MATRIX: Soil/Bulk

LABORATORY CONTROL DUPLICATES

INSTRU UNITS:		te	LAB ID: PREPARED: ANALYZED:					
METHO: ANALY:		OFCU T	REF	REPORTING	SPIKE	RECOVERY	REC LIMITS (%)	RPD
ANALT Cr Cu Mo Ni Pb Se T1 V Zn	Chromium Copper Molybdenum Nickel Lead Selenium Thallium Vanadium Zinc	RESULT 49.0 48.0 45.9 48.3 53.9 17.7 194 49.8 46.6	RESULT 47.7 46.3 43.6 46.8 52.7 16.4 187 48.2 44.6	LIMIT 2 2 2 2 4 20 50 2	VALUE	(%)	LOW HIGH RPD (%) 2.69 3.61 5.14 3.15 2.25 7.62 3.67 3.27 4.39	LIMIT (%) 25 25 25 25 25 25 25 25 25 25
SAMPLI INSTRI UNITS METHOI		Sample Duplica	te	LAB ID: PREPARED: ANALYZED:	IFS_LCR_Z 06/08/98		INSTR RUN: ICP\9805081; BATCH ID: IFS050698-Z DILUTION: 1.000000	
ANALY Sb:		RESULT 45.8	REF RESULT 44.1	REPORTING LIMIT 5	SPIKE VALUE	RECOVERY (%)	REC LIMITS (%) LOW HIGH RPD (%) 3.78	RPD LIMIT (%) 25

MATRIX SPIKE SAMPLES

SAMPLE TYPE: Spike-Sample INSTRUMENT: TJA Enviro 3			LAB ID: PREPARED:	MD04229-05	A	INSTR BATCH		P\98043011 \$043098-0	5500/7/5
JNITS: mg/kg HETHOD:				05/01/98		DILUTI		000000	
111051		REF	REPORTING	SPIKE	RECOVERY	REC LIM	ITS (%)		RPD
NALYTE	RESULT	RESULT	LIMIT	VALUE	(%)	LOW	HIGH	RPD (%)	LIMIT (%)
g Silver	8.51	ND	0.5	10.0	85.1	70	130	• •	, ,
a Barium	388	193	5	200	97.5	70	130		
e Beryllium	9,21	ND	0.5	10.0	92.1	70	130		
d Cadmilum	8.72	ND	0.5	10.0	87.2	70	130		
o Cobalt	95.6	6.52	ī	100	89.1	70	130		
r Chromium	119	31.9	Ž	100	87.1	70	130		
и Соррег	102	12.2	- 2	100	89.8	70	130		
o Molybdenum	84.4	ND	2	100	84.4	60	140		
i Nickel	107	18.8	Ž	100	88.2	60 70	130		
b Lead	92.2	6.98	4	100	85.2	70	130		
l Thallium	310	ND	50	400	77.5	70	130		
Vanad 1 um	123	34.9	Ž	100	88.1	70	130		
in Zinc	109	22.9	5	100	86.1	70	130		~
AMPLE TYPE: Spike-Sample	e/Matrix		LAB ID:	MS04229-05	 iA	INSTR	RUN: IC	P\98043011	5500/6/5
NSTRUMENT: TJA Enviro :	36		PREPARED:			BATCH	ID: IF	S043098-Q	-
NITS: mg/kg ETHOD:				05/01/98		DILUTI	ON: 1.	000000	
		REF	REPORTING	SPIKE	RECOVERY	REC LIM	ITS (%)		RPD

UNITS: mg/kg METHOD:				ANALYZED	: : 05/01/98		DILUTIO		3043098-Q 300000	
			REF	REPORTING	SPIKE	RECOVERY	REC LIM	ITS (%)		RPD
ANAL'	YTE	RESULT	RESULT	LIMIT	VALUE	(%)	LOW	HIĞH	RPD (%)	LIMIT (%)
Ag	Silver	9.34	ND	0.5	10.0	93.4	70	130		•
Ва	8arium	390	193	5	200	98.5	70	130		
8e	Beryllium -	9.96	ND	0.5	10.0	99.6	70	130		
Cd	Cadmium	9.74	ND	0.5	10.0	97.4	70	130		
Co	Cobalt	106	6.52	1	100	99.5	70	130		
Cr	Chromium	129	31.9	2	100	97.1	70	130		
Cu	Copper	107	12.2	2	100	94.8	70	130		
Ma	Molybdenum	92.7	ИD	2	100	92.7	60	140		
Ni	Nicke1	118	18.8	2	100	99.2	70	130		
Pb	Lead	103	6.98	4	100	96.0	70	130		
11	Thallium	346	ND	50	400	86.5	70	130		
٧	Vanad1um	132	34.9	2	100	97.1	70	130		

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QUALITY CONTROL REPORT

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ANALYSIS: Metals Scan by ICP

MATRIX: Soil/Bulk

MATRIX SPIKE SAMPLES

SAMPLE TYPE: INSTRUMENT:	Spike-Sample/Matr TJA Enviro 36	xıx		LAB ID: PREPARED:	MS04229-05	6A	INSTR F		P\98043011 S043098-0	5500/6/5	1
UNITS: METHOD:	mg/kg				05/01/98		DILUTIO		000000		Į
ANALYTE Zn	Zinc	RESULT 115	REF RESULT 22.9	REPORTING LIMIT 5	SPIKE VALUE 100	RECOVERY (%) 92.1	REC LIMI LOW 70	TS (%) HIGH 130	RPD (%)	RPD LIMIT (%)	Į

MATRIX SPIKE DUPLICATES

SAMPLI	E TYPE: Spiked Sample UMENT: TJA Enviro 36	Duplicate		LAB ID: PREPARED:	MR04229-0	5A	INSTR RUN: BATCH ID:	ICP\98043011 IFS043098-0	5500/8/6
UNITS	: mg/kg	1			05/01/98		DILUTION:	1.000000	
(ILTIO	•		REF	REPORTING	SPIKE	RECOVERY	REC LIMITS		RPD .
ANALY	TE	RESULT	RESULT	LIMIT	VALUE	(%)	LOW HIC		LIMIT (%)
Ag	Silver	8.51	9.34	0.5				9.30	25
Вā	8a <i>rium</i>	388	390	5				0.514	25
Ве	Beryllium	9.21	9.96	0.5				7.82	25
Cd	Cadmium	8,72	9.74	0.5				11.1	25 25
Co	Cobalt	95.6	106	1				10.3	25
Cr	Chromium	119	129	ž				8.06	25
Cu	Copper	102	107	ž				4.78	25
Мо	Molybdenum	84.4	92.7	2				9.37	25
Ni	Nickel	107	118	2				9.78	25
Pb	Lead	92.2	103	ā				11.1	25
ΤĪ	Thallium	310	346	50				11.0	25
v i	Vanadium	123	132	2				7.06	25
Žn	Zinc	109	115	5				5.36	25

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QUALITY CONTROL REPORT

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ANALYSIS: Volatile GC/MS

MATRIX: Soil/Bulk

METHOD BLANK SAMPLES

AMPLE TYPE: Blank-Method/Me NSTRUMENT: HP mass spec fo NITS: ug/kg ETHOD:		LAB ID: BLNK_0501 PREPARED: ANALYZED: 05/01/98				INSTR RUN: GCMS12\980501020000/1/ BATCH ID: MS12S050198 DILUTION: 1.00				
NALYTE	RESULT	REF RESULT	REPORTING LIMIT	SPIKE VALUE	RECOVERY (%)	REC LIM:	TS (%)	RPD (%)	RPD LIMIT (%)	
,2-DCA-d4 (surr)	128	KESULI	LIMII	100	128	60	140	KFD (A)	CINII (N	
oluene-d8 (surr)	112			100	112	65	135			
-BFB (surr)	93.1			100	93.1	65	135			
,1-Dichloroethene	מא		5							
enzene	ND		5							
richloroethene	ND		5							
o luene	ND		5							
hlorobenzene	ND		5							
romobenzene	ND		5							
romochloromethane romodichloromethane	ND ND		5							
	ND NO		10							
romoform romomethane	ON ON		5							
-Butylbenzene	ND ND		10 5							
ec-Buty Ibenzene	ON		5							
ert-Buty Ibenzene	ОN		5							
arbon Tetrachloride	ОЙ		5							
hloroethane	ЙĎ		10							
hloroform	ND		5							
hloromethane	ND		10							
-Chlorotoluene	ND		5							
-Ch loroto luene	ND		5 5							
Ibromochloromethane	ND		5							
,2-0ibromo-3-chloropropan	ND		5							
,2-Dibromoethane	ND		5							
ibromomethane	ND		5							
,2-Dichlorobenzene ,3-Dichlorobenzene	ND ON		5 5							
.4-Dichlorobenzene	NO NO		5 5							
ichlorodifluoromethane	ЙD		10							
.1-Dichloroethane	ND		5							
.2-Dichloroethane	ND		5							
is-1,2-Dichloroethene	ND		Š							
rans-1,2-Dichloroethene	ND		5							
.2-Dichloropropane	ND		5							
.3-Dichloropropane	ND		5							
2,2-Dichloropropane	ND		5							
.1-Dichloropropene	ND		5 5							
thy lbenzene	ND		5							
lexachlorobutadiene	DИ		5							
sopropy lbenzene	ND		5 5							
-Isopropyltoluene	ND		.5							
lethylene Chloride Laphthalene	ЖD		10							
iaphtha iene i-Propy lbenzene	ND ND		5 5							
i-rropy ibenzene Stvrene	ND ND		5 5							
.1,1,2-Tetrach loroethane	ND		5							
1,1,2,2-Tetrachloroethane	ND		5							
etrach loroethene	NĎ		5 5							
,2,3-Trichlorobenzene	ND		5							
1,2,4-Trichlorobenzene	ND		5							
.1.1-Trichloroethane	ND		5							
.1.2-Trichloroethane	ND		5							
richlorofluoromethane	ND		5							
.2.3-Trichloropropane	ND		5							
1,2,4-Trimethylbenzene	ND ND		5							
1,3,5-Trimethylbenzene	ND ND		.5							
/inyl Chloride	ND ND		10							
(ylenes, Total Trichlorotrifluoroethane	ND NO		10							
Acetone	ND ND		5							
	ND		100							
Acrylonitrile	ND		10							

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QUALITY CONTROL REPORT

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ANALYSIS: Volatile GC/MS

MATRIX: Soil/Bulk

METHOD BLANK SAMPLES

SAMPLE TYPE: Blank-Method/Media INSTRUMENT: HP mass spec for V UNITS: ug/kg			LAB ID: PREPARED	BLNK_0501 : : 05/01/98		INSTR RUN: GCMS12\S BATCH ID: MS12S050 DILUTION: 1.00	
METHOD:			MINETELD	. 00/01/00		5120110111 21.00	
		REF	REPORTING	SPIKE	RECOVERY	REC LIMITS (%)	RPD
ANALYTE	RESULT	RESULT	LIMIT	VALUE	(%)	LOW HIGH RPD	(%) LIMIT (%)
Carbon Disulfide	סא		10				
trans-1,4-Dichloro-2-buten	ND		100				
cis-1,3-Dichloropropene	ND		5				
trans-1,3-Dichloropropene	ND		5				
2-Hexanone	ND		50				
Methyl Iodide	ЙĎ		200				
4-Methy1-2-pentanone	ND						
Vinyl Acetate	ND		50 50				
2-Chloroethyl Vinyl Ether	ÑĎ		ĨÕ				
o-Xy lene	ND		10				
m.p-Xylene	ND		10				
MTBE	ND		5				

LABORATORY CONTROL SAMPLES

SAMPLE TYPE: Labora INSTRUMENT: HP mas UNITS: ug/kg METHOD:	tory Cont s spec for	rol Spike r Volatiles	~~~*	LAB ID: LCD_0503 PREPARED: ANALYZED: 05/03/98			INSTR RUN: GCMS12\980501020000/7/1 BATCH ID: MS12S050198 DILUTION: 50.0		
ANALYTE 1,2-DCA-d4 Toluene-d8 p-BFB 1,1-Dichloroethene Benzene Trichloroethene Toluene Chlorobenzene	(surr) (surr) (surr)	RESULT 102 104 105 200 2490 2740 2390 3210	REF RESULT 128 112 93.1 ND ND ND ND ND	REPORTING LIMIT	SPIKE VALUE 100 100 2500 2500 2500 2500 2500	RECOVERY (%) 102 104 105 8.00 99.60 109.6 95.60 128.4	REC LIMITS LOW HI 60 14 65 13 65 13 65 13 65 13 65 13 65 13 65 13	GH RPD (%) 5 5 5 5 5 5 5 5	RPD LIMIT (%)
SAMPLE TYPE: Labora	tory Cont			LAR ID.	LCS OFO1	~_	INSTO DIIN	GCMS12\Q804	501020000/2/3

SAMPLE TYPE: Labora INSTRUMENT: HP mas UNITS: ug/kg METHOD:	tory Cont s spec fo	rol Spike r Volatiles		LAB ID: PREPARED ANALYZED	LCS_0501 : : 05/01/98		INSTR BATCH DILUTI	ID: MS1	25050198	1020000/2/1
			REF	REPORTING	SPIKE	RECOVERY	REC LIM	ITS (%)		RPD
ANALYTE 1,2-DCA-d4 Toluene-d8 p-BFB	(surr) (surr) (surr)	RESULT 113 104 99.7	RESULT 128 112 93.1	LIMIT	VALUE 100 100 100	(%) 113 104 99.7	LOW 60 65 65	HIGH 140 135 135	RPD (%)	LIMÎT (%)
1,1-Dichloroethene	, ,	45.8	ND	5	50.0	91.6	65	135		
Benzene		58.9	ND	5	50.0	118	65	135		
Trichloroethene		58.0	ND	5	50.0	116	65	135		
Toluene Chlorobenzene		62.6 67.2	ND ND	5 5	50.0 50.0	125 134	65 65	135 135		

SAMPLE TYPE: Labora INSTRUMENT: HP mas: UNITS: ug/kg METHOD:		rol Spike r Volatiles		LAB ID: PREPARED ANALYZED	LCS_0503 : : 05/03/98		INSTR BATCH DILUTI	ID: MS1	28050198	01020000/6/	1
ANALYTE 1,2-DCA-d4 Toluene-d8 p-BFB 1,1-Dichloroethene Benzene Trichloroethene Toluene	(surr) (surr) (surr)	RESULT 114 100 105 2400 2410 2660 2660	REF RESULT 128 112 93.1 ND ND ND	REPORTING LIMIT	SPIKE VALUE 100 100 2500 2500 2500 2500	RECOVERY (%) 114 100 105 96.00 96.40 106.4 106.4	REC LIM LOW 60 65 65 65 65 65	ITS (%) HIGH 140 135 135 135 135 135	RPD (%)	RPD LIMIT (%)	

WORK ORDER: 9804229

QUALITY CONTROL REPORT

PAGE QR-11

ANALYSIS: Volatile GC/MS

MATRIX: Soil/Bulk

LABORATORY CONTROL SAMPLES

SAMPLE TYPE: Laboratory Conf INSTRUMENT: HP mass spec fo UNITS: ug/kg METHOD:	trol Spike or Volatiles		LAB ID: PREPARED: ANALYZED:	LCS_0503 05/03/98			D: MS1	2S050198	1020000/6/	1
ANALYTE Chlorobenzene	RESULT 3230	REF RESULT ND	REPORTING LIMIT	SPIKE VALUE 2500	RECOVERY (%) 129.2	REC LIMI LOW 65	TS (%) HIGH 135	RPD (%)	RPD LIMIT (%)	

LABORATORY CONTROL DUPLICATES

SAMPLE TYPE: Labora INSTRUMENT: HP mas UNITS: ug/kg METHOD:	tory Contr s spec for	rol Sample O r Volatiles	Ouplicate	LAB ID: LCR_0503 PREPARED: ANALYZED: 05/03/98			INSTR RUN: GCMS12\980501020000/8/6 BATCH ID: MS12S050198 DILUTION: 50.0			
ANALYTE 1,2-DCA-d4 Toluene-d8 p-BFB	(surr) (surr) (surr)	RESULT 102 104 105	REF RESULT 114 100 105	REPORTING LIMIT	SPIKE VALUE	RECOVERY (%) 11.1 3.92 0	REC LIM LOW 60 65 65	HIGH 140 135 135) RPD (%)	RPD LIMIT (%)
1,1-Dichloroethene Benzene Trichloroethene Toluene Chlorobenzene		2000 2490 2740 2390 3210	2400 2410 2660 2660 3230						18.18 3.265 2.963 10.69 0.6211	40 30 30 30 30

SAMPLE SURROGATES

SAMPLE TYPE: INSTRUMENT: UNITS: METHOD:	Sample-Client HP mass spec for ug/kg	Volatiles		LAB ID: PREPARED: ANALYZED:	9804229-05/ 05/03/98		BATCH ID: N	CMS12\980501020000/4/ S12S050198 .00
ANALYTE 1.2-DCA-d4 Toluene-d8 p-BFB	(surr) (surr) (surr)	RESULT 505 494 472	REF RESULT	REPORTING LIMIT	SPIKE VALUE 500 500 500	RECOVERY (%) 101 98.8 94.4	REC LIMITS (% LOW HIGH 60 140 65 135 65 135	
SAMPLE TYPE: INSTRUMENT: UNITS: METHOD:	Sample-Client HP mass spec for ug/kg	Volatiles		LAB ID: PREPARED: ANALYZED:			BATCH ID: M DILUTION: 5	CM\$12\980501020000/5/ \$12\$050198 00
ANALYTE 1,2-DCA-d4 Toluene-d8 p-BFB	(surr) (surr) (surr)	RESULT D D D	REF RESULT	REPORTING LIMIT	SPIKE VALUE 50000 50000	RECOVERY (%) 0 ! 0 ! 0 !	REC LIMITS (% LOW HIGH 60 140 65 135 65 135	
SAMPLE TYPE: INSTRUMENT: UNITS: METHOD:	Sample-Client HP mass spec for ug/kg	Volatiles		LAB ID: PREPARED: ANALYZED:	9804229-07/ 05/03/98	1	BATCH ID: M	CMS12\980501020000/9/ S12S050198 0.0
ANALYTE 1,2-DCA-d4 Toluene-d8 p-BFB	(surr) (surr) (surr)	RESULT 103 127 117	REF RESULT	REPORTING LIMIT	SPIKE VALUE 100 100 100	RECOVERY (%) 103 127 117	REC LIMITS (% LOW HIGH 60 140 65 135 65 135	

----- End of Quality Control Report -----

RYSD

9804229 4217 Chain-of-Custody Record Page of Date: Project No 4710 REMARKS ANALYSES CAM 17 Meta 8260 Additional comments Samplers (Signatures) EPA Method 8270 TPH as gasoline Do Not surmetals until you talk with lenviter patters on. Some samples may need to be composited. TPH as diesel Sample Number 1A 14/22 10925 5-07 14/22 0950 5-08 3A 4/22 1002 5-01 4/22 1025 5-02 (\$-09 5-10 4/22 1100 5-05 14/22 1125 5-04 4/22 1420 5-06 HOLD 4/22/1445 5-09 HOL 0A 4/27/1500 5-10 Turnaround time: Results to: Total No. of containers Jennifer Patterson STANDARN Date. Method of shipment

Lab Pickup

Laboratory comments and Log No:

4/23/92 Composite Sold Sold

Sold Sold Sold

Sold of Composite - For

metals analysis on S

Time. pri J. Patterson Date: Relinquished by: Date: Relinquished by: Relinquished by: Charlie Crocker 4-22-99 Signature: Printed name: RICK CILMORE 8:30 Printed name Printed name: Company

1 COMM (/) X

Received by Company: Received by: Time: Received by: Metals off hold Printed name:

Company:

7/2 Z Signature

7/830 Printed name. 4/ZZ Signature Geomatrix Consultants 100 Pine St 10th Floor San Francisco, CA 94111 (415) 434-9400

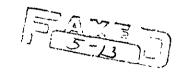
CHANGE ORDER REQUEST

AMERICAN ENVIRONMENTAL NETWORK (AEN)
3440 VINCENT ROAD
PLEASANT HILL, CA 94523
925
PHONE (510) 930-9090

DATE/TIME

AEN REP.

AEN PROJ NO.



COMPANY <u>Geomatrix</u>

CONTACT <u>Tom Graf</u>

PROJECT 4710 4217

925

FAX (510) 930-0256

ADDITIONAL ANALYSIS	CHANGE	D ANALYSIS	OTHER
Standa	nd T	4 T	
	- Pb		
11 A	5-07,08	Comp	
13A	S-09 10	Comp	

ACCEPTED - The above specifications of this Change Order are satisfactory and are hereby accepted

DATE OF ACCEPTANCE ______ SIGNATURE _____

American Environmental Network

Certificate of Analysis

DOHS Certification: 1172

AIHA Accreditation: 11134

PAGE 1

GEOMATRIX CONSULTANTS 100 PINE ST., SUITE 1000 SAN FRANCISCO, CA 94111

ATTN: JEN PATTERSON CLIENT PROJ. ID: 4710

C.O.C. NUMBER: 11029

REPORT DATE: 06/02/98

DATE(S) SAMPLED: 04/22/98

DATE RECEIVED: 04/22/98

AEN WORK ORDER: 9804232

PROJECT SUMMARY:

On April 22, 1998, this laboratory received 10 (4 soil and 6 water) sample(s).

Client requested 9 sample(s) be analyzed for chemical parameters; one sample was placed on hold. Results of analysis are summarized on the following page(s). Please see quality control report for a summary of QC data pertaining to this project.

Samples will be stored for 30 days after completion of analysis, then disposed of in accordance with State and Federal regulations. Samples may be archived by prior arrangement.

If you have any questions, please contact Client Services at (510) 930-9090.

Reviewed by:

William Int

SAMPLE ID: GGW1

AEN LAB NO: 9804232-01 AEN WORK ORDER: 9804232 CLIENT PROJ. ID: 4710

ANALYTE	METHOD/ CAS#	RESULT	REPORTING LIMIT UNITS	DATE ANALYZED
#Digestion, Metals by ICP	EPA 3010		Prep Date	04/28/98
EPA 8260A - Water Benzene Bromobenzene Bromochloromethane Bromodichloromethane Bromoform Bromomethane n-Butylbenzene sec-Butylbenzene tert-Butylbenzene Carbon Tetrachloride Chlorobenzene Chlorotoluene Chlorotoluene Chlorotoluene 2-Chlorotoluene 0ibromochloromethane 1.2-Dibromo-3-chloropropane 1.2-Dibromoethane Dibromomethane 1.2-Dichlorobenzene 1.3-Dichlorobenzene 1.4-Dichlorobenzene 1.1-Dichloroethane 1.1-Dichloroethane 1.1-Dichloroethane 1.2-Dichloroethene cis-1.2-Dichloroethene trans-1,2-Dichloroethene trans-1,2-Dichloroethene trans-1,2-Dichloropropane 1.1-Dichloropropane 1.3-Dichloropropane 1.3-Dichloropropane 1.1-Dichloropropane	EPA 8260A 71-43-2 108-86-1 74-97-5 75-27-4 75-25-2 74-83-9 104-51-8 135-98-8 98-06-6 56-23-5 108-90-7 75-00-3 67-66-3 74-87-3 95-49-8 106-43-4 124-48-1 96-12-8 106-46-7 75-71-8 75-71-8 75-71-8 75-71-8 75-35-4 156-60-5 78-87-5 142-28-9 594-20-7 563-58-6 100-41-4 87-68-3 99-87-6 75-09-2 91-20-3 103-65-1		50 ug/L 50 ug/L	05/06/98 05/06/98

SAMPLE ID: GGW1 AEN LAB NO: 9804232-01 AEN WORK ORDER: 9804232 CLIENT PROJ. ID: 4710

ANALYTE	METHOD/ CAS#	RESULT	REPORTING LIMIT	DATE UNITS ANALYZED
Styrene 1.1.2-Tetrachloroethane 1.1.2.2-Tetrachloroethane Tetrachloroethene Toluene 1.2.3-Trichlorobenzene 1.2.4-Trichlorobenzene 1.1.1-Trichloroethane 1.1.2-Trichloroethane Trichloroethene Trichlorofluoromethane 1.2.3-Trichloropropane 1.2.4-Trimethylbenzene 1.3.5-Trimethylbenzene Vinyl Chloride Xylenes, Total Acetone 2-Butanone Carbon Disulfide cis-1.3-Dichloropropene trans-1,3-Dichloropropene 2-Hexanone 4-Methyl-2-pentanone Vinyl Acetate 2-Chloroethyl Vinyl Ether	100-42-5 630-20-6 79-34-5 127-18-4 108-88-3 87-61-6 120-82-1 71-55-6 79-00-5 79-01-6 75-69-4 96-18-4 95-63-6 108-67-8 75-01-4 1330-20-7 67-64-1 78-93-3 75-15-0 10061-01-5 10061-02-6 591-78-6 108-05-4 110-75-8	ND ND ND ND ND ND ND ND ND ND ND ND ND N	50 ug/l 50 ug/l 50 ug/l 50 ug/l 50 ug/l 50 ug/l 50 ug/l 50 ug/l 50 ug/l 100 ug/l 100 ug/l 100 ug/l 50 ug/l 500 ug/l 500 ug/l 500 ug/l 500 ug/l 500 ug/l	L 05/06/98 05/06/98 05/06/98 05/06/98 05/06/98 05/06/98 05/06/98 05/06/98 05/06/98 05/06/98 05/06/98 05/06/98 05/06/98 05/06/98 05/06/98 05/06/98 05/06/98 05/06/98
CCR 17 Metals Silver Arsenic Barium Beryllium Cadmium Cobalt Chromium Copper Mercury Molybdenum Nickel Lead Antimony Selenium Thallium Yanadium Zinc	EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010	ND 0.034 * 0.23 * ND ND 0.005 * ND ND ND 0.03 * 0.01 * ND ND ND ND	0.01 mg/L 0.002 mg/L 0.005 mg/L 0.005 mg/L 0.01 mg/L 0.01 mg/L 0.002 mg/L 0.01 mg/L 0.01 mg/L 0.01 mg/L 0.05 mg/L 0.05 mg/L 0.05 mg/L	05/05/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98

GEOMATRIX CONSULTANTS

SAMPLE ID: GGW1

AEN LAB NO: 9804232-01 AEN WORK ORDER: 9804232 CLIENT PROJ. ID: 4710

DATE SAMPLED: 04/22/98

DATE RECEIVED: 04/22/98 REPORT DATE: 06/02/98

METHOD/ REPORTING DATE ANALYTE CAS# RESULT LIMIT UNITS ANALYZE						
· ·	ANALYTE	METHOD/ CAS#	RESULT		UNITS	DATE ANALYZED

ND = Not detected at or above the reporting limit * = Value at or above reporting limit

GEOMATRIX CONSULTANTS

SAMPLE ID: GGW1-1.0 AEN LAB NO: 9804232-02 AEN WORK ORDER: 9804232 CLIENT PROJ. ID: 4710

DATE SAMPLED: 04/22/98 DATE RECEIVED: 04/22/98 **REPORT DATE: 06/02/98**

ANALYTE	METHOD/ CAS#	RESULT	REPORTING LIMIT UNITS	DATE ANALYZED
#Digestion for ICP/AA	EPA 3050A	-	Prep Date	04/30/98
Arsenic	EPA 6010	ND	10 mg/kg	05/08/98
Selenium	EPA 6010	ND	20 mg/kg	05/08/98
CCR 17 Metals Silver Barium Beryllium Cadmium Cobalt Chromium Copper Mercury Molybdenum Nickel Lead Antimony Thallium Vanadium Zinc	EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 7471 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010	ND 180 * ND ND 8 * 31 * 12 * ND ND 21 * ND ND ND 31 *	0.5 mg/kg 0.5 mg/kg 1 mg/kg 2 mg/kg 2 mg/kg 0.06 mg/kg 2 mg/kg 4 mg/kg 5 mg/kg 50 mg/kg	05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98

ND = Not detected at or above the reporting limit
 * = Value at or above reporting limit

SAMPLE ID: GGW2

AEN LAB NO: 9804232-03 AEN WORK ORDER: 9804232 CLIENT PROJ. ID: 4710 DATE SAMPLED: 04/22/98

DATE RECEIVED: 04/22/98 REPORT DATE: 06/02/98

ANALYTE	METHOD/ CAS#	RESULT	REPORTING LIMIT	GUNITS	DATE ANALYZED
#Sample Filtration	GFF 0.7 um	-		Filtr Date	04/24/98
TPH as Gas	5030/GC-FID	ND	0.05	mg/L	04/30/98
#Silica gel Cleanup	EPA 3630M	-		C1eanup	05/01/98
#Extraction for TPH	EPA 3510	-		Extrn Date	05/01/98
TPH as Diesel	GC-FID	ND	0.05	mg/L	05/02/98
#Digestion, Metals by ICP	EPA 3010	-		Prep Date	04/28/98
EPA 8260A - Water Benzene Bromobenzene Bromochloromethane Bromodichloromethane Bromomethane Bromomethane n-Butylbenzene sec-Butylbenzene tert-Butylbenzene Carbon Tetrachloride Chlorobenzene Chlorotoluene Chlorotoluene 4-Chlorotoluene 0ibromochloromethane 1.2-Dibromo-3-chloropropane 1.2-Dibromoethane 1.2-Dichlorobenzene 1.3-Dichlorobenzene 1.4-Dichlorobenzene 1.4-Dichloroethane 1.1-Dichloroethane 1.1-Dichloroethane 1.1-Dichloroethane 1.2-Dichloroethene 1.1-Dichloroethene 1.1-Dichloroethene 1.1-Dichloroethene 1.1-Dichloroethene 1.1-Dichloroethene	EPA 8260A 71-43-2 108-86-1 74-97-5 75-27-4 75-25-2 74-83-9 104-51-8 135-98-8 98-06-6 56-23-5 108-90-7 75-00-3 67-66-3 74-87-3 95-49-8 106-43-4 124-48-1 96-12-8 106-93-4 74-95-3 95-50-1 541-73-1 106-46-7 75-71-8 75-34-3 107-06-2 75-35-4 156-59-2 156-60-5 78-87-5	ND ND ND ND ND ND ND ND ND ND ND ND ND N	55550555555555555555555555555555555555	ug/L ug/L ug/L ug/L ug/L ug/L ug/L ug/L	05/01/98 05/01/98

SAMPLE ID: GGW2 AEN LAB NO: 9804232-03 AEN WORK ORDER: 9804232 CLIENT PROJ. ID: 4710

ANALYTE	METHOD/ CAS#	RESULT	REPORTING LIMIT UNITS	DATE ANALYZED
1.3-Dichloropropane 2.2-Dichloropropane 1.1-Dichloropropene Ethylbenzene Hexachlorobutadiene Isopropylbenzene p-Isopropyltoluene Methylene Chloride Maphthalene n-Propylbenzene Styrene 1.1.2-Tetrachloroethane 1.2.2-Tetrachloroethane Tetrachloroethene Toluene 1.2.3-Trichlorobenzene 1.1.1-Trichloroethane 1.1.2-Trichloroethane Trichlorofluoromethane Trichlorofluoromethane 1.2.3-Trichloropropane 1.2.4-Trimethylbenzene 1.3.5-Trimethylbenzene 1.3.5-Trimethylbenzene 1.3.5-Trimethylbenzene 2.9utanone Carbon Disulfide cis-1.3-Dichloropropene trans-1.3-Dichloropropene 2-Hexanone 4-Methyl-2-pentanone inyl Acetate 2-Chloroethyl Vinyl Ether	142-28-9 594-20-7 563-58-6 100-41-4 87-68-3 98-82-8 99-87-6 75-09-2 91-20-3 103-65-1 100-42-5 630-20-6 79-34-5 127-18-4 108-88-3 87-61-6 120-82-1 71-55-6 79-00-5 79-01-6 75-69-4 96-18-4 95-63-6 108-67-8 75-01-4 1330-20-7 67-64-1 78-93-3 75-15-0 10061-01-5 10061-02-6 591-78-6 108-05-4 110-75-8	ND ND DD ND	5 ug/L 10 ug/L 10 ug/L 10 ug/L 10 ug/L 10 ug/L 10 ug/L 10 ug/L 10 ug/L 10 ug/L 10 ug/L 10 ug/L 10 ug/L 10 ug/L 10 ug/L 10 ug/L 10 ug/L	05/01/98 05/01/98
CCR 17 Metals Silver Arsenic Barıum Eeryllium Gadmium Cobalt Chromium	EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010	ND ND 0.15 * ND ND 0.006 *	0.002 mg/L 0.005 mg/L	05/01/98 05/05/98 05/01/98 05/01/98 05/01/98 05/01/98

GEOMATRIX CONSULTANTS

SAMPLE ID: GGW2 AEN LAB NO: 9804232-03 AEN WORK ORDER: 9804232 CLIENT PROJ. ID: 4710 DATE SAMPLED: 04/22/98

DATE RECEIVED: 04/22/98 REPORT DATE: 06/02/98

ANALYTE	METHOD/ CAS#	RESULT	REPORTING LIMIT UNIT	DATE S ANALYZED
Copper Mercury Molybdenum Nickel Lead Antimony Selenium Thallium Vanadium Zinc	EPA 6010 EPA 7470 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010	ND ND 0.05 * 0.03 * ND ND ND ND ND ND	0.01 mg/L 0.0002 mg/L 0.01 mg/L 0.01 mg/L 0.04 mg/L 0.02 mg/L 0.01 mg/L 0.05 mg/L 0.005 mg/L 0.001 mg/L	05/01/98 04/26/98 05/01/98 05/01/98 05/01/98 05/05/98 05/01/98 05/01/98 05/01/98

ND = Not detected at or above the reporting limit * = Value at or above reporting limit

GEOMATRIX CONSULTANTS

SAMPLE ID: GGW2-2.0 AEN LAB NO: 9804232-04 AEN WORK ORDER: 9804232 CLIENT PROJ. ID: 4710 DATE SAMPLED: 04/22/98 DATE RECEIVED: 04/22/98 REPORT DATE: 06/02/98

ANALYTE	METHOD/ CAS#	RESULT	REPORTING LIMIT UNITS	DATE ANALYZED
#Digestion for ICP/AA	EPA 3050A	-	Prep Date	04/30/98
Arsenic	EPA 6010	10 *	10 mg/kg	05/08/98
Selenium	EPA 6010	ND	20 mg/kg	05/08/98
CCR 17 Metals Silver Barium Beryllium Cadmium Cobalt Chromium Copper Mercury Molybdenum Nickel Lead Antimony Thallium Vanadium Zinc	EPA 6010 EPA 6010	ND 150 * ND ND 6 * 47 * 10 * ND ND 25 * ND ND ND ND 36 *	0.5 mg/kg 5 mg/kg 0.5 mg/kg 0.5 mg/kg 1 mg/kg 2 mg/kg 2 mg/kg 0.06 mg/kg 2 mg/kg 4 mg/kg 5 mg/kg 5 mg/kg 5 mg/kg	05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98

ND = Not detected at or above the reporting limit
* = Value at or above reporting limit

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GEOMATRIX CONSULTANTS

SAMPLE ID: GGW3

AEN LAB NO: 9804232-05 AEN WORK ORDER: 9804232 CLIENT PROJ. ID: 4710 DATE SAMPLED: 04/22/98

DATE RECEIVED: 04/22/98 REPORT DATE: 06/02/98

ANALYTE	METHOD/ CAS#	RESULT	REPORTING LIMIT	UNITS	DATE ANALYZED
#Sample Filtration	0.45 um	-		Filtr Date	04/24/98
TPH as Gas	5030/GC-FID	0.22 *	0.05	mg/L	04/30/98
#Digestion, Metals by ICP	EPA 3010	-		Prep Date	04/28/98
EPA 8260A - Water Benzene Bromobenzene Bromochloromethane Bromodichloromethane Bromoform Bromomethane n-Butylbenzene sec-Butylbenzene tert-Butylbenzene Carbon Tetrachloride Chlorobenzene Chlorotoluene Chloroform Chloromethane 2-Chlorotoluene 4-Chlorotoluene Dibromochloromethane 1.2-Dibromo-3-chloropropane 1.2-Dibromoethane 1.2-Dichlorobenzene 1.3-Dichlorobenzene 1.4-Dichlorobenzene 1.4-Dichloroethane 1.1-Dichloroethane 1.1-Dichloroethene cis-1,2-Dichloroethene trans-1,2-Dichloroethene trans-1,2-Dichloropropane 1.3-Dichloropropane 1.3-Dichloropropane 1.1-Dichloropropane 1.1-Dichloropropane 1.1-Dichloropropane 1.1-Dichloropropane 1.1-Dichloropropane 1.1-Dichloropropane 1.1-Dichloropropane 1.1-Dichloropropane	EPA 8260A 71-43-2 108-86-1 74-97-5 75-27-4 75-25-2 74-83-9 104-51-8 135-98-8 98-06-6 56-23-5 108-90-7 75-00-3 67-66-3 74-87-3 95-49-8 106-43-4 124-48-1 96-12-8 106-93-4 74-95-3 95-50-1 541-73-1 106-46-7 75-71-8 75-34-3 107-06-2 75-35-4 156-59-2 156-60-5 78-87-5 142-28-9 594-20-7 563-58-6 100-41-4 87-68-3 98-82-8	X X X X X X X X X X X X X X X X X X X	10555550505555555555555555555555555555	ug/L ug/L ug/L ug/L ug/L ug/L ug/L ug/L	05/01/98 05/01/98

SAMPLE ID: GGW3

AEN LAB NO: 9804232-05 AEN WORK ORDER: 9804232 CLIENT PROJ. ID: 4710 DATE SAMPLED: 04/22/98 DATE RECEIVED: 04/22/98 REPORT DATE: 06/02/98

ANALYTE	METHOD/ CAS#	RESULT	REPORTING LIMIT UN	DATE ITS ANALYZED
p-Isopropyltoluene Methylene Chloride Naphthalene n-Propylbenzene Styrene l.1.2-Tetrachloroethane I.1.2.Tetrachloroethane Tetrachloroethene Toluene l.2.3-Trichlorobenzene l.1.1-Trichlorobenzene l.1.2-Trichloroethane Trichloroethene Trichloroethene Trichloroethene Trichlorofluoromethane l.2.3-Trichloropropane l.2.4-Trimethylbenzene l.3.5-Trimethylbenzene l.3.5-Trimethylbenzene Vinyl Chloride Xylenes. Total Acetone 2-Butanone Carbon Disulfide cis-1.3-Dichloropropene trans-1,3-Dichloropropene 2-Hexanone 4-Methyl-2-pentanone Vinyl Acetate 2-Chloroethyl Vinyl Ether	99-87-6 75-09-2 91-20-3 103-65-1 100-42-5 630-20-6 79-34-5 127-18-4 108-88-3 87-61-6 120-82-1 71-55-6 79-00-5 79-01-6 75-69-4 96-18-4 95-63-6 108-67-8 75-01-4 1330-20-7 67-64-1 78-93-3 75-15-0 10061-01-5 10061-02-6 591-78-6 108-05-4 110-75-8	ND ND ND ND ND ND ND ND ND ND ND ND ND N	5 ug/L 10 ug/L	05/01/98 05/01/98
CCR 17 Metals Silver Arsenic Barium Beryllium Cadmium Cobalt Chromium Copper Mercury Molybdenum Nickel Lead Antimony	EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 7470 EPA 6010 EPA 6010 EPA 6010	ND ND 0.52 * ND ND 0.006 * ND ND ND ND ND	0.002 mg/L 0.005 mg/L	05/01/98 05/05/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 04/26/98 05/01/98 05/01/98 05/01/98

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GEOMATRIX CONSULTANTS

SAMPLE ID: GGW3

AEN LAB NO: 9804232-05 AEN WORK ORDER: 9804232 CLIENT PROJ. ID: 4710

DATE SAMPLED: 04/22/98 DATE RECEIVED: 04/22/98 REPORT DATE: 06/02/98

ANALYTE	METHOD/ CAS#	RESULT	REPORTING RESULT LIMIT UNITS				
Selenium	EPA 6010	ND	0.01 m	g/L	05/05/98		
Thallium	EPA 6010	ND	0.05 m		05/01/98		
Vanadium	EPA 6010	ND	0.005 m	g/L	05/01/98		
Zinc	EPA 6010	0.10 *	0.01 m	g/L	05/01/98		

ND = Not detected at or above the reporting limit * = Value at or above reporting limit

SAMPLE ID: GGW3-1.5 AEN LAB NO: 9804232-06 AEN WORK ORDER: 9804232 CLIENT PROJ. ID: 4710

DATE SAMPLED: 04/22/98 DATE RECEIVED: 04/22/98 REPORT DATE: 06/02/98

ANALYTE	METHOD/ CAS#	RESULT	REPORTING LIMIT UNITS	DATE ANALYZED
#Digestion for ICP/AA	EPA 3050A	-	Prep Date	04/30/98
Arsenic	EPA 6010	12 *	10 mg/kg	05/08/98
Selenium	EPA 6010	ND	20 mg/kg	05/08/98
CCR 17 Metals Silver Barium Beryllium Cadmium Cobalt Chromium Copper Mercury Molybdenum Nickel Lead Antimony Thallium Vanadium Zinc	EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 7471 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010	ND 1,600 * ND 3.0 * 11 * 88 * 32 * 0.32 * ND 29 * 390 * ND 40 * 900 *	0.5 mg/kg 5 mg/kg 0.5 mg/kg 0.5 mg/kg 1 mg/kg 2 mg/kg 2 mg/kg 0.06 mg/kg 4 mg/kg 5 mg/kg 50 mg/kg 5 mg/kg 5 mg/kg	05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98

ND = Not detected at or above the reporting limit
* = Value at or above reporting limit

SAMPLE ID: GGW4

AEN LAB NO: 9804232-07 AEN WORK ORDER: 9804232 CLIENT PROJ. ID: 4710

DATE SAMPLED: 04/22/98 DATE RECEIVED: 04/22/98 REPORT DATE: 06/02/98

ANALYTE	METHOD/ CAS#	RESULT	REPORTING LIMIT UNITS	DATE ANALYZED
#Digestion, Metals by ICP	EPA 3010	-	Prep Date	04/28/98
EPA 8260A - Water Benzene Bromobenzene Bromochloromethane Bromodichloromethane Bromoform Bromomethane n-Butylbenzene sec-Butylbenzene carbon Tetrachloride Chlorobenzene Chlorotoluene Chlorotoluene Chlorotoluene 0 ibromochloromethane 1.2-Dibromo-3-chloropropane 1.2-Dibromoethane 0 ibromomethane 1.2-Dichlorobenzene 1.3-Dichlorobenzene 1.4-Dichlorobenzene 0 ichlorodifluoromethane 1.1-Dichloroethane 1.1-Dichloroethane 1.1-Dichloroethane 1.1-Dichloroethene cis-1.2-Dichloroethene trans-1.2-Dichloroethene 1.3-Dichloropropane 1.3-Dichloropropane 1.3-Dichloropropane 1.3-Dichloropropane 1.1-Dichloropropane	EPA 8260A 71-43-2 108-86-1 74-97-5 75-27-4 75-25-2 74-83-9 104-51-8 135-98-8 98-06-6 56-23-5 108-90-7 75-00-3 67-66-3 74-87-3 95-49-8 106-43-4 124-48-1 96-12-8 106-93-4 74-95-3 95-50-1 541-73-1 106-46-7 75-71-8 75-34-3 107-06-2 75-35-4 156-59-2 156-60-5 78-87-5 142-28-9 594-20-7 563-58-6 100-41-4 87-68-3 98-82-8 99-87-6 75-09-2 91-20-3 103-65-1		500 ug/L 500 ug/L	05/06/98 05/06/98

SAMPLE ID: GGW4

AEN LAB NO: 9804232-07 AEN WORK ORDER: 9804232 CLIENT PROJ. ID: 4710 DATE SAMPLED: 04/22/98 DATE RECEIVED: 04/22/98 REPORT DATE: 06/02/98

ANALYTE	METHOD/ CAS#	RESULT	REPORTING LIMIT UNITS	DATE ANALYZED
Styrene 1.1.2.Tetrachloroethane 1.1.2.2-Tetrachloroethane Tetrachloroethene Toluene 1.2.3-Trichlorobenzene 1.2.4-Trichlorobenzene 1.1.1-Trichloroethane 1.1.2-Trichloroethane Trichloroethene Trichlorofluoromethane 1.2.3-Trichloropropane 1.2.4-Trimethylbenzene 1.3.5-Trimethylbenzene Vinyl Chloride Xylenes, Total Acetone 2-Butanone Carbon Disulfide cis-1.3-Dichloropropene trans-1.3-Dichloropropene 2-Hexanone 4-Methyl-2-pentanone Vinyl Acetate 2-Chloroethyl Vinyl Ether	100-42-5 630-20-6 79-34-5 127-18-4 108-88-3 87-61-6 120-82-1 71-55-6 79-00-5 79-01-6 75-69-4 96-18-4 95-63-6 108-67-8 75-01-4 1330-20-7 67-64-1 78-93-3 75-15-0 10061-01-5 10061-02-6 591-78-6 108-10-1 108-05-4 110-75-8		500 ug/L 500 ug/L 500 ug/L 500 ug/L 500 ug/L 500 ug/L 500 ug/L 500 ug/L 500 ug/L 500 ug/L 500 ug/L 500 ug/L 1000 ug/L	05/06/98 05/06/98 05/06/98 05/06/98 05/06/98 05/06/98 05/06/98 05/06/98 05/06/98 05/06/98 05/06/98 05/06/98 05/06/98 05/06/98 05/06/98 05/06/98 05/06/98 05/06/98
CCR 17 Metals Silver Arsenic Barıum Beryllium Cadmıum Cobalt Chromıum Copper Mercury Molybdenum Nickel Lead Antımony Selenium Thallium Vanadıum Zinc	EPA 6010 EPA 6010	ND 0.008 * 0.04 * ND ND ND ND ND ND ND ND ND ND ND ND ND	0.01 mg/L 0.002 mg/L 0.005 mg/L 0.005 mg/L 0.01 mg/L 0.01 mg/L 0.002 mg/L 0.01 mg/L 0.01 mg/L 0.01 mg/L 0.04 mg/L 0.02 mg/L 0.05 mg/L 0.05 mg/L	05/01/98 05/05/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98

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GEOMATRIX CONSULTANTS

SAMPLE ID: GGW4

AEN LAB NO: 9804232-07 AEN WORK ORDER: 9804232 CLIENT PROJ. ID: 4710 **DATE SAMPLED:** 04/22/98

DATE RECEIVED: 04/22/98

REPORT DATE: 06/02/98

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ANALYTE	METHOD/ CAS#	RESULT	REPORTING LIMIT	UNITS	DATE ANALYZED

ND = Not detected at or above the reporting limit
* = Value at or above reporting limit

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GEOMATRIX CONSULTANTS

SAMPLE ID: GGW4-1.5 AEN LAB NO: 9804232-08 AEN WORK ORDER: 9804232 CLIENT PROJ. ID: 4710

DATE SAMPLED: 04/22/98 DATE RECEIVED: 04/22/98 **REPORT DATE: 06/02/98**

ANALYTE	METHOD/ CAS#	RESULT	REPORTING LIMIT UNITS	DATE ANALYZED
#Digestion for ICP/AA	EPA 3050A	-	Prep Date	04/30/98
Arsenic	EPA 6010	ИD	10 mg/kg	05/08/98
Selenium	EPA 6010	ND	20 mg/kg	05/08/98
CCR 17 Metals Silver Barium Beryllium Cadmium Cobalt Chromium Copper Mercury Molybdenum Nickel Lead Antimony Thallium Vanadium Zinc	EPA 6010 EPA 6010	ND 260 * 0.5 * 0.6 * 18 * 41 * 24 * 0.07 * ND * ND * ND * ND 43 *	0.5 mg/kg 5 mg/kg 0.5 mg/kg 0.5 mg/kg 1 mg/kg 2 mg/kg 2 mg/kg 0.06 mg/kg 4 mg/kg 5 mg/kg 50 mg/kg 50 mg/kg	05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98

ND = Not detected at or above the reporting limit
* = Value at or above reporting limit

SAMPLE ID: GGW13 duplicate 66 w3

AEN LAB NO: 9804232-09 AEN WORK ORDER: 9804232 CLIENT PROJ. ID: 4710 DATE SAMPLED: 04/22/98 DATE RECEIVED: 04/22/98 REPORT DATE: 06/02/98

ANALYTE	METHOD/ CAS#	RESULT	REPORTING LIMIT	UNITS	DATE ANALYZED
#Sample Filtration	GFF 0.7 um	-		Filtr Date	04/24/98
TPH as Gas	5030/GC-FID	0.24 *	0.05	mg/L	04/30/98
#Silica gel Cleanup	EPA 3630M	-		Cleanup	05/01/98
#Extraction for TPH	EPA 3510	-		Extrn Date	05/01/98
TPH as Di e sel	GC-FID	ND	0.05	mg/L	05/02/98
#Digestion, Metals by ICP	EPA 3010	-		Prep Date	04/28/98
Benzene Bromobenzene Bromochloromethane Bromodichloromethane Bromoform Bromomethane n-Butylbenzene sec-Butylbenzene tert-Butylbenzene Carbon Tetrachloride Chlorobenzene Chlorotehane Chlorotoluene Chlorotoluene Dibromochloromethane 1.2-Dibromo-3-chloropropane 1.2-Dibromoethane 0ibromomethane 1.2-Dichlorobenzene 1.3-Dichlorobenzene 1.4-Dichlorobenzene 1.4-Dichloroethane 1.1-Dichloroethane 1.1-Dichloroethane 1.2-Dichloroethane 1.1-Dichloroethane 1.1-Dichloroethane 1.1-Dichloroethane 1.1-Dichloroethane 1.1-Dichloroethene trans-1,2-Dichloroethene trans-1,2-Dichloroethene	EPA 8260A 71-43-2 108-86-1 74-97-5 75-27-4 75-25-2 74-83-9 104-51-8 135-98-8 98-06-6 56-23-5 108-90-7 75-00-3 67-66-3 74-87-3 95-49-8 106-43-4 124-48-1 96-12-8 106-93-4 74-95-3 95-50-1 541-73-1 106-46-7 75-71-8 75-34-3 107-06-2 75-35-4 156-59-2 156-60-5 78-87-5	ND ND DD DD DD DD ND ND ND ND ND ND ND N	55055555555555555555555555555555555555	ug/L ug/L ug/L ug/L ug/L ug/L ug/L ug/L	05/02/98 05/02/98

SAMPLE ID: GGW13 duplinate 664-5 AEN LAB NO: 9804232-09 AEN WORK ORDER: 9804232 CLIENT PROJ. ID: 4710

DATE SAMPLED: 04/22/98 DATE RECEIVED: 04/22/98 REPORT DATE: 06/02/98

ANALYTE	METHOD/ CAS#	RESULT	REPORTING LIMIT UNITS	DATE ANALYZED
1.3-Dichloropropane 2.2-Dichloropropane 1.1-Dichloropropene Ethylbenzene Hexachlorobutadiene Isopropylbenzene p-Isopropyltoluene Methylene Chloride Naphthalene n-Propylbenzene Styrene 1.1.2-Tetrachloroethane 1.1.2-Tetrachloroethane Tetrachloroethene Toluene 1.2.3-Trichlorobenzene 1.1.1-Trichloroethane 1.1.2-Trichloroethane Trichlorofluoromethane Trichlorofluoromethane 1.2.3-Trichloropropane 1.2.4-Trimethylbenzene 1.3.5-Trimethylbenzene 1.3.5-Trimethylbenzene 2.4-Trimethylbenzene 1.3.5-Trimethylbenzene 1.3.5-Trimethylbenzene 1.3.5-Trimethylbenzene 2.4-Trichloropropane 1.2.4-Trichloropropane 1.2.4-Trimethylbenzene 2.4-Trimethylbenzene 2.5-Trimethylbenzene 2.6-Trimethylbenzene 2.6-Trimethylbenzene 2.7-Trichloropropene 2.8-Trimethylbenzene 3.5-Trimethylbenzene	142-28-9 594-20-7 563-58-6 100-41-4 87-68-3 98-82-8 99-87-6 75-09-2 91-20-3 103-65-1 100-42-5 630-20-6 79-34-5 127-18-4 108-88-3 87-61-6 120-82-1 71-55-6 79-00-5 79-01-6 75-69-4 96-18-4 95-63-6 108-67-8 75-01-4 1330-20-7 67-64-1 78-93-3 75-15-0 10061-01-5 10061-02-6 591-78-6 108-05-4 110-75-8	ND ND DD DD DD DD DD ND ND ND ND ND ND N	55555555555555555555555555555555555555	05/02/98 05/02/98
CCR 17 Metals Silver Arsenic Barium Beryllium Cadmium Cobalt Chromium	EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010	ND ND 0.41 * ND ND 0.006 *	0.005 mg/L 0.005 mg/L 0.01 mg/L 0.002 mg/L 0.005 mg/L 0.005 mg/L 0.01 mg/L	05/01/98 05/05/98 05/01/98 05/01/98 05/01/98 05/01/98 05/01/98

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GEOMATRIX CONSULTANTS

SAMPLE ID: GGW13 duplicate 6643

AEN LAB NO: 9804232-09 AEN WORK ORDER: 9804232 CLIENT PROJ. ID: 4710

DATE SAMPLED: 04/22/98 DATE RECEIVED: 04/22/98 **REPORT DATE: 06/02/98**

ANALYTE	METHOD/ CAS#	RESULT	REPORTING LIMIT UNITS	DATE S ANALYZED
Copner Mercury Molybdenum Nickel Lead Antimony Selenium Thallium Vanadium Zinc	EPA 6010 EPA 7470 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010	ND ND ND ND ND ND ND ND ND	0.01 mg/L 0.0002 mg/L 0.01 mg/L 0.01 mg/L 0.04 mg/L 0.02 mg/L 0.01 mg/L 0.05 mg/L 0.05 mg/L 0.01 mg/L	05/01/98 04/26/98 05/01/98 05/01/98 05/01/98 05/01/98 05/05/98 05/01/98 05/01/98

ND = Not detected at or above the reporting limit
 * = Value at or above reporting limit

AEN (CALIFORNIA) QUALITY CONTROL REPORT

AEN JOB NUMBER: 9804232 CLIENT PROJECT ID: 4710

Quality Control and Project Summary

All laboratory quality control parameters were found to be within established limits.

<u>Definitions</u>

Laboratory Control Sample (LCS)/Method Spikes(s): Control samples of known composition. LCS and Method Spike data are used to validate batch analytical results.

Matrix Spike(s): Aliquot of a sample (aqueous or solid) with added quantities of specific compounds and subjected to the entire analytical procedure. Matrix spike and matrix spike duplicate QC data are advisory.

Method Blank: An analytical control consisting of all reagents, internal standards, and surrogate standards carried through the entire analytical process. Used to monitor laboratory background and reagent contamination.

Not Detected (ND): Not detected at or above the reporting limit.

Relative Percent Difference (RPD): An indication of method precision based on duplicate analyses.

Reporting Limit (RL): The lowest concentration routinely determined during laboratory operations. The RL is generally 1 to 10 times the Method Detection Limit (MDL). Reporting limits are matrix, method, and analyte dependent and take into account any dilutions performed as part of the analysis.

Surrogates: Organic compounds which are similar to analytes of interest in chemical behaviour, but are not found in environmental samples. Surrogates are added to all blanks, calibration and check standards, samples, and spiked samples. Surrogate recovery is monitored as an indication of acceptable sample preparation and instrument performance.

- D: Surrogates diluted out.
- I. Interference.
- 1. Indicates result outside of established laboratory QC limits.

QUALITY CONTROL REPORT

PAGE QR-2

ANALYSIS: Arsenic

MATRIX: Water

METHOD BLANK SAMPLES

LAB ID: IFW_PBW_W INSTR RUN: ICP\980505212400/10/ PREPARED: BATCH ID: IFW050598-W

SAMPLE TYPE: Blank-Method/Media blank INSTRUMENT: TJA Enviro 36 BATCH ID: IFW050598-W DILUTION: 0.200

ANALYZED: 05/05/98 UNITS: mg/L

METHOD:

SPIKE RECOVERY REC LIMITS (%) RPD VALUE (%) LOW HIGH RPD (%) LIMIT (%) REF REPORTING RESULT RESULT LIMIT ANALYTE

0.0050 Arsenic ND

LABORATORY CONTROL SAMPLES

INSTR RUN: ICP\980505212400/12/10 SAMPLE TYPE: Spike-Method/Media blank INSTRUMENT: TJA Enviro 36 UNITS: mg/L LAB ID: IFW_LCD_W PREPARED:

BATCH ID: IFW050598-W DILUTION: 0.200 ANALYZED: 05/05/98

METHOD: REPORTING RFF

RECOVERY REC LIMITS (%) RPD (%) LOW HIGH RPD (%) LIMIT (%) 93.6 80 120 SPIKE VALUE LIMIT VALUE RESULT RESULT ANALYTE 0.0749 ND Arsenic

INSTR RUN: ICP\980505212400/11/10 SAMPLE TYPE: Spike-Method/Media blank

LAB ID: IFW_LCS_W PREPARED: INSTRUMENT: TJA Enviro 36

BATCH ID: IFW050598-W DILUTION: 0.200 ANALYZED: 05/05/98 UNITS:

mg/L METHOD:

REF REPORTING SPIKE RECOVERY REC LIMITS (%) RPD
RESULT RESULT LIMIT VALUE (%) LOW HIGH RPD (%) LIMIT (%)
0.0751 ND 0.0050 0.0800 93.9 80 120 ANALYTE

Arsenic

LABORATORY CONTROL DUPLICATES

LAB ID: IFW_LCR_W PREPARED: INSTR RUN: ICP\980505212400/13/11 SAMPLE TYPE: Method Spike Sample Duplicate

INSTRUMENT: TJA Enviro 36

BATCH ID: IFW050598-W DILUTION: 0.200 ANALYZED: 05/05/98

mg/L UNITS: METHOD:

SPIKE RECOVERY REC LIMITS (%)
VALUE (%) LOW HIGH RPD (%) LIMIT (%)
0.267 15 REPORTING REF RESULT LIMIT 0.0050 ANALYTE RESULT

0.0751 0.0749 Arsenic

QUALITY CONTROL REPORT

PAGE QR-3

ANALYSIS: Extractable TPH

MATRIX: Water

METHOD BLANK SAMPLES

SAMPLE TYPE: Blank-Me INSTRUMENT: HP 5890 UNITS: mg/L METHOD:	thod/Media blank		LAB ID: PREPARED: ANALYZED:		1	INSTR BATCH DILUTI	ID: DS	_C\9805010 LW050198-1 000000		•
ANALYTE Diesel Motor Oil n-Pentacosane	RESULT ND ND surr) 92.3	REF RESULT	REPORTING LIMIT 0.05 0.2	SPIKE VALUE 100	RECOVERY (%)	REC LIM LOW	130	RPD (%)	RPD LIMIT (%)	

LABORATORY CONTROL SAMPLES

SAMPLE TYPE: Lab INSTRUMENT: HP UNITS: mg/ METHOD:	5890	ol Spike		LAB ID: PREPARED: ANALYZED:	LCDW-0501- 05/01/98 05/04/98	1	INSTR RUN: BATCH ID: DILUTION:	GC_C\9805010 DSLW050198-	
ANALYTE Diesel n-Pentacosane	(surr)	RESULT 1.97 91.2	REF RESULT ND 92.3	REPORTING LIMIT 0.05	SPIKE VALUE 2.00 100	RECOVERY (%) 98.5 91.2	REC LIMITS LOW HI 60 13 60 13	GH RPD (%)	RPO LIMIT (%)
SAMPLE TYPE: Lab INSTRUMENT: HP UNITS: mg/ METHOD:	5890	ol Spike			LCSW-0501- 05/01/98 05/04/98	1	INSTR RUN: BATCH ID: DILUTION:	GC_C\9805016 DSLW050198- 1.000000	
ANALYTE Diesel n-Pentacosane	(surr)	RESULT 1.72 81.5	REF RESULT ND 92.3	REPORTING LIMIT 0.05	SPIKE VALUE 2.00 100	RECOVERY (%) 86.0 81.5	60 13	GH RPD (%)	RPD LIMIT (%)

LABORATORY CONTROL DUPLICATES

INSTRUMENT:	Laboratory Contro HP 5890 mg/L	l Sample D	Ouplicate	LAB ID: PREPARED: ANALYZED:		I	INSTR I BATCH I DILUTIO	ID: DS[C\9805010 W050198-1 000000	00000/18/16
ANALYTE Diesel Motor Oil n-Pentacosane	ı (surr)	RESULT 1.97 ND 91.2	REF RESULT 1.72 ND 81.5	REPORTING LIMIT 0.05 0.2	SPIKE VALUE	RECOVERY (%)	REC LIMI LOW	ITS (%) HIGH 130	RPD (%) 13.6 0	RPD LIMIT (%) 20

SAMPLE SURROGATES

SAMPLE TYPE: Sample-Client INSTRUMENT: HP 5890 UNITS: mg/L METHOD:			9804232-03D 05/01/98 05/02/98	INSTR RUN: GC_C\980501000 BATCH ID: DSLW050198-1 DILUTION: 1.000000	000/25/
ANALYTE n-Pentacosane (surr)	REF RESULT RESUL 107.8	REPORTING T LIMIT	SPIKE RECOVER VALUE (%) 100 108		RPD .IMIT (%)

American Environmental Network

WORK ORDER: 9804232

QUALITY CONTROL REPORT

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ANALYSIS: Extractable TPH

MATRIX: Water

SAMPLE SURROGATES

SAMPLE TYPE: Sample-Client LAB ID: 9804232-090 INSTR RUN: GC C\9805010000000/26/
INSTRUMENT: HP 5890 PREPARED: 05/01/98 BATCH ID: DSLW050198-1
UNITS: mg/L ANALYZED: 05/02/98 DILUTION: 1.000000
METHOO:

REF REPORTING SPIKE RECOVERY REC LIMITS (%) RPD

REF REPORTING SPIKE RECOVERY REC LIMITS (%) RPO
ANALYTE RESULT RESULT LIMIT VALUE (%) LOW HIGH RPD (%) LIMIT (%)
n-Pentacosane (surr) 100.9 101 60 130

QUALITY CONTROL REPORT

PAGE QR-5

ANALYSIS: Mercury

MATRIX: Water

METHOD BLANK SAMPLES

LAB ID: HGW_BLNK INSTR RUN: HG\980426133000/1/ PREPARED: BATCH ID: HGW042698 ANALYZED: 04/26/98 DILUTION: 1.000000 SAMPLE TYPE: Blank-Method/Media blank

INSTRUMENT: Coleman Hg Analyzer 50D

UNITS: METHOD:

SPIKE RECOVERY REC LIMITS (%) RPD REF REPORTING ANALYTE RESULT LOW HIGH RPD (%) LIMIT (%) RESULT LIMIT VALUE (%)

Mercury in water ND 0.0002

LABORATORY CONTROL SAMPLES

 SAMPLE TYPE:
 Spike-Method/Media blank
 LAB ID:
 HGW LCD
 INSTR RUN:
 HG\980426133000/3/1

 INSTRUMENT:
 Coleman Hg Analyzer 50D
 PREPARED:
 BATCH ID:
 HGW042698

 UNITS:
 mg/L
 ANALYZED:
 04/26/98
 DILUTION:
 1.000000

METHOD:

REF REPORTING SPIKE RECOVERY REC LIMITS (%) RPD
RESULT LIMIT VALUE (%) LOW HIGH RPD (%) LIMIT (%)
ND 0.0002 0.00200 102 80 120 ANALYTE RESULT Mercury in water 0.00203

ND -----_____

INSTR RUN: HG\980426133000/2/1

SAMPLE TYPE: Spike-Method/Media blank LAB ID: HGW LCS INSTR RUN: HG\9804261 INSTRUMENT: Coleman Hg Analyzer 50D PREPARED: BATCH ID: HGW042698 UNITS: mg/L ANALYZED: 04/26/98 DILUTION: 1.000000

METHOD:

REF REPORTING SPIKE RECOVERY REC LIMITS (%) LOW HIGH RPD (%) LIMIT (%) 80 120 LIMIT 0.0002 RESULT RESULT VALUE (%)

Mercury in water ND 0.00200 0.00200

LABORATORY CONTROL DUPLICATES

SAMPLE TYPE: Method Spike Sample Duplicate
INSTRUMENT: Coleman Hg Analyzer 50D
PREPARED:
BATCH ID: HGW042698
DILUTION: 1 000000

BATCH ID: HGW042698 DILUTION: 1.000000 UNITS: ANALYZED: 04/26/98 mg/L METHOD:

REF REPORTING
ANALYTE RESULT RESULT LIMIT
Mercury in water 0.00203 0.00200 0.0002 SPIKE RECOVERY REC LIMITS (%) RPD VALUE (%) LOW HIGH RPD (%) LIMIT (%)
1.49 20

MATRIX: Soil/Bulk

METHOD BLANK SAMPLES

LAB ID: HGS_BLNK SAMPLE TYPE: Blank-Method/Media blank INSTR RUN: HG\980503123000/1/

BATCH ID: HGS053598 DILUTION: 1.000000 INSTRUMENT: Coleman Hg Analyzer 500

ANALYZED: 05/03/98 UNITS: mg/kg

METHOD:

RECOVERY REC LIMITS (%) RPD REF REPORTING SPIKE ANALYTE RESULT Mercury in soil EPA 7471 ND LOW HIGH RPD (%) LIMIT (%) LIMIT RESULT VALUE (%)

0.06

QUALITY CONTROL REPORT

PAGE QR-6

ANALYSIS: Mercury

MATRIX: Soil/Bulk

LABORATORY CONTROL SAMPLES

SAMPLE TYPE: Spike-Method/N INSTRUMENT: Coleman Hg And UNITS: mg/kg		LAB ID: HGS_LCD PREPARED: ANALYZED: 05/03/98				INSTR RUN: HG\980503123000/3/1 BATCH ID: HGS053598 DILUTION: 1.000000		
METHOO: ANALYTE Mercury in soil EPA 7471	RESULT		REPORTING LIMIT 0.06	SPIKE VALUE 0.400	RECOVERY (%) 95.8	REC LIMITS (%) RPD LOW HIGH RPO (%) LIMIT (%) 75 125		
SAMPLE TYPE: Spike-Method/i INSTRUMENT: Coleman Hg And UNITS: mg/kg METHOD:	Media blank alyzer 50D		LAB ID: PREPARED ANALYZED		,,, ,, 	INSTR RUN: HG\980503123000/2/1 BATCH ID: HGS053598 DILUTION: 1.000000		
ANALYTE Mercury in soil EPA 7471	RESULT 0.383	REF RESULT ND	REPORTING LIMIT 0.06	SPIKE VALUE 0.400	RECOVERY (%) 95.8	REC LIMITS (%) RPD LOW HIGH RPD (%) LIMIT (%) 75 125		

LABORATORY CONTROL DUPLICATES

SAMPLE TYPE: Method Spike Samp INSTRUMENT: Coleman Hg Analyz UNITS: mg/kg		LAB ID: HGS_L PREPARED: ANALYZED: 05/03	BATCH ID	INSTR RUN: HG\980503123000/4/2 BATCH ID: HGS053598 DILUTION: 1.000000			
METHOD: ANALYTE	REF RESULT RESULI	REPORTING SPIN		'S (%) RPD HIGH RPD (%) LIMIT (%)			
Mercury in soil EPA 7471	0.383 0.383	0.06	(11)	0 25			

QUALITY CONTROL REPORT

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ANALYSIS: Metals Scan by ICP

MATRIX: Water

METHOD BLANK SAMPLES

INSTRUMENT: TJA Enviro 3 UNITS: mg/L METHOD:	/Media blank 6		LAB ID: PREPARED: ANALYZED:	IFW_PBW_K 05/01/98		INSTR RUN: ICP\980501200300/1/ BATCH ID: IFW0402898-K DILUTION: 1.000000
ANALYTE Ag Silver Ba Barium Be Beryllium Cd Cadmium Co Cobalt Cr Chromium Cu Copper Mo Molybdenum Ni Nickel Pb Lead Sb Antimony Tl Thallium V Vanadium Zn Zinc	RESULT ND ND ND ND ND ND ND ND ND ND ND ND ND	REF RESULT	REPORTING LIMIT 0.005 0.01 0.002 0.005 0.005 0.01 0.01 0.01 0.01 0.04 0.02 0.05 0.005		(%)	REC LIMITS (%) RPD LOW HIGH RPD (%) LIMIT (%)
SAMPLE TYPE: Blank-Method INSTRUMENT: TJA Enviro 3 UNITS: mg/t	I/Media blank		LAB ID: PREPARED:			INSTR RUN: ICP\980505212400/1/ BATCH ID: IFW050598-W
METHOD:						DILUTION: 1.000000

LABORATORY CONTROL SAMPLES

	E TYPE: Spike-Methoument: TJA Enviro			LAB ID: IFW_LCD_K INSTR RUN: ICP\980501200300/3/1 PREPARED: BATCH ID: IFW0402898-K						
UNITS				ANALYZE	DILUTION: 1.000000					
METHO	υ:		REF	REPORTING	SPIKE	RECOVERY	REC LIM	175 (%)		RPD
ANALY	TE	RESULT	RESULT	LIMIT	VALUE	(%)	LOW	HIGH	RPD (%)	LIMÎT (%)
Ag	Silver	0.0256	ND	0.005	0.0250	102	80	120		, ,
₿a	8arium	1.04	ИD	0.01	1.00	104	80	120		
Ве	Beryllium	0.0262	ND	0.002	0.0250	105	80	120		
Cd	Cadmium	0.0542	ND	0.005	0.0500	108	80	120		
Co	Cobalt	0.262	ИÐ	0.005	0.250	105	80	120		
Çr	Chromium	0.103	ND	0.01	0.100	103	80	120		
Cu	Copper	0.1285	ND	0.01	0.125	103	80	120		
Мо	Molybdenum	0.208	ND	0.01	0.200	104	80	120		
Νi	Nickel	0.254	ND	0.01	0.250	102	80	120		
Pb	Lead	0.531	ND	0.04	0.500	106	80	120		
Sb	Ant imony	0.497	ИD	0.02	0.500	99.4	80	120		
11	Thallium	0.495	ND	0.05	0.500	99.0	80	120		
Ā	Vanadium	0.259	ND	0.005	0.250	104	80	120		
Zn	Zinc	0.261	ΝD	0.01	0.250	104	80	120		

QUALITY CONTROL REPORT

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ANALYSIS: Metals Scan by ICP

MATRIX: Water

LABORATORY CONTROL SAMPLES

SAMPLE TYPE: Spike-M INSTRUMENT: TJA Env	ethod/Media blank iro 36		LAB ID: PREPARED	IFW_LCS_K		INSTR RUN: ICP\98 BATCH IO: IFW040	30501200300/2/1 02898-K
METHOD: ANALYTE Ag Silver Ba Barium Be Beryllium Cd Cachmium Co Cobalt Cr Chromium Cu Copper Mo Molybdenum Ni Nickel Pb Lead Sb Antimony Tl Thallium V Vanadium Zn Zinc	RESULT 0.0257 1.01 0.0258 0.0506 0.253 0.101 0.121 0.198 0.246 0.505 0.473 0.481 0.251	REF RESULT ND ND ND ND ND ND ND ND ND ND ND ND ND	REPORTING LIMIT 0.005 0.01 0.002 0.005 0.01 0.01 0.01 0.01 0.01 0.01 0.0	SPIKE VALUE 0.0250 1.00 0.0250 0.0500 0.250 0.100 0.125 0.200 0.250 0.500 0.500 0.500 0.250	RECOVERY (%) 103 101 103 101 101 101 96.8 99.0 98.4 101 94.6 96.2 100 102	REC LIMITS (%) LOW HIGH RP 80 120	RPO (%) LIMIT (%)
SAMPLE TYPE: Spike-M INSTRUMENT: TJA Env UNITS: mg/l	ethod/Media blank iro 36		LAB ID: PREPARED	IFW_LCD_W : 05/05/98		INSTR RUN: ICP\98 BATCH ID: IFW050	30505212400/3/1 0598-W
ANALYTE Ag Silver Ba Barium Be Beryllium Cd Cadmium Co Cobalt Cr Chromium Cu Copper Mo Molybdenum Ni Nickel Pb Lead Sb Antimony T1 Thallium V Vanadium Zn Zinc	RESULT 0.0230 0.958 0.0230 0.0480 0.240 0.0962 0.115 0.186 0.232 0.480 0.437 0.481 0.238	REF RESULT ND ND ND ND ND ND ND ND ND ND ND ND ND	REPORTING LIMIT 0.005 0.01 0.002 0.005 0.005 0.01 0.01 0.01 0.01 0.04 0.02 0.05 0.005 0.005	SPIKE VALUE 0.0250 1.00 0.0250 0.0500 0.100 0.125 0.200 0.250 0.500 0.500 0.500 0.250	RECOVERY (%) 92.0 95.8 92.0 96.0 96.2 92.0 93.0 92.8 96.0 87.4 96.2 95.2	REC LIMITS (%) LOW HIGH RP 80 120	RPD D (%) LIMIT (%)
SAMPLE TYPE: Spike-M INSTRUMENT: TJA Env UNITS: mg/L METHOD:	ethod/Media blank iro 36		LAB ID: PREPARED ANALYZED	IFW_LCS_W : 05/05/98		INSTR RUN: ICP\98 BATCH ID: IFW050 DILUTION: 1.00	
ANÅLYTE Ag Silver	RESULT 0.0220 0.950 0.0224 0.0472 0.236 0.0912 0.113 0.182 0.229 0.471 0.430 0.487 0.235	REF RESULT ND ND ND ND ND ND ND ND ND ND ND ND ND	REPORTING LIMIT 0.005 0.01 0.002 0.005 0.005 0.01 0.01 0.01 0.01 0.02 0.02 0.05 0.005	SPIKE VALUE 0.0250 1.00 0.0250 0.0500 0.250 0.100 0.125 0.200 0.250 0.500 0.500 0.500 0.250	RECOVERY (%) 88.0 95.0 89.6 94.4 91.2 90.4 91.6 94.2 86.0 97.4 94.0 92.8	REC LIMITS (%) LOW HIGH RP 80 120	RPD D (%) LIMIT (%)

QUALITY CONTROL REPORT

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ANALYSIS: Metals Scan by ICP

MATRIX: Water

LABORATORY CONTROL DUPLICATES

SAMPLI	E TYPE: Method Spike (UMENT: TJA Enviro 36 : mg/L	Sample Duplic	ate	LAB ID:	IFW_LCR_K		INSTR RUN:	ICP\98050120 IFW0402898~K	0300/4/2
UNITS METHO	: mg/L D:								'
			REF	REPORTING	SPIKE	RECOVERY	REC LIMITS (%)	RPD
ANALY 4~	Silver	RESULT	RESULT	LIMIT	VALUE	(%)	LOW HIG	H RPD (%)	
Ag Ba	Barium	0.0256	0.0257	0.005				0.390	20
3e		1.04	1.01	0.01				2.93	20
d.	Cachmium	0.0262 0.0542	0.0258	0.002				1.54	20 20
io Co	Caba 1+	0 000	0.0506	0.005				6.87	20
ir	Chromium	0.262	0.253	0.005				3.50	20
Cu	Copper	0.262	0.101	0.01				1.96	20 20 20 20
lo lo	Molybdenum	0.125 0.208	0.121 0.198	0.01 0.01				3.25	20
11	Nicke 1	0.254	0.196	0.01				4.93	20
b	المسا	0 501	0 505	0 0 0				3.20	20 20 20
Sb	Antimony	0.331	0.303	0.04				5.02	20
1	Thallium	0.437	0.4/3	0.02				4.95	20
	Vine and Jum	0.453 0.250	0.401	0.05				2.07	20 20
;				v.uua				3.14	20
'n	Antimony Thallium Vanadium Zinc TYPE: Method Spike S								20 2400/4/2
Zn 									
Zn 	E TYPE: Method Spike S UMENT: TJA Enviro 36 : mg/L		ate	LAB ID: PREPARED: ANALYZED:	IFW_LCR_W 05/05/98		INSTR RUN: BATCH ID: DILUTION:		
Zn SAMPLI INSTRI JN ITS METHOL	E TYPE: Method Spike S UMENT: TJA Enviro 36 : mg/L D:	Sample Duplic	ate REF	LAB ID: PREPARED: ANALYZED: REPORTING	IFW_LCR_W 05/05/98 SPIKE	RECOVERY	INSTR RUN: BATCH ID: DILUTION: REC LIMITS (ICP\98050521 IFW050598-W 1.000000	2400/4/2 RPD
Zn SAMPLI INSTRI JNITS METHOI	E TYPE: Method Spike S UMENT: TJA Enviro 36 : mg/L D:	Sample Duplic	REF RESULT	LAB ID: PREPARED: ANALYZED: REPORTING LIMIT	IFW_LCR_W 05/05/98 SPIKE	RECOVERY	INSTR RUN: BATCH ID: DILUTION:	ICP\98050521 IFW050598-W 1.000000 %) H RPD (%)	2400/4/2 RPD LIMIT (%)
IN SAMPLI INSTRI INITS METHOI MALY	E TYPE: Method Spike S UMENT: TJA Enviro 36 : mg/L D: TE Silver	Sample Duplic RESULT 0.0230	REF RESULT 0.0220	LAB ID: PREPARED: ANALYZED: REPORTING LIMIT 0.005	IFW_LCR_W 05/05/98 SPIKE	RECOVERY	INSTR RUN: BATCH ID: DILUTION: REC LIMITS (ICP\98050521 IFW050598-W 1.000000 %) H RPD (%) 4.44	2400/4/2 RPD LIMIT (%) 20
Zn SAMPLI INSTRI INITS HETHOI NALY NALY	E TYPE: Method Spike S UMENT: TJA Enviro 36 : mg/L D: TE Silver Barium	RESULT 0.0230 0.958	REF RESULT 0.0220 0.950	LAB ID: PREPARED: ANALYZED: REPORTING LIMIT 0.005 0.01	IFW_LCR_W 05/05/98 SPIKE	RECOVERY	INSTR RUN: BATCH ID: DILUTION: REC LIMITS (ICP\98050521 IFW050598-W 1.000000 %) H RPD (%) 4.44 0.839	RPD LIMIT (%) 20 20
Zn SAMPLI INSTRI INITS METHOI ANALY Ag Ba Ba	E TYPE: Method Spike S UMENT: TJA Enviro 36 : mg/L D: TE Silver Barium Beryllium	RESULT 0.0230 0.958 0.0230	REF RESULT 0.0220 0.950 0.0224	LAB ID: PREPARED: ANALYZED: REPORTING LIMIT 0.005 0.01 0.002	IFW_LCR_W 05/05/98 SPIKE	RECOVERY	INSTR RUN: BATCH ID: DILUTION: REC LIMITS (ICP\98050521 IFW050598-W 1.000000 %) H RPD (%) 4.44 0.839 2.64	RPD LIMIT (%) 20 20
Zn SAMPLI INSTRI INITS METHOL ANALY Ag Ba Be Cd	E TYPE: Method Spike S UMENT: TJA Enviro 36 : mg/L D: TE Silver Barium Beryllium Cadmium	RESULT 0.0230 0.958 0.0230 0.0480	REF RESULT 0.0220 0.950 0.0224 0.0472	LAB ID: PREPARED: ANALYZED: REPORTING LIMIT 0.005 0.01 0.002 0.005	IFW_LCR_W 05/05/98 SPIKE	RECOVERY	INSTR RUN: BATCH ID: DILUTION: REC LIMITS (ICP\98050521 IFW050598-W 1.000000 %) H RPD (%) 4.44 0.839 2.64 1.68	RPD LIMIT (%) 20 20
Zn SAMPLI INSTRI INITS METHOL ANALY Ag Sa Se Cd Co	E TYPE: Method Spike S UMENT: TJA Enviro 36 : mg/L D: TE Silver Barium Beryllium Cachnium Cobalt	RESULT 0.0230 0.958 0.0230 0.0480 0.240	REF RESULT 0.0220 0.950 0.0224 0.0472 0.236	LAB ID: PREPARED: ANALYZED: REPORTING LIMIT 0.005 0.01 0.002 0.005 0.005	IFW_LCR_W 05/05/98 SPIKE	RECOVERY	INSTR RUN: BATCH ID: DILUTION: REC LIMITS (ICP\98050521 IFW050598-W 1.000000 %) H RPD (%) 4.44 0.839 2.64 1.68	RPD LIMIT (%) 20 20
AMPLI INSTRI INITS IETHOI MALY Ba Be Id Io	E TYPE: Method Spike S UMENT: TJA Enviro 36: D: mg/L D: TE Silver Barlium Beryllium Cadmium Cobalt Cohromium	RESULT 0.0230 0.958 0.0230 0.0480 0.240 0.0962	REF RESULT 0.0220 0.950 0.0224 0.0472 0.236 0.0912	LAB ID: PREPARED: ANALYZED: REPORTING LIMIT 0.005 0.01 0.002 0.005 0.005 0.005 0.005	IFW_LCR_W 05/05/98 SPIKE	RECOVERY	INSTR RUN: BATCH ID: DILUTION: REC LIMITS (ICP\98050521 IFW050598-W 1.000000 %) H RPD (%) 4.44 0.839 2.64 1.68 1.68 5.34	RPD LIMIT (%) 20 20 20 20 20 20 20
AMPLI NSTRI INITS METHO MALY Sa Se Cd Co	E TYPE: Method Spike S UMENT: TJA Enviro 36: D: TE Silver Barium Beryllium Cadmium Cobalt Chromium Copper	RESULT 0.0230 0.958 0.0230 0.0480 0.240 0.0962 0.115	REF RESULT 0.0220 0.950 0.0224 0.0472 0.236 0.0912 0.113	LAB ID: PREPARED: ANALYZED: REPORTING LIMIT 0.005 0.01 0.002 0.005 0.01 0.005 0.01	IFW_LCR_W 05/05/98 SPIKE	RECOVERY	INSTR RUN: BATCH ID: DILUTION: REC LIMITS (ICP\98050521 IFW050598-W 1.000000 %) H RPD (%) 4.44 0.839 2.64 1.68 1.68 5.34 1.75	RPD LIMIT (%) 20 20 20 20 20 20 20
Zn SAMPLI INSTRI INSTRI INITS METHOI ANALY Ag Sa Cd Co Cr Cu Ho	E TYPE: Method Spike S UMENT: TJA Enviro 36 : mg/L D: TE Silver Barium Beryllium Cadmium Cobalt Chromium Copper Molybdenum	RESULT 0.0230 0.958 0.0230 0.0480 0.240 0.0962 0.115 0.186	REF RESULT 0.0220 0.950 0.0224 0.0472 0.236 0.0912 0.113 0.182	LAB ID: PREPARED: ANALYZED: REPORTING LIMIT 0.005 0.01 0.002 0.005 0.005 0.005 0.01	IFW_LCR_W 05/05/98 SPIKE	RECOVERY	INSTR RUN: BATCH ID: DILUTION: REC LIMITS (ICP\98050521 IFW050598-W 1.000000 %) H RPD (%) 4.44 0.839 2.64 1.68 1.68 1.68 5.34 1.75 2.17	RPD LIMIT (%) 20 20 20 20 20 20 20
Zn SAMPLI INSTRI INITS JETHOI ANALY Ag Sa Sa Cd Cd Cu	E TYPE: Method Spike S UMENT: TJA Enviro 36: : mg/L D: TE Silver Barium Beryllium Cadmium Cobalt Chromium Copper Molybdenum Nickel	RESULT 0.0230 0.958 0.0230 0.0480 0.240 0.0962 0.115 0.186 0.232	REF RESULT 0.0220 0.950 0.0224 0.0472 0.236 0.0912 0.113 0.182 0.229	LAB ID: PREPARED: ANALYZED: REPORTING LIMIT 0.005 0.01 0.002 0.005 0.005 0.005 0.01 0.01	IFW_LCR_W 05/05/98 SPIKE	RECOVERY	INSTR RUN: BATCH ID: DILUTION: REC LIMITS (ICP\98050521 IFW050598-W 1.000000 %) H RPD (%) 4.44 0.839 2.64 1.68 1.68 1.68 5.34 1.75 2.17 1.30	RPD LIMIT (%) 20 20 20 20 20 20 20 20 20 20
Zn SAMPLI INSTRI INITS HETHOI ANALY Ag Ba Be Cd Co Cr Cu Ho Ho Ho Ho Ho Ho Ho Ho Ho Ho Ho Ho Ho	E TYPE: Method Spike S UMENT: TJA Enviro 36: D: TE Silver Barium Beryllium Cadmium Cobalt Chromium Copper Molybdenum Nickel Lead	RESULT 0.0230 0.958 0.0230 0.0480 0.240 0.0962 0.115 0.186 0.232 0.480	REF RESULT 0.0220 0.950 0.0224 0.0472 0.236 0.0912 0.113 0.182 0.229 0.471	LAB ID: PREPARED: ANALYZED: REPORTING LIMIT 0.005 0.01 0.002 0.005 0.005 0.01 0.001 0.01 0.01	IFW_LCR_W 05/05/98 SPIKE	RECOVERY	INSTR RUN: BATCH ID: DILUTION: REC LIMITS (ICP\98050521 IFW050598-W 1.000000 %) H RPD (%) 4.44 0.839 2.64 1.68 1.68 5.34 1.75 2.17 1.30 1.89	RPD LIMIT (%) 20 20 20 20 20 20 20 20 20 20 20 20 20
Zn SAMPLI INSTRI JNITS	E TYPE: Method Spike S UMENT: TJA Enviro 36: mg/L D: TE Silver Barium Beryllium Cadmium Cobalt Chromium Copper Molybdenum Nickel Lead Antimony	RESULT 0.0230 0.958 0.0230 0.0480 0.240 0.0962 0.115 0.186 0.232 0.480 0.437	REF RESULT 0.0220 0.950 0.0224 0.0472 0.236 0.0912 0.113 0.182 0.229 0.471 0.430	LAB ID: PREPARED: ANALYZED: REPORTING LIMIT 0.005 0.01 0.005 0.01 0.01 0.01 0.01 0.01 0.01 0.01 0.01 0.01 0.01 0.01 0.01 0.01 0.01 0.04 0.02	IFW_LCR_W 05/05/98 SPIKE	RECOVERY	INSTR RUN: BATCH ID: DILUTION: REC LIMITS (ICP\98050521 IFW050598-W 1.000000 %) H	RPD LIMIT (%) 20 20 20 20 20 20 20 20 20 20 20 20 20
Zn SAMPLI INSTRI INITS HETHOI ANALY Ag Ba Cd Co Cr Cu Ho Ho Ho Ho Ho Ho Ho Ho Ho Ho Ho Ho Ho	E TYPE: Method Spike S UMENT: TJA Enviro 36: D: TE Silver Barium Beryllium Cadmium Cobalt Chromium Copper Molybdenum Nickel Lead	RESULT 0.0230 0.958 0.0230 0.0480 0.240 0.0962 0.115 0.186 0.232 0.480	REF RESULT 0.0220 0.950 0.0224 0.0472 0.236 0.0912 0.113 0.182 0.229 0.471	LAB ID: PREPARED: ANALYZED: REPORTING LIMIT 0.005 0.01 0.002 0.005 0.005 0.01 0.001 0.01 0.01	IFW_LCR_W 05/05/98 SPIKE	RECOVERY	INSTR RUN: BATCH ID: DILUTION: REC LIMITS (ICP\98050521 IFW050598-W 1.000000 %) H RPD (%) 4.44 0.839 2.64 1.68 1.68 5.34 1.75 2.17 1.30 1.89	RPD LIMIT (%) 20 20 20 20 20 20 20 20 20 20 20 20 20

MATRIX SPIKE SAMPLES

	E TYPE: Spike-Sample UMENT: TJA Enviro 3			LAB ID: PREPAREC	MD04232-0	1I	INSTR RUN: ICP\980501200300/7/5 BATCH ID: IFW0402898-K				
UNITS METHO				ANALYZED: 05/01/98			DILUTION: 1.000000				
			REF	REPORTING	SPIKE	RECOVERY	REC LIM	IITS (%)		RPD	
analy		RESULT	RESULT	LIMIT	VALUE	(%)	LOW	HIGH	RPD (%)	LIMIT (%)	
Ag 💮	Silver	0.0251	ОN	0.005	0.0250	100	80	120	*** ****		
8a	Barium	1.25	0.228	0.01	1.00	102	80	120			
le	Beryllium	0.0258	ND	0.002	0.0250	103	80	120			
`d	Cadmium	0.0514	ND	0.005	0.0500	103	80	120			
o`	Cobalt	0.256	0.00530	0.005	0.250	100	80	120			
r	Chromium	0.0998	ND	0.01	0.100	99.8	80	120			
u	Copper	0.124	ND	0.01	0.125	99.2	80	120			
lo	Molybdenum	0.231	0.0285	0.01	0.200	101	80	120			
i	Nickel	0.256	0.0107	0.01	0.250	98.1	80	120			
b,	Lead	0.502	ND	0.04	0.500	100	80	120			
b	Antimony	0.481	ND	0.02	0.500	96.2	80	120			
1	Thallium	0.526	ND	0.05	0.500	105	80	120			
٧	Vanadium	0.255	ND	0.005	0.250	102	80	120			

QUALITY CONTROL REPORT

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ANALYSIS: Metals Scan by ICP

MATRIX: Water

MATRIX SPIKE SAMPLES

	E TYPE: Spike-Sampl			LAB ID: MS04232-01I							
UNITS METHO		30		PREPARED: ANALYZED: 05/01/98			DILUTION: 1.000000				
MC I NO	ν.		REF	REPORTING	SPIKE	RECOVERY	REC LIM			RPD	
ANALY	TE	RESULT	RESULT	LIMIT	VALUE	(%)	ΓÓΜ	HIGH	RPD (%)	LIMIT (%)	
Ag	Silver	0.0252	ND	0.005	0.0250	101	80	120			
8a	Barium	1.23	0.228	0,01	1.00	100	80	120			
Вe	Beryllium	0.0249	ND	0.002	0.0250	99.6	80	120			
Cd	Cacimium	0.0479	ИD	0.005	0.0500	95.8	80	120			
Co	Cobalt	0.250	0.00530	0.005	0.250	97.9	80	120			
Cr	Chromium	0.0981	ND	0.01	0.100	98.1	80	120			
Ču	Copper	0.121	ИĎ	0.01	0.125	96.8	80	120			
Mo	Molybdenum	0.224	0.0285	0.01	0.200	97.8	80	120			
Ni	Nickel	0.250	0.0107	0.01	0.250	95.7	80	120			
Pb	Lead	0.494	ND	0.04	0.500	98.8	80	120			
Sb	Antimony	0.470	ND	0.02	0.500	94.0	80	120			
ΤĪ	Thallium	0.489	ND	0.05	0.500	97.8	80	120			
Ý	Vanadium	0.248	NĎ	0.005	0.250	99.2	80	120			_

MATRIX SPIKE DUPLICATES

INSTRU UNITS:	mg/L	e Ouplicate 6		LAB ID: PREPARED: ANALYZED:	MR04232-01 05/01/98	1 I	INSTR BATCH DILUTI	ID: IF	P\98050120 W0402898-K 000000	
METHOL ANALYI Ag Ba Be Cd Co Cr Cu Mo Ni Pb Sb Ti		RESULT 0.0251 1.25 0.0258 0.0514 0.256 0.0998 0.124 0.231 0.256 0.502 0.481 0.526 0.255	REF RESULT 0.0252 1.23 0.0249 0.0479 0.250 0.0981 0.121 0.224 0.250 0.494 0.470 0.489	REPORTING LIMIT 0.005 0.01 0.002 0.005 0.005 0.01 0.01 0.01 0.01 0.01 0.01 0.01 0.01 0.05 0.05	SPIKE VALUE	RECOVERY (%)	REC LIM LOW	ITS (%) HIGH	RPD (%) 0.398 1.61 3.55 7.05 2.37 1.72 2.45 3.08 2.37 1.61 2.31 7.29 2.78	RPD LIMIT (%) 20 20 20 20 20 20 20 20 20 20 20

MATRIX: Soil/Bulk

METHOD BLANK SAMPLES

SAMPLE INSTRU UNITS:	: mg/kg		LAB ID: SAND_PBS PREPARED: ANALYZED: 05/01/98				INSTR RUN: ICP\980430115500/1/ BATCH ID: IFS043098-Q DILUTION: 1.000000				
ANALYTAG Ag Ba Be Cd Co Cr Cu Mo Ni Pb		RESULT ND ND ND ND ND ND ND ND ND ND ND ND ND	REF RESULT	REPORTING LIMIT 0.5 5 0.5 0.5 2 2 2 2 4	SPIKE VALUE	RECOVERY (%)	REC LIMIT LOW	TS (%) HIGH	RPD (%)	RPD LIMIT (%)	

QUALITY CONTROL REPORT

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ANALYSIS: Metals Scan by ICP

MATRIX: Soil/Bulk

METHOD BLANK SAMPLES

INSTRUMENT: T	lank-Method/Medi JA Enviro 36 g/kg	a blank		LAB ID: PREPARED: ANALYZED:	SAND_PBS 05/01/98		INSTR RUN: ICP\980430115500/1/ BATCH ID: IFS043098-Q DILUTION: 1.000000			
ANALYTE fl Thall V Vanad Zn Z	ium	RESULT ND ND ND	REF RESULT	REPORTING LIMIT 50 2 5	SPIKE VALUE		REC LIMITS (%) RPD LOW HIGH RPD (%) LIMIT (%)			
(NSTRUMENT: T	lank-Method/Medi			LAB ID: PREPARED:	IFS_P8S_Z 05/08/98		INSTR RUN: ICP\980508123100/1/ BATCH ID: IFS050698-Z DILUTION: 1.000000			
ANALYTE Ag Sil As Arse Ba Baryll Cd Cadm Co Cob Cr Chrom Cu Cop Mo Molybde Ni Nic Pb L Se Selen Ti Thali V Vanad	ium ium ium alt ium per num kel ead ium	RESULT ND ND ND ND ND ND ND ND ND ND ND ND ND	REF RESULT	REPORTING LIMIT 0.5 10 5 0.5 2 2 2 2 2 2 2 2 50 50	SPIKE VALUE	(%)	REC LIMITS (%) RPD LOW HIGH RPD (%) LIMIT (%)			
SAMPLE TYPE: BINSTRUMENT: TUNITS: m	lank-Method/Medi JA Enviro 36 g/kg	a blank		LAB ID: PREPARED:		~~~	INSTR RUN: ICP\980508123100/15/ BATCH ID: IFS050698-Z DILUTION: 1.000000			
ANALYTE Sb Antim	ony	RESULT ND	REF RESULT	REPORTING LIMIT 5	SPIKE VALUE	RECOVERY (%)	REC LIMITS (%) RPD LOW HIGH RPD (%) LIMIT (%)			

LABORATORY CONTROL SAMPLES

	E TYPE: Spike-Metho UMENT: TJA Enviro : mg/kg			LAB ID: SAND_LCD PREPARED: ANALYZED: 05/01/98				INSTR RUN: ICP\980430115500/3/1 BATCH ID: IFS043098-Q DILUTION: 1.000000				
METHO				ANALIZED	. 03/01/30		DILOII	ON. 1.	300000			
			REF	REPORTING	SPIKE	RECOVERY	REC LIM	ITS (%)		RPD		
ANALY		RESULT	RESULT	LIMIT	VALUE	(%)	LOW	HIĠH	RPD (%)	LIMIT (%)		
Ag	Silver	9.76	ND	0.5	10.0	97.6	75	125				
Ва	Barium	209	ИD	5	200	105	75	125				
Вe	Beryllium	9.41	ИD	0.5	10.0	94.1	75	125				
Cd	Cadmium	9.72	ND	0.5	10.0	97.2	75	125				
Co	Cobalt	103	ND	1	100	103	75	125				
Çr	Chromium	103	ďЯ	2	100	103	75	125				
Cu	Copper	99.5	ND	2	100	99.5	75	125				
Мо	Molybdenum	94.9	ND	2	100	94.9	75	125				
Ni	Nickel	102	ДN	2	100	102	75	125				
РЬ	Lead	101	ND	4	100	101	75	125				
11	Thallium	398	ND	50	400	99.5	75	125				
Ã	Vanadium	102	ИD	2	100	102	75	125				
Zn	Zinc	98.9	ND	5	100	98.9	75	125		_		

QUALITY CONTROL REPORT

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ANALYSIS: Metals Scan by ICP

MATRIX: Soil/Bulk

LABORATORY CONTROL SAMPLES

							~~~~~~~	
SAMPLE TYPE: Spike-Method/Med INSTRUMENT: TJA Enviro 36 UNITS: mg/kg METHOD:	dia blank		LAB ID: PREPARED: ANALYZED:	SAND_LCS 05/01/98		INSTR BATCH DILUTI	RUN: ICP\98043 ID: IFS043098 ON: 1.000000	0115500/2/1 -Q
METHOD:  ANALYTE Ag Silver Ba Barium Be Beryllium Cd Cadmium Co Cobalt Cr Chromium Cu Copper Mo Molybdenum Ni Nickel Pb Lead Tl Thallium V Vanadium Zn Zinc	RESULT 9.89 210 9.37 9.89 103 104 99.7 90.5 102 103 401 103 98.5	REF RESULT ND ND ND ND ND ND ND ND ND ND ND	REPORTING LIMIT 0.5 5 0.5 0.5 2 2 2 2 4 50 2 5	SPIKE VALUE 10.0 200 10.0 10.0 100 100 100 100 100 100 100	RECOVERY (%) 98.9 105 93.7 98.9 103 104 99.7 90.5 102 103 100 103 98.5	REC LIM LOW 75 75 75 75 75 75 75 75 75 75	HTS (%) HIGH RPD (9 125 125 125 125 125 125 125 125 125 125	RPD LIMIT (%)
SAMPLE TYPE: Spike-Method/Med INSTRUMENT: TJA Enviro 36 UNITS: ma/ka	dia blank		LAB ID: PREPARED: ANALYZED:	IFS_LCD_Z 05/08/98		INSTR BATCH DILUTI	RUN: ICP\98050 ID: IFS050698 ON: 1.000000	8123100/4/2 -Z
ANALYTE Ag Silver As Arsenic Ba Barium Be Beryllium Cd Cadmium Co Cobalt Cr Chromium Cu Copper Mo Molybdenum Ni Nickel Pb Lead Se Selenium Tl Thallium V Vanadium Zn Zinc	RESULT 4.56 7.51 101 4.65 5.10 49.3 49.0 48.0 45.9 48.3 53.9 17.7 194 49.8 46.6	REF RESULT ND ND ND ND ND ND ND ND ND ND ND ND	REPORTING LIMIT 0.5 10 5 0.5 1 2 2 2 2 4 20 50 5	SPIKE VALUE 5.00 10.0 100 5.00 5.50 50.0 50.0 50.0	RECOVERY (%) 91.2 75.1 101 93.0 92.7 98.6 98.0 96.0 91.8 96.6 98.0 88.5 92.4 99.6 93.2	REC LIM LOW 75 75 75 75 75 75 75 75 75 75 75	ITS (%) HIGH RPD (%) 125 125 125 125 125 125 125 125 125 125	RPD (%)
SAMPLE TYPE: Spike-Method/Med INSTRUMENT: TJA Enviro 36 UNITS: mg/kg METHOD:	dia blank		LAB ID: PREPARED: ANALYZED:	IFS_LCD_Z 06/08/98		INSTR BATCH DILUTI	RUN: ICP\98050 ID: IFS050698 ON: 1.000000	8123100/18/16 -Z
ANALYTE Sb Antimony	RESULT 45.8		LIMIT 5					
SAMPLE TYPE: Spike-Method/Med INSTRUMENT: TJA Enviro 36 UNITS: mg/kg METHOD:	dia blank		LAB ID: PREPARED: ANALYZED:	IFS_LCS_Z 05/08/98		INSTR BATCH DILUTI	RUN: ICP\98050 ID: IFS050698 ON: 1.000000	8123100/3/2 -Z
ANALYTE Ag Silver As Arsenic Ba Barium Be Beryllium Cd Cadmium Co Cobalt Cr Chromium Cu Copper Mo Molybdenum Ni Nickel	RESULT 4.42 7.55 98.7 4.49 5.06 47.7 47.7 46.3 43.6 46.8	REF RESULT ND ND ND ND ND ND ND ND	LIMIT 0.5 10 5 0.5 0.5	5.00 5.50		REC LIM LOW 75 75 75 75 75 75 75 75 75	ITS (%) HIGH RPD (%) 125 125 125 125 125 125 125 125 125 125	RPD &) LIMIT (%)

QUALITY CONTROL REPORT

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ANALYSIS: Metals Scan by ICP

MATRIX: Soil/Bulk

#### LABORATORY CONTROL SAMPLES

SAMPLE INSTRUM UNITS:	ENT: TJA Enviro 36 mg/kg			LAB ID: PREPARED; ANALYZED:	1FS_LCS_Z 05/08/98	INSTR RUN: ICP\980508123100/3/2 BATCH ID: IFS050698-Z DILUTION: 1.000000			
METHOD: ANALYTE Pb Se T1 V Zn		RESULT 52.7 16.4 187 48.2 44.6	REF RESULT ND ND ND ND	REPORTING LIMIT 4 20 50 2 5	SPIKE VALUE 55.0 20.0 210 50.0 50.0	RECOVERY (%) 95.8 82.0 89.0 96.4 89.2	REC LIMITS (%) RPD LOW HIGH RPD (%) LIMIT (%) 75 125 75 125 75 125 75 125 75 125		
SAMPLE INSTRUM UNITS: METHOD:	mg/kg	Media blank		LAB ID: PREPARED: ANALYZED:	IFS_LCS_Z 06/08/98		INSTR RUN: ICP\980508123100/17/16 BATCH ID: IFS050698-Z DILUTION: 1.000000		
ANALYTE Sb	Ant imony	RESULT 44.1	REF RESULT ND	REPORTING LIMIT 5	SPIKE VALUE 50.0	RECOVERY (%) 88.2	REC LIMITS (%) RPD LOW HIGH RPD (%) LIMIT (%) 75 125		

#### LABORATORY CONTROL DUPLICATES

SAMPLINSTRI UNITS METHO	Sample Duplica	te	LAB ID: PREPARED: ANALYZED:			INSTR RUN: ICP\980430115500/4/2 BATCH ID: IFS043098-Q DILUTION: 1.000000			
ANALY Ag Ba Be Cd Co Cr Cr Cu Mo Ni Pb Tl	RESULT 9.76 209 9.41 9.72 103 103 99.5 94.9 102 101 398 102 98.9	REF RESULT 9.89 210 9.37 9.89 103 104 99.7 90.5 102 103 401 103 98.5	REPORTING LIMIT 0.5 5 0.5 1 2 2 2 2 2 4 50 2 5	SPIKE VALUE	RECOVERY (%)	REC LIMITS (%) LOW HIGH RPD (%) L: 1.32 0.477 0.426 1.73 0 0.966 0.201 4.75 0 1.96 0.751 0.976 0.405	RPD IMIT (%) 25 25 25 25 25 25 25 25 25 25 25		
SAMPLI INSTRI UNITS METHO	Sample Duplica	<b></b> te	LAB ID: PREPARED: ANALYZED:	IFS_LCR_Z 05/08/98		INSTR RUN: ICP\98050812310 BATCH ID: IFS050698-Z DILUTION: 1.000000	0/5/3		
ANALY	DECIII T	REF	REPORTING	SPIKE	RECOVERY	REC LIMITS (%)	RPD		

			REF	REPORTING	SPIKE	RECOVERY	REC LIM	ITS (%)	RPD
ANALY		RESULT	RESULT	LIMIT	VALUE	(%)	LOW	HIGH RPD (%)	LIMIT (%)
Ag	Silver	4.56	4.42	0.5		` .		3,12	25`
As	Arsenic	7.51	7.55	10				0.531	25
βa	Barium	101	98.7	5				2.30	25
Вę	Beryllium	4.65	4.49	0.5				3.50	25 25
Cd	Cachn i um	5.10	5.06	0.5				0.787	25
Со	Cobalt	49.3	47.7	1				3,30	25
Cr	Chromium	49.0	47.7	2				2.69	25
Сu	Copper	48.0	46.3	2				3.61	25 25
Мо	Molybdenum	45.9	43.6	2				5,14	25
Ni	Nickel	48.3	46.8	2				3.15	25
Рb	Lead	53.9	52.7	4				2.25	25
Se	Selenium	17.7	16.4	20				7.62	25
1)	Thallium	194	187	50				3.67	25
٧	Vanadium	49.8	48.2	2				3.27	25
Zn	Zinc	46.6	44.6	5				4.39	25

#### American Environmental Network

WORK ORDER: 9804232

QUALITY CONTROL REPORT

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ANALYSIS: Metals Scan by ICP

MATRIX: Soil/Bulk

#### LABORATORY CONTROL DUPLICATES

INSTR RUN: ICP\980508123100/19/17 BATCH ID: IFS050698-Z DILUTION: 1.000000 SAMPLE TYPE: Method Spike Sample Duplicate LAB ID: IFS_LCR_Z PREPARED:

INSTRUMENT: TJA Enviro 36

UNITS: mg/kg ANALYZED: 06/08/98 METHOD:

RECOVERY REC LIMITS (%)
(%)
LOW HIGH RPD (%) LIMIT (%)
3.78
25 REF REPORTING SPIKE RESULT RESULT ANALYTE LIMIT VALUE

Sb Antimony 45.8 44.1

# QUALITY CONTROL REPORT

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ANALYSIS: Selenium

MATRIX: Water

METHOD BLANK SAMPLES

SAMPLE TYPE: Blank-Method/Media blank LAB ID: IFW PBW INSTR RUN: ICP\980505212400/14/
INSTRUMENT: TJA Enviro 36 PREPARED: BATCH ID: IFW050598-W
UNITS: mg/L ANALYZED: 05/05/98 DILUTION: 0.200

METHOD:

REPORTING REF ANALYTE

SPIKE RECOVERY REC LIMITS (%)
VALUE (%) LOW HIGH RPD (%) LIMIT (%) RESULT RESULT LIMIT ND 0.01

LABORATORY CONTROL SAMPLES

SAMPLE TYPE: Spike-Method/Media blank INSTR RUN: ICP\980505212400/16/14

LAB ID: IFW_LCD_W PREPARED: BATCH ID: IFW050598-W DILUTION: 0.200 INSTRUMENT: TJA Enviro 36

UNITS: mg/L ANALYZED: 05/05/98 METHOD:

REF REPORTING SPIKE RECOVERY REC LIMITS (%) RPD
RESULT RESULT LIMIT VALUE (%) LOW HIGH RPD (%) LIMIT (%)
0.0906 ND 0.01 0.100 90.6 80 120 ANALYTE

Selenium ------

INSTR RUN: ICP\980505212400/15/14 BATCH IO: IFW050598-W DILUTION: 0.200 LAB ID: IFW_LCS_W PREPARED: SAMPLE TYPE: Spike-Method/Media blank

INSTRUMENT: TJA Enviro 36

ANALYZED: 05/05/98 UNITS: mg/L METHOD:

RECOVERY REC LIMITS (%) RPD REPORTING RESULT ANALYTE

VALUE (%) LOW HIGH RPD (%) LIMIT (%) 0.100 86.6 80 120 LIMIT 0.01 RESULT RESULT RESULT 0.0866 ND Selenium

LABORATORY CONTROL DUPLICATES

LAB ID: IFW_LCR_W INSTR RUN: ICP\980505212400/17/15
PREPARED: BATCH ID: IFW050598-W
ANALYZED: 05/05/98 DILUTION: 0.200 SAMPLE TYPE: Method Spike Sample Duplicate

INSTRUMENT: TJA Enviro 36

UNITS:

mg/L METHOD:

RECOVERY REC LIMITS (%)
(%) LOW HIGH RPD (%) LIMIT (%)
4.51 15 SPIKE REF REPORTING RESULT RESULT LIMIT 0.0906 0.0866 0.01 ANALYTE (%) Selenium

# QUALITY CONTROL REPORT

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ANALYSIS: Volatile GC/MS

MATRIX: Water

## METHOD BLANK SAMPLES

SAMPLE TYPE: Blank-Method/Med INSTRUMENT: HP mass spec for JNITS: ug/L GETHOD:		LAB ID: PREPARED: ANALYZED:	BLNK_0501 05/01/98		INSTR RUN: GCMS12\980501000000/1/ BATCH ID: MS12W050198 DILUTION: 1.00				
	REF	REPORTING	SPIKE	RECOVERY	REC LIMITS (%)	RPD			
ANALYTE	RESULT RESULT	LIMIT	VALUE	(%)	LOW HIGH RPD (	%) LIMIT (%)			
1,2-DCA-d4 (surr)	88.3		100	88.3	65 140				
[o]uene~d8 (surr)	102		100	102	70 135 70 135				
o-BFB (surr) L.1-Dichloroethene	102	c	100	102	70 133				
Benzene	ND ND	5 5							
richloroethene	ND	5 5 5							
To luene	ND	5							
Ch lorobenzene	מא	5							
romobenzene	ND	5							
Promoch Loromethane	ND	5							
Promodichloromethane	ND	5 5							
Bromomethane	ND ND	10							
n-Buty Ibenzene	ND								
sec-Buty Ibenzene	ND	5 5 5							
tert-Butylbenzene	ND	5							
Carbon Tetrachloride	ND	5							
Ch loroethane	מא	10							
Chloroform	ND	.5							
Chloromethane	ND	10							
?-Chlorotoluene I-Chlorotoluene	ND ND	5 5							
)ibromochloromethane	ND ND	5							
1.2-Dibromo-3-chloropropan	ND								
.2-Dibromoethane	ND	5 5							
) i bromomethane	ND	5							
,2-Dichlorobenzene	ND	5							
.3-Dichlorobenzene	NO	5							
i,4-Dichlorobenzene	ND	5 10							
lichlorodifluoromethane 1,1-Dichloroethane	ND ND	5							
.2-Dichloroethane	ND	5							
is-1,2-Dichloroethene	ND	5							
rans-1,2-Dichloroethene	ND	5							
1,2-Dichloropropane	ND	5							
L.3-Dichloropropane	ND	5							
2,2-Dichloropropane	ND	5							
l,1-Dichloropropene	ND	5							
Ethylbenzene	ND	5							
Hexachlorobutadiene Isopropylbenzene	ND ND	5 5							
p-Isopropy to an incompany to luene	שא ND	5							
Methylene Chloride	ND	10							
Naphtha lene	ND	5							
n-Propy Ibenzene	ND	5							
Styrene	ND	5							
1,1,1,2-Tetrachloroethane	ND	5							
1,1,2,2-Tetrachloroethane	NO	5							
Tetrachloroethene 1.2.3-Trichlorobenzene	ND ND	5 5							
1,2,4-Trichlorobenzene	ND GN	5 5							
1,1,1-Trichloroethane	ND	5							
1,1,2-Trichloroethane	ND	5							
Trichlorofluoromethane	ND	5							
1,2,3-Trichloropropane	ND	5							
1,2,4-Trimethylbenzene	ND	5							
1.3.5-Trimethylbenzene	ND	. 5							
Vinyl Chloride	ND	10							
Xylenes, Total	ND	10							
T_iah lawahaif l:									
Trichlorotrifluoroethane	ND	5 100							
Trichlorotrifluoroethane Acetone Acrylonitrile	NU NO ND	100 10							

# QUALITY CONTROL REPORT

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ANALYSIS: Volatile GC/MS

MATRIX: Water

# METHOD BLANK SAMPLES

dia blank		LAB ID:	BLNK_0501		INSTR	RUN: GC		1000000/1/
· Volatiles			05/01/98					
RESULT ND NO NO ND ND ND ND ND ND ND ND ND ND ND ND ND	REF RESULT	REPORTING LIMIT 10 100 5 5 50 200 50 50 10 10	SPIKE VALUE	(%)	LOW	HIĠH	RPD (%)	RPO LIMIT (%)
dia blank Volatiles		LAB ID: PREPARED:	BLNK_0505		INSTR BATCH	RUN: GCI	4S12\98050 L2W050598	5020000/1/
T 101 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	REF	REPORTING 555555555555555555555555555555555555	SPIKE VALUE 100 100 100	RECOVERY (%) 103 101 99.0	REC LIN LOW 65 70 70	HITS (%) HIGH 140 135 135	RPD (%)	RPD LIMIT (%)
	A SULT NO NO NO NO NO NO NO NO NO NO NO NO NO	REF RESULT RESULT ND ND ND ND ND ND ND ND ND ND ND ND ND	Volatiles	A	A	Tablank	Tablank	

# QUALITY CONTROL REPORT

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ANALYSIS: Volatile GC/MS

MATRIX: Water

#### **METHOD BLANK SAMPLES**

SAMPLE TYPE: Blank-Method/Med INSTRUMENT: HP mass spec for			LAB ID: PREPARED:	BLNK_0505		INSTR RUN: GCMS12\980505020000/1/ BATCH ID: MS12W050598
UNITS: ug/L METHOD:			ANALYZED:	05/05/98		DILUTION: 1.00
		REF	REPORTING	SPIKE	RECOVERY	REC LIMITS (%) RPD
ANALYTE		RESULT	LIMIT	VALUE	(%)	LOW HIGH RPD (%) LIMIT (%)
Naphtha lene	ND		5			
n-Propy Ibenzene	ND		5			
Styrene	ND		5			
1,1,1,2-Tetrachloroethane	ND		5			
1,1,2,2-Tetrachloroethane	ND		5			
Tetrachloroethene	ND		5			
1,2,3-Trichlorobenzene	ND		5			•
1,2,4-Trichlorobenzene	ND		5			
1,1,1-Trichloroethane	ND		5			
1,1,2-Trichloroethane	ND		5			
Trichlorofluoromethane	ND		5			
1,2,3-Trichloropropane	ND		5 5 5			
1,2,4-Trimethy ibenzene	ND		5			
1,3,5-Trimethylbenzene	ND		5			
Vinyl Chloride	ND		10			
Xylenes, Total	ND		10			
Trichlorotrifluoroethane	ND		5			
Acetone	ND		100			
Acrylonitrile	ND		10			
2-Butanone	ND		100			
Carbon Disulfide	ND		10			
trans-1,4-Dichloro-2-buten	NO		100			
cis-1,3-Dichloropropene	ND		5			
trans-1,3-Dichloropropene	ND		5			
2-Hexanone	NO		50			
Methyl Iodide	ND		200			
4-Methy1-2-pentanone	ND		50			
Vinyl Acetate	ND		50			
2-Chloroethyl Vinyl Ether	ND		ĭŏ			
o-Xylene	NO		io			
m,p-Xylene	ND		10			
MTBE	ND		5			

#### LABORATORY CONTROL SAMPLES

SAMPLE TYPE: Labora INSTRUMENT: HP mas UNITS: ug/L METHOD:		rol Spike r Volatiles		LAB ID: PREPARED ANALYZED	INSTR RUN: GCMS12\980505020000/4/1 BATCH ID: MS12W050598 DILUTION: 1.00					
ANALYTE 1,2-DCA-d4 Toluene-d8 p-BFB 1,1-Dichloroethene	(surr) (surr) (surr)	RESULT 105 102 97.3 44.7	REF RESULT 103 101 99.0 ND	REPORTING LIMIT	SPIKE VALUE 100 100 100 50.0	RECOVERY (%) 105 102 97.3 89.4	REC LIM LOW 65 70 70 70	HIGH 135 130 130 130	RPD (%)	RPO LIMIT (%)
Benzene Trichloroethene Toluene Chlorobenzene		54.2 55.4 53.2 63.8	ND ND NO ND	5 5 5 5	50.0 50.0 50.0 50.0	108 111 106 128	70 70 70 70	130 130 130 130		

# QUALITY CONTROL REPORT

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ANALYSIS: Volatile GC/MS

MATRIX: Water

#### MATRIX SPIKE SAMPLES

	olke-Sample/Mai mass spec for n/L	trix r Volatiles		LAB ID: PREPARED ANALYZED	INSTR RUN: GCMS12\980501000000/7/4 BATCH ID: MS12W050198 DILUTION: 1.00						
METHOD:	,, -		ANAC12ED: 03/02/30				DILOTION. LIGO				
			REF	REPORTING	SPIKE	RECOVERY	REC LIM	IITS (%)		RPD	
ANALYTE		RESULT	RESULT	LIMIT	VALUE	(%)	LOW	HIGH	RPD (%)	LIMIT (%)	
1,2-DCA-d4	(surr)	100	106		100	100	65	135	• •	• •	
To luene-d8	(surr)	107	105		100	107	70	130			
p-8FB	(surr)	91.3	95.4		100	91.3	70	130			
1,1-Dichloroeth	nene	48.7	В	5	50.0	97.4	70	130			
Benzene		51.3	ND	5	50.0	103	70	130			
Trichloroethene	;	56.2	ND	5	50.0	112	70	130			
Toluene		48.9	ИВ	5	50.0	97.8	70	130			
Chlorobenzene		62.5	ND	5	50.0	125	70	130			

#### MATRIX SPIKE DUPLICATES

SAMPLE TYPE: Spiked INSTRUMENT: HP mas	d Sample Du	uplicate		LAB ID: PREPARED	INSTR RUN: GCMS12\980501000000/9/7 BATCH ID: MS12W050198					
INSTRUMENT: HP mass spec for Volatiles UNITS: ug/L METHOD:				DILUTION: 1.00						
ANALYTE 1,2-DCA-d4 Toluene-d8 p-BFB	(surr) (surr) (surr)	RESULT 151 93.4 104	REF RESULT 100 107 91.3	REPORTING LIMIT	SPIKE VALUE 100 100 100	RECOVERY (%) 151 ! 93.4 104	REC LIM LOW 65 70 70	ITS (%) HIGH 135 130 130	RPD (%)	RPD LIMIT (%)
1.1-Dichloroethene Benzene Trichloroethene Toluene Chlorobenzene		45.8 55.3 58.0 55.8 66.6	48.7 51.3 56.2 48.9 62.5	5 5 5 5 5	50.0 50.0 50.0 50.0 50.0				6.14 7.50 3.15 13.2 6.35	20 20 20 20 20

#### SAMPLE SURROGATES

SAMPLE TYPE INSTRUMENT: UNITS: METHOD:	: Sample-Client HP mass spec for ug/L	mass spec for Volatiles			LAB ID: 9804232-03F PREPARED: ANALYZED: 05/01/98				INSTR RUN: GCMS12\980501000000/4/ BATCH ID: MS12W050198 DILUTION: 1.00		
ANALYTE 1,2-DCA-d4 Toluene-d8 p-BF8	(surr) (surr) (surr)	RESULT 106 105 95.4	REF RESULT	REPORTING LIMIT	SPIKE VALUE 100 100 100	RECOVERY (%) 106 105 95.4	65 70	S (%) HIGH 135 130	RPD (%)	RPD LIMIT (%)	
SAMPLE TYPE INSTRUMENT: UNITS: METHOD:	: Sample-Client HP mass spec for ug/L	Volatiles	***********	LAB ID: PREPARED: ANALYZED:	9804232-05F 05/01/98	:	INSTR RU BATCH ID DILUTION	: MS1	2W050198	1000000/5/	
ANALYTE		RESULT	REF RESULT	REPORTING LIMIT	SPIKE VALUE	RECOVERY (%) 121		S (%) HIGH 135	RPD (%)	RPD LIMIT (%)	

# QUALITY CONTROL REPORT

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ANALYSIS: Volatile GC/MS

MATRIX: Water

#### SAMPLE SURROGATES

SAMPLE TYPE: INSTRUMENT: UNITS: METHOD:	Sample-Client HP mass spec for ug/L	Volatiles		LAB ID: PREPARED: ANALYZED:	98 <b>04232-0</b> 91 05/02/98	F	INSTR RI BATCH II DILUTIO	D: MS1	2W050198	000000/6/
ANALYTE 1,2-DCA-d4 Toluene-d8 p-BFB	(surr) (surr) (surr)	RESULT 132 95.5 96.1	REF RESULT	REPORTING LIMIT	SPIKE VALUE 100 100	RECOVERY (%) 132 95.5 96.1	REC LIMI' LOW 65 70 70	TS (%) HIGH 135 130 130	RPD (%)	RPD LIMIT (%)
SAMPLE TYPE: INSTRUMENT: UNITS: METHOD:	Sample-Client HP mass spec for ug/L	Volatiles		LAB ID: PREPARED: ANALYZED:	9804232-016 05/06/98	в	INSTR REBATCH IN	D: MS1	2W050598	5020000/5/
ANALYTE 1.2-DCA-d4 Toluene-d8 p-BFB	(surr) (surr) (surr)	RESULT 1180 971 960	REF RESULT	REPORTING LIMIT	SPIKE VALUE 1000 1000 1000	RECOVERY (%) 118.0 97.1 96.0	REC LIMI LOW 65 70 70	TS (%) HIGH 135 130 130	RPD (%)	RPD LIMIT (%)
SAMPLE TYPE: INSTRUMENT: UNITS: METHOD:	Sample-Client HP mass spec for ug/L	Volatiles		LAB ID: PREPARED: ANALYZED:	98 <b>0</b> 4232-071 05/06/98	 H	INSTR RUBATCH II	D: MS1	\$12\98050! 2W050598	5020000/6/
ANALYTE 1,2-DCA-d4 Toluene-d8 p-BFB	(surr) (surr) (surr)	RESULT 8570 9640 10100	REF RESULT	REPORTING LIMIT	SPIKE VALUE 10000 10000 10000	RECOVERY (%) 85.70 96.40 101.00	REC LIMIT LOW 65 70 70	TS (%) HIGH 135 130 130	RPD (%)	RPD LIMIT (%)

PAGE QR-21

# QUALITY CONTROL DATA

METHOD: EPA 8020, 5030 GCFID

AEN JOB NO: 9804232 AEN LAB NO: E043003 DATE ANALYZED: 04/30/98

INSTRUMENT: E MATRIX: WATER

# Method Blank

	CAS #	Result (ug/L)	Reporting Limit (ug/L)
Benzene	71-43-2	ND	0.5
Toluene	108-88-3	ND	0.5
Ethylbenzene	100-41-4	ND	0.5
Xylenes, Total	1330-20-7	ND	2
HCs as Gasoline		ND mg/L	0.05 mg/L

#### American Environmental Network

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# QUALITY CONTROL DATA

METHOD: EPA 8020, 5030 GCFID

AEN JOB NO: 9804232

INSTRUMENT: E MATRIX: WATER

# Surrogate Standard Recovery Summary

Date Analyzed	Client Id.	Lab Id.	Percent Recovery Fluorobenzene
04/30/98	GGW2	03	95
04/30/98 04/30/98	GGW3	05	95
04/30/98	GGW13	09	95
QC Limits:			70-130

DATE ANALYZED: 04/30/98

SAMPLE SPIKED: INSTRUMENT: E

## Laboratory Control Sample Recovery

						QC Lim	its
Analyte	Spike Added (ug/L)	LCS Result (ug/L)	LCD Result (ug/L)	LCS Percent Recovery	RPD	Percent Recovery	RPD
Benzene Toluene Ethylbenzene Total Xylenes	200 200 200 600	216 210 208 621	221 213 211 620	108 105 104 104	2 1 1 <1	70-130 70-130 70-130 70-130	20 20 20 20

			12553	×45,	A	Sala		9804232
	CHAIN-OF-CUSTO	Y RECOP	RD	Nº	11029	Date:	4-9	22-98 Page / of /
	Project No.: 47/0			ANALYS				REMARKS
	Samplers (Signatures):	EPA Mathod 8010 EPA Method 8020 EPA Method 8020 (BTEX only) EPA Method 8240	EPA Method 8270 TPH as gasoline	VOCS by 8260		Hold	Cooled Soil (S), Water (W), or Vapor (V)	Additional Comments  Noun Page 10 Additional Comments  Notes by 8260
1 2 4 1 1	1315 GGWZ 1150 GGWZ-2.0 1515 GGWZ* 0950 GGWZ-1.5			X   X   X   X   X   X   X   X   X   X			Y W 1/2 Y S A Y W 1/2 Y S A Y W 1/2 Y S A	W9 724 GGW-1 & GGW-4.  1 1 Metalr by CAM-17  W8 (field filtered-plastic HNO3)  1 1 TPHg TPHd by  1 1 Midified 8015
T T BC	0915 GGW4-1.5 1530 GGW13 1525 Blank	Turnaround time:	Stock	Results	ito: Jen Pat-	Total No.	YSA	# Ou not use pastic betters for metals. Filter snapk from extra amount analyse of
	Printed Name:  Company:  Received by (signature):  Printed Name:  Printed Name:  Printed Name:  Printed Name:  Printed Name:	Printed Na Received to Printed Ma Received to Printed Ma Romany:  Printed Ma Company:	ed by signature):  Of LMON  (Signature):  A Color  (Color  (Co	12E /8	Printed Nar  8-30 Company:	ne: y (signature):	Date: Time:  Date:	Method of Shipment:  Pick-up  Laboratory Comments and Log No.:  PEN J. Patturan - Hold  GGW3 & GGW13 12 Plastic  alz-do not us; for meta  F, 141 & presure duplicate  Oiesy ambro for 6663 &  GGWATRIX Consultants  100 Pine Street, 10th Floor  San Francisco, California 94111  415 434 9400

Forms(PF) 012 (Revised 12/95)

# American Environmental Network

# Certificate of Analysis

DOHS Certification: 1172

AIHA Accreditation: 11134

PAGE 1

GEOMATRIX CONSULTANTS 100 PINE ST., SUITE 1000 SAN FRANCISCO, CA 94111

ATTN: JENNIFER PATTERSON CLIENT PROJ. ID: 4710

C.O.C. NUMBER: 11007

REPORT DATE: 05/26/98

DATE(S) SAMPLED: 04/27/98

DATE RECEIVED: 04/28/98

AEN WORK ORDER: 9804273

#### PROJECT SUMMARY:

On April 28, 1998, this laboratory received 6 water sample(s).

Client requested sample(s) be analyzed for chemical parameters. Results of analysis are summarized on the following page(s). Please see quality control report for a summary of QC data pertaining to this project.

Samples will be stored for 30 days after completion of analysis, then disposed of in accordance with State and Federal regulations. Samples may be archived by prior arrangement.

If you have any questions, please contact Client Services at (510) 930-9090.

Reviewed by:

SAMPLE ID: MW-4

AEN LAB NO: 9804273-01 AEN WORK ORDER: 9804273 CLIENT PROJ. ID: 4710

DATE SAMPLED: 04/27/98 DATE RECEIVED: 04/28/98 REPORT DATE: 05/26/98

ANALYTE	METHOD/ CAS#	RESULT	REPORTING LIMIT UNITS	DATE ANALYZED
#Digestion, Metals by ICP	EPA 3010	-	Prep Date	05/05/98
EPA 8260A - Water Benzene Bromobenzene Bromochloromethane Bromodichloromethane Bromoform Bromomethane n-Butylbenzene sec-Butylbenzene tert-Butylbenzene Carbon Tetrachloride Chlorobenzene Chlorotoluene Chlorotoluene Chlorotoluene 4-Chlorotoluene 0ibromochloromethane 1,2-Dibromo-3-chloropropane 1,2-Dibromoethane Dibromomethane 1,2-Dichlorobenzene 1,3-Dichlorobenzene 1,4-Dichlorobenzene 1,4-Dichloroethane 1,1-Dichloroethane 1,1-Dichloroethane 1,2-Dichloroethene cis-1,2-Dichloroethene trans-1,2-Dichloroethene 1,2-Dichloropropane 1,3-Dichloropropane	EPA 8260A 71-43-2 108-86-1 74-97-5 75-27-4 75-25-2 74-83-9 104-51-8 135-98-8 98-06-6 56-23-5 108-90-7 75-00-3 67-66-3 74-87-3 95-49-8 106-43-4 124-48-1 96-12-8 106-93-4 74-95-3 95-50-1 541-73-1 106-46-7 75-34-3 107-06-2 75-35-4 156-59-2 156-60-5 78-87-5 142-28-9 594-20-7 563-58-6 100-41-4 87-68-3 98-82-8 99-87-6 75-09-2 91-20-3 103-65-1	* * * * * * * * * * * * * * * * * * *	55555555555555555555555555555555555555	05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98

SAMPLE ID: MW-4

AEN LAB NO: 9804273-01 AEN WORK ORDER: 9804273 CLIENT PROJ. ID: 4710 DATE SAMPLED: 04/27/98 DATE RECEIVED: 04/28/98 REPORT DATE: 05/26/98

ANALYTE	METHOD/ CAS#	RESULT	REPORTING LIMIT UN	DATE NITS ANALYZED
Styrene 1,1,1,2-Tetrachloroethane 1,1,2,2-Tetrachloroethane Tetrachloroethene Toluene 1,2,3-Trichlorobenzene 1,2,4-Trichloroethane 1,1,2-Trichloroethane Trichloroethene Trichlorofluoromethane 1,2,3-Trichloropropane 1,2,4-Trimethylbenzene 1,3,5-Trimethylbenzene Vinyl Chloride Xylenes, Total Acetone 2-Butanone Carbon Disulfide cis-1,3-Dichloropropene trans-1,3-Dichloropropene 2-Hexanone 4-Methyl-2-pentanone Vinyl Acetate 2-Chloroethyl Vinyl Ether	100-42-5 630-20-6 79-34-5 127-18-4 108-88-3 87-61-6 120-82-1 71-55-6 79-00-5 79-01-6 75-69-4 96-18-4 95-63-6 108-67-8 75-01-4 1330-20-7 67-64-1 78-93-3 75-15-0 10061-01-5 10061-02-6 591-78-6 108-10-1 108-05-4 110-75-8		5 ug/L 5 ug/L 5 ug/L 5 ug/L 5 ug/L 5 ug/L 5 ug/L 5 ug/L 10 ug/L 100 ug/L 100 ug/L 100 ug/L 50 ug/L 50 ug/L 50 ug/L	05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98
CCR 17 Metals Silver Arsenic Barium Beryllium Cadmium Cobalt Chromium Copper Mercury Molybdenum Nickel Lead Antimony Selenium Thallium Vanadium Zinc	EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010	ND 0.018 * 0.15 * ND ND ND ND ND 0.02 * 0.01 * ND ND ND ND	0.005 mg/L 0.005 mg/L 0.01 mg/L 0.002 mg/L 0.005 mg/L 0.005 mg/L 0.01 mg/L 0.01 mg/L 0.01 mg/L 0.01 mg/L 0.01 mg/L 0.02 mg/L 0.05 mg/L 0.05 mg/L 0.05 mg/L	05/06/98 05/05/98 05/05/98 05/05/98 05/05/98 05/05/98 05/05/98 05/05/98 05/05/98 05/05/98 05/05/98 05/05/98 05/05/98

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# GEOMATRIX CONSULTANTS

SAMPLE ID: MW-4

AEN LAB NO: 9804273-01 AEN WORK ORDER: 9804273 CLIENT PROJ. ID: 4710

**DATE SAMPLED: 04/27/98** 

DATE RECEIVED: 04/28/98 REPORT DATE: 05/26/98

ANALYTE	METHOD/ CAS#	RESULT	REPORTING LIMIT	UNITS	DATE ANALYZED

ND = Not detected at or above the reporting limit
 * = Value at or above reporting limit

SAMPLE ID: MW-1

AEN LAB NO: 9804273-02 AEN WORK ORDER: 9804273 CLIENT PROJ. ID: 4710 DATE SAMPLED: 04/27/98 DATE RECEIVED: 04/28/98 REPORT DATE: 05/26/98

ANALYTE	METHOD/ CAS#	RESULT	REPORTING LIMIT UN	DATE NITS ANALYZED
EPA 8260A - Water	EPA 8260A			
Benzene	71-43-2	ND	5 ug/L	05/21/98
Bromobenzene	108-86-1	ND	5 ug/L	05/21/98
Bromoch1oromethane	74-97-5	ND	5 ug/L 5 ug/l	05/21/98
Bromodichloromethane	75-27-4	ND	5 ug/L 5 ug/L	05/21/98
Bromoform	75-25-2	ND	5 ua/l	05/21/98
Bromomethane	74-83-9	ND	10 ug/L 5 ug/L 5 ug/L 5 ug/L 5 ug/L 5 ug/L	05/21/98
n-Butylbenzene	104-51-8	ND	5 ug/L	05/21/98
sec-Butylbenzene	135-98-8	8 7	5 ug/L	05/21/98
tert-Butylbenzene	98-06-6	NĎ	5 ug/L	05/21/98
Carbon Tetrachloride	56-23-5	ND	5 ug/L	05/21/98
Chlorobenzene	108-90-7	ND	5 ug/L	05/21/98
Chloroethane	75-00-3	ND	10 ug/L	05/21/98
Chloroform	67-66-3	ND	5 ug/L	05/21/98
Chloromethane	74-87-3	ND	10 ug/L	05/21/98
2-Chlorotoluene	95-49-8	ND	5 ug/L	05/21/98
4-Chlorotoluene	106-43-4	ND	5 ug/L	05/21/98
Dibromochloromethane	124-48-1	ND	5 ug/L	05/21/98
1,2-Dibromo-3-chloropropane	96-12-8	ND	5 ug/L 5 ug/L 5 ug/L 5 ug/L 5 ug/L 5 ug/L 5 ug/L	05/21/98
1,2-Dibromoethane	106-93-4	ND	5 ug/L	05/21/98
Dibromomethane	74-95-3	ND	5 ug/L	05/21/98
1,2-Dichlorobenzene	95-50-1	ND	5 ug/L	05/21/98
1,3-Dichlorobenzene	541-73-1	ND	5 ug/L	05/21/98
1,4-Dichlorobenzene	106-46-7	ND	5 ug/L	05/21/98
Dichlorodifluoromethane	75-71-8	ND	10 ug/L	05/21/98
1,1-Dichloroethane	75-34-3	18 *	r 5 ug∕L	05/21/98
1,2-Dichloroethane	107-06-2	ND	5 ug/L	05/21/98
1,1-Dichloroethene	75-35 <b>-</b> 4	ND	5 ug/L	05/21/98
cis-1,2-Dichloroethene	156-59-2	ND	5 ug/L	05/21/98
trans-1,2-Dichloroethene	156-60-5	10 *	5 ug/L 5 ug/L 5 ug/L 5 ug/L 5 ug/L 5 ug/L	05/21/98
1,2-Dichloropropane	78-87-5	ND		05/21/98
1,3-Dichloropropane	142-28-9	ND	5 ug/L	05/21/98
2.2-Dichloropropane	594-20-7	ND	5 ug/L	05/21/98
1.1-Dichloropropene	563-58-6	ND	5 ug/L	05/21/98
Ethylbenzene	100-41-4	ND	5 ug/L	05/21/98
Hexachlorobutadiene	87-68-3	ИD	5 ug/L	05/21/98
Isopropylbenzene	98-82-8	8 *	f 5 ug/L	05/21/98
p-Isopropyltoluene	99-87-6	ND	5 ug/L	05/21/98
Methylene Chloride	75-09-2	ND	10 ug/L	05/21/98
Naphthalene	91-20-3	ND	5 ug/L	05/21/98
n-Propylbenzene	103-65-1	9 *	' 5 ug/L	05/21/98
Styrene	100-42-5	ND	5 ug/L	05/21/98
1.1.1.2-Tetrachloroethane	630 <b>-</b> 20-6	ND	5 ug/L	05/21/98

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### GEOMATRIX CONSULTANTS

SAMPLE ID: MW-1

AEN LAB NO: 9804273-02 AEN WORK ORDER: 9804273 CLIENT PROJ. ID: 4710

**DATE SAMPLED: 04/27/98** 

DATE RECEIVED: 04/28/98 **REPORT DATE: 05/26/98** 

ANALYTE	METHOD/ CAS#	RESULT	REPORTING LIMIT UNITS	DATE ANALYZED
1.1.2.2-Tetrachloroethane Tetrachloroethene Toluene 1.2.3-Trichlorobenzene 1.2.4-Trichloroethane 1.1.2-Trichloroethane 1.1.2-Trichloroethane Trichloroethene Trichlorofluoromethane 1.2.3-Trichloropropane 1.2.4-Trimethylbenzene 1.3.5-Trimethylbenzene vinyl Chloride Xylenes, Total Acetone 2-Butanone Carbon Disulfide cis-1,3-Dichloropropene trans-1,3-Dichloropropene 2-Hexanone 4-Methyl-2-pentanone Vinyl Acetate 2-Chloroethyl Vinyl Ether	79-34-5 127-18-4 108-88-3 87-61-6 120-82-1 71-55-6 79-00-5 79-01-6 75-69-4 96-18-4 95-63-6 108-67-8 75-01-4 1330-20-7 67-64-1 78-93-3 75-15-0 10061-01-5 10061-02-6 591-78-6 108-05-4 110-75-8	ND ND ND ND ND ND ND ND ND ND ND ND ND N	5 ug/L 5 ug/L 5 ug/L 5 ug/L 10 ug/L	05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98

ND = Not detected at or above the reporting limit
* = Value at or above reporting limit

SAMPLE ID: MW-2

AEN LAB NO: 9804273-03 AEN WORK ORDER: 9804273 CLIENT PROJ. ID: 4710

**DATE SAMPLED: 04/27/98** DATE RECEIVED: 04/28/98 REPORT DATE: 05/26/98

ANALYTE	METHOD/ CAS#	RI RESULT	EPORTING LIMIT UNITS	DATE ANALYZED
#Digestion, Metals by ICP	EPA 3010	-	Prep Date	05/05/98
EPA 8260A - Water Benzene Bromobenzene Bromochloromethane Bromodichloromethane Bromoform Bromomethane n-Butylbenzene sec-Butylbenzene tert-Butylbenzene chlorobenzene Chlorobenzene Chlorotoluene Chlorotoluene d-Chlorotoluene 1.2-Dibromo-3-chloropropane 1.2-Dibromoethane Dibromomethane 1.2-Dichlorobenzene 1.3-Dichlorobenzene 1.4-Dichlorobenzene 1.4-Dichloroethane 1.1-Dichloroethane 1.1-Dichloroethane 1.1-Dichloroethene cis-1.2-Dichloroethene trans-1.2-Dichloroethene trans-1.2-Dichloropropane 1.3-Dichloropropane 1.1-Dichloropropane 1.3-Dichloropropane 1.3-Dichloropropane 1.3-Dichloropropane 1.3-Dichloropropane 1.3-Dichloropropane 1.3-Dichloropropane 1.1-Dichloropropane	EPA 8260A 71-43-2 108-86-1 74-97-5 75-27-4 75-25-2 74-83-9 104-51-8 135-98-8 98-06-6 56-23-5 108-90-7 75-00-3 67-66-3 74-87-3 95-49-8 106-43-4 124-48-1 96-12-8 106-93-4 74-95-3 95-50-1 541-73-1 106-46-7 75-71-8 75-34-3 107-06-2 75-35-4 156-59-2 156-60-5 78-87-5 142-28-9 594-20-7 563-58-6 100-41-4 87-68-3 98-82-8 99-87-6 75-09-2 91-20-3	NDDDDDDDDDDDDDDDDDDDXNNNNNNNNNNNNNNNNN	//L/L/L/L/L/L/L/L/L/L/L/L/L/L/L/L/L/L/	05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98

SAMPLE ID: MW-2

AEN LAB NO: 9804273-03 AEN WORK ORDER: 9804273 CLIENT PROJ. ID: 4710 DATE SAMPLED: 04/27/98 DATE RECEIVED: 04/28/98 REPORT DATE: 05/26/98

ANALYTE	METHOD/ CAS#	RESULT	REPORTING LIMIT UNITS	DATE ANALYZED
Styrene 1.1.1.2-Tetrachloroethane 1.1.2.2-Tetrachloroethane Tetrachloroethene Toluene 1.2.3-Trichlorobenzene 1.2.4-Trichlorobenzene 1.1.1-Trichloroethane 1.1.2-Trichloroethane Trichloroethene Trichlorofluoromethane 1.2.3-Trichloropropane 1.2.4-Trimethylbenzene 1.3.5-Trimethylbenzene Vinyl Chloride Xylenes, Total Acetone 2-Butanone Carbon Disulfide cis-1.3-Dichloropropene trans-1.3-Dichloropropene 2-Hexanone 4-Methyl-2-pentanone Vinyl Acetate 2-Chloroethyl Vinyl Ether	100-42-5 630-20-6 79-34-5 127-18-4 108-88-3 87-61-6 120-82-1 71-55-6 79-00-5 79-01-6 75-69-4 96-18-4 95-63-6 108-67-8 75-01-4 1330-20-7 67-64-1 78-93-3 75-15-0 10061-01-5 10061-02-6 591-78-6 108-10-1 108-05-4 110-75-8	ND ND ND 120 * ND ND ND ND ND ND ND ND ND ND ND ND ND	5 ug/L 5 ug/L 5 ug/L 5 ug/L 5 ug/L 5 ug/L 5 ug/L 10 ug/L	05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98
CCR 17 Metals Silver Arsenic Barium Beryllium Cadmium Cobalt Chromium Copper Mercury Molybdenum Nickel Lead Antimony Selenium Thallium Vanadium Zinc	EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010 EPA 6010	ND 0.007 * 19 * ND ND ND ND ND ND ND ND ND ND ND ND ND	0.01 mg/L 0.002 mg/L 0.005 mg/L 0.005 mg/L 0.01 mg/L 0.01 mg/L 0.0002 mg/L 0.01 mg/L	05/06/98 05/05/98 05/05/98 05/05/98 05/05/98 05/05/98 05/05/98 05/05/98 05/05/98 05/05/98 05/05/98 05/05/98 05/05/98

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### GEOMATRIX CONSULTANTS

SAMPLE ID: MW-2

AEN LAB NO: 9804273-03 AEN WORK ORDER: 9804273 CLIENT PROJ. ID: 4710

**DATE SAMPLED: 04/27/98** DATE RECEIVED: 04/28/98 REPORT DATE: 05/26/98

ANALYTE	METHOD/ CAS#	RESULT	REPORTING LIMIT	UNITS	DATE ANALYZED

ND = Not detected at or above the reporting limit
 * = Value at or above reporting limit

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# GEOMATRIX CONSULTANTS

SAMPLE ID: MW-3

AEN LAB NO: 9804273-04 AEN WORK ORDER: 9804273 CLIENT PROJ. ID: 4710

**DATE SAMPLED: 04/27/98** DATE RECEIVED: 04/28/98 REPORT DATE: 05/26/98

ANALYTE	METHOD/ CAS#	RESULT	REPORTING LIMIT UNITS	DATE ANALYZED
EPA 8260A - Water Benzene Bromobenzene	EPA 8260A 71-43-2 108-86-1	ND ND	30 ug/L 30 ug/L	05/21/98 05/21/98
Bromochloromethane	74-97-5	ND	30 ug/L	05/21/98
Bromodichloromethane	75-27-4	ND	30 ug/L	05/21/98
Bromoform	75-25-2	ND	30 ug/L	05/21/98
Bromomethane	74-83-9	ND	50 ug/L	05/21/98
n-Butylbenzene	104-51-8	ND	30 ug/L	05/21/98
sec-Butylbenzene	135-98-8	ND	30 ug/L	05/21/98
tert-Butylbenzene	98-06-6	ND	30 ug/L	05/21/98
Carbon Tetrachloride	56-23-5	ND	30 ug/L	05/21/98
Chlorobenzene	108-90-7	ND	30 ug/L	05/21/98
Chloroethane	75-00-3	ND	50 ug/L	05/21/98
Chloroform	67-66-3	ND	30 ug/L	05/21/98
Chloromethane 2-Chlorotoluene 4-Chlorotoluene	74-87-3 95-49-8 106-43-4	ND ND ND	50 ug/L 30 ug/L 30 ug/L 30 ug/L	05/21/98 05/21/98 05/21/98
Dibromochloromethane	106-93-4	ND	30 ug/L	05/21/98
1,2-Dibromo-3-chloropropane		ND	30 ug/L	05/21/98
1,2-Dibromoethane		ND	30 ug/L	05/21/98
Dibromomethane	74-95-3	ND	30 ug/L	05/21/98
1,2-Dichlorobenzene	95-50-1	ND	30 ug/L	05/21/98
1,3-Dichlorobenzene	541-73-1	ND	30 ug/L	05/21/98
1,4-Dichlorobenzene	106-46-7	ND	30 ug/L	05/21/98
Dichlorodifluoromethane	75-71-8	ND	50 ug/L	05/21/98
1,1-Dichloroethane	75-34-3	ND	30 ug/L	05/21/98
1,2-Dichloroethane	107-06-2	ND	30 ug/L	05/21/98
1.1-Dichloroethene cis-1.2-Dichloroethene trans-1.2-Dichloroethene	75-35-4 156-59-2 156-60-5 78-87-5	ND 140 * ND ND	30 ug/L	05/21/98 05/21/98 05/21/98 05/21/98
1,2-Dichloropropane 1,3-Dichloropropane 2,2-Dichloropropane 1,1-Dichloropropene	142-28-9 594-20-7 563-58-6	ND ND ND ND	30 ug/L 30 ug/L 30 ug/L 30 ug/L	05/21/98 05/21/98 05/21/98
Ethylbenzene	100-41-4	ND	30 ug/L	05/21/98
Hexachlorobutadiene	87-68-3	ND	30 ug/L	05/21/98
Isopropylbenzene	98-82-8	ND	30 ug/L	05/21/98
p-Isopropyltoluene Methylene Chloride Naphthalene	99-87-6 75-09-2 91-20-3 103-65-1	ND ND ND ND	30 ug/L 50 ug/L 30 ug/L 30 ug/L	05/21/98 05/21/98 05/21/98 05/21/98
n-Propylbenzene Styrene 1,1,1,2-Tetrachloroethane	100-42-5 630-20-6	ND ND	30 ug/L 30 ug/L 30 ug/L	05/21/98 05/21/98

SAMPLE ID: MW-3

AEN LAB NO: 9804273-04 AEN WORK ORDER: 9804273 CLIENT PROJ. ID: 4710

**DATE SAMPLED: 04/27/98** DATE RECEIVED: 04/28/98 REPORT DATE: 05/26/98

ANALYTE	METHOD/ CAS#	RESULT	REPORTING LIMIT UNIT	DATE FS ANALYZED
1.1.2.2-Tetrachloroethane Tetrachloroethene Toluene 1.2.3-Trichlorobenzene 1.2.4-Trichlorobenzene 1.1.1-Trichloroethane 1.1.2-Trichloroethane Trichloroethene Trichlorofluoromethane 1.2.3-Trichloropropane 1.2.4-Trimethylbenzene 1.3,5-Trimethylbenzene Vinyl Chloride Xylenes. Total Acetone 2-Butanone Carbon Disulfide cis-1.3-Dichloropropene trans-1.3-Dichloropropene 2-Hexanone 4-Methyl-2-pentanone Vinyl Acetate 2-Chloroethyl Vinyl Ether	79-34-5 127-18-4 108-88-3 87-61-6 120-82-1 71-55-6 79-00-5 79-01-6 75-69-4 96-18-4 95-63-6 108-67-8 75-01-4 1330-20-7 67-64-1 78-93-3 75-15-0 10061-01-5 10061-02-6 591-78-6 108-10-1 108-05-4 110-75-8	ND	30 ug/L 30 ug/L 30 ug/L 30 ug/L 30 ug/L 30 ug/L 30 ug/L 30 ug/L 30 ug/L 30 ug/L 50 ug/L	05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98 05/21/98

Reporting limits elevated due to high levels of target compounds. Sample run at dilution.

ND = Not detected at or above the reporting limit
 * = Value at or above reporting limit

SAMPLE ID: MW-5 duplicate MW-3

AEN LAB NO: 9804273-05 AEN WORK ORDER: 9804273 CLIENT PROJ. ID: 4710 DATE SAMPLED: 04/27/98 DATE RECEIVED: 04/28/98

**REPORT DATE:** 05/26/98

ANALYTE	METHOD/ CAS#	RESULT	REPORTING LIMIT UNITS	DATE ANALYZED
EPA 8260A - Water	EPA 8260A			
Benzene	71-43-2	ND	30 ug/L	05/15/98
Bromobenzene	108-86-1	ND	30 ug/L	05/15/98
Bromochloromethane	74-97-5	ND	30 ug/L	05/15/98
Bromodichloromethane	75-27-4	ND	30 ug/L	05/15/98
Bromoform	75-25-2	ND	30 ug/L	05/15/98
Bromomethane	74-83-9	ND	50 ug/L	05/15/98
n-Butylbenzene	104-51-8	ND	30 ug/L	05/15/98
sec-Butylbenzene	135 <b>-</b> 98-8	ND	30 ug/L	05/15/98
tert-Butylbenzene	98-06-6	ND	30 ug/L	05/15/98
Carbon Tetrachloride	56 <b>-</b> 23-5	ND	30 ug/L	05/15/98
Chlorobenzene	108-90-7	ND	30 ug/L	05/15/98
Chloroethane	75-00-3	ND	50 ug/L	05/15/98
Chloroform	67-66-3	ND	30 ug/L	05/15/98
Chloromethane	74-87-3	ND	50 ug/L	05/15/98
2-Chlorotoluene	95-49-8	ND	30 ug/L	05/15/98
4-Chlorotoluene	106-43-4	NĎ	30 ug/L	05/15/98
Dibromochloromethane	124-48-1	ND	30 ug/L	05/15/98
1.2-Dibromo-3-chloropropane	96-12-8	ND	30 ug/L	05/15/98
1,2-Dibromoethane	106-93-4	ND	30 ug/L	05/15/98
Dibromomethane	74-95-3	ND	30 ug/L	05/15/98
1,2-Dichlorobenzene	95-50-1	ND	30 ug/L	05/15/98
1,3-Dichlorobenzene	541-73-1	ND	30 ug/L	05/15/98
1,4-Dichlorobenzene	106-46-7	ND	30 ug/L	05/15/98
Dichlorodifluoromethane	75-71-8	ND	50 ug/L	05/15/98
1,1-Dichloroethane	75-34-3	ND	30 ug/L	05/15/98
1,2-Dichloroethane	107-06-2	ND	30 ug/L	05/15/98
1,1-Dichloroethene	75-35-4	ND	30 ug/L	05/15/98
cis-1,2-Dichloroethene	156-59-2	140 *	30 ug/L	05/15/98
trans-1,2-Dichloroethene	156-60-5	ND	30 ug/L	05/15/98
1,2-Dichloropropane	78-87-5	ND	30 ug/L	05/15/98
1,3-Dichloropropane	142-28-9	ND	30 ug/L	05/15/98
2,2-Dichloropropane	594-20-7	ND	30 ug/L	05/15/98
1,1-Dichloropropene	563-58-6	ND	30 ug/L	05/15/98
Ethylbenzene	100-41-4	ND	30 ug/L	05/15/98
Hexachlorobutadiene	87-68-3	ND	30 ug/L	05/15/98
Isopropylbenzene	98-82-8	ND	30 ug/L	05/15/98
p-Isopropyltoluene	99-87-6	ND	30 ug/L	05/15/98
Methylene Chloride	75-09-2	ND	50 ug/L	05/15/98
Naphthalene	91-20-3	ND	30 ug/L	05/15/98
n-Propylbenzene	103-65-1	NĎ	30 ug/L	05/15/98
Styrene	100-42-5	ND	30 ug/L	05/15/98
1,1,1,2-Tetrachloroethane	630-20-6	ND	30 ug/L	05/15/98

SAMPLE ID: MW-5 duplicate MW-3

AEN LAB NO: 9804273-05 AEN WORK ORDER: 9804273 CLIENT PROJ. ID: 4710

**DATE SAMPLED:** 04/27/98 DATE RECEIVED: 04/28/98 **REPORT DATE: 05/26/98** 

ANALYTE	METHOD/ CAS#	R RESULT	EPORTING LIMIT UNITS	DATE ANALYZED
1.1.2.2-Tetrachloroethane Tetrachloroethene Toluene	79-34-5 127-18-4 108-88-3	ND 30 * 220 *	30 ug/L 30 ug/L 30 ug/L	05/15/98 05/15/98 05/15/98
1,2,3-Trichlorobenzene 1,2,4-Trichlorobenzene 1,1,1-Trichloroethane	87-61-6 120-82-1 71-55-6	ND ND ND	30 ug/L 30 ug/L 30 ug/L	05/15/98 05/15/98 05/15/98
1.1.2-Trichloroethane Trichloroethene Trichlorofluoromethane	79-00-5 79-01-6 75-69-4	ND 190 * ND	30 ug/L 30 ug/L 30 ug/L	05/15/98 05/15/98 05/15/98
1,2,3-Trichloropropane 1,2,4-Trimethylbenzene 1,3,5-Trimethylbenzene Vinyl Chloride	96-18-4 95-63-6 108-67-8 75-01-4	ND ND ND 110 *	30 ug/L 30 ug/L 30 ug/L	05/15/98 05/15/98 05/15/98
Xylenes, Total Acetone 2-Butanone	1330-20-7 67-64-1 78-93-3	ND ND ND	50 ug/L 50 ug/L 500 ug/L 500 ug/L	05/15/98 05/15/98 05/15/98 05/15/98
Carbon Disulfide cis-1,3-Dichloropropene trans-1,3-Dichloropropene	75-15-0 10061-01-5 10061-02-6	ND ND ND	50 ug/L 30 ug/L 30 ug/L 30 ug/L	05/15/98 05/15/98 05/15/98
2-Hexanone 4-Methyl-2-pentanone Vinyl Acetate	591-78-6 108-10-1 108-05-4	ND ND ND	300 ug/L 300 ug/L 300 ug/L 300 ug/L	05/15/98 05/15/98 05/15/98
2-Chloroethyl Vinyl Ether	110-75-8	ND	50 ug/L 50 ug/L	05/15/98

Reporting limits elevated due to high levels of target compounds. Sample run at dilution.

ND = Not detected at or above the reporting limit
 * = Value at or above reporting limit

#### AEN (CALIFORNIA) QUALITY CONTROL REPORT

AEN JOB NUMBER: 9804273 CLIENT PROJECT ID: 4710

### Quality Control and Project Summary

All Samples:

Samples for EPA 8260 were analyzed out of hold time. Results are estimated.

All other laboratory quality control parameters were found to be within established limits.

#### Definitions

Laboratory Control Sample (LCS)/Method Spikes(s): Control samples of known composition. LCS and Method Spike data are used to validate batch analytical results.

Method Blank: An analytical control consisting of all reagents, internal standards, and surrogate standards carried through the entire analytical process. Used to monitor laboratory background and reagent contamination.

Not Detected (ND): Not detected at or above the reporting limit.

Relative Percent Difference (RPD): An indication of method precision based on duplicate analyses.

Reporting Limit (RL): The lowest concentration routinely determined during laboratory operations. The RL is generally 1 to 10 times the Method Detection Limit (MDL). Reporting limits are matrix, method, and analyte dependent and take into account any dilutions performed as part of the analysis.

Surrogates: Organic compounds which are similar to analytes of interest in chemical behaviour, but are not found in environmental samples. Surrogates are added to all blanks, calibration and check standards, samples, and spiked samples. Surrogate recovery is monitored as an indication of acceptable sample preparation and instrument performance.

- D: Surrogates diluted out.
- I. Interference.
- 1. Indicates result outside of established laboratory QC limits.

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WORK ORDER: 9804273

# QUALITY CONTROL REPORT

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**ANALYSIS:** Mercury

MATRIX: Water

METHOD BLANK SAMPLES

SAMPLE TYPE: Blank-Method/Media blank LAB ID: HGW PBW INSTR RUN: HG\980505123000/1/

INSTRUMENT: Coleman Hg Analyzer 50D PREPARED: BATCH ID: HGW050498

ANALYZED: 05/05/98 UNITS: mg/L DILUTION: 1.000000

METHOD:

REPORTING SPIKE REF REC LIMITS (%) ANALYTE RESULT RESULT LIMIT VALUE (%) LOW HIGH RPD (%) LIMIT (%)

0.0002 Mercury in water

LABORATORY CONTROL SAMPLES

LAB ID: HGW_LCD INSTR RUN: HG\980505123000/3/1

SAMPLE TYPE: Spike-Method/Media blank INSTRUMENT: Coleman Hg Analyzer 50D BATCH ID: HGW050498 DILUTION: 1.000000 PREPARED:

ANALYZED: 05/05/98 UNITS: mg/L

METHOD:

REF SPIKE RECOVERY REPORTING REC LIMITS (%) RESULT RESULT LIMIT VALUE LOW HIGH RPD (%) LIMIT (%) Mercury in water 0.00216 ND 0.0002 0.00200 108 80 120

SAMPLE TYPE: Spike-Method/Media blank INSTRUMENT: Coleman Hg Analyzer 50D HGW_LCS LAB ID: INSTR RUN: HG\980505123000/2/1

BATCH ID: HGW050498 DILUTION: 1.000000 PREPARED:

. . . . . .

ANALYZED: 05/05/98 UNITS: mg/L

METHOD:

REPORTING RECOVERY REC LIMITS (%) REF RESULT RESULT VALUE LOW ANALYTE LIMIT HIGH RPD (%) LIMIT (%) 0.00210 0.0002 0.00200 120 Mercury in water ND

LABORATORY CONTROL DUPLICATES

SAMPLE TYPE: Method Spike Sample Duplicate LAB_ID:_ HGW_LCR INSTR RUN: HG\980505123000/4/2

INSTRUMENT: Coleman Hg Analyzer 50D PREPARED: BATCH ID: HGW050498 DILUTION: 1.000000

UNITS: ANALYZED: 05/05/98 mg/L

METHOD: SPIKE RECOVERY

REF REPORTING REC LIMITS (%) RPD RESULT RESULT LOW HIGH RPD (%) ANALYTE LIMIT VALUE (1) LIMIT (%)

20` 0.00216 0.00210 0.0002 2.82 Mercury in water

MATRIX SPIKE SAMPLES

SAMPLE TYPE: Spike-Sample/Matrix INSTRUMENT: Coleman Hg Analyzer 50D LAB ID: MD04273-03I INSTR RUN: HG\980505123000/8/6

BATCH ID: HGW050498 DILUTION: 1.000000 PREPARED:

ANALYZED: 05/05/98 UNITS: mg/L

METHOD:

REF REPORTING RECOVERY REC LIMITS (%) RESULT ANALYTE RESULT LIMIT VALUE (X)HIGH RPD (な) LOW

LIMIT (%) 0.00190 Mercury in water ND 0.0002 0.00200 120

SAMPLE TYPE: Spike-Sample/Matrix INSTRUMENT: Coleman Hg Analyzer 50D LAB ID: MS04273-03I INSTR RUN: HG\980505123000/7/6 PREPARED:

BATCH ID: HGW050498 DILUTION: 1.000000 UNITS: mg/L ANALYZED: 05/05/98

METHOD: REC LIMITS (%) REF REPORTING RECOVERY

RPD RESULT **RESULT** VALUE LOW 80 **ANALYTE** LIMIT HIGH RPD (%) LIMIT (%) 0.00184 ND 0.00200 92.0 Mercury in water 0.0002 120

QUALITY CONTROL REPORT

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**ANALYSIS:** Mercury

MATRIX: Water

MATRIX SPIKE DUPLICATES

METHOD:

SAMPLE TYPE: Spiked Sample Duplicate INSTRUMENT: Coleman Hg Analyzer 50D UNITS: mg/L

ANALYTE Mercury in water

RESULT RESULT 0.00190 0.00184

REPORTING LIMIT 0.0002

LAB ID: MR04273-03I INSTR RUN: HG\980505123000/9/7
PREPARED: BATCH ID: HGW050498
ANALYZED: 05/05/98 DILUTION: 1.000000 ANALYZED: 05/05/98 SPIKE

VALUE

RECOVERY REC LIMITS (%)
(%) LOW HIGH RPD (%)
3.21

RPD LIMIT (%) 20

. QUALITY CONTROL REPORT

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ANALYSIS: Metals Scan by ICP

MATRIX: Water

# METHOD BLANK SAMPLES

SAMPLE INSTRU UNITS: METHOD	MENT: TJA En mg/L	Method/Media blank viro 36		LAB ID: PREPARED: ANALYZED:	IFW_BLNK_W 05/05/98	• • • • • • • • • •	INSTR F BATCH I DILUTIO	ID: IFI	N98050521 050598-W	2400/1/
ANALYT Ag As Bae Cd Co Cr Cu Mo Ni Pb Sb Se T1 V Zn		RESULT ND ND ND ND ND ND ND ND ND ND ND ND ND	REF RESULT	REPORTING LIMIT 0.005 0.04 0.01 0.005 0.005 0.005 0.01 0.01 0.01 0	SPIKE VALUE	RECOVERY	REC LIMI LOW	ITS (%)	RPD (%)	RPD LIMIT (%)

# LABORATORY CONTROL SAMPLES

		<b></b>							<b></b>	<b>.</b>	
SAMPL INSTRI UNITS METHO		d/Media blank 36		LAB ID: PREPARED ANALYZED	INSTR RUN: ICP\980505212400/3/1 BATCH ID: IFW050598-W DILUTION: 1.00						
ANALY Ag As Bae CCo Cr Cw Mi Pbb Se T1 Vzn	TE Silver Arsenic Barium Beryllium Cadmium Cobalt Chromium Coper Molybdenum Nickel Lead Antimony Selenium Thallium Vanadium Zinc	RESULT 0.0230 0.0749 0.958 0.0230 0.0480 0.240 0.0962 0.115 0.186 0.232 0.480 0.437 0.0906 0.481 0.238 0.232	REF RESULT ND ND ND ND ND ND ND ND ND ND ND ND ND	REPORTING LIMIT 0.005 0.04 0.01 0.002 0.005 0.005 0.01 0.01 0.01 0.01 0.04 0.02 0.07 0.05 0.005 0.005	SPIKE VALUE 0.0250 0.0800 1.00 0.0250 0.0500 0.250 0.100 0.125 0.200 0.250 0.500 0.500 0.500 0.500	RECOVERY 92.0 93.6 95.8 96.0 96.0 96.2 92.0 93.0 93.0 94.0 95.2 96.2	REC LIM 80 80 80 80 80 80 80 80 80 80 80 80	ITS (%) HIGH 120 120 120 120 120 120 120 120 120 120	RPD (*)	RPD LIMIT (%)	•••
SAMPL INSTR UNITS METHO		d/Media blank 36		LAB ID: PREPARED ANALYZED	IFW_LCS_W : 05/05/98	•••••	INSTR BATCH DILUTI	ID: IFW	\98050521  050598-W  0	2400/2/1	••

METHOD:	ilig/ L			ANAL I ZEI	J. 03/03/30		DILOTI	ON: I.	30	
TIL TITLE			REF	REPORTING	SPIKE	RECOVERY	REC LIM	ITS (%)		RPD
ANALYTE		RESULT	RESULT	LIMIT	VALUE	(%)	LOW	HIGH	RPD (な)	LIMÏT (な)
Ag	Silver	0.0220	ND	0.005	0.0250	88.0	80	120		
As	Arsenic	0.0751	ND	0.04	0.0800	93.9	80	120		
Вa	Barium	0.950	ND	0.01	1.00	95.0	80	120		
Be	Beryllium	0.0224	ND	0.002	0.0250	89.6	80	120		
Çd	Cadmium	0.0472	ND	0.005	0.0500	94.4	80	120		
Co	_ Cobalt	0.236	МD	0.005	0.250	94.4	80	120		
Cr	Chromium	0.0912	ND	0.01	0.100	91.2	80	120		
Cu	Copper	0.113	ND	0.01	0.125	90.4	80	120		

QUALITY CONTROL REPORT

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ANALYSIS: Metals Scan by ICP

MATRIX: Water

### LABORATORY CONTROL SAMPLES

INSTRU UNITS:	mg/L	Media blank		LAB ID: PREPARED: ANALYZED:	IFW_LCS_W 05/05/98		INSTR BATCH DILUTI	ID: IFV	\98050521 1050598·W	2400/2/1
METHOD  ANALYT  Mo N1 Pb Sb Se T1 V Zn		RESULT 0.182 0.229 0.471 0.430 0.0866 0.487 0.235 0.232	REF RESULT ND ND ND ND ND ND ND ND ND	REPORTING LIMIT 0.01 0.01 0.04 0.02 0.07 0.05 0.005 0.001	SPIKE VALUE 0.200 0.250 0.500 0.500 0.100 0.500 0.250	RECOVERY (%) 91.0 91.6 94.2 86.0 86.6 97.4 94.0 92.8	REC LIM LOW 80 80 80 80 80 80 80 80	ITS (%) HIGH 120 120 120 120 120 120 120 120 120 120	RPD (*)	RPD LIMIT (%)

# LABORATORY CONTROL DUPLICATES

INSTRI UNITS	ke Sample Duplica 36	ite	LAB ID: PREPARED: ANALYZED:	IFW_LCR_W 05/05/98			CP\98050521 FW050598-W .000000	2400/4/2
METHO ANALY Ag Ba Be Cd Cor Cu Mo Ni Pb Sb Se T1 V Zn	RESULT 0.230 0.0749 0.958 0.0230 0.0480 0.240 0.115 0.186 0.232 0.480 0.437 0.0906 0.481 0.238 0.232	REF RESULT 0.0220 0.0751 0.950 0.0224 0.0472 0.236 0.0912 0.113 0.182 0.229 0.429 0.430 0.0866 0.487 0.235 0.232	REPORTING LIMIT 0.005 0.04 0.01 0.002 0.005 0.005 0.01 0.01 0.01 0.01 0.02 0.02 0.05 0.05	SPIKE VALUE	RECOVERY	REC LIMITS (% LOW HIGH	RPD (%) 165 ! 0.267 0.839 2.64 1.68 1.68 5.34 1.75 2.17 1.30 1.89 1.61 4.51 1.24 1.27	RPD (*) 20 20 20 20 20 20 20 20 20 20 20 20 20

QUALITY CONTROL REPORT

QUALITY CONTINUE NEIGHT

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ANALYSIS: Volatile GC/MS

MATRIX: Water

# METHOD BLANK SAMPLES

WORK ORDER: 9804273

# QUALITY CONTROL REPORT

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ANALYSIS: Volatile GC/MS

MATRIX: Water

# METHOD BLANK SAMPLES

SAMPLE TYPE: Blank-Method/Media blank INSTRUMENT: HP mass spec for Volatiles UNITS: ug/L METHOD:	LAB ID PREPAR ANALYZI	ED: — ED: 05/15/98	INSTR RUN: GCMS12\980515020000/1/ BATCH ID: MS12W051598 DILUTION: 1.00
ANALYTE RESULT Acetone ND Acrylonitrile ND 2-Butanone ND Carbon Disulfide ND trans-1,4-Dichloro-2-buten ND cis-1,3-Dichloropropene ND trans-1,3-Dichloropropene ND 4-Methyl Iodide ND 4-Methyl -2-pentanone ND Vinyl Acetate ND 2-Chloroethyl Vinyl Ether ND m,p-Xylene ND MTBE ND	REF REPORTING RESULT LIMIT 100 10 100 100 55 50 200 50 10 10 10 10 55	SPIKE RECOVERY VALUE (*)	REC LIMITS (%) RPD LOW HIGH RPD (%) LIMIT (%)
SAMPLE TYPE: Blank-Method/Media blank INSTRUMENT: HP mass spec for Volatiles UNITS: ug/L METHOD:	LAB ID PREPARI ANALYZI	: BLNK_0521 ED: ED: 05/21/98	INSTR RUN: GCMS13\980521000000/1/ BATCH ID: MS13W052198 DILUTION: 1.00
ANALYTE (surr) 112 Toluene-d8 (surr) 95.3 p-BFB (surr) 97.0 1.1-Dichloroethene ND Trichloroethene ND Trichloroethene ND Toluene ND Toluene ND Chlorobenzene ND Bromochloromethane ND Bromodichloromethane ND Bromomethane ND Bromomethane ND Bromomethane ND Bromomethane ND Carbon Tetrachloride ND Chloroform ND Chloroform ND Chloromethane ND Chlorotoluene ND Chlorotoluene ND Chlorotoluene ND 1.2-Dibromo-3-chloropropan ND 1.2-Dichlorobenzene ND 1.3-Dichlorobenzene ND 1.4-Dichlorobenzene ND 1.1-Dichloroethane ND 1.2-Dichlorobenzene ND 1.2-Dichlorobenzene ND 1.2-Dichlorobenzene ND 1.3-Dichlorobethane ND 1.2-Dichlorobethane ND 1.2-Dichlorobethane ND 1.2-Dichlorobethane ND 1.2-Dichloroethane ND 1.2-Dichloropropane ND 1.3-Dichloropropane ND	REF REPORTING LIMIT 55555555555555555555555555555555555	SPIKE RECOVERY VALUE (%) 100 112 100 95.3 100 97.0	REC LIMITS (%) RPD LOW HIGH RPD (%) LIMIT (%) 65 140 70 135 70 135

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QUALITY CONTROL REPORT

ANALYSIS: Volatile GC/MS

MATRIX: Water

# METHOD BLANK SAMPLES

WORK ORDER: 9804273

SAMPLE TYPE: Blank-Method/Me (NSTRUMENT: HP mass spec fo	dia blank r Volatiles		LAB ID: PREPARED	BLNK_0521		BATCH ID	: MS:	L3W052198	21000000/1/
JNITS: ug/L ÆTHOD:			ANALYZED	: 05/21/98		DILUTION	: 1.0	00	
WALYTE	RESULT	REF Result	REPORTING LIMIT	SPIKE VALUE	RECOVERY (な)	REC LIMIT		DDD (#)	RPD
1-Dichloropropene	ND ND	IVESUE !	5	4VEGE	(4)	LOW	HIGH	RPD (X)	LIMIT (X
thylbenzene	ND		5						
lexachlorobutadiene	ND		5						
sopropylbenzene	ND		5						
-Isopropyltoluene	ND		5						
ethylene Chloride	ND		10						
aphthalene	ND		10						
-Propylbenzene	ND		Ş						
tyrene	ND		2						
.1.1.2-Tetrachloroethane	ND		5						
.1.2.2-Tetrachloroethane	ND		5						
etrach)oroethene	ND		Ş						
.2.3-Trichlorobenzene	ND		Ş						
.2.4-Trichlorobenzene	ND Nati		ភភភភភភភភភភភភភភភភភភភភភភភភភភភភភភភភភភភភភ						
.1.1-Trichloroethane	ND		D E						
.1.2-Trichloroethane	ND		5						
richlorofluoromethane	ND ON		5						
,2,3-Trichloropropane	ND		ž						
.2.4-Trimethylbenzene	ND		5						
3.5-Trimethylbenzene	ND		5						
inyl Chloride	ND		10						
lenes. Total	ND		10 10						
richlorotrifluoroethane	ND		5						
cetone	ND		100						
crylonitrile	ND		100						
Butanone	ND		100						
arbon Disulfide	ND		100						
rans-1,4-Dichloro-2-buten	ND								
is-1,3-Dichloropropene	ND		100						
ans-1,3-Dichloropropene	ND		5 5						
Hexanone	ND		50						
thyl Iodide	ND ND		20						
Methyl -2-pentanone	ON		200						
inyl Acetate	ND		50						
Chloroethyl Vinyl Ether	ND ND		50 10						
·Xylene	ND ND		10						
,p-Xylene	ND DN		10						
,p-xyrene IBE	ND ND		10						
J DEL	NU		5						

# LABORATORY CONTROL SAMPLES

SAMPLE TYPE: Labora INSTRUMENT: HP mas UNITS: ug/L METHOD:	rol Spike r Volatiles		LAB ID: LCS_0515 PREPARED: ANALYZED: 05/15/98				INSTR RUN: GCMS12\980515020000/2/1 BATCH ID: MS12W051598 DILUTION: 1.00				
ANALYTE 1,2-DCA-d4 Toluene-d8 p-BFB 1,1-Dichloroethene Benzene Trichloroethene Toluene Chlorobenzene	(surr) (surr) (surr)	RESULT 105 105 105 52.9 51.8 48.7 52.2 55.3	REF RESULT 117 103 95.5 ND ND ND ND	REPORTING LIMIT 5 5 5 5 5 5	SPIKE VALUE 100 100 100 50.0 50.0 50.0 50.0	RECOVERY (%) 105 105 105 106 104 97.4 104 111	REC LIM LOW 65 70 70 70 70 70 70	HIGH 135 130 130 130 130 130 130 130	RPD (%)	RPD LIMIT (%)	

# QUALITY CONTROL REPORT

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ANALYSIS: Volatile GC/MS

MATRIX: Water

### LABORATORY CONTROL SAMPLES

UNITS: ug/L	itory Conti ss spec foi	rol Spike r Volatiles		LAB ID: LCS_0521 INSTR RUN: GCMS13\98052 PREPARED: BATCH ID: MS13W052198 ANALYZED: 05/21/98 DILUTION: 1.00						1000000/2/1
METHOD:  ANALYTE 1.2-DCA-d4 Toluene-d8 p.BFB 1.1-Dichloroethene Benzene Trichloroethene Toluene Chlorobenzene	(surr) (surr) (surr)	RESULT 90.2 101 94.2 48.6 53.9 53.0 53.3 57.7	REF RESULT 112 95.3 97.0 ND ND ND ND ND	REPORTING LIMIT 5 5 5 5 5	SPIKE VALUE 100 100 50.0 50.0 50.0 50.0 50.0	RECOVERY (*) 90.2 101 94.2 97.2 108 106 107	REC LIM LOW 65 70 70 70 70 70 70	HIGH 135 130 130 130 130 130 130 130	RPD (%)	RPD LIMIT (%)

# MATRIX SPIKE SAMPLES

SAMPLE TYPE: Spike INSTRUMENT: HP ma:	Sample/Ma	trix r Volatiles		LAB ID: PREPARED:	MD04273-010	D	INSTR BATCH		MS13\98052 L3W052198	21000000/8/3
UNITS: ug/L METHOD:	ss spec ro	i voiatiles		ANALYZED:	05/21/98		DILUTI			
ANALYTE		RESULT	REF RESULT	REPORTING LIMIT	SPIKE VALUE	RECOVERY	REC LIM LOW	ITS (%) HIGH	RPD (X)	RPD LIMIT (%)
1,2-DCA-d4 Toluene-d8	(surr) (surr)	84.8 100	78.4 110		100 100	84.8 100	65	135 130		
p-BFB 1.1-Dichloroethene	(surr)	102 51.2	98.4 ND	5	100 50.0	102 102	70 70 70 70 70 70	130 130		
Benzene Trichloroethene		50.4 48.1	ND ND	5 5	50.0 50.0	101 96.2	70 70	130 130		
Toluene Chlorobenzene		51.0 53.1	ND ND	5 5	50.0 50.0	102 106	70 70	130 130		
		· · · · · · · · · · · · · · · · · · ·				- · · · · · · · · · · · · · · · · · · ·		• • • • • • • • • •		•••••
SAMPLE TYPE: Spike	-Sample/Ma	trix		LAB ID:	MS04273-01	D	INSTR		4S13\98052	1000000/7/3

INSTRUMENT: HP mas UNITS: ug/L METHOD:	s spec for	r Volatiles	PREPARED: ANALYZED: 05/21/98				BATCH ID: MS13W052198 DILUTION: 1.00				
ANALYTE 1.2-DCA-d4 Toluene-d8 p-BFB	(surr) (surr) (surr)	RESULT 84.5 101 _103	REF RESULT 78.4 110 98.4	REPORTING LIMIT	SPIKE VALUE 100 100 100	RECOVERY (%) 84.5 101 103	REC LIM LOW 65 70 70	HIGH 135 130 130	RPD (%)	RPD LIMIT (%)	
1,1-Dichloroethene Benzene Trichloroethene Toluene Chlorobenzene		56.3 54.1 51.0 52.9 57.0	ND ND ND ND ND	5 5 5 5 5 5	50.0 50.0 50.0 50.0 50.0	113 108 102 106 114	70 70 70 70 70	130 130 130 130 130			

### MATRIX SPIKE DUPLICATES

		. <b></b>								
SAMPLE TYPE: Spiked	Sample Du	uplicate	LAB ID: MR04273-01D			INSTR RUN: GCMS13\980521000000/9/7 BATCH ID: MS13W052198				
UNITS: ug/L	s spec for	Volatiles		PREPARED: ANALYZED:	05/21/98		DILUTI			
METHOD:			REF	REPORTING	SPIKE	RECOVERY	REC LIM	TTS (2)		RPD
ANALYTE		RESULT	RESULT	LIMIT	VALUE	(%)	LOW	HIGH	RPD (%)	LIMÏT (X)
1.2-DCA-d4 Toluene-d8	(surr) (surr)	84.8 100	84.5 101		100 100	84.8 100	65 70	135 130		
p-BFB	(surr)	102	103		100	102	70	130		
1.1-Dichloroethene		51.2	56.3	5	50.0				9.49	20

QUALITY CONTROL REPORT

ANALYSIS: Volatile GC/MS

MATRIX: Water

# MATRIX SPIKE DUPLICATES

WORK ORDER: 9804273

SAMPLE TYPE: Spiked Sample D INSTRUMENT: HP mass spec fo JNITS: ug/L	Ouplicate or Volatiles		LAB ID: PREPAREC ANALYZEC	MR04273-0 ): ): 05/21/98	10	INSTR RUN: GC BATCH ID: MS DILUTION: 1.	13W052198	21000000/9/7
1ETHOD:		REF	REPORTING	SPIKE	RECOVERY	REC LIMITS (%)		RPD
NALYTE	RESULT	RESULT	LIMIT	VALUE	(%)	LOW HIGH	RPD (%)	LIMIT (%)
Benzene	50.4	54.1	5	50.0			7.08	20
[richloroethene	48.1	51.0	5555555	50.0			5.85	20
[o] uene	51.0	52.9	5	50.0			3.66	20
Chlorobenzene	53.1	57.0	5	50.0			7.08	20
Promobenzene	ND	ND	5				0	
3romochloromethane	ND	ND	5				Õ	
3romodichloromethane	ND	ND	þ				0	
Bromoform	ND	ND					Ŏ	
Bromomethane	ND	ND	10 5 5 5 5				Õ	
n-Butylbenzene	ND	ND	5				Ŏ	
sec-Butylbenzene	ND	ND	ຼີ				0	
tert-Butylbenzene	ND ND	ND	D D				Õ	
Carbon Tetrachloride	ND ND	ND ND	10				0 0	
Chloroethane Chloroform	ND ND	ND ND	10 5					
Chloromethane	ND ND	ND ND	10				0	
	ND ND	ND ND	10				0	
2-Chlorotoluene 1-Chlorotoluene	ND	ND ND	5				Ö	
)ibromochloromethane	ND	ND	5				Ö	
1,2-Dibromo-3-chloropropan	ND	NO	i i				ŏ	
L,2-Dibromoethane	ND	ND	5				ŏ	
)ibromomethane	ND	ND	555555555				ŏ	
1.2.Dichlorobenzene	ND	ND	ž				ŏ	
1.3-Dichlorobenzene	ND	ND	5				ŏ	
L.4-Dichlorobenzene	ND	ND	5				Ŏ	
Dichlorodifluoromethane	ND	ND	10				Ŏ	
L.1-Dichloroethane	ЙĎ	ND	15				ŏ	
1.2-Dichloroethane	Й	ND	ž				ŏ	
cis-1,2-Dichloroethene	ND	ND	š				ŏ	
trans-1,2-Dichloroethene	ND	ND	š				ŏ	
L.2-Dichloropropane	ÖN	NĎ	š				ŏ	
1,3-Dichloropropane	ND	ND	5				ŏ	
2.2-Dichloropropane	ND	NĎ	5665655555555				ŏ	
L.1-Dichloropropene	ND	NĎ	Š				ŏ	
thylbenzene	NĎ	ND	š				ŏ	
lexachlorobutadiene	ND	ND	š				ŏ	
[sopropy]benzene	ND	ND	Š				ŏ	
-Isopropyltoluene	ND	ND	Š				ŏ	
lethylene Chloride	ND	ND	10				Ŏ	
laphthalene	ND	ND	- <u>Š</u>				ŏ	
n-Propylbenzene	ÑĎ	NĎ	5				Ö O	
Styrene	ND	ND	555555				Ŏ	
L.1.1.2-Tetrachloroethane	ND	ND	5				Ŏ	
L.1.2.2-Tetrachloroethane	ND	ND	5				Ŏ	
[etrach]oroethene	ND	ND	5				Ŏ	
1.2.3-Trichlorobenzene	ND	ND	5				Ŏ	
1,2,4-Trichlorobenzene	ND	ND	5				Ŏ	
L.1.1-Trichloroethane	ND	ND	5 5 5 5 5 5				Ō	
l,1,2·Trichloroethane	ND	ND	5				0	
[richlorofluoromethane	ND	ND	5				0	
1,2,3.Trichloropropane	ND	ND	5				0	
l,2,4.Trimethylbenzene	ND	ND	5				Ō	
l,3,5-Trimethylbenzene	ND	ND	5				Ō	
/inyl Chloride	ND	ND	10				Ŏ	
(ylenes, Total	ND	ND	10				Õ	
richlorotrifluoroethane	ND	ND	5				Ō	
Acetone	ND	ND	100				ŏ	
Acrylonitrile	ND	ND	10				Ŏ	
2-Butanone	ND	ND	100				Ō	
Carbon Disulfide	ND	ND	10				Ō	

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# QUALITY CONTROL REPORT

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ANALYSIS: Volatile GC/MS

MATRIX: Water

# MATRIX SPIKE DUPLICATES

SAMPLE TYPE: Spiked Sample Du INSTRUMENT: HP mass spec for UNITS: ug/L	plicate Volatiles		LAB ID: PREPARED: ANALYZED:	MR04273-03 05/21/98	lD		MS13\98052 13W052198 00	1000000/9/7
METHOD:  ANALYTE trans-1,4-Dichloro-2-buten cis-1,3-Dichloropropene trans-1,3-Dichloropropene 2-Hexanone Methyl Iodide 4-Methyl-2-pentanone Vinyl Acetate 2-Chloroethyl Vinyl Ether o-Xylene m.p-Xylene MTBE	RESULT ND ND ND ND ND ND ND ND ND ND ND ND ND	REF RESULT ND ND ND ND ND ND ND ND ND ND ND ND ND	REPORTING LIMIT 100 5 5 50 200 50 10 10	SPIKE VALUE	RECOVERY (な)	REC LIMITS (%) LOW HIGH	RPD (%) 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	RPD LIMIT (%)

# SAMPLE SURROGATES

SAMPLE TYPE: INSTRUMENT: UNITS: METHOD:	Sample-Client HP mass spec for ug/L	Volatiles	• • • • • • • • • •	LAB ID: PREPARED: ANALYZED:	9804273-05/ 05/15/98	A	DILUTION: 5.	12W051598	
ANALYTE 1,2-DCA-d4 Toluene-d8 p-BFB	(surr) (surr) (surr)	RESULT 599 505 526	REF RESULT	REPORTING LIMIT	SPIKE VALUE 500 500 500	RECOVERY (%) 120 101 105	REC LIMITS (%) LOW HIGH 65 135 70 130 70 130	RPD (%)	RPD LIMIT (%)
SAMPLE TYPE: INSTRUMENT: UNITS: METHOD:	Sample-Client HP mass spec for ug/L	Volatiles		LAB ID: PREPARED: ANALYZED:	9804273-01 05/21/98	)	INSTR RUN: GC BATCH ID: MS DILUTION: 1.	13W052198	.000000/3/
ANALYTE 1,2·DCA-d4 Toluene·d8 p-BFB	(surr) (surr) (surr)	RESULT 78.4 110 98.4	REF RESULT	REPORTING LIMIT	SPIKE VALUE 100 100 100	RECOVERY (%) 78.4 110 98.4	REC LIMITS (%) LOW HIGH 65 135 70 130 70 130	RPD (%)	RPD LIMIT (*)
INSTRUMENT: UNITS:		• • • • • • • • • • • • • • • • • • • •			9804273-02	)	INSTR RUN: GC BATCH ID: MS DILUTION: 1.	13W052198	1000000/4/
METHOD: ANALYTE 1,2-DCA-d4 Toluene-d8 p-BFB	(surr) (surr) (surr)	RESULT 77.3 104 99.0	REF RESULT	REPORTING LIMIT	SPIKE VALUE 100 100 100	RECOVERY (%) 77.3 104 99.0	REC LIMITS (%) LOW HIGH 65 135 70 130 70 130	RPD (%)	RPD LIMIT (%)
SAMPLE TYPE INSTRUMENT: UNITS: METHOD:	Sample-Client HP mass spec for ug/L	Volatiles		LAB ID: PREPARED:	9804273-03 05/21/98	D	INSTR RUN: GC BATCH ID: MS DILUTION: 1.	13W052198	
ANALYTE 1.2-DCA-d4 Toluene-d8 p-BFB	(surr) (surr) (surr)	RESULT 76.9 101 97.6	REF RESULT	REPORTING LIMIT	SPIKE VALUE 100 100 100	RECOVERY (%) 76.9 101 97.6	REC LIMITS (%) LOW HIGH 65 135 70 130 70 130	RPD (%)	RPD LIMIT (%)

American Environmental Network

WORK ORDER: 9804273 QUALITY CONTROL REPORT

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ANALYSIS: Volatile GC/MS

MATRIX: Water

# SAMPLE SURROGATES

	Sample-Client HP mass spec for ug/L		• • • • • • • • • •	LAB ID: PREPARED: ANALYZED:	9804273-04		INSTR BATCH DILUTI	RUN: GCI ID: MS:	13W052198	1000000/6/
ANALYTE 1,2-DCA-d4 Toluene-d8 p-BFB	(surr) (surr) (surr)	RESULT 486 489 524	REF RESULT	REPORTING LIMIT	SPIKE VALUE 500 500 500	RECOVERY (%) 97.2 97.8 105	REC LIM LOW 65 70 70	ITS (%) HIGH 135 130 130	RPD (%)	RPD LIMIT (%)

----- End of Quality Control Report -----

9804273

CHAIN-OF-CUSTODY RECORD	2 W=±11007	Date: 4/27/98   Page   of
Project No.: 4710	d Hanadyses	REMARKS
Samplers (Signatures) A Method 8010  A Method 8020  A Method 8240  A Method 8270  A Method 8270	TPH as desei CAM 17 method 8260 Hold-TPHG 8260 hold	Cooled Soil (S), water (W), or vapor (V), or vapor (V), and containers  Number of containers  Soil (S), water (W), or vapor (W),
4/21/18/1030 MW-4		
115 MD-1		
1150 MW-2		1/3   7 \   /   /   /
1400 MW-3		
1430 MW-5		
V 1500 MW-6		XWY8 GA-CH
		wec wec
Turnaround time:	Results to: Patterson	Total No. of containers: 3 50
Relinquished by (signature). Date: Relinquished by (s		
With Carlet		Laboratory Comments and Log No.:
Printed Name: Time: Printed Name:	Time: Printed Name:	Time:
Company: Company:	Company:	
Received by (signature), Date: Received by (signature)	ure): Date: Received by (signature):	Date:
Printed Name: Time: Printed Name:	Time: Printed Name:	Time:
PAUL C RICHEDA 1400		Geomatrix Consultants
Company: Company:	Company:	100 Pine Street, 10th Floor San Francisco, California 94111 415 434 9400

# American Environmental Network

# Certificate of Analysis

DOHS Certification: 1172

AIHA Accreditation: 11134

PAGE 1

GEOMATRIX CONSULTANTS 100 PINE ST., SUITE 1000 SAN FRANCISCO, CA 94111

ATTN: WILL CALDICOTT CLIENT PROJ. ID: 4710.01

C.O.C. NUMBER: 10166

REPORT DATE: 05/22/98

DATE(S) SAMPLED: 05/15/98

DATE RECEIVED: 05/15/98

AEN WORK ORDER: 9805174

#### PROJECT SUMMARY:

On May 15, 1998, this laboratory received 2 water sample(s).

Client requested sample(s) be analyzed for chemical parameters. Results of analysis are summarized on the following page(s). Please see quality control report for a summary of QC data pertaining to this project.

Samples will be stored for 30 days after completion of analysis, then disposed of in accordance with State and Federal regulations. Samples may be archived by prior arrangement.

If you have any questions, please contact Client Services at (510) 930-9090.

Reviewed by:

SAMPLE ID: MW-4

AEN LAB NO: 9805174-01 AEN WORK ORDER: 9805174 CLIENT PROJ. ID: 4710.01 DATE SAMPLED: 05/15/98 DATE RECEIVED: 05/15/98 REPORT DATE: 05/22/98

ANALYTE	METHOD/ CAS#	RESULT	REPORTING LIMIT UNITS	DATE ANALYZED
EPA 8260A - Water	EPA 8260A			
Benzene	71 <b>-</b> 43-2	ND	5 ug/L	05/19/98
Bromobenzene	108-86-1	ND	5 ug/L	05/19/98
Bromochloromethane	74-97-5	ND	5 ug/L	05/19/98
Bromodichloromethane	75-27 <b>-</b> 4	ND	5 ug/L	05/19/98
Bromoform	75-25-2	ND	5 ug/L	05/19/98
Bromomethane	74-83 <b>-</b> 9	ND	1 <u>0</u> ug/L	05/19/98
n-Butylbenzene	104-51-8	ND	5 ug/L	05/19/98
sec-Butylbenzene	135-98-8	ND	5 ug/L	05/19/98
tert-Butylbenzene	98-06-6	ND	5 ug/L	05/19/98
Carbon Tetrachloride	56-23-5	ND	5 ug/L	05/19/98
Chlorobenzene	108-90-7	ND	5 ug/L	05/19/98
Chloroethane	75-00-3	ND	10 ug/L	05/19/98
Chloroform	67-66-3	ND	5 ug/L	05/19/98
Chloromethane	74-87-3	ND ND	10 ug/L	05/19/98
2-Chlorotoluene	95-49-8	ND	5 ug/L	05/19/98
4-Chlorotoluene	106-43-4	ND	5 ug/L	05/19/98 05/19/98
Dibromochloromethane	124-48-1 96-12-8	DN DN	5 ug/L 5 ug/L	05/19/98
1,2-Dibromo-3-chloropropane 1,2-Dibromoethane	106-93-4	ND	5 ug/L	05/19/98
Dibromomethane	74-95-3	ND ND	5 ug/L	05/19/98
1,2-Dichlorobenzene	95-50-1	ND	5 ug/L	05/19/98
1,3-Dichlorobenzene	541 <i>-</i> 73-1	ND	5 ug/L	05/19/98
1,4-Dichlorobenzene	106-46-7	ND	5 ug/L	05/19/98
Dichlorodifluoromethane	75-71-8	ND	10 ug/L	05/19/98
1,1-Dichloroethane	75-34-3	ND	5 ug/L	05/19/98
1,2-Dichloroethane	107-06-2	ND	5 ug/L	05/19/98
1,1-Dichloroethene	75-35-4	ND	5 ug/L	05/19/98
cis-1,2-Dichloroethene	156-59-2	ND	5 ug/L	05/19/98
trans-1,2-Dichloroethene	156-60-5	ND	5 uğ/L	05/19/98
1,2-Dichloropropane	78 <b>-87-</b> 5	ND	5 ug/L	05/19/98
1.3-Dichloropropane	142-28-9	ND	5 ug/L	05/19/98
2.2-Dichloropropane	594-20-7	ND	5 ug/L	05/19/98
1,1-Dichloropropene	563-58-6	ND	5 ug/L	05/19/98
Ethylbenzene	100-41-4	ND	5 ug/L	05/19/98
Hexachlorobutadiene	87-68-3	ND	5 ug/L	05/19/98
Isopropylbenzene	98-82-8	ND	5 ug/L	05/19/98
p-Isopropyltoluene	99-87 <b>-</b> 6	ND	5 ug/L	05/19/98
Methylene Chloride	75-09-2	ND	10 ug/L	05/19/98
Naphthalene	91-20-3	ND	5 ug/L	05/19/98
n-Propylbenzene	103-65-1	ND	5 ug/L	05/19/98
Styrene	100-42-5	ND	5 ug/L	05/19/98
1,1,1,2-Tetrachloroethane	630-20-6	ND	5 ug/L	05/19/98

SAMPLE ID: MW-4 AEN LAB NO: 9805174-01 AEN WORK ORDER: 9805174 CLIENT PROJ. ID: 4710.01

**DATE SAMPLED:** 05/15/98 DATE RECEIVED: 05/15/98 **REPORT DATE: 05/22/98** 

ANALYTE	METHOD/ CAS#	RESULT	REPORTING LIMIT UNITS	DATE ANALYZED
1,1,2,2-Tetrachloroethane Tetrachloroethene	79-34-5 127-18-4	ND ND	5 ug/L 5 ug/L	05/19/98 05/19/98
Toluene	108-88-3	ND	5 110/1	05/19/98
1.2.3-Trichlorobenzene	87-61-6	ND	5 ug/L 5 ug/L 5 ug/L 5 ug/L 5 ug/L 5 ug/L	05/19/98
1,2,4-Trichlorobenzene	120-82-1	ND	5 ug/L	05/19/98
1.1.1-Trichloroethane	71-55-6	ND	5 ug/L	05/19/98
1.1.2-Trichloroethane	79-00-5	ND	5 ug/L	05/19/98
Trichloroethene	79-01-6	ND ND	5 ug/L	05/19/98 05/19/98
Trichlorofluoromethane 1,2,3-Trichloropropane	75-69-4 96-18-4	ND ND	5 ug/L 5 ug/l	05/19/98
1,2,4-Trimethylbenzene	95-63-6	ND	5 ug/L 5 ug/L 5 ug/L	05/19/98
1,3,5-Trimethylbenzene	108-67-8	ND	5 ug/L	05/19/98
Vinyl Chloride	75-01-4	ND	10 ug/L	05/19/98
Xylenes, Total	1330-20-7	ND	10 ug/L	05/19/98
Acetone	67-64-1	ND	100 ug/L	05/19/98
2-Butanone	78-93-3	ND	100 ug/L	05/19/98
Carbon Disulfide	75-15-0	ND ND	10 ug/L 5 ug/L	05/19/98 05/19/98
cis-1,3-Dichloropropene trans-1,3-Dichloropropene	10061-01-5 10061-02-6	ND ND	5 ug/L 5 ug/L	05/19/98
2-Hexanone	591-78-6	ND	50 ug/L	05/19/98
4-Methy1-2-pentanone	108-10-1	ND	50 ug/L	05/19/98
Vinyl Acetate	108-05-4	ND	50 ug/L	05/19/98
2-Cȟloroethyl Vinyl Ether	110-75-8	ND	10 ug/L	05/19/98

ND = Not detected at or above the reporting limit
* = Value at or above reporting limit

### AEN (CALIFORNIA) QUALITY CONTROL REPORT

AEN JOB NUMBER: 9805174 CLIENT PROJECT ID: 4710.01

### Quality Control and Project Summary

All laboratory quality control parameters were found to be within established limits.

#### Definitions

Laboratory Control Sample (LCS)/Method Spikes(s): Control samples of known composition. LCS and Method Spike data are used to validate batch analytical results.

Matrix Spike(s): Aliquot of a sample (aqueous or solid) with added quantities of specific compounds and subjected to the entire analytical procedure. Matrix spike and matrix spike duplicate QC data are advisory.

Method Blank: An analytical control consisting of all reagents, internal standards, and surrogate standards carried through the entire analytical process. Used to monitor laboratory background and reagent contamination.

Not Detected (ND): Not detected at or above the reporting limit.

Relative Percent Difference (RPD): An indication of method precision based on duplicate analyses.

Reporting Limit (RL): The lowest concentration routinely determined during laboratory operations. The RL is generally 1 to 10 times the Method Detection Limit (MDL). Reporting limits are matrix, method, and analyte dependent and take into account any dilutions performed as part of the analysis.

Surrogates: Organic compounds which are similar to analytes of interest in chemical behaviour, but are not found in environmental samples. Surrogates are added to all blanks, calibration and check standards, samples, and spiked samples. Surrogate recovery is monitored as an indication of acceptable sample preparation and instrument performance.

- D: Surrogates diluted out.
- I: Interference.
- !: Indicates result outside of established laboratory QC limits.

# QUALITY CONTROL REPORT

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ANALYSIS: Volatile GC/MS

MATRIX: Water

# METHOD BLANK SAMPLES

SAMPLE TYPE: Blank-Method/Med INSTRUMENT: HP mass spec for UNITS: ug/L	ia blank Volatiles	LAB ID: PREPARED:	BLNK_0519 05/19/98	********		MS13\9805190000000/1/ 13W051998
METHOD:				0.50015501		
ANALYTE 1.2-DCA-d4 (surr) Toluene-d8 (surr) p-BFB (surr)	REF RESULT RESULT 83.8 101 98.5	REPORTING LIMIT	SPIKE VALUE 100 100 100	RECOVERY (%) 83.8 101 98.5	REC LIMITS (%) LOW HIGH 65 140 70 135 70 135	RPD (*) LIMIT (*)
1.1-Dichloroethene Benzene Trichloroethene	ND ND ND	555555555				
Toluene Chlorobenzene Bromobenzene	ND ND ND	5 5 5				
Bromochloromethane Bromodichloromethane Bromoform	ND ND ND	5 5 5				
Bromomethane n-Butylbenzene sec-Butylbenzene	ND ND ND	10 5 5 5 5				
tert-Butylbenzene Carbon Tetrachloride	ND ND ND					
Chloroethane Chloroform Chloromethane	ND ND	10505555555555555555555555555555555555				
2-Chlorotoluene 4-Chlorotoluene Dibromochloromethane	ND ND ND	5 5 5				
1.2-Dibromo-3-chloropropan 1.2-Dibromoethane Dibromomethane	ND ND ND	5 5 5				
1,2-Dichlorobenzene 1,3-Dichlorobenzene 1,4-Dichlorobenzene	ND ND ND	5 5 5				
Dichlorodifluoromethane 1,1-Dichloroethane 1,2-Dichloroethane	ND ND ND	10				
cis-1,2-Dichloroethene trans-1,2-Dichloroethene 1,2-Dichloropropane	ND ND ND	555555555555555555555555555555555555555				
1,3-Dichloropropane 2,2-Dichloropropane 1,1-Dichloropropene	ND ND ND	5 5 5				
Ethylbenzene Hexachlorobutadiene	ND ND	5 5				
Isopropylbenzene p-Isopropyltoluene Methylene Chloride	ND ND ND	5 10				
Naphthalene n-Propylbenzene Styrene	ND ND ND					
1,1.1,2-Tetrachloroethane 1,1,2,2-Tetrachloroethane Tetrachloroethene	ND ND ND	5 5 5 5 5 5 5				
1.2.3-Trichlorobenzene 1.2.4-Trichlorobenzene 1.1.1-Trichloroethane	ND ND ND	5 5 5				
1,1,2-Trichloroethane Trichlorofluoromethane 1,2,3-Trichloropropane	ND ND ND	5 5 5 5				
1.2.4-Trimethylbenzene 1.3.5-Trimethylbenzene	ND ND ND	5 5 5 10				
Vinyl Chloride Xylenes, Total Trichlorotrifluoroethane	ND ND	10 5				
Acetone Acrylonitrile 2-Butanone	ND ND ND	100 10 100				

QUALITY CONTROL REPORT

PAGE QR-3

ANALYSIS: Volatile GC/MS

MATRIX: Water

# METHOD BLANK SAMPLES

SAMPLE TYPE: Blank-Method/Mei INSTRUMENT: HP mass spec for UNITS: ug/L		LAB ID: PREPARED ANALYZED	BLNK_0519 : 05/19/98		INSTR RUN: GC BATCH ID: MS DILUTION: 1.	13W051998	9000000/1/
METHOD:  ANALYTE Carbon Disulfide trans-1.4-Dichloro-2-buten cis-1.3-Dichloropropene trans-1.3-Dichloropropene 2-Hexanone Methyl Iodide 4-Methyl-2-pentanone Vinyl Acetate 2-Chloroethyl Vinyl Ether o-Xylene m.p-Xylene MTBE	RE RESULT RESU ND ND ND ND ND ND ND ND ND ND ND ND ND		SPIKE VALUE	RECOVERY (%)	REC LIMITS (%) LOW HIGH	RPD (X)	RPD LIMIT (%)

### LABORATORY CONTROL SAMPLES

SAMPLE TYPE: Labora INSTRUMENT: HP mas UNITS: ug/L METHOD:	tory Contr s spec for	ol Spike Volatiles		LAB ID: PREPARED: ANALYZED:	LCS_0519 05/19/98		INSTR BATCH DILUTI	ID: MSI	13W051998	9000000/2/1
ANALYTE 1.2-DCA-d4 Toluene-d8 p-BFB 1.1-Dichloroethene Benzene Trichloroethene Toluene Chlorobenzene	(surr) (surr) (surr)	RESULT 84.7 98.8 98.8 54.5 52.3 51.3 53.9 56.4	REF RESULT 83.8 101 98.5 ND ND ND ND ND	REPORTING LIMIT 5 5 5 5 5	SPIKE VALUE 100 100 50.0 50.0 50.0 50.0 50.0	RECOVERY (*) 84.7 98.8 98.8 109 105 103 108 113	REC LIM LOW 65 70 70 70 70 70 70	ITS (%) HIGH 135 130 130 130 130 130 130 130	RPD (%)	RPD LIMIT (%)

# SAMPLE SURROGATES

SAMPLE TYPE: INSTRUMENT: UNITS: METHOD:	Sample-Client HP mass spec for ug/L	Volatiles	• • • • • • • • •	LAB ID: PREPARED: ANALYZED:	9805174-01/ 05/19/98	A		ID: MS	L3W051998	9000000/8/
ANALYTE 1,2-DCA-d4 Toluene-d8 p-BFB	(surr) (surr) (surr)	RESULT 76.3 99.8 89.4	ref Result	REPORTING LIMIT	SPIKE VALUE 100 100 100	RECOVERY (*) 76.3 99.8 89.4	REC LIM LOW 65 70 70	ITS (%) HIGH 135 130 130	RPD (%)	RPD LIMIT (%)

..... End of Quality Control Report .....

Time:

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Company:

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Geomatrix Consultants

100 Pine Street, 10th Floor

GEOMATRIX San Francisco, California 94111 415 434 9400

Forms(PF) 012 (Revised 12/95)

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### FRIEDMAN & BRUYA, INC.

### **ENVIRONMENTAL CHEMISTS**

James E. Bruya, Ph.D. Charlene Jensen, M.S. Bradley T. Benson, B.S. Kurt Johnson, B.S.

3012 16th Avenue West Seattle, WA 98119-2029 TEL: (206) 285-8282 FAX: (206) 283-5044 e-mail: fbi@isomedia.com

May 19, 1998

Will Caldicott, Project Manager Geomatrix Consultants, Inc. 100 Pine Street, Suite 1000 San Francisco, CA 94111-5112

Dear Mr. Caldicott:

Included are the results from the testing of material submitted on May 15, 1998 from your 4710.01 project. Any samples that may remain are currently scheduled for disposal in 30 days. If you would like us to return your samples or arrange for long term storage at our offices, please contact us as soon as possible.

We appreciate this opportunity to be of service to you and hope you will call if you should have any questions.

Sincerely,

FRIEDMAN & BRUYA, INC.

Charlene Jensen

Chemist

Enclosures GMC0519R.DOC

### FRIEDMAN & BRUYA, INC.

### **ENVIRONMENTAL CHEMISTS**

Date of Report: 05/19/98 Date Received: 05/15/98

Project: 4710.01

Date Extracted: 05/18/98 Date Analyzed: 05/19/98

# RESULTS FROM THE ANALYSIS OF THE WATER SAMPLE FOR FINGERPRINT CHARACTERIZATION BY CAPILLARY GAS CHROMATOGRAPHY USING A FLAME IONIZATION DETECTOR (FID) AND ELECTRON CAPTURE DETECTOR (ECD)

### Sample ID

### GC Characterization

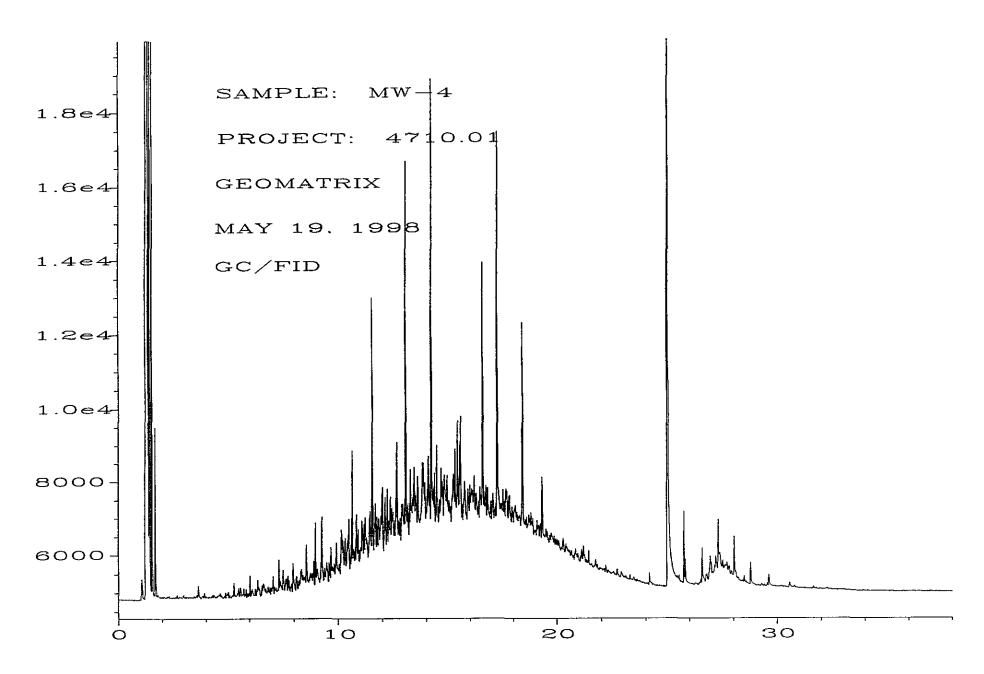
MW-4

The GC trace using the flame ionization detector (FID) showed the presence of medium boiling compounds. The patterns displayed by these peaks are indicative of diesel fuel.

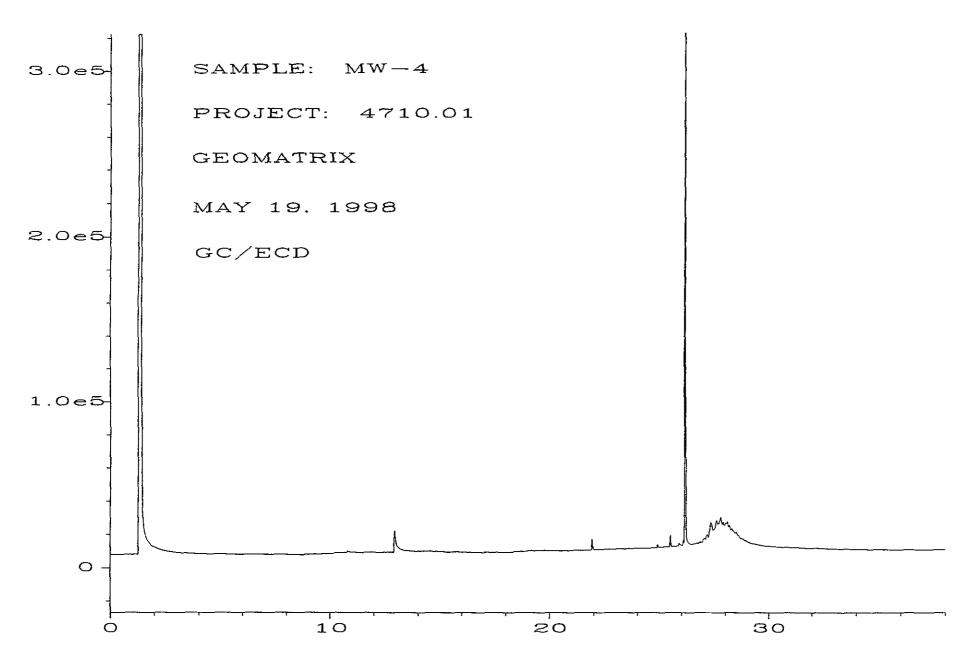
The medium boiling compounds appeared as a pattern of peaks eluting from n-C9 to n-C26. A dominant pattern of n-alkanes was not seen. The medium boiling material appears to have undergone evaropative and biological degradation.

The large peak seen near 25 minutes on the GC/FID trace is pentacosane, added as a quality assurance check for this GC analysis. There is a second surrogate present that is seen on the GC/ECD trace at about 26 minutes which is dibutyl chlorendate.

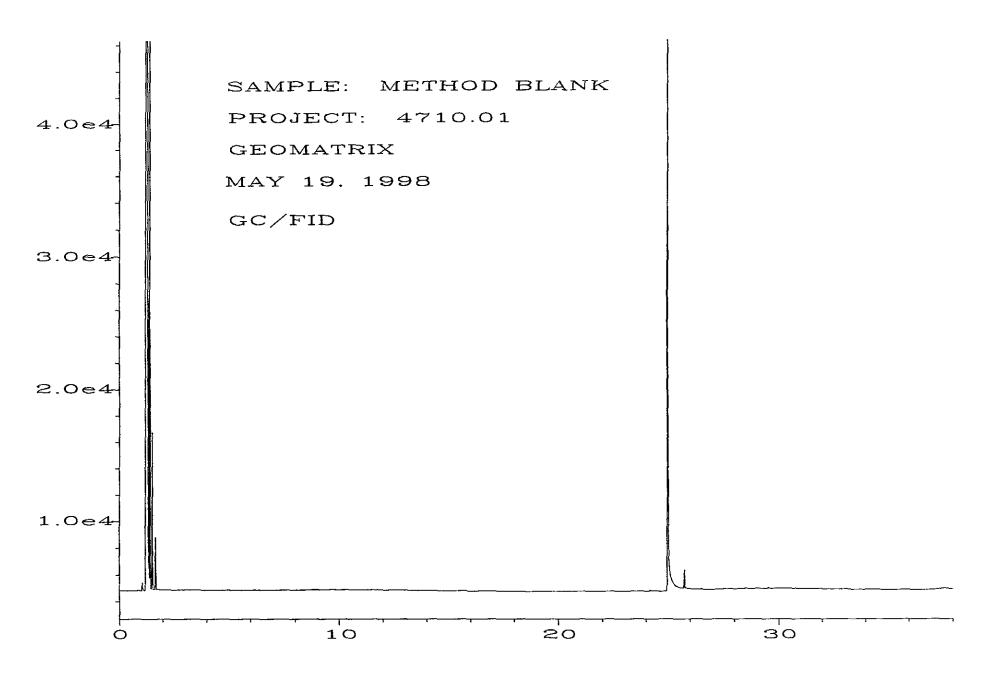
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Company: Friedray & Bross	Сотрапу		Company			100 Pine Street 10th Floor San Francisco California 94711 445 434 9400



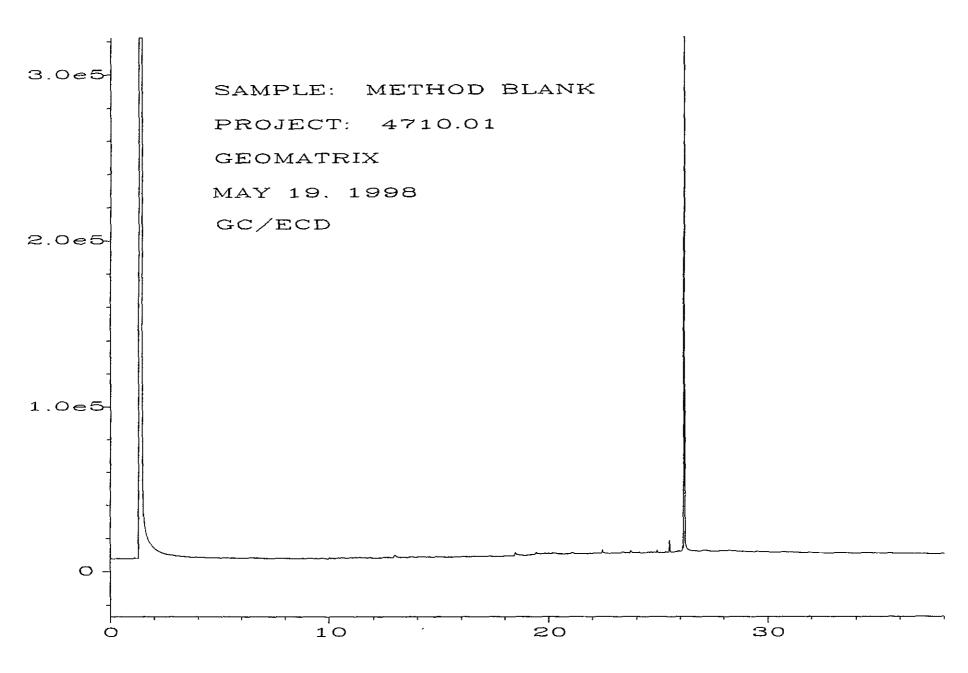
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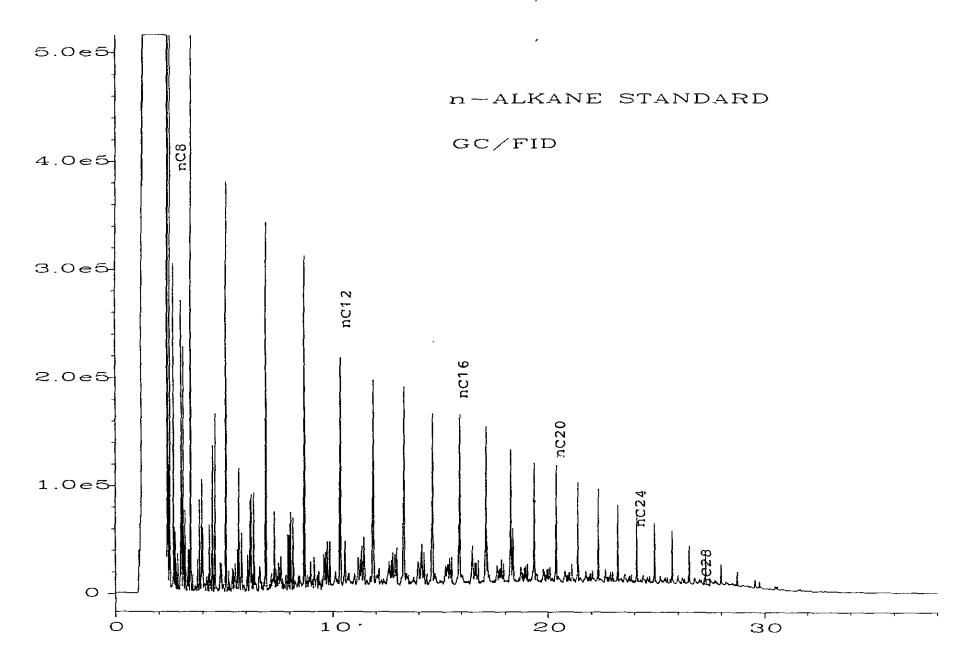
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### APPENDIX G

# ESTIMATION OF CHEMICAL EMISSIONS AND AMBIENT AND INDOOR AIR CONCENTRATIONS



## APPENDIX G

# ESTIMATION OF CHEMICAL EMISSIONS AND AMBIENT AND INDOOR AIR CONCENTRATIONS

The vapor emission rates of volatile chemicals of potential concern (COPCs) in soil were estimated using the VOC emissions model provided in the Preliminary Endangerment Assessment (PEA) guidance manual (Cal-EPA, 1994a). In addition, the VOC emissions model was modified to accommodate emissions from groundwater. This model is a simple screening model that assumes that all phases are in equilibrium (vapor, liquid, solid phases) and that the source remains constant over time (i.e., does not account for source depletion). The estimated emission rates calculated for soil and groundwater were used to calculate indoor and ambient air concentrations using a box model. The box model is a simple mass balance equation that is based on the concept of a theoretically enclosed space over the area of interest. These models are described in the following paragraphs.

## Soil Vapor Emissions

The VOC emissions model is not considered valid if "free phase" chemicals are present in the soil matrix. The presence of free phase chemicals can be assessed by comparing the concentration of the volatile COPCs in soil to the soil saturation concentration ( $C_{sat}$ ), which is based on the adsorptive limit of the soil and the theoretical dissolution limit of each chemical in the available soil moisture. When the concentration in soil is greater than or equal to the  $C_{sat}$ , free phase material is considered to be present. The  $C_{sat}$  can be calculated using the equation presented in the U.S. EPA Soil Screening Guidance (U.S. EPA, 1996) and the default parameters presented in the PEA guidance manual (Cal-EPA, 1994a). The  $C_{sat}$  was calculated for each volatile COPC using the following equation; chemical-specific input parameters, including their source and rationale, are provided in Table 6 of the main text.

$$C_{sat} = \frac{S}{\rho b} (Kd \times \rho b + Pw + H' \times Pa)$$

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Where:

 $C_{sat}$  = Soil saturation concentration for volatile COPCs ( $mg_{chemical}/kg_{soil}$ )

S = Chemical-specific solubility in water  $(mg_{chemical}/L_{water}; chemical-specific)$ 

ρb = Dry soil bulk density (1.5 kg_{soil}/L_{soil}; Cal-EPA, 1994a)

Kd = Soil-water partition coefficient [cm 3 _{water}/g_{soil}; equal to the chemical-specific organic carbon partition coefficient (Koc; L_{water}/kg_{soil} or cm 3 _{water}/g_{soil} chemical-specific) multiplied by the fraction of organic carbon in soil (foc; 0.02; Cal-EPA, 1994a)].

Pw = Water-filled soil porosity (0.15 cm³_{water}/ cm³_{soil}; Cal-EPA, 1994a)

H' = Dimensionless Henry's Law Constant [cm³_{water}/cm³_{air}; equal to the Henry's Law Constant (chemical-specific) multiplied by 41 (conversion factor) Cal-EPA, 1994a]

Pa = Air-filled soil porosity (0.284 cm³_{air}/ cm³_{soil}; Cal-EPA, 1994a)

As shown in the attached calculation spreadsheets, the concentrations of COPCs in soil are less than the Csat value. Therefore, use of the VOC emissions model is valid for concentrations in soil at the site.

The VOC emissions model uses chemical-specific physical characteristics (e.g., diffusivity in air) and assumptions regarding soil properties at the site (e.g., air-filled porosity) to predict the emission rate of volatile chemicals at the soil surface. The emission rate for each volatile COPC in soil was calculated using the following equation; chemical-specific parameters are presented in Table R-3 of the main text.

$$Ei = \frac{2 \times A \times Dei \times Pa \times Kas \times Cs \times 0.001 \, kg \, / \, g}{\sqrt{\pi \times \alpha \times T}}$$

Where:

Ei = Emission rate over area of interest  $(mg_{chemical}/sec)$ 

A = Emission area (12,000,000 cm²; equivalent to 1200 m²; site-specific)

Dei = Chemical-specific effective diffusivity [cm³/cm-sec.; calculated as follows: Di x (Pa³.333/Pt²); where Di is the chemical-specific diffusivity in air (cm²/sec), Pa is equal to the air-filled porosity, and Pt equals the total soil porosity]

Pa = Air-filled soil porosity (0.284 cm³_{air}/ cm³_{soil}; Cal-EPA, 1994a)

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Pt = Total porosity (0.434 cm³_{water}/ cm³_{soil}; Cal-EPA, 1994a)

Kas = Soil/air partition coefficient [g_{soil}/cm³_{air}; calculated as follows: (H/Kd) x 41; where H is the Henry's Law Constant (chemical specific) and Kd is soilwater partition coefficient (defined above)]

Cs = Bulk soil concentration of COPC  $(mg_{chemical}/kg_{soil})$ 

 $\pi = 3.14$ 

 $\alpha = \text{cm}^3/\text{cm-sec}$ ; calculated as follows:  $\alpha = \frac{(\text{Dei} \times \text{Pa})}{\text{Pa} + [\rho \times (1 - \text{Pa})/\text{Kas}]}$ 

 $\rho$  = Particle density (2.65  $g_{soil}/cm_{soil}^3$ ; Cal-EPA, 1994a)

T = Exposure duration (sec; equal to 1 year for noncarcinogenic health risks and 25 years for carcinogenic health risks)

## **Groundwater Vapor Emissions**

As presented in the PEA guidance manual, the VOC emissions model estimates vapor emission rates from volatile chemicals in soil. For purposes of this evaluation, the VOC emissions model was modified to estimate groundwater vapor emissions by replacing the components of the model that estimate the concentration in soil vapor based on a soil concentration (i.e., measured soil concentration and soil-to-air partitioning coefficient) with components that estimate the concentration in soil vapor based on a groundwater concentration (i.e., measured groundwater concentration and Henry's Law Constant). The emission rate for each volatile COPC in groundwater was calculated using the following equation; the chemical-specific parameters are presented in Table 6 of the main text.

$$Ei = \frac{2 \times A \times Dei \times Pa \times H' \times Cw \times 0.001 L_{water}/cm^{3}_{soil}}{\sqrt{\pi \times \alpha \times T}}$$

Where:

Ei = Emission rate over area of interest (mg_{chemical}/sec)

A = Emission area (12,000,000 cm²; equivalent to 1200 m²; site-specific)

Dei = Chemical-specific effective diffusivity [cm³/cm-sec.; calculated as follows: Di x (Pa³ 333/Pt²); where Di is the chemical-specific diffusivity in air

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(cm²/sec) and Pa is equal to the air-filled porosity and Pt equals the total soil porosity]

Pa = Air-filled soil porosity (0.284 cm³_{air}/ cm³_{soil}; Cal-EPA, 1994a)

Pt = Total porosity (0.434 cm³_{pore space}/ cm³_{soil}; Cal-EPA, 1994a)

H' = Dimensionless Henry's Law Constant [(cm³_{water}/ cm³_{air}) equal to the Henry's Law Constant (chemical-specific) multiplied by 41 (conversion factor; Cal-EPA, 1994a).

Cw = Groundwater concentration of COPC  $(mg_{chemical}/L_{water})$ 

 $\pi = 3.14$ 

 $\alpha = \text{cm}^3/\text{cm-sec}$ ; calculated as follows:  $\alpha = \frac{(\text{Dei} \times \text{Pa})}{\text{Pa} + [\rho \times (1 - \text{Pa})/\text{Kas}]}$ 

 $K_{as}$  = Soil/air partition coefficient [ $g_{soil}$ /cm³ ar; calculated as follows: (H'/Kd) where Kd is soil-water partition coefficient]

 $K_d$  = Soil-water partition coefficient [cm³_{water}/g_{soil}; equal to the chemical-specific organic carbon partition coefficient (Koc; L_{water}/kg_{soil} or cm³_{water}/g_{soil} chemical-specific) multiplied by the fraction of organic carbon in soil [foc; 0.02 unitless; Cal-EPA, 1994a])].

 $\rho$  = Particle density (2.65  $g_{soil}/cm_{soil}^3$ ; Cal-EPA, 1994a)

T = Exposure interval (sec; equal to 1 year for noncarcinogenic health risks and 25 years for carcinogenic health risks)

# Indoor Air Concentrations

The box model is typically used to estimate ambient air concentrations assuming vapors enter the box from the subsurface and are removed from the box by the wind. For purposes of this evaluation, the box model was modified to estimate indoor air concentrations by assuming that vapors enter the box via emissions through the cracks in the floor and are removed by ventilation. With these modifications, indoor air concentrations were calculated using the following equation, which was taken from the U.S. EPA's "Air/Superfund National Technical Guidance Study Series Assessing Potential Indoor Air Impacts at Superfund Sites" (1992); the calculations are presented in the attached calculation spreadsheets.

$$Cin = \frac{Ei \times F}{ACH/CF \times V}$$



## Where:

Cin = Indoor air concentration  $(mg/m^3)$ 

Ei = Emission rate from groundwater (mg/sec; calculated previously)

F = Fraction of floor area that is cracked (0.001 unitless; upper end of range for a house [Grimsrud et al., 1983])

ACH = Air exchanges per hour (1.5 hr⁻¹, U.S EPA, 1992; upper end of the range reported for a house)

CF = Conversion factor (3600 sec/hr)

V = Volume of air in building (3600 m³; site-specific)



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n-Butylbenzene	1 2E+07	0.064	0.284	0.434	5 12E-03	1 25E-03	2 48E+03	0 02	1 03E-03	6.5	2 65	790E-07	3 15E+07	2 65E-02	1	i	
sec-Bussibenzene	1 2E+α7	0.064	0.284	11 434	5 L2E-03	1 14E-02	891E+02	0.02	2 62E-u2	6	2.45	2.00E-05	3 LSE+07	t 23E-ut	1	}	
I I-Dichloroethane	NA	, NA	0 284	0 434	NA.	NA NA	NA NA	- 002	<u>NA</u>	NA	2 65	NA	3 15E+07	NA	+	1	
cis-1 2-Dichloroethene trans-1.2-Dichloroethene	NA NA	NA NA	0 284	0 434	NA NA	i NA i NA	NA NA	0 02	NA	NA NA	2 65 2 65	NA NA	3 15E+07	NA NA	1	+	
Ethylbenzene	1 2E+07	NA 1 0 075	0 284	0 434	6 00E-03	7 88E-03	3 63E+02	- 0 02	4 45E-02	0 03	2 65	3 97E-05	3 ISE+07	87tE-04	1		
Isopropylben/ene	1 2E+07	0 068	0 284	0 434	5 44E-03	5 58E-03	2 82E+03	0 02	4 06E-03	4	2 65	3 30E-06	3 15E+07	3 33E-02	i - 1	E I	ļ
Isopropy lialuene	1 2E+07	0.068	0 284	0 434	5 44E-03	1 10E-02	2 41E+03	0 02	9 36E-03	03	2 65	7 61E-06	3 15E+07	3 79E-03	<b>}</b>	}	ı
Naphthalene	1 2E+07	11 (159	0 284	0 434	4 72E-03	4 83E-04	2 00E+03	002	4 95E-04	51	2 65	3 50E-07	3 15E+07	138E-02	1 1		
n-Propylbenzene	1 2E+07	0.068	0 284	0 434	5 44E-03	1 08E-02	7 41E+02	0.02	2 99E-02	59	2 65	2 42E-05	3 15E+07	33E-01		1	
Tetrachloroethene	NA	NA	0 284	U 434	NA	NA	NA.	0 02	NA.	NA	2 65	NA	3 15E+07	NA	1 1	į	j
Toluene	NA.	NA	0 284	0 434	NA	NA	NA.	0 02	NA.	NA	2 65	NA	3 15E+07	NA	1 1	İ	
Trichloroethene	NA.	NA NA	0 284	0 434	NA	NA NA	NA.	0 02	NA	NA	2 65	NA	3 13E+07	NA	1 1	1	
1 2 4-Tranethylbenzene	1 2E+07	0 068	0 284	Ö 434	5 44E-03	5 70E-03	3 72E+03	0 02	3 15E-03	19	2 65	2 56E-06	3 15E+07	139E-02		i	
1,3 5-Trimethylbenzene	1 2E+07	0 068	0 284	0 434	5 44E-03	6 73E-03	6.618+02	0 02	2 09E-02	14	2 65	1 69E-05	3 15E+07	2 64E-02	! !	;	
Vmyl Chloride	NA	NA.	0 284	U 434	NA.	NA NA	NA.	0 02	NA NA	NA	2 65	NA NA	3 15E+07	NA	i i	ĺ	
Xylene	1 2E+07	0.087	0 284	0 434	6 96E-03	6 73E-03	3 %6E+02	0 02	3 57E-02	11	2 65	3 70E-05	3 15E+07	3 08E-02	1	ĺ	
ł	ļ	}	I	}			-	]		ì	i		ı İ		}	}	
Step 3 Calculate Indoor A	br Concentrations	ł	i	i		[			[		j		i j		] [	1	;
i	1		1	ļ			İ				1		į			ĺ	
	C2n = (E1 * F)/((ACH/	CF) * V)					1		ļ.		ì						
	1	į	1			İ	j .				i		1		1 1	ļ	
ł	ļ	ļ		Volume of Air in		indoor Air	}			1	}		i 1		1	}	
1	Emission Rate (Ei)	Frem of Flr	Rate (ACH)	Bldg (V)	Conversion	Conc (Cin)		1			1				1	1	
Chemical	(mg/sec)	that is Crkd (F)	(ltr)*1	(m³)	Factor (sec/hr)	(mg/m²)			!		1		:		] ]		
l	1	1	1					·					[		1 1	ļ	
n-Buty Ibenzenc	2 65E-02	0.001	1.5	3600	3600	1 76E-05									1	Į	
sec-Butylbenzene	t 23E-01	0.001	1.5	3600	3600	8 22E-05		· ·		}	1				1	į	
1.1-Dichloroethanc cis-1 2-Dichloroethene	NA NA	0 001	15	3600 3600	3600 3600	NA NA		l –			1				1 1	-	
trans-1 2-Dichlorocthene	NA NA	1000	15	3600	3600	NA NA		ļ <del></del> -							+ 1	-	-
Ethylbenzene	8 71E-04	0.001	15	3600	3600	5 80E-07	1				1		ł		! !		
Isopropy lbenzene	3 33E-02	0.001	15	3600	3600	2 22E-05							ł		f - 1	ł	İ
Isopropyltoluene	3 79E-03	0 001	15	3600	3600	2 53E-06							1		1 1	1	
Naphthalene	1 38E-02	0.001	1.5	3600	3600	9 20E-06			<del></del> -		†		. [		1 1	†	
n-Propylbonzone	t 33E-01	0.001	15	3600	3600	8 90E-05	-						†		1	-	
Tetrachloroethene	NA	0.001	1.5	3600	3600	NA						: f	:		1	ļ	
Foluenc	NA	0.001	1.5	3600	3600	NA						'	i		j †	1	
Trichloroethene	NA.	0.001	1.5	3600	3600	NA	1						† 1		; 1	1	İ
1,2,4-Trimethy lbenzenc	1 39E-02	0.001	15	3600	3600	9 28E-06	_	Γ 1	Γ			İ	1				
1.3.5-Trimethy benzene	2 64E-02	0.001	15	3600	3600	1 76E-05	-						į		į į	1	
Vary1 Chlorado	NA.	0.001	15	3600	3600	NA						i	[		1	į	j
Xylene	3 08E-02	0.001	1.5	3600	3600	2 05E-05						[	ĺ			[	
	·		<u> </u>														



Human Health Risk Asses	Ameni			,					<del></del>			<del></del>					$\overline{}$
Ris Properts		•				•				:							
Emersydic, California		•							!	ļ				`	•		<b>\</b>
								•	į					,			
Current and Future On-5	otte Commercial Work	er: Reasonable Mas	imum Exposori	(RME) Scenario			;	1				•				I	- 1
Initioar Air Concentration	is for Chemicals in Gr	oundwarter-Noncar	cinogenic and C	arcinogenic Health	Risks			! [	! !							· .	
Step 1 Estimate Llux from				:			;		1	:					1	;	1
					2 *A* Dei * Pa * C	w . H CL)/(≤ . ız	ć, £λ,ά 2		ļ		:	;			1		i
				Where		i	H' = H * 41		Į	Į.	Ì	, .		į.	1		ŀ
	•			pa = (Dei * Pa)/(Pa Dei = Di * (Pa ^{3.33}			Kd = Koc foc		1	!	:	!		i	1		
1	•	1		Kas = (H/Kd)*41	/na:) !	1	MI = NOC-10¢	-	1	}	}	1		i	ì	j	h
			;	(13/10) 41			;	İ	†	1	Ì			,		!	' I
		•	•	1		i	1		1	1	Soil-Water	'		•	1	;	- 1
	Max GW	1	1	i		İ	Ì	Dimensionless	Organic Carbon	Fraction of	Partition	Soil-Air			1	'	
}	Conc	Emissions	Diffusivity	Air-Filled Soil	Total	Effective	Henry's Law	Henry's Law	Partition	Organic	Coefficia	Parution	Particle	1	1	Conversion	
ĺ	(Cu)	Arca (A)	is Air (Di)	Porosity (Pa)	Porosity (P1)	Diffusivity (Dei)		Constant (H')	Coefficient (Koc)	Carbon (foc)	, (Kd)	Coefficat (Kas)	* - * -	Alpha (a)	Time (T)	Factor (CF)	Raic (E,)
Chemical	(mg/l)	(cm²)	(cm²/scc)	(cm³/cm³)	(cm /cm )	(cm /cm-sec)	(atm-m /mole)	(cm²/cm²)	(ml/g)	(Unitless)	(cm /g)	(g/cm²)	(g/cm²)	(cm /cm-sec)	(scc)	(l/cm )	(mg/sec)
n Butylbenzene	ND	1 2E+07	0 064	11 284	0 434	5 128-03	1 25E-113	5 13E-02	2 48 €+03	0.02	49 66	1 032E-03	265	7 90E-07	3 15E+07	1 00E-03	NA 5.tog oz
sec-Butylbenzene	0.015	1 2E+07	0.064	0 284	0 434 0 434	5 12E-03 5 93E-03	1 14E-02	4 67E-01 2 30E-01	8 91 E+02 3 16 E+01	0 02 0 02	17 82 0 63	2 623E-02 3 646E-03	2 65 2 65	2 00E-05 3 07E-04	3 15E+07	1 00E-03 1 00E-03	5 49E-03 6 41E-03
1 1-Dichloroethane cis-1,2-Dichloroethene	0 12 0 14	1 2E+07 1 2E+07	0 074	0 284	0 434	5 89E-03	5 62E-03 4 08E-03	1 67E-01	3 55E+01	0 02	0.71	2 356E-01	2 65	2 00E-04	3 15E+07	1 00E-03	6 67E-03
trans-1 2-Dichloroethene	0018	1 2E+07	0.071	0 284	0 434	5 65E-03	9 38E-03	3 85E-01	5 25E+01	0 02	1 105	3 663E-01	2 65	2 94E-04	3 15E+07	1 00E-03	1 56E-03
Ethythenzene	0.007	1 2E+07	0 075	0 284	0 434	6 00E-03	7 88E-03	3 23E-01	3 63E+02	0.02	7 26	4 456E-02	2 65	3 97E-05	3 15E+07	1 00E-03	1 17E-03
Isopropy lbenzene	8 816	1 2E+07	0.068	0 284	11 434	5 44E-03	5 58E-03	1 29E-01	2 82E+03	4:42	56 36	4 059E-03	2 65	3 30E-06	3 15E+07	1 00E-03	7 50E-03
Isopropy Itoluene	ND	1 2E+07	0.064	0 284	0.434	5 12E-03	1 10E-02	4 51E-01	2 41E+03	0.02	48 20	9 357E-03	2 65	7 16E-06	3 15E+07	1 00E-03	NA
Naphthalene	0.005	I 2E+07	0.059	0 284	0 434	4 72E-03	4 83E-04	1 98E-02	2 00€+03	0 02	40 00	4 951E-04	2 65	3 50E-07	3 15E+07	1 00E-03	5 41E-04
n-Propy Ibenzene	0.009	1 2E+07	0 068	0 284	0 434	5 44E-03	1 08E-02	4 43E-01	7 41E+02	0.02	14 83	2 987E-02	2 65	2 42E-05	3 15E+07	1 00E-03 1 00E-03	3 02E-03 2 51E-02
Tetrachloroothene	0 12 0 25	1 2E+07	0 072	0 284	0 434 0 434	5 76E-03 6 96E-03	1 84E-02 6 64E-03	7 54E-01 2 72E-01	1 55E+02 1 82E+02	0 02 0 02	3 10 3 64	2 434E-01 7 479E-02	2 65 2 65	2 02E-04 7 70E-05	3 15E+07	1 00E-03	3 70E-02
Frichloroethene	019	1 2E+07	0.079	0 284	0 434	6 32E-03	1 03E-02	4 22E-01	1 66E+02	0 02	3 32	1 272E-01	2 65	1 18E-04	3 15E+07	1 00E-03	3 20E-02
1,2,4-Trimethylbenzene	0 12	J 2E+07	0.068	0 284	0 434	5 44E-113	5 70E-03	2 34E-01	3 72E+03	0 02	74 31	3 145E-03	2 65	2 56E-06	3 15E+07	1 00E-03	6 53E-02
1.3.5-Tramethy lbenzene	0.011	1 2E+07	0 068	0 284	0 434	5 44E-03	6,73E-03	2 76E-01	661E+02	0.02	13 21	2 088E-02	2 65	1 69E-05	3 158+07	1 00E-03	2 75E-03
Vinv1 Chloride	0.11	1 2E+07	0 106	0 284	0 434	8 48E-03	2 70E-02	Î HE+(X)	1 86E+01	0.02	0.37	2 976E+00	2 65	2 61E-03	3 15E+07	1 00E-03	1 38E-02
Xylene	0 075	l 2E+117	0 087	0 284	0 434	6 96E-03	6 73E-03	2 76E-01	3 86E+02	0 02	7 73	3 571E-02	2 65	3 70E-05	3 15E+07	1 (X)E-03	1 62E-02
	t		į.	C VEASIMAC	LI (CENSIA)						}				1		
Step 2 Calculate Indoor A	ir Concentrations-None	aremogenic ricaim n 	(SVEY	$C_{\mu} = (E_i * F)/((AC_i)$	n/Cr)*Y)	}	•		<del> </del>	1	+				1		
			Air Exchange	Volume of Air in		Indoor Air	†		-	ł	1	1 1		ţ		l l	ļ
	Emission Rate (E _i )	Frein of Fir	Rate (ACH)	Bldg (V)	Conversion	Cone (C _{sn} )	1										1
Chemical	(mg/scc)	that is Crkd (F)	(hr)-i	(m')	Factor (sec/hr)	(mg/m³)			İ	ļ		1		}			
	()		1,	\/		1 (	f		-	İ				i I	Ì		
n-Butylbenzene	NA.	0 001	15	36190	3600	NA.	1		<del> </del> -	t	ţ	t l		1	Į.	}	[
sec-Buty lbenzene	5 49E-03	100 0	15	3600	3600	3 66E-06	j			1	ŀ	1		İ	İ		
1 I-Dichlorocthane	6 41E-03	0 001	15	3600	3600	4 28E-06	] -		Γ.		]	1		1		] ;	Į
cis-1,2-Dichloroethene	6 67E-03	0.001	1.5	3600	3600	4 45E-06	]		]_		]	į j		1	ļ		1
trans-1 2-Dichloroethene	1 56E-03	0 001	1.5	3600	3600	1 04E-06			Į	ļ	1	į i		Į	Į		1
Ethylbenzene	1 47E-03	0.001	1.5	3600	3600	9 83E-07	İ	-	-	1	[						1
Isopropylbenzene	7 50E-03	0 001	15	3600	3600	5 00E-06		—	<del> </del> -	į	1			Į	-		j
isopropy Itoluene Naphthalene	NA 5 41E-04	0 001	15	3600 3600	3600 3600	NA 3 61E-07			<del> </del> -	}	į			1	-		l
n-Propylbenzene	3 02E-03	0.001	15	3600	3600	2 01E-06	1			†	†	† j		1	ţ		
Tetrachloroethene	2 51E-02	0 001	15	3600	3600	1 67E-05	1		<b>]</b> "		1	, )		1	1	† į	}
Tolsene	3 70E-02	6 001	15	3600	3600	2 46E-05	]	I	1	Ī		] ]		i	į		ł
Trichloroethene	3 20E-02	0 001	15	3600	3600	2 13E-05	1		L			]		1	1	j	- 1
1 2,4- Francilis Ibenzenc	6 53E-02	100 0	1.5	3600	3600	4 35E-05	1 1		-	· I	ļ			1			
1,3,5-3 rimethy lbenzene	2 75E-03	0.001	15	36(X)	3600	1 83E-06	1		}-	}	ĺ	} i		}	}	1	ŀ
Vinyl Chloride	1 38E-02 1 62E-02	0 001	15	3600 3600	3600 3600	9 22E-06 1 08E-05				+		1					ŀ
Xylene	1 020-02	0 001	15	30,00	3000	1 001:-03	{ i	·	·			! !		į	<u> </u>		- 1
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Human Health Risk Asse	asment		•											•		:	•
Rix Property				: .		'	: .		Į.								
Emeryville, California								:	:		,	•		-			
C			p	ADMES C				<u>.</u>	1	1	•						:
Current and Future On-S	site Commercial Work	er: Reasonable (12)	mann Exposure	(KWE) SCHILLE		•	:			•							
Step 1 Ashmate Flux from	n Cremedwaler-Careino	genie Health Risks		Emission Rate = (	2 *A* Dei * Pa * C	'x.*H' * CF)/(# * «	* T)^0,5	 	İ	1					•		
.,		<b>4</b>		,					i					•			_
		_		Where				•	i								
			1	ta = (Det * Pa)/(Pa			H' = H * 41		1		;	,		;	+		:
1				Det = Dt * (Pa 3.33	/Pt ² )		Kd = Koc*foc	Į	1								į l
			•	Kas = (H/Kd)*41						,		,		:	!		!
1					,		1		1					,	ļ	1	!
1	7			1		į	:		1	•	Soil-Water	1 - 1			1	i	!
	/ GW \	١				}		Dimensionless	Organic Carbon	Fraction of	Partition	Sou-Air		1	1		!
	Conc	Emissions	Diffusivity	Air-Filled Soil	Total	Effective	Henry's Law	Henry's Law	Partition	Organic	Coefficit	Partition	Particle			Conversion	
	(Cn)	Arca (A)	is Air (Di)	Porosity (Pa)	Porosity (Pt)	Diffusivity (Dei)		Constant (H")	Coefficient (Koc)	Carbon (foc)	(Kd)	Coefficht (Kas)		Alpha (a)	Time (T)	Factor (CF)	
( hemical	(mg/l)	(cm²)	(cm²/sec)	(cm'/cm')	(cm³/cm³)	(cm /cm-sec)	(aun-m /mole)	(cm³/cm³)	(m½g)	(Unitless)	(cm /g)	(g/cm ')	(g/cm ')	(cm'/cm-sec)	(scc)	(i/cm )	(mg/sec)
	ND	1.35,07	0.061	0.294	0 434	5 12E-03	1 25E-03	5 13E-02	2 48E+113	0 02	49 66	1 03E 403	2 65	7 90E-07	788E+08	[   LOUE-03	NA.
n-Butylbenzene sec-Butylbenzene	, 0:015	1 2E+07	0.064	0 284	0 434	5 12E-03	1 25E-03 1 14E-02	4 67E-01	891E+02	0.02	1782	2 62E-02	2 65	2 00E-05	7 88E+08	L 00E-03	NA 1 108-03
1.1-Dichloroethane	0.12	1 26+07	0.034	0.284	0.434	5 12E-03	1 62E-03	2.30E-01	3 (6E+0)	0.02	1163	3 65E-01	265	3 07E-04	788E+08	[ 00E-03	1 2xE-03
cis-1,2-Dichlorouthene	014	1 2E+07	0 074	0 284	0 434	5 89E-03	4 08E-03	1 67E-01	3 55E+01	0 02	071	2 36E-01	2 65	2 00E-04	7 88E+08	1 00E-03	1 33E-03
trans-1 2-Dichloroethene	0.018	1 2E+07	0 074	0 284	0 434	5 65E-03	9 38E-03	3 85E-01	5 25E+01	0.02	105	3 66E-01	2 65	2 94E-04	7 88E+08	1 00E-03	3 13E-04
Ethylbenzene	0 007	1 2E+07	0.075	0 284	0 434	6 00E-03	7 88E-03	3 23E-01	3 63E+02	0 02	7 26	4 45E-02	2 65	3 97E-05	7 88E+08	1 00E-03	2 95E-04
Isopropy Ibenzene	0.036	1 2E+07	0 068	0 284	0 434	5 44E-03	5 58E-03	2 29E-01	2 82E+03	0 02	56 36	4 06E-03	2 65	3 30E-06	7 88E+08	1 00E-03	1 50E-03
Isopropy Itoluene	ND	1 2E+07	0 064	0 284	0 434	5 12E-03	1 10E-02	4 51E-01	2 41E+03	0 02	48 20	9 36E-03	2 65	7 16E-06	788E+08	1 00E-03	NA
Naphthalene	0.005	1 2E+07	0 059	0 284	0 434	4 72E-03	4 83E-14	1 98E-02	2 (K)E+03	0.02	40 00	4 95E-04	2 65	3 50E-07	788E+08	t 00E-03	1 08E-04
n-Propylbenzene	0.009	1 2E+07	0.068	0 284	0 434	5 44E-03	1 08E-02	4 43E-01	7.41E+02	0.02	14 83	2.99E-02	2 65	2 42E-05	7 88E+08	L00E-03	6 03E-04
Tetrachloroethene	0 12	1 2E+07	0 072	0 284	0 434	5 76E-03	1 84E-02	7 54E-01	1 55E+02	0.02	3 10	2 43E-01	2 65	2 02E-04	788E+08	J 00E-03	5 02E-03
Toluene	0.25	1 2E+07	0.087	0.284	0 434	6 96E-03	6 64E-03	2 72E-01	1 82E+02	0 02	3 64	7 48E-02	2 65	7 70E-05	7 88E+08	1 00E-03	7 39E-03
Trichloroethene	0.19	1 2E+07	0 079	U 284	0 434	6 32E-03	1 03E-02	4 22E-01	1 66E+02	0 02	3 32	î 27E-UL	2 65	1 18E-04	788E+08	1 00E-03	6 39E-03
1.2.4-Tranethylbenzene	e 12	1 2E+07	0.068	0 284	(1434	5 44E-03	5 70E-03	2 34E-01	3 72E+03	0.02	7431	3 15E-03	2 65	2 56E-06	788E+08	1 00E-03	1 31E-02
1.3 5-Trunethylbenzene	110.0	[ 2E+07	0 068	0 284	0 434	5 44E-03	6 73E-03	2 76E-01	6 61E+02	0 02	13 21	2 09E-02	2 65	3 69E-05	788E+08	1 00E-03	5 49E-04
Vinyl Chloride	0.11	1 2E+07	0 106	0 284	0 434	8 48E-03	2 70E-02	1 11E+00	1 86E+01	0.02	0.37	2 98E+(X)	2 65	2 61E-03	788E+08	1 00E-03	2 77E-03
Xylene	0 075	1 2E+07	0.087	0 284	0 434	6 96E-03	6 73E-03	2 76E-01	3 86E+02	0 02	7 73	3 57E-02	2 65	3 70E-05	7 88E+08	1 00E-03	3 24E-03
	İ				-	į	_	u	<u> </u>	1	[			ļ			
Step 2 Calculate Indoor A	ir Concentrations (arci	magenic Health Risks		$C_m = (E_i + F)/((AC)$	H/CF)*V)	1	1					!					
	1			La	-			_				ļ				}	
	Emission		Air Exchange	Volume of Air in		Indoor Au	l					[					1
	Rate (E,)	Frem of Fir	Rate (ACH)	Bidg (V)	Conversion	Conc (C _m )									-		
Chemical	(mg/sec)	that is Crkd (F)	(lur)	(m')	Factor (scc/hr)	(mg/m³)	[		ļ. <u></u> .					1	[		
l	ļ.	Į	Į	į į		Į	(		ļ	Ę.	ļ	į į			ļ	ļ	(
n-Butylbenzene	NA	0.001	1.5	3600	3600	NA	1	<u>-</u>	<u> </u>	1		1			ļ		
sec-Butylbenzene	1 10E-03	0.001	1.5	3600	36110	7 33E-07			ļ	1	1			ļ		1	
I,I-Dichloroethane	t 28E-03	0.003	15	3600	3600	8 55E-07			<u></u>	1	1	1			1	1	[
cis-1,2-Dichloroethene	1 33E-03	0.001	15	3600	3600	8 89E-07	ļ .	<u>.</u> . <u>-</u>	<del> </del> -	1					1	+	
trans-1,2-Dichloroethene	3 13E-04	0 001	1.5	3600	3600	2 09E-07			ļ. <i>.</i>	1				1	-	1	
Ethylbenzene	2 95E-04	0 001	1.5	3600	3600	1 97E-07			<del> </del>	1		1		-	1	1	
Isopropy locn/ene	J 50E-03	0 001	. 15	3600	3600 3600	1.00E-06	!		<del> </del> -	l		F -		ł	}	1	<u> </u>
Isopropyltoluene	NA 1 me o t	0 001	15	3600	3600 3600	NA 7 22E-08	-					‡			1		
Naphthalene	1 08E-04 6 03E-04	100 0 100 0	15	3600 3600	3600	4 02E-07	ł		ļ ·			1			ł	1	
n-Propylbenzene Tetrachloroethene	5 02E-03	1	15	3600	3600	3 35E-06	†		ļ -	1		1 1			ł	1	;
Tolucac	7 39E-03	0 001	15	3600	3600	4 93E-06	<b>}</b>		<del>├</del> -	·	ļ	1		}	}	1	<b>\</b>
Trichloroethene	6 39E-03	1 0.001	15	3600	3600	4 93E-06 4 26E-06	-		<del> </del> -	1	ł	1		1	1	1	
1,2 4-Trimethylbenzene	1 31E-02	0.001	15	3600	3600	8 71E-06	1		<del> </del>	<u> </u>		1		t			
1,3 5-Trimethy loenzene	5 49E-04	0 001	15	3600	3600	3 66E-07	į ·		·			}		1	1		
Vinyl Chloride	2 77E-03	0 001	15	3600	36(8)	1 84E-06	<u> </u>	-	t	Ì	l I				1	! 	1
Xviene	3 24E-03	0.001	15	3600	3600	2 16E-06	ļ i		ţ			1 !			t	t	
	1 22.27.22										·····			· · · · · · · · · · · · · · · · · · ·			



# APPENDIX H

EQUATIONS USED FOR ESTIMATING PHYSICAL CONSTANTS OF SELECTED CHEMICALS OF POTENTIAL CONCERN



### APPENDIX H

# EQUATIONS USED FOR ESTIMATING PHYSICAL CONSTANTS OF SELECTED CHEMICALS OF POTENTIAL CONCERN

The diffusivity in air coefficient (Di) was calculated for several chemicals of potential concern using the following equation from U.S. EPA's Superfund Exposure Assessment Manual (1988b), the calculations are shown in the attached calculation spreadsheet.

$$Di = \frac{0.001 \times T^{1.75} \sqrt{\frac{1}{MWi} + \frac{1}{MWa}}}{Pa[(\sum Vi)^{1/3} + (\sum Va)^{1/3}]^2}$$

Where:

Di = Diffusivity in air  $(cm^2/sec)$ 

T = Absolute temperature (°Kelvin)

MWi = Molecular weight of toxic substance (g/mole)

MWa = Molecular weight of air (g/mole)

Pa = Absolute pressure (atm.)

 $\sum Vi$  = Molecular diffusion volumes of toxic substance (cm³/mole)

 $\sum Va = Molecular diffusion volumes of air (cm³/mole)$ 

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ssment		-			1			
			:	}				
·		•	†	 	;			•
in Air Coefficier	nts Using Equat	ion in U.S. EPA G	uidance Manus	ls (Superfund	Exposure Asses	sment Manual)		•
C atom	16.5	•			į	:	1	
H atom	1.98	•	$Di = (0.001T^{4})$	1.75*(1/MWi +	- 1/MWa)^0.5) /	(Pa * [(Vi)^1/3 + (Va)	· ^1/3)]^2)	Í
O atom	5.48	t 	1		,			r 1
'N atom	5.69		,	1	• 	!	ı	
Cl atom	19.5	• I	i	İ	i	1	•	
Aromatic	-20.2		1		1	•	İ	
Heterocyc	-20.2						1	
MWi	# Carbons	# Hydrogens	# Oxygen	# Nitrogen	# Chlorine	# Aromatic Rings	# Heterocyclic	Vi
(g/mole)	İ			ļ			İ	(cm³/mole)
134.22	10	14	0	0	0	1	0	172.5
134.22	10	14	0	0	0	1	0	172.5
120.19	9	12	0	0	0	1	0	152.1
134.22	10	14	0	0	0	1	0	172.5
120.19	9	12	0	0	0	1	0	152.1
120.19	9	12	0	0	0	1	0	152.1
120.19	9	12	0	0	0	1	0	152.1
1						j		Ì
						ļ		r [
Constant	Temp	MWi	MWa	Pa	Vi	Va	Di	7
_		(g/mole)	(g/mole)	(atm)	(cm³/mole)	(cm³/mole)	(cm ² /sec)	
0.001				1				
: !			1	† - i	1	1	1	1
1 1			l	1	!	!	0.068	-
			1	<u>1</u>		Į.	j .	1
		_	1	1	1	l .	0.068	
1 . 1			L	<u> </u>	1	1	1	
! 1				i i	1	i		] 
	C atom H atom O atom N atom Cl atom Aromatic Heterocyc  MWi (g/mole) 134.22 120.19 134.22 120.19 120.19 120.19	C atom 16.5 H atom 1.98 O atom 5.48 N atom 5.69 Cl atom 19.5 Aromatic -20.2 Heterocyc -20.2  MWi #Carbons (g/mole) 134.22 10 134.22 10 120.19 9 134.22 10 120.19 9 120.19 9 120.19 9 120.19 9 120.19 9 120.19 9 0.001 298 0.001 298 0.001 298 0.001 298 0.001 298 0.001 298	C atom 16.5 H atom 1.98 O atom 5.48 N atom 5.69 Cl atom 19.5 Aromatic -20.2 Heterocyc -20.2  MWi # Carbons # Hydrogens  (g/mole) 134.22 10 14 120.19 9 12 134.22 10 14 120.19 9 12 120.19 9 12 120.19 9 12 120.19 9 12 120.19 9 12 120.19 9 12 120.19 9 12 120.19 9 12 120.19 9 12 120.19 9 12 120.19 9 12 120.19 9 12 120.19 9 12 120.19 9 12 120.19 9 12	Catom   16.5   Hatom   1.98   Di = (0.00117')   Oatom   5.48   Natom   5.69   Clatom   19.5     Aromatic   -20.2   Heterocyc   -20.2     MWi	in Air Coefficients Using Equation in U.S. EPA Guidance Manuals (Superfund  C atom 16.5 H atom 1.98 O atom 5.48 N atom 5.69 Cl atom 19.5 Aromatic -20.2 Heterocyc -20.2  MWi #Carbons #Hydrogens #Oxygen #Nitrogen  (g/mole) 134.22 10 14 0 0 134.22 10 14 0 0 134.22 10 14 0 0 120.19 9 12 0 0 134.22 10 14 0 0 120.19 9 12 0 0 134.22 10 14 0 0 120.19 9 12 0 0 120.19 9 12 0 0 120.19 9 12 0 0 120.19 9 12 0 0 120.19 9 12 0 0 120.19 9 12 0 0 120.19 9 12 0 0 120.19 9 12 0 0 120.19 9 12 0 0 120.19 9 12 0 0 120.19 9 12 0 0 120.19 9 12 0 0 120.19 9 12 0 0 120.19 9 12 0 0 120.19 9 12 0 0 120.19 9 12 0 0 120.19 9 12 0 0 120.19 9 12 0 0 120.19 9 12 0 0 120.19 9 12 0 0 120.19 9 12 0 0 120.19 9 12 0 0 120.19 9 12 0 0 120.19 9 12 0 0 120.19 9 12 0 0 120.19 9 12 0 0 120.19 9 12 0 0 120.19 9 12 0 0 120.19 9 12 0 0 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.19 120.1	Table   Catom   16.5     Hatom   1.98   Di = (0.001T^\1.75*(1/MWi + 1/MWa)^\0.5)/   Oatom   5.48   Natom   5.69     Clatom   19.5   Aromatic   -20.2     Heterocyc   -20.2     MWi	in Air Coefficients Using Equation in U.S. EPA Guidance Manuals (Superfund Exposure Assessment Manual)  C atom	Catom   16.5



# APPENDIX I

HUMAN HEALTH RISK ASSESSMENT CALCULATION SPREADSHEETS



Human Health Risk Asses	ssment							. :		
Rix Property										
Emeryville, California								•		
Current and Future On-S	ite Commercial Worker:	: Reasonable M	aximum Exposur	e (RME) Scenari	jo	•		:		
				D-APPACDV/DW	** A TT\	Hazard Quotient	A A D D /D (T)			
Inhalation of Soil Vapors	4		AADD = (Cin+)	Ra*EF*ED)/(BW	*A1)	Hazara Quotieni	- AADD/KIDI		:	
	Indoor Air (Soil) Concentration (Cin)	Inhalation Rate (IR _a )	Exposure Frequency (EF)	Exposure Duration (ED)	Body Weight (BW)	Averaging Time (AT)	Annual Average Daily Dose (AADD)	Inhalation Reference Dose (RfDi)	Hazard Quotient	
Chemical	(mg/m³)	(m³/day)	(days/year)	(years)	(kg)	(days)	(mg/kg-day)	(mg/kg-day)		
n-Butylbenzene	1.76E-05	20	250	25	70	9125	3.45E-06	1.0E-02	3E-04	
sec-Butylbenzene	8.22E-05	20	250	25	70	9125	1.61E-05	1.0E-02	2E-03	
1,1-Dichloroethane	NA	20	250	25 25	70	9125	NA	1.0E-02	NA I	
cis-1,2-Dichloroethene	i NA	20	250	25	70	9125	NA	1.0E-02	NA	
trans-1,2-Dichloroethene	NA NA	20	250	25	70	9125	NA	1.0E-02	NA	
Ethylbenzene	5.80E-07	20	250	25	70	9125	1.14E-07	2.9E-01	4E-07	
Isopropylbenzene	2 22E-05	20	250	25	70	9125	4 34E-06	2.6E-03	2E-03	
Isopropyltoluene	2 53E-06	20	250	25	70	9125	4.95E-07	NA NA	NA ]	
Naphthalene	9.20E-06	20	250	25	70	9125	1 80E-06	4.0E-02	5E-05	
n-Propyibenzene	8.90E-05	20	250	25	70	9125	1 74E-05	1.0E-02	2E-03	
Tetrachloroethene	i NA	20	250	25	70	9125	NA	1.0E-02	NA	
Toluene	NA	20	250	2 <u>5</u> 2 <u>5</u>	70	9125	NA	1.0E-02	NA	
Trichloroethene	NA .	20	250	25	70	9125	NA	1.0E-02	NA	
1,2,4-Trimethylbenzene	9.28E-06	20	250	25	70	9125	1.82E-06	1.7E-03	1E-03	
1,3,5-Trimethylbenzene	1 76E-05	20	250	25	70	9125	3.45E-06	1.7E-03	2E-03	
Vinyl Chloride	NA	20	250	25	70	9125	NA	1 7E-03	NA	
Xylene	2.05E-05	20	250	25	70	9125	4.02E-06	2.0E-01	2E-05	
		•			-			Total	9E-03	
İ	\ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \		1		}	1		1		

### HQ-ECR



Human Health Risk Asses	ssment								,
Rix Property						•			
Emeryville, California	•							i	
Current and Future On-S	iite Commercial Worker	: Reasonable M	aximum Exposur	e (RME) Scenari	o				
Inhalation of Groundwate	er Vapors		AADD = (Cin*l	Ra*EF*ED)/(BW	*AT)	Hazard Quotient	= AADD/RfDi	,	;
	Indoor Air (Soil) Concentration (Cin)	Inhalation Rate (IR _a )	Exposure Frequency (EF)	Exposure Duration (ED)	Body Weight (BW)	Averaging Time (AT)	Annual Average Daily Dose (AADD)	Inhalation Reference Dose (RfDi)	Hazard Quotient
Chemical	(mg/m³)	(m³/day)	(days/year)	(years)	(kg)	(days)	(mg/kg-day)	(mg/kg-day)	!
n-Butylbenzene	NA :	20	250	1	70 70	9125	NA	1.00E-02	NA
sec-Butylbenzene	3 66E-06	20	250	1	70	9125	2.87E-08	1.00E-02	3E-06
1,1-Dichloroethane	4.28E-06	20	250	1	70	9125	3 35E-08	1.40E-01	2E-07
cis-1,2-Dichloroethene	4.45E-06	20	250	1	70	9125	3 48E-08	1 00E-02	3E-06
trans-1,2-Dichloroethene	1.04E-06	20	250	ı	70	9125	8.16E-09	2 00E-02	4E-07
Ethylbenzene	9.83E-07	20	250	1	70	9125	7 70E-09	2.90E-01	3E-08
Isopropylbenzene	5 00E-06	20	250	1	70	9125	3.92E-08	1 10E-01	4E-07
Isopropyltoluene	NA	20	250	1	70	9125	NA	NA	NA
Naphthalene	3 61E-07	20	250	1	70	9125	2.82E-09	8.60E-04	3E-06
n-Propylbenzene	2.01E-06	20	250	1	70	9125	1.57E-08	1.00E-02	2E-06
Tetrachloroethene	1.67E-05	20	250	i	70	9125	1 31E-07	1.10E-01	1E-06
Toluene	2.46E-05	20	250	j	70	9125	1.93E-07	1.10E-01	2E-06
Trichloroethene	2.13E-05	20	250	1	70 70	9125	1.67E-07	6.00E-03	3E-05
I,2,4-Trimethylbenzene	4.35E-05	$\bar{2}0$	250	Ī	70	9125	3.41E-07	1.70E-03	2E-04
1,3,5-Trimethylbenzene	1.83E-06	20	250	1	70	9125	1 43E-08	1.70E-03	8E-06
Vinyl Chloride	9.22E-06	20	250	Ì	70	9125	7.22E-08	NA NA	NA
Xylene	1.08E-05	20	250	i	70	9125	8.46E-08	2.00E-01	4E-07
		-	]			]		Total	3E-04



Human Health Risk Asses	sment								
Rix Property	•		•	:		•			•
Emeryville, California			1			•	ı	•	•
Current and Future On-S	ite Commercial Worker	: Reasonable M	aximum Exposur	e (RME) Scenar	o			· :	• •
Inhalation of Groundwate	er Vapors		LADD = (Cin*)	Ra*EF*ED)/(BW	*AT)	Excess Cancer F	Risk = LADD*SFi	; !	1
}	Indoor Air (Soil) Concentration (Cin)	Inhalation Rate (IR _a )	Exposure Frequency (EF)	Exposure Duration (ED)	Body Weight (BW)	Averaging Time (AT)	Lifetime Average Daily Dose (LADD)	Inhalation Slope Factors (SFi)	Excess Cancer Risk
Chemical	(mg/m³)	(m³/day)	(days/year)	(years)	(kg)	(days)	(mg/kg-day)	(mg/kg-day)	
n-Butylbenzene	NA	20	250	25	70	25550	NA	. NA	NA
sec-Butylbenzene	7.33E-07	20	250	25	70	25550	5 12E-08	NA ,	NA
1,1-Dichloroethane	8.55E-07	20	250	25	70	25550	5.98E-08	NA	NA
cis-1,2-Dichloroethene	8 89E-07	20	250	25	70	25550	6.21E-08	NA	NA
trans-1,2-Dichloroethene	2.09E-07	20	250	25	70	25550	1.46E-08	NA	NA
Ethylbenzene	1.97E-07	20	250	25	70	25550	1 37E-08	NA	NA
Isopropylbenzene	1 00E-06	20	250	25	70	25550	6.99E-08	NA	NA
Isopropyltoluene	, NA	20	250	25	70	25550	NA	NA	NA
Naphthalene	7.22E-08	20	250	25	70	25550	5.04E-09	NA	NA
n-Propylbenzene	4.02E-07	20	250	25	70	25550	2.81E-08	NA	NA
Tetrachloroethene	3.35E-06	20	250	25	70	25550	2 34E-07	2 10E-02	5E-09
Toluene	4.93E-06	20	250	25	70	25550	3 44E-07	NA	NA
Frichloroethene	4.26E-06	20	250	25	70	25550	2.98E-07	1.00E-02	3E-09
1,2,4-Trimethylbenzene	8 71E-06	20	250	25	70	25550	6.09E-07	NA	NA
1,3,5-Trimethylbenzene	3.66E-07	20	250	25	70	25550	2 56E-08	NA	NA /
Vinyl Chloride	1.84E-06	20	250	25	70	25550	1 29E-07	2 70E-01	3E-08
Xylene	2.16E-06	20	250	25	70	25550	151E-07	NA	NA /
		-			<u> </u>			Total	4E-08



# APPENDIX J

# LEAD SPREAD MODEL OUTPUT SHEETS FOR HUMAN HEALTH RISK ASSESSMENT

# LEAD RISK ASSESSMENT SPREADSHEET

CALIFORNIA DEPARTMENT OF TOXIC SUBSTANCES CONTROL
Current and Future On-Site Worker - Reasonable Maximum Exposure Scenario

INPUT			OUTPUT			**			
MEDIUM	LEVEL			per	centile	s		PRG-99	PRG-95
LEAD IN AIR (ug/m^3)	0.15	]	50th	90th	95th	98th	99th	(ug/g)	(ug/g)
LEAD IN SOIL (ug/g)	(430.0	BLOOD Pb, ADULT (ug/di)	NA	NA	NA	NA	NA	NA	NA
LEAD IN WATER (ug/l)	15 -	BLOOD Pb, CHILD (ug/dl)	NA	NA	NA	NA	NA	NA	NA
PLANT UPTAKE? 1=YES 0=NO	1 ·	BLOOD Pb, PICA CHILD (ug/dl)	NA	NA	NA	NA	NA	NA	NA
RESPIRABLE DUST (ug/m^3)	50	BLOOD Ph, INDUSTRIAL (ug/dl)	2.2	3.4	3.8	4.4	4.9	4262.5	6306.5

#### **EXPOSURE PARAMETERS**

ะหอ				
	resi	dential		industrial
units	adults	children	children	adults
			with pica	
days/wk	7	7	7	5
cm^2	3700	2800	2800	5800
mg/cm^2	0.5	0.5	0.5	0.5
(ug/dl)/(ug/day)	0.00011	0.00011	0.00011	0.00011
mg/day	25	55	790	25
(ug/dl)/(ug/day)	0.0176	0.0704	0.0704	0.0176
m^3/day	20	10	10	20
(ug/di)/(ug/day)	0.082	0.192	0.192	0.082
I/day	1.4	0.4	0.4	1.4
(ug/dl)/(ug/day)	0.04	0.16	0.16	0.04
kg/day	2.2	1.3	1.3	2.2
(ug/di)/(ug/day)	0.04	0.16	0.16	0.04
ug/kg	20.1	20.1	20.1	10.0
ug/kg	193.5	193.5	193.5	· · · · · · · · · · · · · · · · · · ·
	units  days/wk  cm^2 mg/cm^2 (ug/dl)/(ug/day)  mg/day (ug/dl)/(ug/day)  l/day (ug/dl)/(ug/day)  l/day (ug/dl)/(ug/day)  kg/day (ug/dl)/(ug/day)  ug/kg	resinal adults    days/wk   7     cm^2   3700     mg/cm^2   0.5     (ug/dl)/(ug/day)   0.00011     mg/day   25     (ug/dl)/(ug/day)   0.0176     m^3/day   20     (ug/dl)/(ug/day)   0.082     I/day   1.4     (ug/dl)/(ug/day)   0.04     kg/day   2.2     (ug/dl)/(ug/day)   0.04     ug/kg   20.1	residential   adults   children     days/wk   7   7   7	residential   adults   children   children   with pica   days/wk   7   7   7   7   7   7   7   7   7

## PATHWAYS, ADULTS

	Reside	ntial	Indu	strial	
Pathway	Blood Pb ug/dl	percent of total	Blood Pb ug/dl	percent of total	Concentration in medium
SOIL CONTACT:	0.08	NA	0.09	4%	430 ug/g
SOIL INGESTION:	0.19	NA	0.14	6%	430 ug/g
INHALATION:	0.28	NA	0.20	9%	0.17 ug/m^3
WATER INGESTION:	0.84	NA	0.84	39%	15 ug/l
FOOD INGESTION:	1.77	NA	0.88	41%	20.1 ug Pb/kg diet

### PATHWAYS, CHILDREN

	Typic	cal	with	pica	
	Blood Pb	percent	Blood Pb	percent	Concentration
Pathway	ug/dl	of total	ug/dl	of total	in medium
SOIL CONTACT:	0.06	NA	0.06	NA	430 ug/g
SOIL INGESTION:	1.66	NA	23.91	NA	430 ug/g
INHALATION:	0.33	NA	0.33	NA	0.17 ug/m^3
WATER INGESTION:	0.96	NA	0.96	NA	15 ug/l
FOOD INGESTION:	4.18	NA	4.18	NA	20.1 ug Pb/kg diet