

WORK PLAN

Shell Service Station 1800 Powell Street Emeryville, California

general contractors

November 2, 1989

Mr. Larry Seto
Department of Environmental Health
Hazardous Materials Program
80 Swan Way, Room 200
Oakland, California 94621

Reference:

Shell Service Station 1800 Powell Street Emeryville, California

Gentlemen:

As authorized by Shell Oil Company, we are forwarding a copy of the Work Plan prepared by GeoStrategies Inc., dated October 27, 1989, for the above referenced location.

Please do not hesitate to call should you have any questions or comments.

Sincerely,

John P. Werfal Project Manager

enclosure

cc: Ms. Wendy Howell, Shell Oil Company

Ms. Diane Lundquist, Shell Oil Company

Mr. Tom Callaghan, Regional Water Quality Control Board



2140 WEST WINTON AVENUE HAYWARD, CALIFORNIA 94545

(415) 352-4800

October 27, 1989

Gettler-Ryan Inc. 1992 National Avenue Hayward, California 94545

Attn:

Mr. John Werfal

Re:

WORK PLAN

Shell Service Station 1800 Powell Street Emeryville, California

Gentlemen:

This work plan has been prepared by GeoStrategies Inc. (GSI) for the Shell Service Station at the above referenced location (Plates 1 and 2). The work plan addresses the need to further assess petroleum hydrocarbon migration from the underground fuel storage complex.

BACKGROUND

Prior to August 1983, five tank backfill wells (S-1 through S-4, and S-11) and six ground-water monitoring wells (S-5 through S-10) were installed at the site. Boring logs and well construction details are not available for these wells.

In October 1988, the existing monitoring network outside of the tank area (Wells S-5 through S-10) was sampled and analyzed for Total Petroleum Hydrocarbons calculated as Gasoline (TPH-Gasoline), and Benzene, Toluene, Ethylbenzene, and Xylenes (BTEX). TPH-Gasoline concentrations ranged from 0.05 to 700 parts per million (ppm). Benzene concentrations ranged from to 37 ppm. Well S-9 separate-phase contained hydrocarbons (floating product). Bi-weekly monitoring of Well S-9 reveals a range of floating product thickness from 1.21 to 1.50 feet in measured thickness, through January 1989.

Gettler-Ryan Inc. October 27, 1989 Page 2

In February, Gettler-Ryan Inc. (G-R) conducted ground-water sampling for Wells S-5 and S-6 contained a sheen of the first quarter of 1989. floating product. and Well S-9 had 1.3 feet of floating TPH-Gasoline and Benzene were detected in all wells sampled. TPH-Gasoline concentrations ranged from 0.05 ppm to 6.5 ppm. Benzene concentrations ranged from 0.0009 ppm to 0.74 ppm. The results of this sampling event are presented in the GeoStrategies (GSI) quarterly report dated April 14. 1989.

In April 1989, G-R conducted second quarter ground-water sampling. Well S-9 contained floating product (1.25 feet in measured thickness). TPH-Gasoline was detected in Wells S-5, S-6, S-8 and S-10, and ranged in concentrations from 2.7 ppm to 13 ppm. Benzene was detected in all wells sampled, and ranged from 0.0010 ppm to 2.4 ppm. The results of this sampling event are presented in the GSI quarterly report dated July 13, 1989.

On October 10, 1989 GSI issued an interim ground-water sampling report summarizing the third quarterly sampling conducted by G-R. Well S-9 contained floating hydrocarbon product (1.20 feet in measured thickness). TPH-Gasoline and Benzene were detected in all wells sampled. Benzene concentrations ranged from 0.0022 ppm to 1.7 ppm. The results of the July sampling event are presented in a G-R report dated August 11, 1989 (Appendix B).

TECHNICAL APPROACH

GSI has reviewed available field and chemical data for this site. No boring logs or well construction details are available for review. Potentiometric data collected during the third quarterly ground-water sampling event indicate that groundwater beneath the site flows to the southwest towards San Francisco Bay (Plate 3). Due to the lack of boring logs and well construction details, we recommend that existing Wells S-6 and S-7 be properly abandoned. The tank backfill wells (S-1 through S-4 and S-11) and Wells S-8 through S-10 should remain as monitoring points along with Well S-5, which was installed to monitor the Pacific Gas and Electric (PG&E) vault box located adjacent to the well. We also recommend that routine pumping of Well S-9 be conducted to reduce the amount of separate-phase product at this location. If separate-phase and dissolved ground-water conditions remain unchanged by the next quarterly sampling, additional remedial measures will be proposed at that time.

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In addition to the well abandonment, we recommend that a ground-water sampling survey be conducted around the perimeter of the site, at the intersection of Powell Street and the Frontage Road, and in the area downgradient from Well S-5 (Plate 2). This in-situ procedure uses a hollow probe, with a detachable end point, which is inserted through the soil to the targeted shallow water-bearing unit. The probe is then withdrawn to permit the inflow of water into the resulting hole. A vacuum adaptor is placed on top of the probe and is used to connect the probe to The ground-water sample is extracted using a vacuum a peristaltic pump. The sample is then analyzed on-site in a mobile syringe mechanism. laboratory. The results should provide data that will assist evaluating subsurface hydrocarbon distribution and therefore, be helpful selecting new monitoring well locations. If this method is not feasible, due to the fine-grained nature of the soils beneath the site and the presence of fill of unknown composition resulting in very slow ground-water sample extraction times, a soil vapor survey will be used as an alternative cost-effective method for delineating soil vapors relative to hydrocarbons.

We further recommend that three new monitoring wells be installed to provide lithologic information and monitoring points for obtaining potentiometric data as well as ground-water quality data. The locations of the proposed wells are shown on Plate 2. The borings will be drilled to the bottom of the first encountered saturated zone. Soil samples will be collected every five feet as a minimum and at significant lithologic changes for lithologic description, field head-space analysis for organic vapor presence using an OVM, and for chemical analysis. Additional samples may be collected for lithologic description and/or chemical analysis.

Soil and ground-water samples will be analyzed in a State-certified laboratory for TPH-Gasoline according to EPA Method 8015 (Modified), and BTEX according to EPA Method 8020.

The rationale for the above proposed scope of work is based on the following criteria:

Floating product has been identified in existing Well S-9. Hydrocarbon product thickness measurements have ranged from 1.20 feet to 1.50 feet (measurements taken from G-R historical monitoring data through August 1989). Therefore, interim remediation (pumping monitoring well S-9) seems appropriate to begin to remove floating product from the shallow aquifer zone.

Gettler-Ryan Inc. October 27, 1989 Page 4

- o Boring logs and well construction details are not available for the existing wells; therefore the validity of potentiometric and ground-water quality data remain questionable. Existing Wells S-6 and S-7 should be abandoned and at least three new wells installed.
- O Dissolved petroleum hydrocarbons have migrated off-site (Plates 4 and 5). A cross-gradient, up-gradient, and down-gradient No Detection (ND) boundary needs to be established so that the hydrocarbon plume configuration can be estimated for the selection and implementation of the appropriate remedial action. An in-situ ground-water sampling survey (if feasible) or a soil vapor survey will assist in selecting optimum locations for monitoring wells both on- and off-site so the lateral extent of the hydrocarbon plume can be evaluated.

GSI recommends that the proposed monitoring wells screened intervals be installed a minimum of one-foot above the equilibrated water level to accommodate for separate-phase product (if present), and potential diurnal and seasonal ground-water fluctuations. Notwithstanding, the well screens will be emplaced so that well designs are compatible with subsurface geologic conditions. No well screens will be installed that potentially could permit cross-contamination of adjacent aquifers.

If chemical analytical results from either soil or ground-water samples show elevated concentration levels, appropriate DHS or RWQCB guidance documents and policies will be used to assess whether additional work is necessary.

Attached to this work plan are the procedures, protocols, and methods that will be used to investigate this site. Additional scopes of work, if necessary, will follow the procedures described in this work plan, or shall be added as appropriate addendum.

Gettler-Ryan Inc. October 27, 1989 Page 5

If you have any questions, please call.

GeoStrategies Inc. by,

David A. Ferreira

Geologist

Jeffrey L. Peterson Senior Hydrogeologist

My 2. Loturon

R.E.A. 1021

№ 1262 CERTIFIED **ENGINEERING GEOLOGIST**

Christopher M. Palmer C.E.G. 1262, R.E.A. 285

JLP/CMP/mlg

Plate 1. Vicinity Map

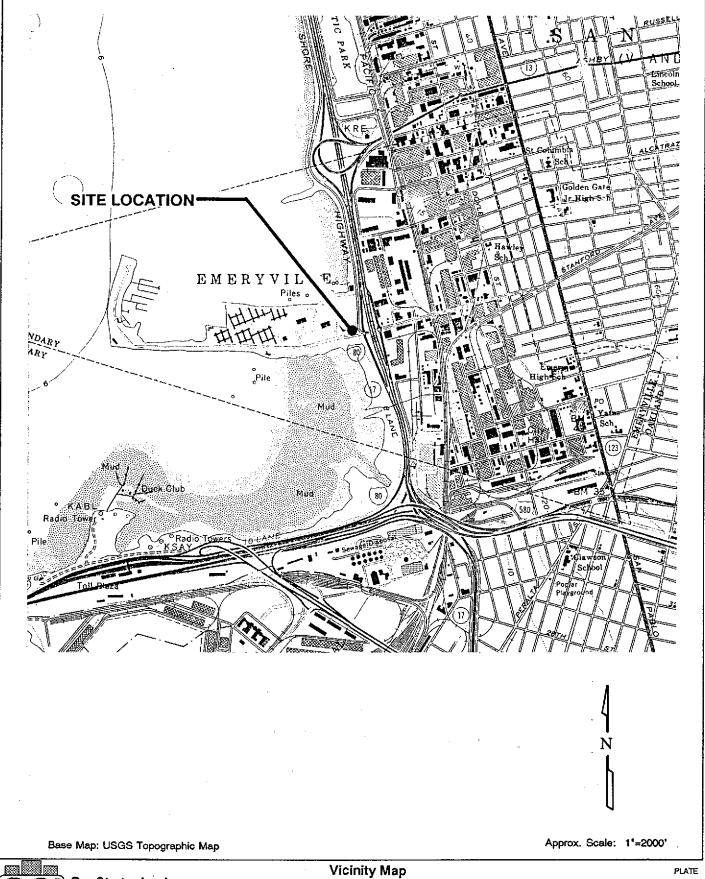
Plate 2. Site Plan

Plate 3. Potentiometric Map

Plate 4. TPH Isoconcentration Map Plate 5. Benzene Isoconcentration Map

Appendix A: Field Methods and Procedures

Appendix B: G-R Groundwater Sampling Report (August 11, 1989)





Vicinity Map Shell Service Station 1800 Powell Street Emeryville, California

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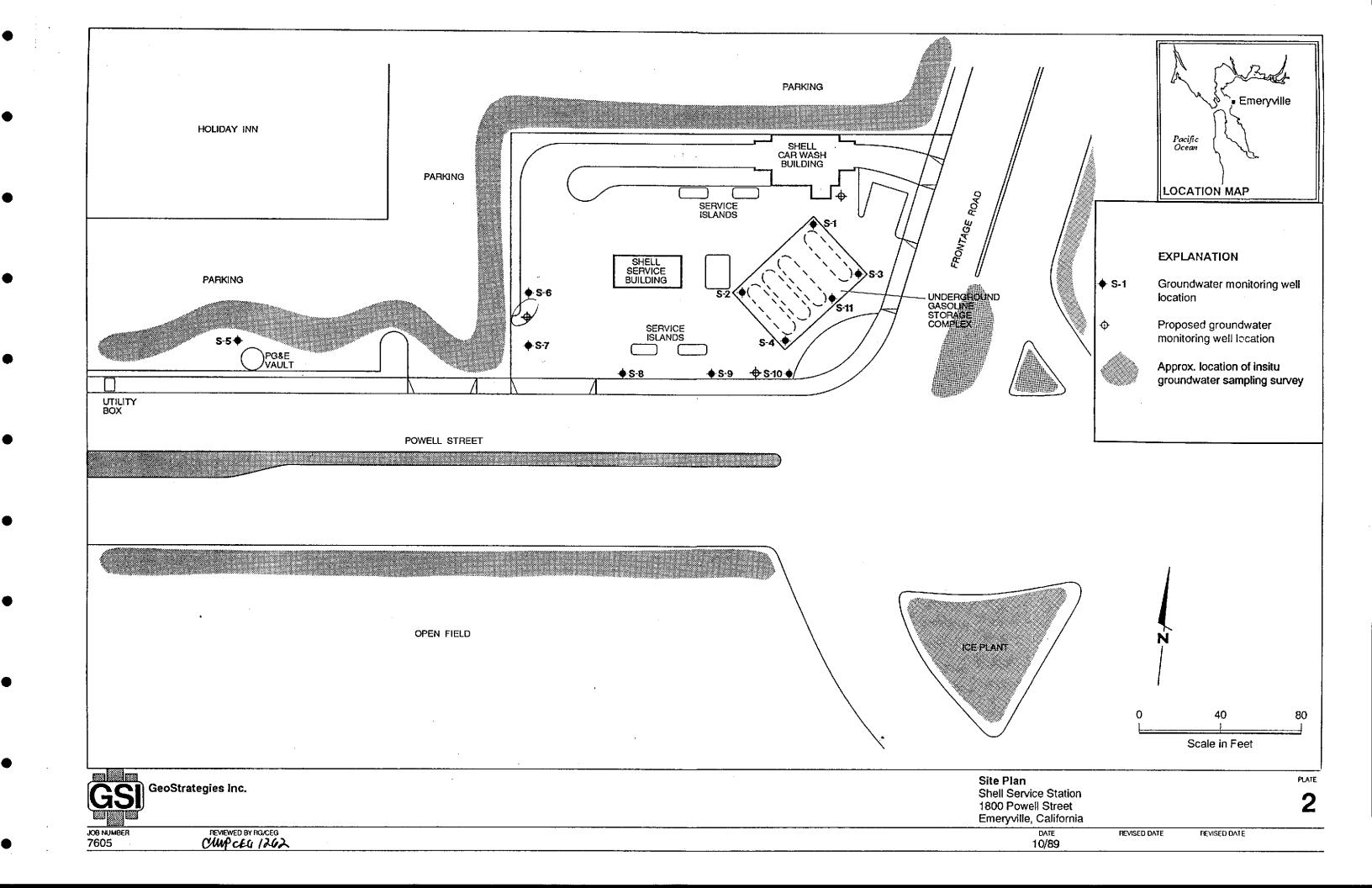
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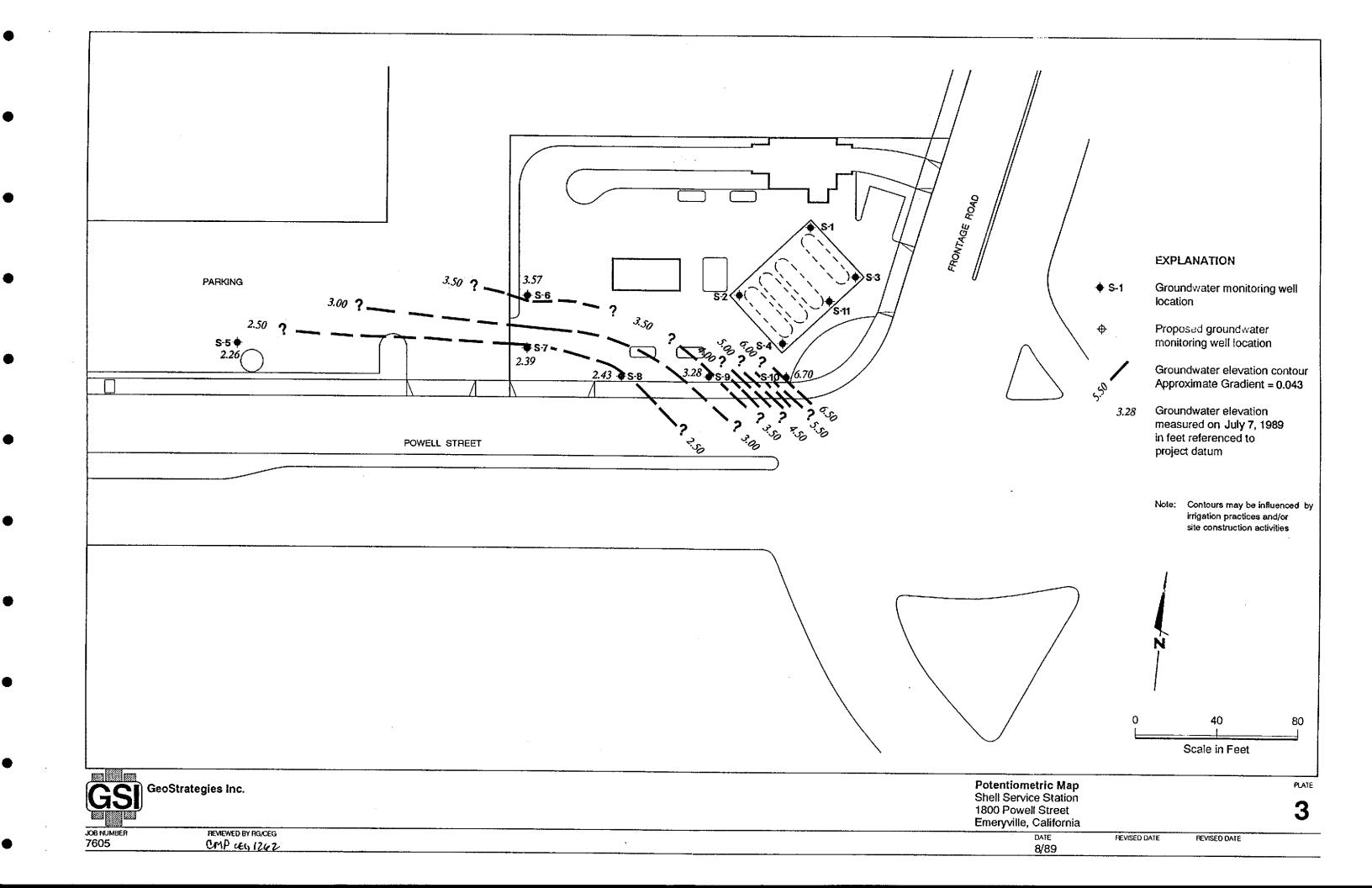
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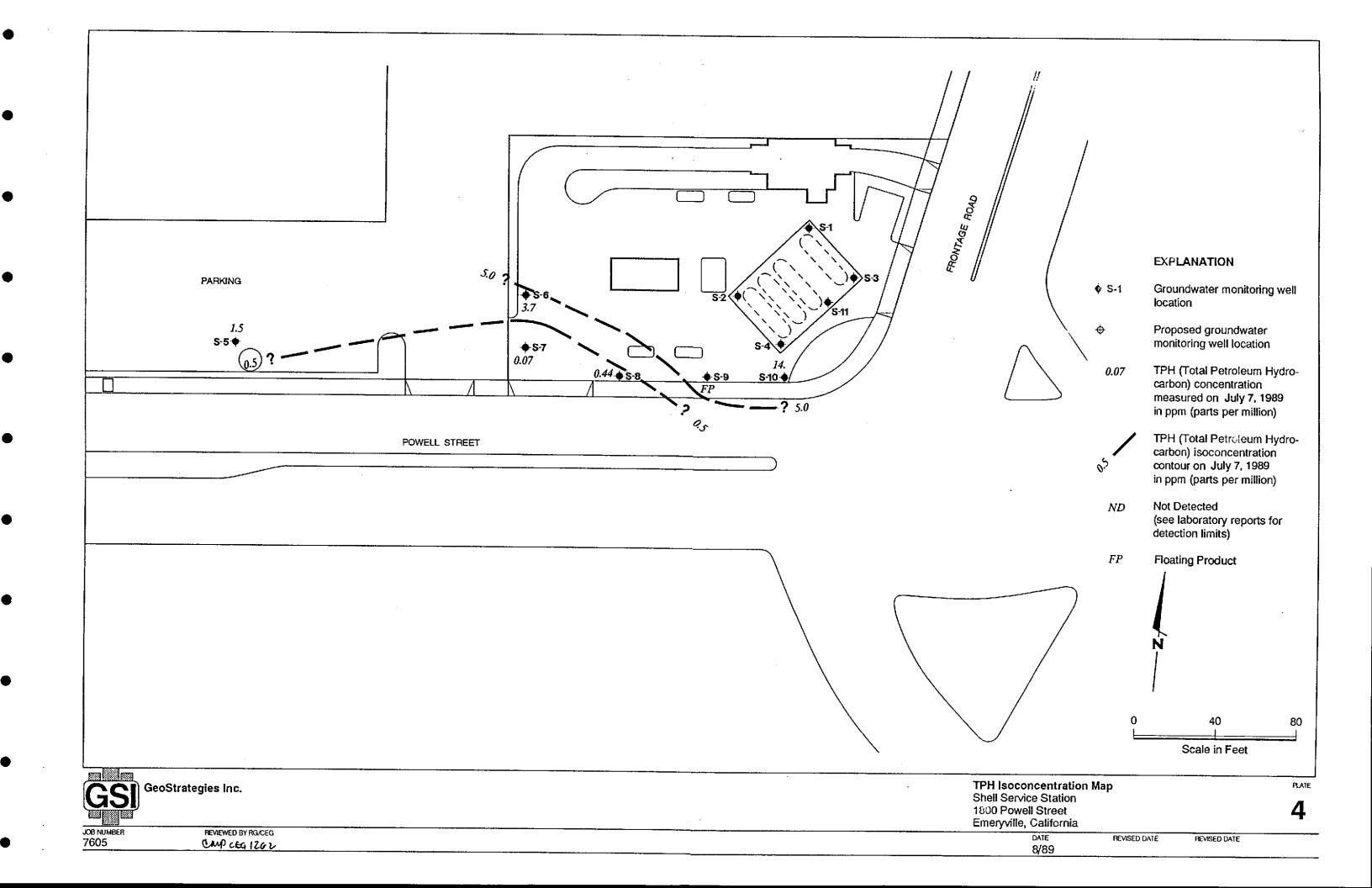
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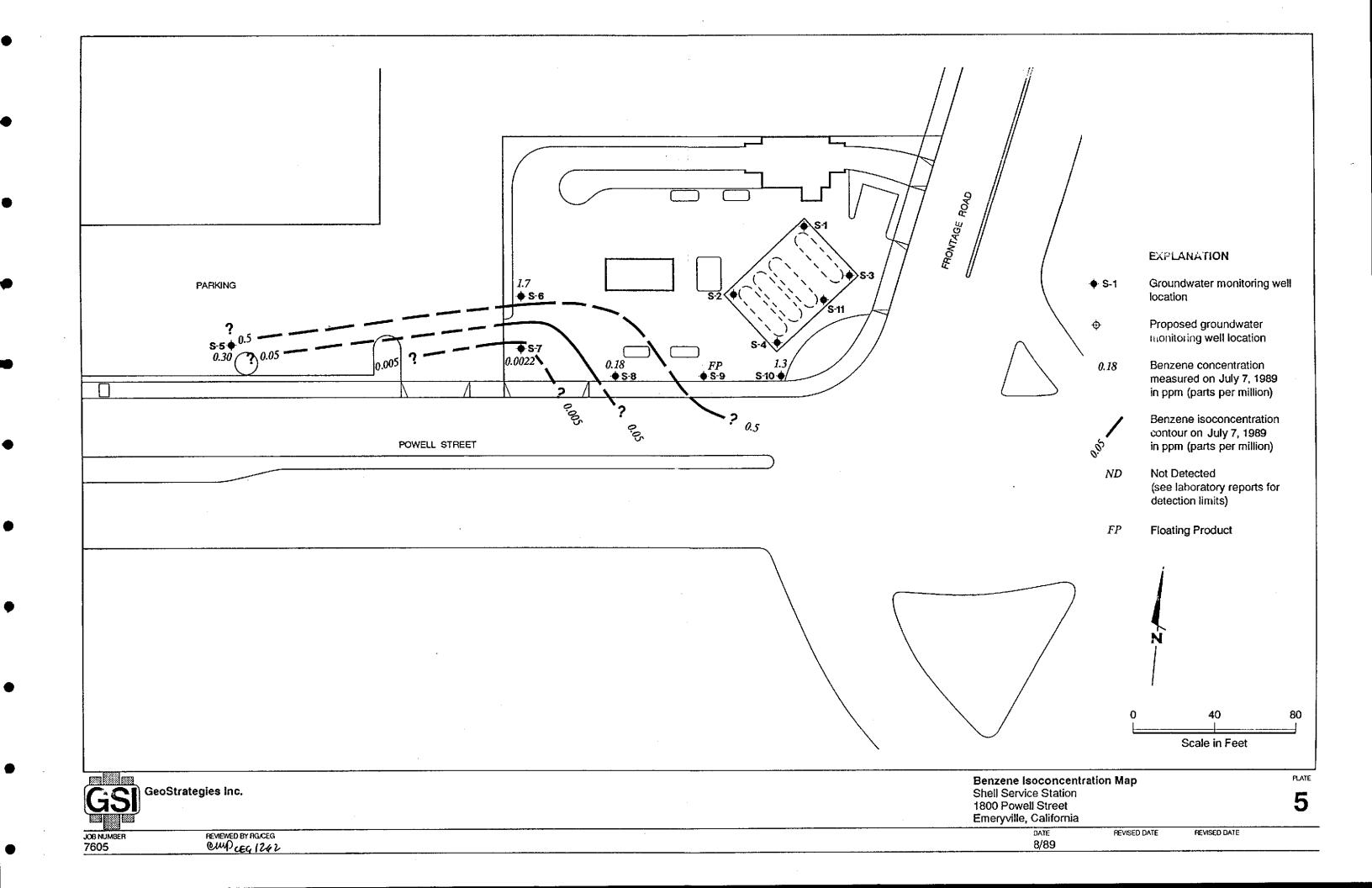
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APPENDIX A

METHODS AND PROCEDURES

FIELD METHODS AND PROCEDURES

EXPLORATION DRILLING

Mobilization

Prior to any drilling activities, GSI will verify that necessary drilling permits have been secured.

Utility locations will be located and drilling will be conducted so as not to disrupt activities at a project site. GSI will obtain and review available public data on subsurface geology and if warranted, the location of wells within a half-mile of the project site will be identified. Drillers will be notified in advance so that drilling equipment can be inspected prior to performing work.

Drilling

The subsurface investigations are typically performed to assess the lateral and vertical extent of petroleum hydrocarbons present in soils and ground water. Drilling methods will be selected to optimize field data requirements as well as be compatible with known or suspected subsurface geologic conditions.

Monitoring wells are installed using a truck-mounted hollow-stem auger drill rig or mud-rotary drill rig. Typically, the hollow-stem rig is used for wells up to 100 feet, if subsurface conditions are Wells greater than 100-feet deep are typically drilled using mud-rotary techniques. When mud rotary drilling is used, an electric log will be performed for additional lithological information. Also during mud rotary drilling, precautions will be taken to prevent mud from circulating contaminants by using a conductor casing to seal off contaminated zones. Samples will be collected for lithologic logging by continuous chip, and where needed by drive sample or core as specified by the supervising geologist.

Soil Sampling

Shallow soil borings will be drilled using a truck-mounted hollow-stem auger drilling rig, unless site conditions favor a different drilling method. Drilling and sampling methods will be consistent with ASTM Method D-1452-80. The auger size will be a minimum 6-inch nominal outside-diameter (O.D). No drilling fluids will be used during this drilling method. The augers and other tools used in the bore hole will be steam cleaned before use and between borings to minimize the possibilities of cross-contamination between borings.

Soil samples are typically collected at 5-foot intervals as a minimum from ground surface to total depth of boring. Additional soil samples will be collected based on significant lithologic changes and/or potential chemical content. Soil samples from each sampling interval will be lithologically described by a GSI geologist (Figure 1). Soil colors will be described using the Munsell Color Chart. Rock units will be logged using appropriate lithologic terms, and colors described by the G.S.A. Rock Color Chart.

Head-space analyses will be performed to check for the evidence of volatile organic compounds. Head-space analyses will be performed using an organic vapor analyzer; either an OVA, HNU, or OVM. Organic vapor concentrations will be recorded on the GSI field log of boring (Figure 1). The selection of soil samples for chemical analysis are typically based on the following criteria:

- 1) Soil discoloration
- 2) Soil odors
- 3) Visual confirmation of chemical in soil
- 4) Depth with respect to underground tanks (or existing grade)
- 5) Depth with respect to ground water
- 6) OVA reading

Soil samples (full brass liners) selected for chemical analysis are immediately covered with aluminum foil and the liner ends are capped to prevent volatilization. The samples are labeled and entered onto a Chain-of-Custody form, and placed in a cooler on blue ice for transport to a State-certified analytical laboratory.

Soil cuttings are stockpiled on-site. Soils are sampled and analyzed for site-specific chemical parameters. Disposition of soils is dependent of chemical analytical results of the samples.

Soil Sampling - cont.

Soil borings not converted to monitoring wells will be backfilled (sealed) to ground surface using either a neat cement or cement-bentonite grout mixture. Backfilling will be tremied by continuously pumping grout from the bottom to the top of the boring where depth exceeds 20' or as required by local permit requirements.

All field and office work, including exploratory boring logs, are prepared under the direction of a registered geologist.

Monitoring Well Installation

Monitoring well casing and screen will be constructed of Schedule 40, flush-joint threaded polyvinylchloride (PVC). The well screen will be factory mill-slotted unless additional open area is required (eg. conversion to an extraction well in a low-yield aquifer). The screen length will be placed adjacent to the aquifer material to a minimum of 2-feet above encountered water. No screen shall be placed in a borehole that potentially creates hydraulic interconnection of two or more aquifer units. Screen slot size and well sand pack will be compatible with encountered aquifer materials, as confirmed by sieve analysis.

Monitoring wells will be completed below grade (Figure 2) unless special conditions exist that require above-grade completion design. In the event a monitoring well is required in an aquifer unit beneath an existing aquifer, the upper aquifer will be sealed off by installing a steel conductor casing with an annular neat cement or cement-bentonite grout seal. This seal will be continuously tremied pumped from the bottom of the annulus to ground surface.

The monitoring well sand pack will be placed adjacent to the entire screened interval and will extend a recommended minimum distance of 2-feet above the top of the screen. No sand pack will be placed that interconnects two or more aquifer units. A minimum 2-foot bentonite pellet or bentonite slurry seal will be placed above the sand pack. Sand pack, bentonite, and cement seal levels will be confirmed by sounding the annulus with a calibrated weighted tape. The remaining annular space above the bentonite seal will be grouted with a bentonite-cement mixture and will be tremie-pumped from the bottom of the annular space to the ground surface. The bentonite content of the grout will not exceed 5 percent by weight. A field log of boring and a field well completion form will be prepared by GSI for each well installed.

Decontamination of drilling equipment before drilling and between wells will consist of steam cleaning, and/or Alconox wash.

Well Development

Monitoring wells will be developed using a submersible pump, bladder pump or bailer. All well developing equipment will be decontaminated prior to development using a steam cleaner and/or Alconox detergent wash. Wells will be developed until discharge water is visibly clear and free of sediment. The adequacy of well development will be assessed by the GSI geologist. Indicator parameters (pH, specific conductance, and temperature) will be monitored and recorded during well development. Field instrument calibrations will be performed according to manufacturer's specifications.

Well Surveying

Monitoring wells will be surveyed to obtain top of box elevations to the nearest ± 0.01 foot. Water level measurements will be recorded to the nearest ± 0.01 foot and referenced to mean sea level (MSL). If additional wells are required, then existing and newly installed wells are surveyed relative to MSL.

GROUND-WATER SAMPLING AND ANALYSIS

Quality Assurance/Quality Control Objectives

The sampling and analysis procedures employed by Gettler-Ryan Inc. (G-R) for ground-water sampling and monitoring follow specific Quality Assurance/Quality Control (QA/QC) guidelines. Ouality objectives have been established by G-R to develop and implement procedures for obtaining and evaluating water quality and field data an accurate, precise, and complete manner so that sampling procedures and field measurements provide information comparable and representative of actual field conditions. Control (QC) is maintained by G-R by using specific field protocols requiring the analytical laboratory to perform internal external QC checks. It is the goal of G-R to provide data that are accurate, precise, complete, comparable, and representative. definitions for accuracy, precision, completeness, comparability, representativeness are as follows:

- Accuracy the degree of agreement of a measurement with an accepted referenced or true value.
- <u>Precision</u> a measure of agreement among individual measurements under similar conditions. Usually expressed in terms of the standard deviation.
- <u>Completeness</u> the amount of valid data obtained from a measurement system compared to the amount that was expected to meet the project data goals.
- <u>Comparability</u> expresses the confidence with which one data set can be compared to another.
- Representativeness a sample or group of samples that reflects the characteristics of the media at the sampling point. It also includes how well the sampling point represents the actual parameter variations which are under study.

As part of the G-R QA/QC program, applicable federal, state, and local reference guidance documents are followed. The procedures outlined in these regulations, manuals, handbooks, guidance documents, and journals are incorporated into the G-R sampling procedures to assure that; (1) ground-water samples are properly collected, (2) ground-water samples are identified, preserved, and transported in a manner such that they are representative of field conditions, and (3) chemical analysis of samples are accurate and reproducible.

Guidance and Reference Documents Used to Collect Groundwater Samples

These documents are used to verify Gettler-Ryan Inc. sampling procedures and consistent with current regulatory guidance. If site specific work and sampling plans are required, those plans will be developed from these documents.

NEIC Manual for Groundwater/Subsurface Investigation at Hazardous Waste Sites

U.S.E.P.A. - 530/SW611

Procedures Manual for Groundwater Monitoring at Solid Waste Disposal Facilities (August, 1977)

U.S.E.P.A. - 600/4-79-020

Methods for Chemical Analysis of Water and Wastes (1983)

U.S.E.P.A. - 600/4-82-029

Handbook for Sampling and Sample Preservation of Water and Wastewater (1982)

U.S.E.P.A. - 600/4-82-057

Test Methods for Organic Chemical Analysis of Municipal and Industrial Wastewater (July, 1982)

U.S.E.P.A. - SW-846#, 3rd Edition

Test Methods for Evaluating Solid Waste - Physical/Chemical Methods (November, 1986)

40 CFR 136.3e, Table II (Code of Federal Regulations) Required Containers, Preservation Techniques, and Holding Times

Resources Conservation and Recover Act (OSWER 9950.1)

Groundwater Monitoring Technical Enforcement Guidance Document (September, 1986)

California Regional Water Quality Control Board (Central Valley Region)

A Compilation of Water Quality Goals (September, 1988); Updates (October, 1988)

California Regional Water Quality Control Board (North Coast, San Francisco Bay, and Central Valley)

Regional Board Staff Recommendations for Initial Evaluations and Investigation of Underground Tanks: Tri-Regional Recommendations (June, 1988)

Guidance and Reference Documents Used to Collect Groundwater Samples (cont.)

Regional Water Quality Control Board (Central Valley Region)

Memorandum: Disposal, Treatment, and Refuse of Soils Contaminated with Petroleum Fractions (August, 1986)

State of California Department of Health Services

Hazardous Waste Testing Laboratory Certification List (March, 1987)

State of California Water Resources Control Board Leaking Underground Fuel Tank (LUFT) Field Manual (May, 1988), and LUFT Field Manual Revision (April, 1989)

State of California Water Resources Control Board

Title 23, (Register #85.#33-8-17-85), Subchapter 16: Underground Tank Regulations; Article 3, Sections 2632 and 2634; Article 4, Section 2647 (October, 1986)

Alameda County Water District

Groundwater Protection Program: Guidelines for Groundwater and Soil Investigations at Leaking Underground Fuel Tank Sites (November, 1988)

American Public Health Association

Standard Methods for the Examination of Water and Wastewaters, 16th Edition

Analytical Chemistry (journal)

Principles of Environmental Analysis, Volume 55, Pages 2212-2218 (December, 1983)

Santa Clara Valley Water District

Guidelines for Preparing or Reviewing Sampling Plans for Soil and Groundwater Investigation of Fuel Contamination Sites (January, 1989)

Santa Clara Valley Water District

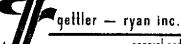
Investigation and Remediation at Fuel Leak sites: Guidelines for Investigation and Technical Report Preparation (March 1989)

American Petroleum Institute

Groundwater Monitoring & Sample Bias; API Publication 4367, Environmental Affairs Department, June 1983

Site Specific (as needed)

General and specific regulatory documents as required.



Because ground-water samples collected by G-R are analyzed to the parts per billion (ppb) range for many compounds, extreme care is exercised to prevent contamination of samples. When volatile or semi-volatile organic compounds are included for analysis, G-R sampling crew members will adhere to the following precautions in the field:

- 1. A clean pair of new, disposable gloves are worn for each well being sampled.
- When possible, samples are collected from known or suspected wells that are least contaminated (i.e. background) followed by wells in increasing order of contamination.

When known or potential organic compounds are being sampled for, the following additional precautions are taken:

- 1. All sample bottles and equipment are kept away from fuels and solvents. When possible, gasoline (used in generators) is stored away from bailers, sample bottles, purging pumps, etc.
- 2. Bailers are made of Teflon or Stainless Steel. Other materials such as plastic may contaminate samples with phthalate esters which interfere with many Gas Chromatography (GC) analyses.
- 3. Volatile organic ground-water samples are collected so that air passage through the sample does not occur or is minimal (to prevent volatiles from being stripped from the samples): sample bottles are filled by slowly running the sample down the side of the bottle until there is a positive convex meniscus over the neck of the bottle; the Teflon side of the septum (in cap) is positioned against the meniscus, and the cap screwed on tightly; the sample is inverted and the bottle lightly tapped. The absence of an air bubble indicates a successful seal; if a bubble is evident, the cap is removed, more sample is added, and the bottle is resealed.
- 4. Extra Teflon seals are brought into the field in case seals are difficult to handle and/or are dropped. Dropped seals are considered contaminated and are not used. When replacing seals or if seals become flipped, care is taken to assure that the Teflon seal faces down.

Sample analysis methods, containers, preservatives and holding times are shown on Table 1.

Laboratory and field handling procedures of samples are monitored by including QC samples for analysis with every submitted sample lot from a project site. QC samples may include any combination of the following:

- A. <u>Trip Blank</u>: Used for purgeable organic compounds only; QC samples are collected in 40 milliliter (ml) samples vials filled in the analytical laboratory with organic-free water. Trip blanks are sent to the project site, and travel with project site samples. Trip blanks are not opened, and are returned from a project site with the project site samples for analysis.
- B. <u>Field Blank</u>: Prepared in the field using organic-free water. These QC samples accompany project site samples to the laboratory and are analyzed for specific chemical parameters unique to the project site where they were prepared.
- C. <u>Duplicates</u>: Duplicated samples are collected "second samples" from a selected well and project site. They are collected as either split samples or second-run samples collected from the same well.
- D. Equipment Blank: Periodic QC sample collected from field equipment rinsate to verify decontamination procedures.

The number and types of QC samples are determined as follows:

- A. Up to 2 wells Trip Blank Only
- B. 2 to 5 Wells 1 Field Blank and 1 Trip Blank
- C. 5 to 10 Wells 1 Field blank, 1 Trip Blank, and 1 Duplicate
- D. More than 10 Wells 1 Field Blank, 1 Trip Blank, and 1 Duplicate per each 12 wells
- E. If sampling extends beyond one day, quality control samples will be collected for each day.

SAMPLE COLLECTION

This section describes the routine procedures followed by G-R while collecting ground-water samples for chemical analysis. These procedures include decontamination, water-level measurements, well purging, physical parameter measurements, sample collection, sample preservation, sample handling, and sample documentation. Critical sampling objectives for G-R are to:

- 1. Collect ground-water samples that are representative of the sampled matrix and,
- 2. Maintain sample integrity from the time of sample collection to receipt by the analytical laboratory.

Sample analyses methods, containers, preservation, and holding times are presented in Table 1.

Decontamination Procedures

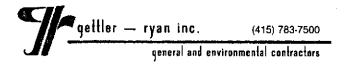
All physical parameter measuring and sampling equipment are decontaminated prior to sample collection using Alconox or equivalent detergent followed by steam cleaning with deionized water. Any sampling equipment surfaces or parts that might absorb specific contaminants, such as plastic pump valves, impellers, etc., are cleaned in the same manner.

Sample bottles, bottle caps, and septa used for sampling volatile organics are thoroughly cleaned and prepared in the laboratory. Sample bottles, bottle caps, and septa are protected from all potential chemical contact before actual usage at a sample location.

During field sampling, equipment placed in a well are decontaminated before purging or sampling the next well. The equipment are decontaminated by cleaning with Alconox or equivalent detergent followed by steam cleaning with deionized water.

Water-Level Measurements

Prior to purging and sampling a well, the static-water levels are measured in all wells at a project site using an electric sounder and/or calibrated portable oil-water interface probe (Figure 3). Both static water-level and separate-phase product thickness are measured to the nearest ±0.01 foot. The presence of separate-phase product is confirmed using a clean, acrylic or polyvinylchloride (PVC) bailer, measured to the nearest ±0.01 foot with a decimal scale tape.



Water-Level Measurements (continued)

The monofilament line used to lower the bailer is replaced between line to preclude the possibility cross-contamination. Field observations (e.g. well integrity, product color, turbidity, water color, odors, etc.) are noted on the G-R Well Sampling Field Data Sheet shown in Figure 3. Before and after each electric sounder. interface probe and bailer washing with Alconox or equivalent decontaminated detergent followed rinsing with deionized water cross-contamination.

As mentioned previously, water-levels are measured in wells with known or suspected lowest dissolved chemical concentrations to the highest dissolved concentrations.

Well Purging

Before sampling occurs, well casing storage water and interstitial water in the artificial sand pack will be purged using (1) a positive displacement bladder pump constructed of inert, non-wetting, Teflon and stainless steel, (2) a pneumatic-airlift pumping system, (3) a centrifigal pumping system, or (4) a Teflon or Stainless steel bailer Methods of purging will be assessed based on well size. (Figure 4). location, accessibility, and known chemical conditions. Individual well purge volumes are calculated from borehole volumes which take into account the sand packed interval in the well annular space. As a general rule, a minimum of 3 and a maximum of 10 borehole volumes will be purged. Wells which dewater or demonstrate slow recharge periods (i.e. low-yield wells) during purging activities may be sampled after fewer purging cycles. If a low-yield (low recovery) well is to be sampled, sampling will not take place until at least 80 percent of the previously measured water column has been replaced by recharge, or as per local requirements. Physical parameter measurements (temperature, pH, and specific conductance) are closely monitored throughout the well purging process and are used by the G-R sampling crew as indicators for assessing sufficient purging. Purging is continued all three physical parameters have stabilized. conductance (conductivity) meters read are to the nearest umhos/cm, and are calibrated daily. pH meters are read to the nearest ± 0.1 pH units and are calibrated daily. Temperature is read to the nearest 0.1 degree F. Calibration of physical parameter meters will follow manufacturers specifications. Monitoring wells will be purged according to the protocol presented in Figure 4. Collected field data during purging activities will be entered on the G-R Well Sampling Field Data Sheet shown in Figure 3. Copies of the G-R Field Data Sheets will be reviewed by the G-R Sampling Manager for accuracy and completeness.

DOCUMENTATION

Sample Container Labels

Each sample container will be labeled by an adhesive label, noted in permanent ink immediately after the sample is collected. Label information will include:

Sample point designation (i.e. well number or code)

Sampler's identification

Project number

Date and time of collection

Type of preservation used

Well Sampling Data Forms

In the field, the G-R sampling crew will record the following information on the Well Sampling Data Sheet for each sample collected:

Project number

Client

Location

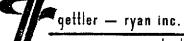
Source (i.e. well number)

Time and date

Well accessibility and integrity

Pertinent well data (e.g. depth, product thickness, static water-level, pH, specific conductance, temperature)

Calculated and actual purge volumes



Chain-of-Custody

A Chain-of-Custody record (Figure 5) shall be completed and accompany sample and every shipment of samples to the analytical laboratory in order to establish the documentation necessary to trace sample possession from time of collections. The record will contain the following information:

- Sample or station number or sample identification (ID)
- Signature of collector, sampler, or recorder
- Date and time of collection
- Place of collection
- Sample type
- Signatures of persons involved in chain of possession
- Inclusive dates of possession

Samples shall always be accompanied by a Chain-of-Custody record. transferring the samples, the individual relinquishing and receiving the samples will sign, date, and note the time on the Chain-of-Custody record. G-R will be responsible for notifying the laboratory coordinator when and how many samples will be sent to the laboratory for analysis, and what types of analyses shall be performed.

SAMPLE ANALYSIS METHODS, CONTAINERS, PRESERVATIONS, AND HOLDING TIMES

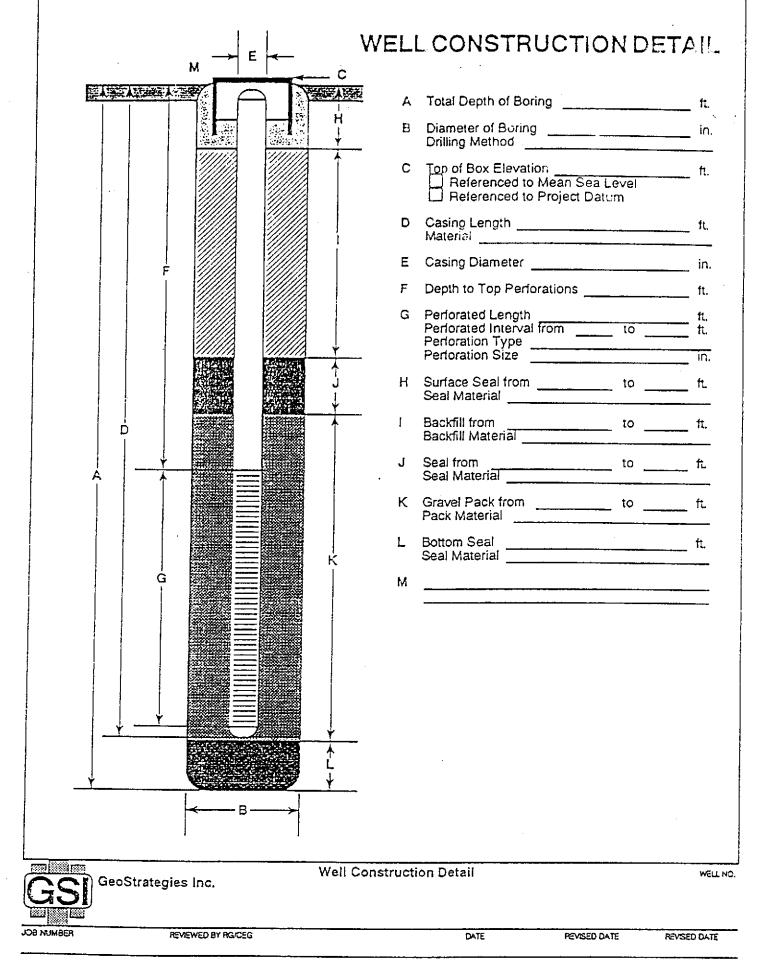
TABLE 1

Parameter	Analytical <u>Method</u>	Reporting Units	Container	Preservation	Maximum Holding <u>Time</u>
Total Petroleum Hydrocarbons (gasoline)	EPA 8015 (modified)	mg/l ug/l	40 ml. vial glass, Teflon	cool, 4 C HC1 to pH<2	14 days (maximum)
Benzene Toluene Ethylbenzene Xylenes (BTEX)	EPA 8020	mg/l ug/l mg/l	50 ml. vial glass, Teflon lined septum 1 l glass, Teflon	cool, 4 C HC1 to pH<2	7 days (w/o preservative) 14 days (w preservative)
Oil & Grease	SM 503E	ug/l	lined septum	H2SO4 to pH<2	28 days (maximum)
Total Petroleum Hydrocarbons (Diesel)	EPA 8015 (modified)	mg/l ug/l	40 ml. vial glass, Teflon lined septum	cool, 4 C	14 days (maximum)
Halogented Volatile Organics (chlorinated solvents)	8010	mg/l ug/l	40 ml. vial glass, Teflon lined septum	cool, 4 C	14 days (maximum)
Non chlorinated solvents	8020	mg/l ug/l	40 ml. vial glass, Teflon lined septum	cool, 4 C HCl to pH<2	14 days (maximum)
Volatile Organics	8240	mg/t ug/l	40 ml. vial glass, Teflon lined septum	cool, 4 C	14 days (maximum)
Semi-Volatile Organics	8270	mg/l ug/l	40 ml. vial glass, Teflon lined septum	cool , 4 C	14 days (maximum)
Specific Conductance (Field test)		umhos/cm			
pH (Field test)		pH units			
Temperature (Field test)		Deg F			



FIELDEXPLORATORYBORINGLOG

Field loc	ation of bo	ring:						Project No.:		Date:	· - · · ·	Boring (vo:
								Client:		······································		-
1								Location:		· · · · · · · · · · · · · · · · · · ·		
								City:	•			Sheet
			÷.					Logged by:		Driller:		of
								Casing installs	ition data:	i		J
Drilling o	nethod:							1				
Hole dia	meler:					· ·		Top of Box Ele	evation:	-	Datum:	
i	Blows/ft. or Pressure (pst)							Water Level				1
οÊ	13 F	Type of Sample	Samole Number	Depth (R.)	Sample	# 3	Sol Group Symbol (USCS)	Time				<u> </u>
Old (mxxt)	50 28	CyT.	28.5	B.	525	Vveli Detail	28.5 S. 1	Date				
	- &			3			ν -			Description		
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Remarks:												
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GETTLER-RYAN INC.

General and Environmental Contractors

WELL SAMPLING FIELD DATA SHEET

COMPANY			JOB #	
LOCATION			DATE	
CITY			TIME	
Well ID.		Well Co	ondition	
Well Diameter	in_		arbon Thickness	
Total Depth Depth to Liquid-	ft.	Volume Factor	2" = 0.17 6" = 1.50 3" = 0.38 8" = 2.60 4" = 0.66 10" = 4.10	0 12" = 5.80
/ # of \		L	=(Estimated Purge Volume)	
Purging Equipment				
Sampling Equipment				
Starting Time		Purging	Flow Rate	gp
Estimated Purge Volume	gal. / (Purging Flow Rate)	gpm. = (Anticipated Purging Time	mi
m:				
Time	pН	Conductivity	y Temperature	Volume
	рН	Conductivity	7 Temperature	Volume
11me	рн	Conductivity	' Temperature	Volume
Time	рн	Conductivity	Temperature	Volume
lime				
Did well dewater?	lf	yes, time	Volume	
Did well dewater?	lf	yes, time	Volume	
Oid well dewater?	If	yes, time Weather Co	Volume	
Did well dewater?	If	yes, time _ Weather Co	Volume onditions	
Did well dewater?	If	yes, time _ Weather Co	Volume onditions ottles Used	

Sampling Crew Reviews Project

```
Sampling Requirments/Schedule
                                               Field Decontemination and
                                              Instrumentation Calibration
                                                Check Integrity of Well
                                               (Inspect for Well Damage)
                                           Heasure and Record Depth to Water
                                                 and Total Well Depth
                                                (Electric Well Sounder)
                                              Check for Floating Product
                                              (Oll/Water Interface Probe)
Floating Product
                                            Floating Product Not
Present
                                           Present
Confirm Product Thickness
                                          Purge Volume Calculation
(Acrylic or PVC Bailer)
                             V = Purge volume (gallons)
Collect Free-Product Sample
                               = 3.14159
                             h = Height of Water Column (feet)
Dissolved Product Sample
                             r = Borehole radius (inches)
Not Required
Record Data on
                             Evacuate water from well equal to the calculated purge volume while
Field Data Form
                             monitoring groundwater stabilization indicator parameters (pK, conductivity, temperature)
                             at intervals of one casing volume.
      Well Dewaters after
                                                           Wall Readily Recovers
      One Purge Volume
      (Low yield well).
      Well Recharges to 80% of
                                                           Record Groundwater Stability
      Initials Heasured Water
                                                           Indicator Parameters from each
      Column Height in Feet
                                                          Additional Purge Volume
      within 24 hrs. of Evacuation.
                                                          Stability indicated when the following criteria are met:
      Measure Groundwater Stability
                                                                         ± 0.1 pH units
± 10%
      Indicator Parameters (pH,
                                                          Conductivity:
      Temp., Conductivity)
                                                                        1.0 degree F
                                                          Tempertaure:
      Collect Sample and Complete
                                          Groundwater Stability
                                                                    Groundwater Stability
      Chain-of-Custody
                                          Achieved
                                                                    Not Achieved
                                          Collect Sample and
                                                                    Continue Purging
                                          Complete
                                                                    Until Stability is
                                          Chain-of-Custody
                                                                    Achieved
     Preserve Sample According
                                          Preserve Sample
                                                                    Collect Sample and
     to Required Chemical Analysis
                                          According to Required
                                                                    Complete Chain-of-
                                          Chemical Analysis
                                                                    Custody
                                                                    Preserve Sample
                                                                    According to Required
                                                                    Chemical Analysis
     Transport to Anayltical
                                         Transport to
                                                                   Transport to
     Laboratory
                                         Analytical Laboratory
                                                                   Analytical Laboratory
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Gettler - R	lyan Inc	EN	VIRONMENTAL DI	VISION .	Chain of Custo
COMPANY	····	•			JOB NO:
JOB LOCATION _				·	
CITY	· · · · · · · · · · · · · · · · · · ·			PHONE I	١٥
AUTHORIZED			DATE _	P.O. NO.	
SAMPLE ID	NO. OF CONTAINERS	SAMPLE MATRIX	DATE/TIME SAMPLED	ANALYSIS REQUIRED	SAMPLE CONDITION LAB ID
	·				
			•		
			•		
RELINQUISHED BY	:		RECE	IVED BY:	
RELINQUISHED BY	4		RECE	IVED BY:	
RELINQUISHED BY	:			IVED BY LAB:	· .
		<u>-</u>	· · · · · · · · · · · · · · · · · · ·		
DESIGNATED LABO	DRATORY:			DHS #:	
REMARKS:					
	<u></u>				
NATE COLUDI CTOR					
ATE COMPLETED			FORE	MAN	FIGURE 5

APPENDIX B

CHEMICAL ANALYTICAL REPORTS

GROUNDWATER SAMPLING REPORTS
CHEMICAL ANALYTICAL REPORTS



August 11, 1989

GROUNDWATER SAMPLING REPORT

Referenced Site:

Shell Service Station 1800 Powell Street Emeryville, California

Sampling Date:

July 7, 1989

This report presents the results of the quarterly groundwater sampling and analytical program conducted by Gettler-Ryan Inc. on July 7, 1989 at the referenced location. The site is occupied by an operating service station located on the northwest corner of Powell Street and I-80. The service station has underground storage tanks containing regular leaded, unleaded and super unleaded gasoline products and diesel.

There are currently six groundwater monitoring wells and five tank backfill wells on site at the locations shown on the attached site map. Groundwater samples were not collected from the tank backfill wells. Prior to sampling, all monitoring wells were inspected for total well depth, water levels, and presence of separate phase product using an electronic interface probe. A clean acrylic bailer was used to visually confirm the presence and thickness of separate phase product. Groundwater depths ranged from 5.92 to 10.44 feet below grade. Separate phase product was observed in well S-9.

Wells that did not contain separate phase product were purged and sampled. Standard sampling procedure calls for a minimum of four case volumes to be purged from each well. Each well was purged while pH, temperature, and conductivity measurements were monitored for stability. In cases where a well dewatered or less than four case volumes were purged, groundwater samples were obtained after the physical parameters had stabilized. The purge water was contained in drums for proper disposal. Details of the final well purging results are presented on the attached Table of Monitoring Data.

Samples were collected, using Teflon bailers, in properly cleaned and laboratory prepared containers. All sampling equipment was thoroughly cleaned after each well was sampled and steam cleaned upon completion of work at the site. The samples were labeled, stored on blue ice, and transported to the laboratory for analysis. A field blank (SF-10) and trip blank, supplied by the laboratory, were included and analyzed to assess quality control. A duplicate sample (SD-8), was submitted without a well designation to assess laboratory performance. Analytical results for the blanks are included in the Certified Analytical Report (CAR's). Chain of custody records were established noting sample identification numbers, time, date, and custody signatures.

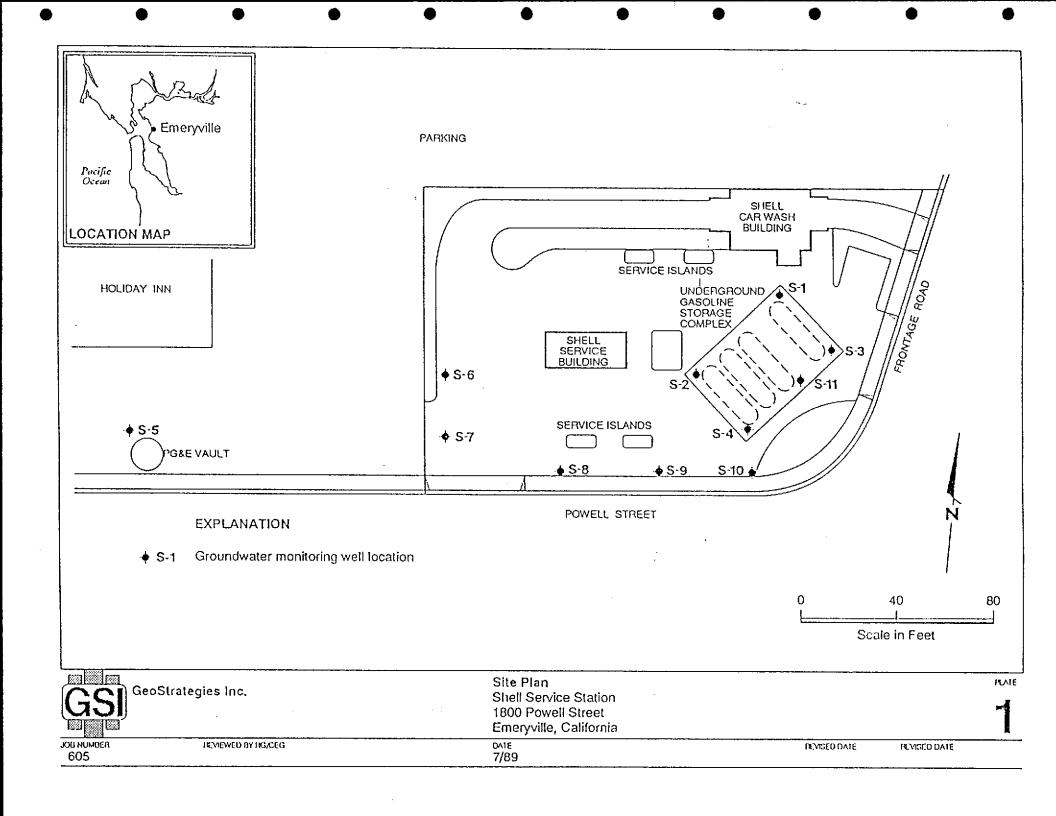
Report 3605-3

The samples were analyzed at International Technology Corporation - Santa Clara Laboratory located at 2055 Junction Avenue, San Jose, California. The laborator assigned a California DHS-HMTL Certification number of 137. The results are present a Certified Analytical Report, a copy of which is attached to this report.

Tom Paulson

Sampling Manager

attachments





ANALYTICAL SERVICES

CERTIFICATE OF ANALYSIS

Gettler-Ryan

1992 National Avenue Hayward, CA 94545 ATTN: John Werfal

Work Order Number: S9-07-061

Date: July 31, 1989

P.O. Number: 3605

This is the Certificate of Analysis for the following samples:

Client Project ID:

GR #3605, Shell, 1800 Powell Street,

Emeryville, CA

Date Received by Lab:

7/10/89 8

Number of Samples:

Sample Type:

Water

The method of analysis for low boiling hydrocarbons is taken from EPA Methods 8015, 8020 and 5030. The sample is examined using the purge and trap technique. Final detection is by gas chromatography using a flame ionization detector as well as a photoionization detector. The result for total low boiling hydrocarbons is calculated as gasoline and includes benzene, toluene, ethyl benzene and xylenes.

Reviewed and Approved

Michael E. Dean Project Manager

MED/an

1 Page Following - Table of Results

American Council of Independent Laboratories International Association of Environmental Testing Laboratories American Association for Laboratory Accreditation

IT ANALYTICAL SERVICES SAN JOSE, CA

Page: 1 of 1
Date: July 31, 1989
Client Project ID: GR #3605, Shell,
1800 Powell Street, Emeryville, CA

Work Order Number: S9-07-061

Lab Sample ID	Client Sample ID	Sample Date	Date Analysis Completed	Sample Condition on Receipt
\$9-07-061-01	s-5	7/7/89	7/21/89	cool pH <2
\$9-07-061-02	s-6	7/7/89	7/21/89	cool pH <2
\$9-07-061-03	s-7	7/7/89	7/21/89	cool pH <2
\$9-07-061-04	s-8	7/7/89	7/20/89	cool pH <2
\$9-07-061-05	S-10	7/7/89	7/21/89	cool pH <2
\$9-07-061-06	SD-8	7/7/89	7/21/89	cool pH >2
\$9-07-061-07	SF-10	7/7/89	7/21/89	cool pH <2
\$9-07-061-08	TRIP BLANK	7/7/89	7/13/89	cool pH <2

Total Petroleum Hydrocarbons - Modified E.P.A. Methods 8015, 8020

ND = None Detected Results - Milligrams per Liter	ND =	None	Detected	Results -	Milligrams pe	er Liter
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Lab Sample ID	Client Sample ID	Low Boiling Hydrocarbons (calculated as Gasoline)	Benzene	Toluene	Ethyl Benzene	Xylenes (total)
S9-07-061-01* Detection Limit	S-5	1.5	0.30	0.008	0.007	0.009
S9-07-061-02*	s-6	3.7	1.7	0.034	0.055	0.20
Detection Limit		0.10	0.001	0.002	0.002	0.006
S9-07-061-03* Detection Limit	s-7	0.07	0.0022 0.0005	ND 0.001	ND 0.001	аи 200.0
S9-07-061-04*	S-8	0.44	0.18	0.005	0.002	0.012
Detection Limit		0.05	0.0005	0.001	0.001	0.003
S9-07-061-05*	s-10	14.	1.3	0.31	0.27	2.4
Detection Limit		2.	0.02	0.05	0.05	0.2
S9-07-061-06* Detection Limit	SD-8	1.4 0.5	0.62 0.005	0.02 0.01	0.01	0.08 0.03
S9-07-061-07	SF-10	ND	ND	ND	ND	ND
Detection Limit		0.05	0.0005	0.001	0.001	0.003
S9-07-061-08	TRIP BLANK	ND	ND	ND	ND	ND
Detection Limit		0.05	0.0005	0.001	0.001	0.003

^{*}Sample contains compounds which caused severe foaming during analysis. A foaming matrix could affect analytical results.

AUTHORIZED	mery ville,	C-44	J-80/58		PHONE NO.	3605	· · · · · · · · · · · · · · · · · · ·
SAMPLE ID	NO. OF CONTAINERS	SAMPLE MATRIX	DATE/TIME SAMPLED	ANALYSIS RE		tA	CONDITION B ID
5-5	3	Liond	7-7/1210	THE (CO.	JBTX	_0K/(ool
5-6			1/1052			' }	
5-7			1-/1127				
2-8			1/4/3		· · · · · · ·		
S-100			10907				
8-02	-						,
SF-10	~	-(/	7:5 / -			1/	
Try black			7:3/			V	
			nco	EIVED BY	1001		
RELINQUISHED B		7-7- 0 01 /		EIVEU BT/	ell -	7-7-89	17:00
RELINOUSHED B	7 0	7-7- 6 4_/_	ILLO	EIVED BY:			
RELINQUISHED B		0-69 11:	15 an	EIVED BY LAB:	\sim \sim		
				sephine_	Delail): 7/10/2	19 11:27
DECICNATED I AS	BORATORY:	IT SC	<u>v</u> ()	DHS#:	137		_
DESIGNATED DV							<u> </u>
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	Jorna	1-1-47				··	

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