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7150 Island Queen Dr.
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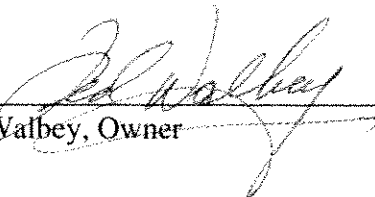
3/14/07

Mr. Barney Chan
Alameda County Health Care Services Agency
Environmental Protection Division
1131 Harbor Bay Parkway, Suite 250
Alameda, CA 94502-6577

Re: Perjury Statement
Former Fiesta Beverages Facility
966 89th Avenue
Oakland, California
ACDEH Fuel Leak Site # RO0000314

Dear Mr Chan,

"I declare under penalty of perjury, that the information and / or recommendations contained in the attached proposal or report is true and correct to the best of my knowledge."



Ted Walbey, Owner

First Quarter 2007 Groundwater Monitoring Event

Former Fiesta Beverages Facility
966 89th Avenue
Oakland, California 94621
ACHCSA Fuel Leak Site # RO0000314

March 8, 2007
BEI Job No. 203004

Prepared for:

Mr. Ted Walbey
Fiesta Beverages
7150 Island Queen Dr.
Sparks, NV 89436

Prepared by:

Blymyer Engineers, Inc.
1829 Clement Avenue
Alameda, CA 94501-1395
(510) 521-3773

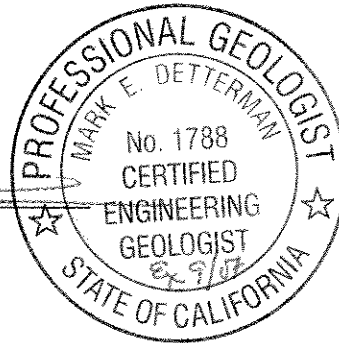
Limitations

Services performed by Blymyer Engineers, Inc. have been provided in accordance with generally accepted professional practices for the nature and conditions of similar work completed in the same or similar localities, at the time the work was performed. The scope of work for the project was conducted within the limitations prescribed by the client. This report is not meant to represent a legal opinion. No other warranty, expressed or implied, is made. This report was prepared for the sole use of the client, Fiesta Beverages.

Blymyer Engineers, Inc.

By:


Mark E. Detterman, CEG
Senior Geologist



And:



Michael S. Lewis, REA
Vice President, Technical Services

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1.0 Introduction and Background

In August 1990, one 500-gallon and one 1,000-gallon gasoline underground storage tanks (USTs) were removed from the subject site (Figures 1 and 2). Soil and groundwater were reported to be impacted from releases from one or both USTs. Overexcavation of the former UST basins occurred in January 1991. The excavations were reported to have reached approximately 15 feet by 8 feet by 14 feet deep and 12 feet by 7 feet by 14 feet deep, respectively, on January 14, 1991. Beginning in April 1991, aeration of the soil occurred onsite. In April 1993, 74.28 tons of soil were transported to the Remco recycling facility.

In June 1993, groundwater monitoring wells MW-1, MW-2, and MW-3 were installed. In general, the wells encountered black to grey to light brown clay to a depth of approximately 15 below grade surface (bgs). At 15 feet bgs, the three bores encountered a 0.5- to 2.0-foot-thick clayey sand. Below this unit a light brown to grey clay was present to a depth of 18 to 21 feet bgs. Underneath this unit, a 1- to 3-foot-thick sand was encountered in bores MW-1 and MW-2, while a clayey silt was encountered in bore MW-3. Below approximately 21 feet bgs, a green-grey or black clay was encountered to the full explored depth of 26.5 feet bgs in bore MW-1 and to 25 feet bgs in bores MW-2 and MW-3. Saturated soil was encountered below a depth of approximately 13 feet bgs (in clay overlaying the uppermost sand unit). The wells were installed with a screened interval between 10 and 25 feet bgs. Groundwater from the three wells was sampled six times between August 1993 and December 1998.

In November 1999, after obtaining appropriate permits, AllCal Property Services, Inc. (AllCal) installed four Geoprobe⁷ soil bores downgradient from the former location of the two USTs. The bores were installed in the public right-of-way across 89th Avenue from the subject site, in an unpaved portion of the roadway. Soil bores SB-1 and SB-2 were logged to a depth of 16 feet bgs. Silty clay was encountered to a depth of approximately 13 to 14 feet bgs. Below that depth, soil consisted of clayey silt that alternated between moist and saturated for several vertical feet. Bore SB-1 also encountered a poorly graded sand at 16 feet bgs. Hydrocarbon odors were present in both bores at a depth of approximately 6 feet bgs and green discolored soil was present at 10 feet bgs in bore SB-1. Discolored soil and gasoline odors were noted in both bores throughout the clayey silt, while brownish colored clay was present in both

bores just above the silt. The groundwater interface appears to have been encountered at an approximate depth of 16 feet bgs in the sand. A sheen was noted at that depth in SB-1. Groundwater samples were obtained from bores SB-1 and SB-2 after pushing the Geoprobe⁷ system to a total depth of 18 feet bgs. Soil bores SB-3 and SB-4 were directly pushed to a total depth of 18 feet bgs in order to obtain grab groundwater samples. Groundwater samples from bores SB-1 and SB-2 contained elevated concentrations of Total Petroleum Hydrocarbons (TPH) as gasoline, and benzene, toluene, ethylbenzene, and total xylenes (BTEX). Significantly lower concentrations of TPH as gasoline and total xylenes were encountered in the groundwater sample from soil bore SB-3, while all analytes were nondetectable in groundwater collected from soil bore SB-4. No soil samples were submitted for laboratory analysis from the four Geoprobe⁷ bores.

After the review of the January 2001 groundwater monitoring report, the Alameda County Health Care Services Agency (ACHCSA) approved the application of a 7% solution of hydrogen peroxide to the wells in an attempt to remediate dissolved constituents. On March 7, 2001, the solution was applied by AllCal and on April 25, 2001, a groundwater monitoring event was conducted to determine if a reduction in dissolved constituents had occurred. Based on the analytical data, a reduction was seen in wells MW-1 and MW-2, with some reductions also seen in well MW-3. This sampling event and subsequent interpretation was complicated by the presumed mis-marking of samples from wells MW-1 and MW-3. No further work at the site is known to have occurred between April 2001 and the March 2003 groundwater monitoring event.

On January 16, 2003, a new case manager, Mr. Amir Gholami, was appointed by the ACHCSA. On September 17, 2003, a workplan for a Geoprobe⁷ investigation of the site was submitted to the ACHCSA. The intent was to attempt to determine the lateral and vertical extent of impacted soil and groundwater in order to better target the residual contamination in future remedial actions to be determined.

Due to the lack of a response from the ACHCSA, on February 17, 2004, Blymyer Engineers issued a *Letter of Intent to Proceed: Geoprobe⁷ Investigation*.

The *Fourth Quarter 2003 Groundwater Monitoring Event* report, dated January 6, 2004, recommended that analysis for fuel oxygenates by EPA Method 8260B be eliminated from the analytical

program. It was reasoned that the data generated to date had been very consistent, and further quantification would not significantly add to the level of understanding at the site. Additionally, the concentration of methyl *tert*-butyl ether (MTBE) can be monitored using EPA Method 8021B for no additional cost, and the resultant concentration of MTBE can be used as a proxy for the approximate concentration of the remaining fuel oxygenates. Based on the lack of response from the ACHCSA, it has been presumed that this was found reasonable and acceptable.

On March 15, 2004, Blymyer Engineers issued a letter entitled *Recommendation for Reduction of Groundwater Monitoring* that provided additional rationale for decreasing the groundwater sampling interval from quarterly to semi-annually. It argued that generation of quarterly analytical data would not significantly improve the level of understanding of impacts to the subsurface at the site, and recommended a reduction of the sampling interval to semi-annual. Based on the lack of response from the ACHCSA, it has been presumed that this was found reasonable and acceptable.

On December 14, 2004, Blymyer Engineers issued to the ACHCSA the *Report on a Geoprobe® Subsurface Investigation* which documented the installation of nine Geoprobe® soil bores at the site. The work further refined the known lateral and vertical extent of soil impacted by the petroleum release at the site. Grab groundwater samples in the upgradient and the eastern cross-gradient directions defined all petroleum compounds in groundwater to concentrations below the San Francisco Bay Regional Water Quality Control Board (RWQCB) Environmental Screening Levels (ESLs). Grab groundwater samples in the downgradient and western cross-gradient directions were unable to define most petroleum compounds to concentrations below the RWQCB ESLs. The installation of additional permanent groundwater monitoring wells was recommended as appropriate at the site in order to allow for groundwater sampling from a “repeatedly accessed location”. It was reasoned that data generated from these locations will assist in determining appropriate remedial actions, and in monitoring remedial progress.

On July 6, 2005, the new case manager for the ACHCSA, Mr. Barney Chan, issued the letter *Fuel Leak Case RO0000314* commenting on the December 14, 2004 report. The ACHCSA determined that the collection of additional data is needed to progress the site towards closure. The letter requested a workplan to clear well MW-1 of several feet of sediment due to the potential for groundwater gradient

biasing, requested further definition of the groundwater and soil plumes through the installation of additional wells and soil bores, requested a conduit study, and requested a Feasibility Study and Remedial Action Plan.

Blymyer Engineers submitted the *Workplan for Remedial Investigation / Feasibility Study*, on October 10, 2005. The Workplan detailed the procedures for the collection of Remediation by Natural Attenuation (RNA) analytical parameters from existing wells as an initial phase of a Remedial Investigation / Feasibility Study (RI/FS), as well as the installation of four additional groundwater monitoring wells, and the destruction and reinstallation of groundwater monitoring well MW-1. On November 18, 2005, the ACHCSA issued the letter *Fuel Leak Case RO0000314* commenting on the Workplan. The ACHCSA requested the following:

- The addition of two wells at specified locations for further plume characterization,
- Use of a maximum of 10 feet of screen in the wells,
- Confirmation of the presence of MTBE by EPA Method 8260 if MTBE concentrations rose significantly, and
- Collection of the RNA parameters.

The ACHCSA requested confirmation that the additional wells would be added by December 19, 2005, and that a RI/FS report would be submitted by February 19, 2006. Confirmation that the additional wells would be included was provided by telephone in December 2005; however, permitting issues delayed installation of the wells. The *Remedial Investigation / Feasibility Study Report* (RI/FS Report), dated September 8, 2006, was submitted to ACHCSA on October 6, 2006.

The RI/FS report documented the destruction of well MW-1, the installation of replacement well MW-1R, and the installation of wells MW-4 through MW-9. The soil and groundwater data collected in the effort achieved vertical delineation, as well as upgradient, lateral, and downgradient delineation of all hydrocarbon compounds in soil and groundwater, with the exception of MTBE in groundwater. MTBE was delineated to below the MCL and the *non-drinking water* ESL goal for the compound, but was

slightly above the *drinking water* goal. Because the site is in an area that is not known to extensively use groundwater as a drinking water source, the numeric remedial goals were predominantly compared to the *non-drinking water* ESL goals; however, the ACHCSA may ultimately apply *drinking water* ESL goals to remedial efforts at the site.

Higher concentrations of TPH as gasoline appear to be relatively isolated near the former source (MW-1, MW-1R, GP-5, and GP-2; the latter based on PID results only). The presence of slightly higher concentrations at GP-6 or GP-8 likely indicates lateral migration through the clay units in the vadose zone in very thin, interbedded coarser grained deposits with more permeability and porosity. A conduit survey indicated that, due to depth of burial, the utility corridors do not appear to be acting as significant conduits in the site vicinity for groundwater movement and therefore contaminant migration. A notable decrease in analyte concentrations in soil is apparent with increasing depth. Generic *non-drinking water* ESL goals for soil were not exceeded for any compound beneath approximately 12 feet bgs.

The distribution of nitrate, methane and dissolved oxygen indicate that the TPH as gasoline groundwater plume is undergoing anaerobic degradation. Specifically, the elevated concentrations of nitrate observed in perimeter wells MW-4 through MW-9, in comparison to the concentration of nitrate in plume core wells MW-1/1R, MW-2 and MW-3, where the concentration is reduced to essentially one-half of its perimeter levels, and the correspondingly high methane concentrations in the plume core area suggest that active anaerobic degradation is occurring. The source of nitrate is likely leaking sewer lines located along 89th Avenue.

For the site as a whole, the limited area of hydrocarbon degradation suggested by the RNA data, collectively with the laboratory notes indicating relatively unmodified gasoline range hydrocarbons are present in soil and groundwater samples, and the continued recontamination of groundwater documented by graphs depicted on Figures 10 through 13 of the *RI/FS Report*, appear to document a release that is undergoing anaerobic microbial degradation, that RNA is oxygen limited, has reached stability with the surrounding area, and will not progress significantly further without remedial efforts.

Six potential remedial options were evaluated for appropriateness at the site; monitored natural attenuation (MNA), groundwater pump and treat, enhanced insitu bioremediation (EIB), air sparging-vapor recovery (ASVR), dual phase extraction, and insitu chemical oxidation (ISCO). A combination of EIB and ISCO was selected as the most appropriate remedial technology for the site due to multiple factors. ISCO was selected for the vicinity of the former tank excavation and would consist of the injection of the commercial oxidation product RegenOx. Chemical oxidation of residual source soil and groundwater containing higher hydrocarbon concentrations is anticipated to eliminate potential residual free-phase hydrocarbons in the tank vicinity. EIB using Oxygen Releasing Compound Advanced (ORC Advanced) was selected for the larger area around and downgradient of the former tank location. Petroleum hydrocarbon compounds are recognized to degrade favorably and rapidly under aerobic (oxygen rich) conditions. To stimulate aerobic bacterial activity and increase the rate of biodegradation within the hydrocarbon plume, non-toxic inorganic chemicals (bionutrients) can be added to the groundwater that release oxygen, nitrogen and phosphate, such as ORC Advanced and bionutrient compounds (typically, nitrogen/phosphorus/potassium (NPK) fertilizer). At sites where stagnant hydrocarbon plumes are present, one or more of the essential bionutrient elements is commonly depleted, and natural attenuation of the hydrocarbon plume due to microbial activity ceases. By determining a site's "bio-needs," the missing elements can be injected into the hydrocarbon plume to boost bioactivity.

At the site, dissolved oxygen in groundwater is depleted to less than 1 mg/L, and based on available information the lack of dissolved oxygen is the limiting factor retarding current biological activity. For EIB, the supply of bio-nutrients is assessed prior to and during remediation. During the course of remediation, if nutrient concentrations are found to be inadequate, then further nutrient addition is performed.

On December 18, 2006, the ACHCSA issued a letter indicating that it was in agreement with the proposed plan of action, namely EIB with localized ISCO, using a combination of ORC Advanced and RegenOx, respectively. The December 18, 2006 letter requested an interim corrective action plan (ICAP) by January 19, 2007, and quarterly monitoring reports by January 30, and April 30, 2007. A request for deadline extension was later submitted to, and approved by, the ACHCSA. The *Interim Corrective Action Plan* was submitted on February 7, 2007, and is pending review by the ACHCSA.

2.0 Groundwater Sample Collection and Analytical Methods

This report documents the First Quarter 2007 groundwater monitoring event at the subject site (Figure 1). This is the first quarterly groundwater monitoring event conducted by Blymyer Engineers, Inc. at the former Fiesta Beverage site in Oakland, California since the ACHCSA requested that the groundwater monitoring interval return to a quarterly interval in their January 2, 2007 letter.

Groundwater samples were collected from monitoring wells MW-1R and MW-2 through MW-9 on February 19, 2007. The groundwater samples were collected by Blaine Tech Services, Inc. (Blaine) in accordance with Blaine *Standard Operating Procedures* for groundwater gauging, purging, and sampling.

A copy is included as Appendix A. Depth to groundwater was measured in all wells at the site. Temperature, pH, conductivity, and turbidity were measured initially, and then after removal of each of three well casing volumes for each well. Dissolved Oxygen (DO) and the Oxygen-Reduction Potential (ORP) were measured prior to purging groundwater. These measurements are generally useful in determining if an adequate supply of oxygen is present in groundwater to allow microbial growth. The groundwater depth measurements and details of the monitoring well purging and sampling are presented on the *Well Monitoring Data Sheets* and *Well Gauging Data Sheets* generated by Blaine and included as Appendix B. Depth-to-groundwater measurements are presented in Table I. All purge and decontamination water was temporarily stored in a Department of Transportation-approved 55-gallon drum for future disposal by the owner.

The groundwater samples were analyzed by McCampbell Analytical, Inc., a California-certified laboratory, on a 5-day turnaround time. Groundwater samples were analyzed for Total Petroleum Hydrocarbons (TPH) as gasoline by Modified EPA Method 8015; and benzene, toluene, ethylbenzene, and total xylenes (BTEX) and MTBE by EPA Method 8021B. Tables II to V summarize current and previous analytical results for groundwater samples. The laboratory analytical report for the current sampling event is included as Appendix C.

3.0 Groundwater Flow Data and Groundwater Sample Analytical Results

Previously surveyed top-of-casing (TOC) elevations were used to construct a groundwater gradient map (Figure 2). Groundwater depths during this monitoring event ranged between 7.77 to 8.61 feet below the top of the casings. Depth to groundwater decreased an average of 0.43 feet. Groundwater predominantly appears to flow to the west during this event, and is generally consistent with historical data, although apparent southern and eastern flow directions have been observed previously. The average groundwater gradient was calculated to be at 0.01 feet/foot for the current monitoring event.

Concentrations of all of the chemical compounds related to gasoline were present in wells MW-1R, MW-2, and MW-3 during the current quarter. Except for MTBE in well MW-5, wells MW-5 through MW-9 did not yield detectable concentrations of petroleum hydrocarbons. With respect to the previous groundwater sampling event in June 2006, concentrations in well MW-2 were generally lower, concentrations in MW-3 were generally higher, and concentrations in MW-1R were generally higher, although more mixed than in comparison to concentrations in well MW-3. Each contained TPH as gasoline and BTEX. While concentrations in each of these wells has increased or decreased, the concentrations are within the range of previous concentrations obtained from groundwater samples collected from wells MW-2 and MW-3. The continued fluctuation in results in the wells suggests a mobilization of residual contamination from soil to groundwater at the site. This is only the second sampling of well MW-1R.

Plots of TPH-G or benzene and groundwater elevation vs. time were generated for well MW-3 (Figures 3 and 4). There is a generally good correlation between a rise in groundwater elevation and a rise in the concentration of both TPH-G and benzene. Previously generated graphs for well MW-1 have documented a good correlation between a rise in groundwater elevation and a decrease in the concentration of both TPH-G and benzene. Consequently there appears to be both recontamination of groundwater upon rise up into contaminated soil as well as drainage from soil to groundwater as groundwater drops in elevation at the site and in the different wells.

The concentration of TPH as gasoline ranged from nondetectable to 280 micrograms per liter (Fg/L) in

well MW-3. Benzene ranged between a concentration of nondetectable and 49 Fg/L (well MW-3). Toluene was present up to a concentration of 11 Fg/L, ethylbenzene up to 18 Fg/L, and total xylenes to up to 23 Fg/L (all in well MW-3).

As is typical, the concentration of benzene in groundwater exceeded the drinking water Maximum Contaminant Level (MCL) in plume core wells MW-1R, MW-2, and MW-3 during the current monitoring and sampling event; however, Blymyer Engineers does not believe that groundwater at this location should be considered as drinking water. Consequently, Blymyer Engineers also references the Environmental Screening Levels (ESL) promulgated by the RWQCB in Table II. Only the RWQCB provides a look-up value for TPH, and for a non-drinking water designation of groundwater.

At the request of the ACHCSA, four quarters of groundwater samples have previously been analyzed for the fuel oxygenates di-isopropyl ether (DIPE), ethyl *tert*-butyl ether (ETBE), methyl *tert*-butyl ether (MTBE), *tert*-amyl methyl ether (TAME), and *tert*-butyl alcohol (TBA), by EPA Method 8260B. Ethanol and methanol have also been analyzed since, and were nondetectable. Due to the consistency of the data, analysis by this EPA method was eliminated as an unnecessary expense. Using EPA Method 8021B, MTBE was detected in wells MW-2 and MW-5, and at a concentration of 7.1 and 5.6 Fg/L, respectively. Although not detected, it is likely that TAME is also present, such as was documented in the June 2003 sampling event (Table III). Of the fuel oxygenates, only MTBE has an MCL, listed at 13 Fg/L.

4.0 Intrinsic Bioremediation Groundwater Sample Analytical Results

Intrinsic bioremediation laboratory analytical parameters were not collected during the current quarter; however, pre-purge field parameters were collected. Tables IV and V present the analytical results of current and previous RNA indicator parameters. Microbial use of petroleum hydrocarbons as a food source is affected by the concentration of a number of chemical compounds dissolved in groundwater at a site. RNA monitoring parameters were established by research conducted by the Air Force Center for Environmental Excellence. The research results were used to develop a technical protocol for documenting RNA in groundwater at petroleum hydrocarbon release sites (Wiedemeier, Wilson, Kampbell, Miller and Hansen, 1995, *Technical Protocol for Implementing the Intrinsic Remediation with Long Term Monitoring for Natural Attenuation of Fuel Contamination Dissolved in Groundwater, Volumes I and II*, U.S. Air Force Center for Environmental Excellence, Brooks Air Force Base, Texas). The protocol focuses on documenting both aerobic and anaerobic degradation processes whereby indigenous subsurface bacteria use various dissolved electron acceptors to degrade dissolved petroleum hydrocarbons.

In the order of preference, the following electron acceptors and metabolic by-products are used and generated, respectively, by the subsurface microbes (aerobes, Mn – Fe reducers, and methanogens) to degrade petroleum hydrocarbons: oxygen to carbon dioxide, nitrate to nitrogen, insoluble manganese (Mn^{4+}) to soluble manganese (Mn^{2+}), insoluble ferric iron (Fe^{3+}) to soluble ferrous iron (Fe^{2+}), sulfate to hydrogen sulfide, and carbon dioxide to methane. With the exception of oxygen, the use of all other electron acceptor pathways by microbes indicates increasingly anaerobic degradation. Aerobic degradation takes place first, and oxygen inhibits anaerobic degradation. As oxygen is consumed and an anoxic zone develops, the Mn – Fe reducers and methanogens begin to grow and release dissolved Mn, dissolved Fe, and methane (Commission on Geosciences, Environment and Resources, *Natural Attenuation for Groundwater Remediation*, 2000). Investigation of each of these electron acceptor pathways was conducted in selected wells at the site as part of the evaluation of RNA chemical parameters. Previous analytical results appear to have documented oxygen and nutrient (nitrate) limited natural biodegradation at the site.

Microbial use of petroleum hydrocarbons as a food source is principally affected by the concentration of dissolved oxygen (DO) in the groundwater present at a site; it is the preferred electron acceptor for the biodegradation of hydrocarbons. DO was present in pre-purge groundwater in concentrations ranging from 0.08 milligrams per liter (mg/L) to 1.98 mg/L. Because standard purge and sample techniques were used in the current quarter in order to decrease labor and equipment costs, only pre-purge DO concentrations were collected due to the elevated potential to entrain oxygen in groundwater during sampling. There is not a reasonable correlation between plume core and plume perimeter wells with respect to DO with the pre-purge readings; however, based on available information the lack of dissolved oxygen is the limiting factor retarding current biological activity.

ORP is another measure of the supply and use of oxygen at a site. The higher the reading in millivolts (mV), the more oxygenated the subsurface environment is, and the lower the readings, the more anaerobic or reducing the subsurface environment is. Due to the pre-purge collection of ORP, a strong correlation is again not observed between plume interior or plume perimeter wells.

For a more in-depth review of recent RNA parameters, please refer to the *Remedial Investigation / Feasibility Study Report*, dated September 8, 2006.

5.0 Conclusions and Recommendations

The following conclusions were generated from the available data discussed above:

- Concentrations of hydrocarbons were present in plume core wells MW-1R, MW-2, and MW-3 this monitoring period. Concentrations generally increased in wells MW-1R and MW-3 and generally decreased in well MW-2. Except for MTBE, detected below the MCL in well MW-5, all fuel petroleum hydrocarbons were not detected above the limits of detection in wells MW-4 through MW-9. Concentrations in the wells are within historic concentration ranges.
- TPH as gasoline and benzene, were over the generic RWQCB ESLs for groundwater (non-drinking water) in wells MW-1R and MW-3, and benzene was over the ESL in well MW-2.
- MTBE was detected in wells MW-2 and MW-5, at a concentrations of 7.1 and 5.6 Fg/L, respectively. TAME has previously been detected in groundwater at the site, thus it is likely that it is present beneath the site. Of fuel oxygenates, only MTBE has an MCL, listed at 13 Fg/L.
- Only DO and ORP of the RNA chemical parameters suite were investigated in order to help determine if biological degradation of the petroleum hydrocarbons at the site is present. Because they were collected prior to well purging due to the use of standard well purging and sampling techniques, good correlations were not observed. Previously, natural degradation of the fuel hydrocarbons beneath the site has been observed to be oxygen limited.
- Groundwater flow appears to be towards the west and the average groundwater gradient was calculated at 0.01 feet/foot for this monitoring event.

The following recommendations were generated from the available data discussed above:

- \$ The next quarterly groundwater sampling event should occur in May 2007.
- \$ Collection of all RNA indicator data should be discontinued as a modest cost saving measure. The collection of additional data is unlikely to contribute to the understanding of biodegradation beneath the site. Collection of RNA indicator data could be resumed thereafter should a need be

documented.

\$ Interim corrective actions should be conducted in accordance with the approved ICAP (approval pending). Future quarterly monitoring events should be used to gauge the effectiveness of corrective actions.

\$ A copy of this letter report should be forwarded to:

Mr. Barney Chan
Alameda County Health Care Services Agency
Environmental Protection Division
1131 Harbor Bay Parkway, Suite 250
Alameda, CA 94502-6577

Tables

**Table I, Summary of Groundwater Elevation Measurements
BEI Job No. 203004, Former Fiesta Beverage
966 89th Avenue, Oakland, California**

Well ID	Date	TOC Elevation (feet)	Depth to Water (feet)	Water Surface Elevation (feet)
MW-1	8/6/1993	18.72	8.96	9.76
	1/12/1996		8.55	10.17
	4/16/1996		7.65	11.07
	7/15/1996		8.76	9.96
	10/16/1996		9.04	9.68
	12/15/1998		8.38	10.34
	1/18/2001		8.49	10.23
	4/25/2001		8.24	10.48
	3/17/03*		8.08	10.64
	6/23/2003		8.63	10.09
	9/18/2003		8.90	9.82
	12/15/2003		8.15	10.57
	6/15/2004		8.67	10.05
	12/15/2004		7.99	10.73
	6/29/2005		7.88	10.84
	5/8/2006		21.70	Destroyed
	2/19/2007	Destroyed		Destroyed

**Table I, Summary of Groundwater Elevation Measurements
BEI Job No. 203004, Former Fiesta Beverage
966 89th Avenue, Oakland, California**

Well ID	Date	TOC Elevation (feet)	Depth to Water (feet)	Water Surface Elevation (feet)
MW-2	8/6/1993	18.44	8.68	9.76
	1/12/1996		8.24	10.20
	4/16/1996		7.41	11.03
	7/15/1996		8.45	9.99
	10/16/1996		8.73	9.71
	12/15/1998		8.05	10.39
	1/18/2001		8.24	10.20
	4/25/2001		7.88	10.56
	3/17/03*		7.08	11.36
	6/23/2003		8.90	9.54
	9/18/2003		8.61	9.83
	12/15/2003		7.97	10.47
	6/15/2004		8.42	10.02
	12/15/2004		8.00	10.44
	6/29/2005		9.51	8.93
	6/12/2006		21.45	8.25
	2/19/2007	8.12		13.33

**Table I, Summary of Groundwater Elevation Measurements
BEI Job No. 203004, Former Fiesta Beverage
966 89th Avenue, Oakland, California**

Well ID	Date	TOC Elevation (feet)	Depth to Water (feet)	Water Surface Elevation (feet)
MW-3	8/6/1993	19.01	9.07	9.94
	1/12/1996		8.65	10.36
	4/16/1996		7.82	11.19
	7/15/1996		8.88	10.13
	10/16/1996		9.16	9.85
	12/15/1998		8.45	10.56
	1/18/2001		8.57	10.44
	4/25/2001		8.29	10.72
	3/17/03*		8.50	10.51
	6/23/2003		9.05	9.96
	9/18/2003		9.11	9.90
	12/15/2003		8.03	10.98
	6/15/2004		8.85	10.16
	12/15/2004		8.84	10.17
	6/29/2005		9.00	10.01
	6/12/2006		22.02	8.62
	2/19/2007	8.12		13.90

**Table I, Summary of Groundwater Elevation Measurements
BEI Job No. 203004, Former Fiesta Beverage
966 89th Avenue, Oakland, California**

Well ID	Date	TOC Elevation (feet)	Depth to Water (feet)	Water Surface Elevation (feet)
MW-1R	6/12/2006	21.73	8.49	13.24
	2/19/2007		7.94	13.79
MW-4	6/12/2006	21.34	8.37	12.97
	2/19/2007		7.77	13.57
MW-5	6/12/2006	22.53	8.75	13.78
	2/19/2007		8.61	13.92
MW-6	6/12/2006	21.97	8.59	13.38
	2/19/2007		7.93	14.04
MW-7	6/12/2006	21.21	8.31	12.90
	2/19/2007		7.85	13.36
MW-8	6/12/2006	20.97	8.37	12.60
	2/19/2007		7.99	12.98
MW-9	6/12/2006	20.98	8.50	12.48
	2/19/2007		8.08	12.90

Notes:

TOC = Top of Casing

* = Initial data set collected under direction of Blymyer Engineers, Inc.

NM = Not measured

¹ = Resurveyed on February 7, or June 22, 2006 by CSS Environmental Services, Inc.

Elevations in feet above mean sea level

Table II, Summary of Groundwater Sample Hydrocarbon Analytical Results
BEI Job No. 203004, Former Fiesta Beverage
966 89th Avenue, Oakland, California

Well ID	Sample Date	Modified EPA Method 8015 (µg/L)	EPA Method 8020 or 8021B (µg/L)				
		TPH as Gasoline	Benzene	Toluene	Ethylbenzene	Total Xylenes	MTBE
MCL		N/A	1	150	700	1,750	13
<i>Drinking Water Source</i> ¹		100	1	40	30	20	5
<i>Non-Drinking Water Source</i> ²		500	46	130	290	100	1,800
MW-1	8/6/1993	17,000	7.1	8.4	9.2	53	NA
	1/12/1996	12,000	1,900	840	370	1,100	NA
	4/16/1996	3,500	700	55	100	180	NA
	7/15/1996	11,000	2,300	450	350	910	NA
	10/16/1996	21,000	4,200	2,200	650	2,600	NA
	12/15/1998	10,000	1,800	520	270	1,100	<350
	1/18/2001	11,000^a	2,000	320	320	1,100	<120
	4/25/2001	2,100^{a, c}	270	46	59	130	<5.0
	3/17/2003*	2,200^a	260	19	36	54	NA ^d
	6/23/2003	6,100^a	930	53	99	200	NA
	9/18/2003	3,800^a	660	13	24	34	NA
	12/15/2003	260^a	19	1.1	<0.5	1.5	NA
	6/15/2004	5,200^a	520	13	38	39	<50
	12/15/2004	2,400^a	370	8.2	13	14	<15
	6/29/2005	5,500^a	750	27	94	140	<100
	5/8/2006	Destroyed	Destroyed	Destroyed	Destroyed	Destroyed	Destroyed
2/19/2007	Destroyed	Destroyed	Destroyed	Destroyed	Destroyed	Destroyed	Destroyed

Table II, Summary of Groundwater Sample Hydrocarbon Analytical Results
BEI Job No. 203004, Former Fiesta Beverage
966 89th Avenue, Oakland, California

Well ID	Sample Date	Modified EPA Method 8015 (µg/L)	EPA Method 8020 or 8021B (µg/L)				
		TPH as Gasoline	Benzene	Toluene	Ethylbenzene	Total Xylenes	MTBE
MCL		N/A	1	150	700	1,750	13
<i>Drinking Water Source</i> ¹		100	1	40	30	20	5
<i>Non-Drinking Water Source</i> ²		500	46	130	290	100	1,800
MW-2	8/6/1993	2,700	1	2	2	8	NA
	1/12/1996	2,700	600	310	94	220	NA
	4/16/1996	190	39	11	10	14	NA
	7/15/1996	700	160	33	34	48	NA
	10/16/1996	190	48	8	10	13	NA
	12/15/1998	200	62	17	5	14	4.4^b
	1/18/2001	300^a	74	26	7	21	7.3
	4/25/2001	<50 ^c	5	2	1	2	<5.0
	3/17/2003*	78^a	26	3	2	4	NA ^d
	6/23/2003	160^a	51	2	1	2	NA
	9/18/2003	<50	2	<0.5	<0.5	<0.5	NA
	12/15/2003	<50	12	<0.5	<0.5	<0.5	NA
	6/15/2004	95^a	15	1.3	2	1	<30
	12/15/2004	<50	11	0.97	1	1	7.8
	6/29/2005	130	29	2	3	3	6.7
	6/13/2006	150^a	59	3	3.4	2.7	11
2/19/2007	51^a	8	1.6	1	2.8	7.1	

Table II, Summary of Groundwater Sample Hydrocarbon Analytical Results
BEI Job No. 203004, Former Fiesta Beverage
966 89th Avenue, Oakland, California

Well ID	Sample Date	Modified EPA Method 8015 (µg/L)	EPA Method 8020 or 8021B (µg/L)				
		TPH as Gasoline	Benzene	Toluene	Ethylbenzene	Total Xylenes	MTBE
MCL		N/A	1	150	700	1,750	13
<i>Drinking Water Source</i> ¹		100	1	40	30	20	5
<i>Non-Drinking Water Source</i> ²		500	46	130	290	100	1,800
MW-3	8/6/1993	5,200	2.1	2.9	3.6	17	NA
	1/12/1996	4,500	280	180	120	470	NA
	4/16/1996	5,400	370	340	160	580	NA
	7/15/1996	1,800	200	220	66	250	NA
	10/16/1996	2,000	340	140	100	300	NA
	12/15/1998	1,400	200	39	72	150	<22
	1/18/2001	1,800^a	240	41	86	120	<10
	4/25/2001	8,300^{a, c}	300	330	200	1,100	<20
	3/17/2003*	2,100^a	240	78	10	280	NA ^d
	6/23/2003	<50	2.5	0.6	0.69	1.4	NA
	9/18/2003	<50	<0.5	<0.5	<0.5	<0.5	NA
	12/15/2003	2,400	300	120	140	260	NA
	6/15/2004	<50	1.1	<0.5	<0.5	<0.5	6.2
	12/15/2004	1,600^a	140	83	83	230	<15
	6/29/2005	230^a	27	6.1	7.2	15	<15
	6/13/2006	68^a	3.1	1.8	<0.5	<0.5	<5.0
2/19/2007	280^a	49	11	18	23	<5.0	

Table II, Summary of Groundwater Sample Hydrocarbon Analytical Results
BEI Job No. 203004, Former Fiesta Beverage
966 89th Avenue, Oakland, California

Well ID	Sample Date	Modified EPA Method 8015 (µg/L)	EPA Method 8020 or 8021B (µg/L)				
		TPH as Gasoline	Benzene	Toluene	Ethylbenzene	Total Xylenes	MTBE
MCL		N/A	1	150	700	1,750	13
<i>Drinking Water Source</i> ¹		100	1	40	30	20	5
<i>Non-Drinking Water Source</i> ²		500	46	130	290	100	1,800
MW-1R	6/13/2006	90^a	24	<0.5	<0.5	1.9	7.0
	2/19/2007	200^a	8	0.80	12	8.7	<5.0
MW-4	6/12/2006	<50	<0.5	<0.5	<0.5	<0.5	5.7
	2/19/2007	<50	<0.5	<0.5	<0.5	<0.5	<5.0
MW-5	6/12/2006	<50	<0.5	<0.5	<0.5	<0.5	<5.0
	2/19/2007	<50	<0.5	<0.5	<0.5	<0.5	5.6
MW-6	6/13/2006	<50	<0.5	<0.5	<0.5	<0.5	<5.0
	2/19/2007	<50	<0.5	<0.5	<0.5	<0.5	<5.0
MW-7	6/12/2006	<50	<0.5	<0.5	<0.5	<0.5	<5.0
	2/19/2007	<50	<0.5	<0.5	<0.5	<0.5	<5.0
MW-8	6/12/2006	<50	<0.5	<0.5	<0.5	<0.5	<5.0
	2/19/2007	<50	<0.5	<0.5	<0.5	<0.5	<5.0
MW-9	6/12/2006	<50	<0.5	<0.5	<0.5	<0.5	5.6
	2/19/2007	<50	<0.5	<0.5	<0.5	<0.5	<5.0

**Table II, Summary of Groundwater Sample Hydrocarbon Analytical Results
BEI Job No. 203004, Former Fiesta Beverage
966 89th Avenue, Oakland, California**

Well ID	Sample Date	Modified EPA Method 8015 (µg/L)	EPA Method 8020 or 8021B (µg/L)				
		TPH as Gasoline	Benzene	Toluene	Ethylbenzene	Total Xylenes	MTBE
MCL		N/A	1	150	700	1,750	13
<i>Drinking Water Source</i> ¹		100	1	40	30	20	5
<i>Non-Drinking Water Source</i> ²		500	46	130	290	100	1,800

- Notes:
- ug/L = micrograms per liter
 - TPH = Total Petroleum Hydrocarbons
 - EPA = Environmental Protection Agency
 - MTBE = Methyl *tert*-Butyl Ether
 - ¹ = From Table A; RWQCB Environmental Screening Levels (ESLs); Groundwater IS a Current or Potential Source of Drinking Water
 - ² = From Table B; RWQCB Environmental Screening Levels (ESLs); Groundwater IS NOT a Current or Potential Source of Drinking Water
 - RWQCB = California Regional Water Quality Control Board, San Francisco Bay Region
 - ESL = Environmental Screening Level
 - N/A = Not applicable
 - NA = Not analyzed
 - RBSL = Risk Based Screening Level
 - <x = Analyte not detected at reporting limit x
 - * = Initial data set collected under direction of Blymyer Engineers, Inc.
 - ^a = Laboratory note indicates the unmodified or weakly modified gasoline is significant.
 - ^b = Confirmed with EPA Method 8260.
 - ^c = Groundwater samples for MW-1 and MW-3 suspected to have been switched (mismarked) in field. First collection of groundwater samples after application of Hydrogen Peroxide on March 7, 2001.
 - ^d = Analysis conducted by EPA Method 8260. See Table III.

Bold results indicate detectable analyte concentrations.

Note: Shaded cell indicates that detected concentration exceeds ESL

Table III, Summary of Groundwater Sample Fuel Oxygenate Analytical Results
BEI Job No. 203004, Former Fiesta Beverage
966 89th Avenue, Oakland, California

Well ID	Sample Date	EPA Method 8260B (ug/L)								
		TAME	TBA	EBD	1,2-DCA	DIPE	Ethanol	ETBE	Methanol	MTBE
<i>Drinking Water Source</i> ¹		NV	12	0.05	0.5	NV	50,000	NV	NV	5
<i>Non-Drinking Water Source</i> ²		NV	18,000	152	204	NV	50,000	NV	NV	1,800
MW-1	3/17/2003	8.3	<5.0	NA	NA	<0.50	NA	<0.50	NA	10.0
	6/23/2003	6.4	<25	NA	NA	<2.5	NA	<2.5	NA	8.0
	9/18/2003	5.3	<25	NA	NA	<2.5	NA	<2.5	NA	8.5
	12/15/03 ³	9.0	<5.0	NA	NA	<0.5	NA	<0.5	NA	12.0
MW-2	3/17/2003	2.1	6.0	NA	NA	<0.50	NA	<0.50	NA	13.0
	6/23/2003	4.5	<5.0	NA	NA	<0.50	NA	<0.50	NA	11.0
	9/18/2003	0.7	<25	NA	NA	<2.5	NA	<2.5	NA	5.0
	12/15/03 ³	3.2	5.2	NA	NA	<0.5	NA	<0.5	NA	13.0
	6/13/2006	4.5	6.5	<5.0	<5.0	<5.0	<50	<0.5	<500	7.6
MW-3	3/17/2003	4.3	8.6	NA	NA	<0.50	NA	<0.50	NA	10.0
	6/23/2003	2.6	<5.0	NA	NA	<0.50	NA	<0.50	NA	5.6
	9/18/2003	3.6	<25	NA	NA	<2.5	NA	<2.5	NA	10.0
	12/15/03 ³	2.7	<5.0	NA	NA	<0.5	NA	<0.5	NA	13.0

**Table III, Summary of Groundwater Sample Fuel Oxygenate Analytical Results
BEI Job No. 203004, Former Fiesta Beverage
966 89th Avenue, Oakland, California**

Well ID	Sample Date	EPA Method 8260B (ug/L)								
		TAME	TBA	EBD	1,2-DCA	DIPE	Ethanol	ETBE	Methanol	MTBE
<i>Drinking Water Source</i> ¹		NV	12	0.05	0.5	NV	50,000	NV	NV	5
<i>Non-Drinking Water Source</i> ²		NV	18,000	152	204	NV	50,000	NV	NV	1,800
MW-4	6/12/2006	NA	NA	NA	NA	NA	NA	NA	NA	6.1

Notes: TAME = Methyl tert-Amyl Ether
TBA = tert-Butyl Alcohol
EDB = 1,2-Dibromoethane
1,2-DCA = 1,2-Dichloroethane
DIPE = Di-isopropyl ether
ETBE = Ethyl tert-butyl ether
MTBE = Methyl tert-butyl ether
(µg/L) = Micrograms per liter
NV = No value
NA = Not analyzed

¹ = From Table A; Environmental Screening Levels (ESLs); Groundwater IS a Current or Potential Source of Drinking Water

² = From Table B; RWQCB Environmental Screening Levels (ESLs); Groundwater IS NOT a Current or Potential Source of Drinking Water

³ = In general after this date, fuel oxygenates were monitored using MTBE detected by EPA Method 8020B, as a proxy for the approximate concentration of the remaining fuel oxygenates.

Bold results indicate detectable analyte concentrations.

Note: Shaded cell indicates that detected concentration exceeds ESL

Table IV, Summary of Groundwater Intrinsic Bioremediation Field Results
BEI Job No. 203004, Former Fiesta Beverage
966 89th Avenue, Oakland, California

Well ID	Sample Date	Field Meter	Field Meter	Field Test Kit	Field Meter	Field Meter
		Dissoved Oxygen (mg/L)	Oxidation Reduction Potential (mV)	Ferrous Iron (Fe 2+)	Field Temperature (o F / o C)	Field pH pH units
MW-1	3/17/2003	NA	NA	NA	60.4 / 60.0 *	7.1 / 7.3
	6/23/2003	0.4	NA	NA	61.0 / 61.0 *	6.9 / 6.9
	9/18/2003	0.4	NA	NA	65.1 / 62.9 *	7.1 / 6.9
	12/15/2003	1.1	NA	NA	13.1 / 13.4	6.8 / 6.7
	6/15/2004	0.1	NA	NA	64.5 / 63.4 *	6.9 / 7.0
	12/15/2004	NA	NA	NA	15.4 / 17.5	7.0 / 6.9
	6/29/2005	0.24 / 0.17	1.0	4.5	19.78 / 21.63	7.15 / 7.08
	5/8/2006	Destroyed	Destroyed	Destroyed	Destroyed	Destroyed
	2/19/2007	Destroyed	Destroyed	Destroyed	Destroyed	Destroyed
MW-2	3/17/2003	NA	NA	NA	66.0 / 64.2 *	7.4 / 7.9
	6/23/2003	0.6	NA	NA	62.1 / 61.8 *	6.8 / 7.1
	9/18/2003	1.3	NA	NA	66.7 / 63.7 *	6.7 / 6.9
	12/15/2003	1.6	NA	NA	13.2 / 13.4	6.6 / 6.6
	6/15/2004	0.1	NA	NA	64.5 / 65.0 *	6.3 / 7.1
	12/15/2004	NA	NA	NA	16.9 / 17.0	7.1 / 7.1
	6/29/2005	0.19 / 0.24	0.7	0.7	18.58 / 21.18	7.12 / 7.13
	6/13/2006	0.80 / 0.42	168.0 / 168.0	0 / 0	17.49 / 17.70	6.97 / 6.98
	2/19/2007	0.2	80	NA	13.6 / 16.3	7.24 / 7.06
MW-3	3/17/2003	NA	NA	NA	63.3 / 60.9 *	7.4 / 7.6
	6/23/2003	0.7	NA	NA	66.4 / 66.9 *	7.3 / 7.2
	9/18/2003	0.4	NA	NA	63.7 / 62.6 *	7.1 / 7.1
	12/15/2003	1.6	NA	NA	14.7 / 15.1	6.5 / 6.4
	6/15/2004	0.0	NA	NA	63.1 / 62.3 *	7.5 / 7.1
	12/15/2004	NA	NA	NA	15.4 / 16.7	7.2 / 7.0
	6/29/2005	0.72 / 0.78	141.7 / -67.6	0.9	17.65 / 18.79	6.94 / 7.02
	6/13/2006	1.01 / 0.41	170.0 / 168.5	0 / 0	17.30 / 17.15	7.02 / 6.98
	2/19/2007	0.08	81	NA	13.7 / 15.6	7.10 / 6.95

Table IV, Summary of Groundwater Intrinsic Bioremediation Field Results
BEI Job No. 203004, Former Fiesta Beverage
966 89th Avenue, Oakland, California

Well ID	Sample Date	Field Meter	Field Meter	Field Test Kit	Field Meter	Field Meter
		Dissoved Oxygen (mg/L)	Oxidation Reduction Potential (mV)	Ferrous Iron (Fe 2+)	Field Temperature (o F / o C)	Field pH pH units
MW-1R	6/13/2006	0.87 / 0.37	172.9 / 172.9	0 / 0	17.31 / 17.36	6.90 / 6.92
	2/19/2007	0.48	8.0	NA	12.2 / 15.8	6.95 / 6.86
MW-4	6/12/2006	0.67 / 0.33	164.3 / 161.0	0.5 / 0	16.90 / 16.79	6.82 / 6.79
	2/19/2007	0.21	98	NA	13.7 / 15.0	7.14 / 7.03
MW-5	6/12/2006	0.61 / 0.31	175.2 / 169.0	0 / 0	18.40 / 18.01	7.01 / 6.94
	2/19/2007	1.98	-114	NA	12.7 / 14.1	6.93 / 6.73
MW-6	6/13/2006	3.10 / 0.81	181.2 / 174.8	0 / 0	17.25 / 17.32	6.94 / 6.83
	2/19/2007	0.21	-30	NA	14.6 / 15.6	6.58 / 6.74
MW-7	6/12/2006	0.59 / 0.27	172.5 / 171.8	0.5 / 0.2	18.14 / 18.00	6.90 / 6.87
	2/19/2007	0.10	110	NA	16.2 / 17.2	7.69 / 7.21
MW-8	6/12/2006	0.37 / 0.33	186.1 / 180.4	0 / 0	18.55 / 18.39	6.85 / 6.85
	2/19/2007	0.11	102	NA	15.2 / 16.6	7.23 / 7.07
MW-9	6/12/2006	2.01 / 1.87	206.0 / 191.0	0 / 0	16.88 / 16.91	6.63 / 6.66
	2/19/2007	0.08	101	NA	15.8 / 16.3	7.56 / 7.23

Notes: mV = Millivolts
mg/L = Milligrams per liter
° F / ° C = degrees Fahrenheit / degrees Centigrade
* = degrees Fahrenheit
2.6 / 2.2 = Initial reading (pre-purge) / Final reading (post-purge)
NA = Not analyzed

**Table V, Summary of Groundwater Intrinsic Bioremediation Analytical Results
BEI Job No. 203004, Former Fiesta Beverage
966 89th Avenue, Oakland, California**

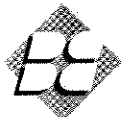
Well ID	Sample Date	Method SM 5310B	Method E300.1		Method RSK 174
		CO ₂	Nitrate (as N)	Sulfate	Methane
		mg/L			µg/L
MW-1	6/29/2005	490	<0.1	5	5,900
	5/8/2006	Destroyed	Destroyed	Destroyed	Destroyed
MW-2	6/29/2005	250	4.1	42	68
	6/13/2006	290	3.2	44	45
MW-3	6/29/2005	230	3.5	33	370
	6/13/2006	220	3.5	33	55
MW-1R	6/13/2006	290	4.3	46	24
MW-4	6/12/2006	260	8.6	44	1.1
MW-5	6/12/2006	240	6.8	45	1.5
MW-6	6/13/2006	290	7.2	50	<0.5
MW-7	6/12/2006	260	6	51	<0.5
MW-8	6/12/2006	330	7.3	46	<0.5
MW-9	6/12/2006	240	8.3	44	1.1

Notes: SM = Standard Method
mg/L = Milligrams per liter
µg/L = Micrograms per liter
CO₂ = Carbon Dioxide

Figures



UNITED STATES GEOLOGICAL SURVEY 7.5' QUADS. "OAKLAND EAST, CA & SAN LEANDRO, CA", BOTH PHOTOREVISED 1981.



BLMYER
ENGINEERS, INC.



SITE LOCATION MAP

FORMER FIESTA BEVERAGE
966 89TH AVE.
OAKLAND, CA

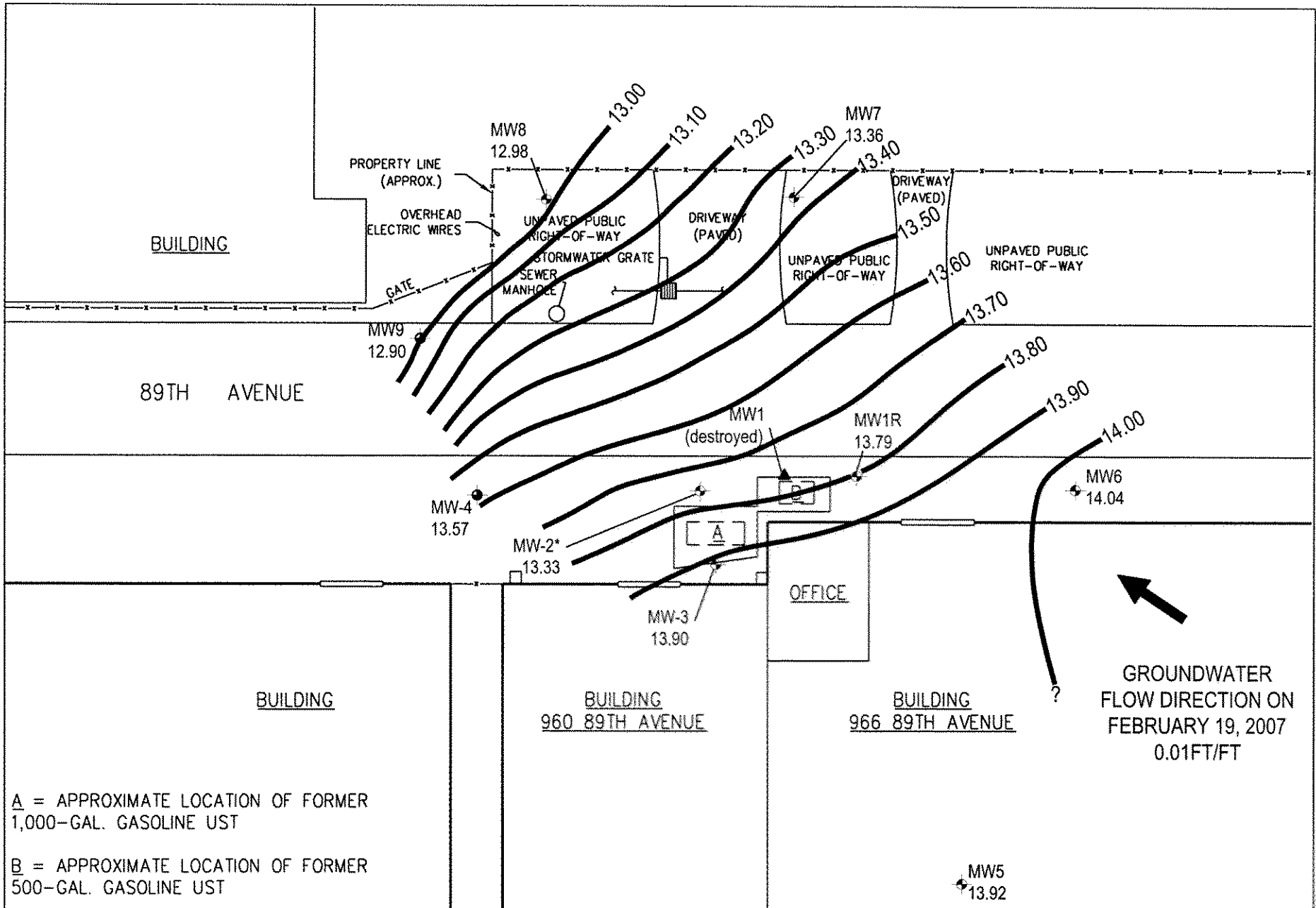
FIGURE

1

BEI JOB NO. 203004

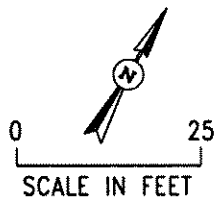
DATE 3-19-03

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A = APPROXIMATE LOCATION OF FORMER 1,000-GAL. GASOLINE UST

B = APPROXIMATE LOCATION OF FORMER 500-GAL. GASOLINE UST



BEI JOB NO. 203004
DATE 3-8-07

LEGEND

- UST UNDERGROUND STORAGE TANK
- ANOMOLOUS DATA; EXCLUDED FROM CONTOURING
- ⊕ GROUNDWATER MONITORING WELL
- ▲ DESTROYED GW MONITORING WELL

GROUNDWATER GRADIENT
FEBRUARY 19, 2007
FORMER FIESTA BEVERAGE
966 89TH AVE.
OAKLAND, CA

FIGURE

2

Figure 3: TPH as Gasoline & Groundwater Elevation vs. Time in Well MW-3

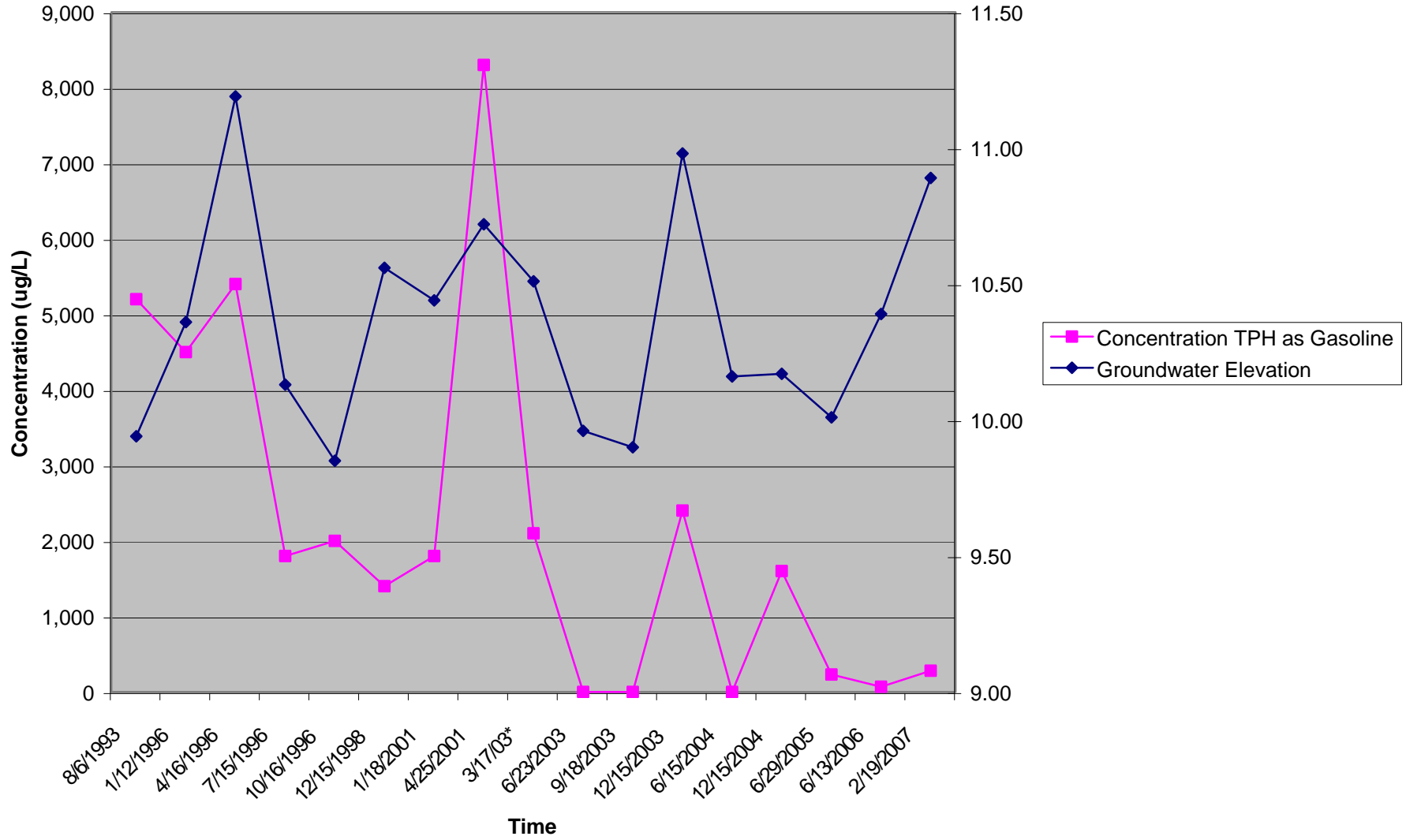
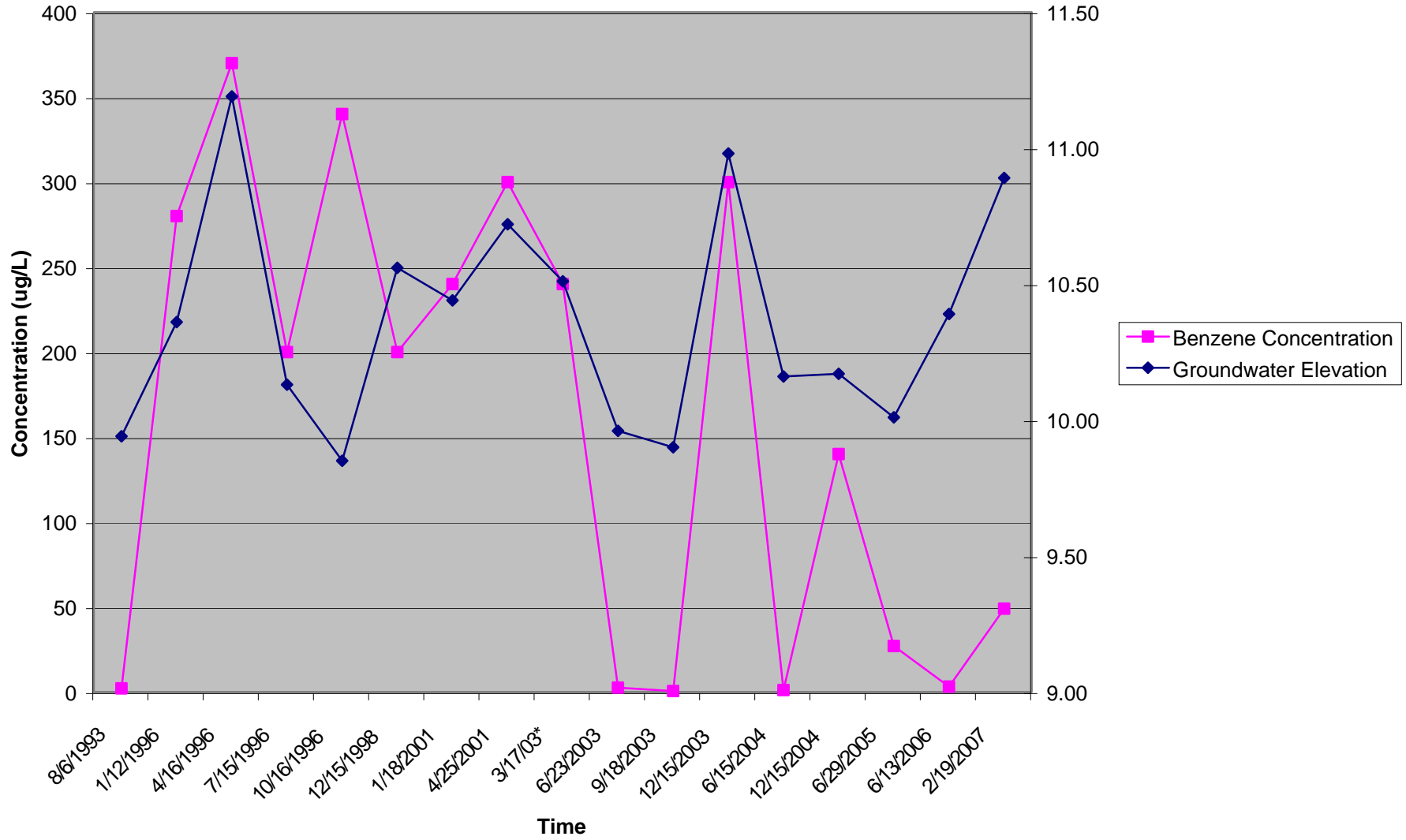


Figure 4: Benzene & Groundwater Elevation vs. Time in Well MW-3



Appendix A

Standard Operating Procedures

Blaine Tech Services, Inc.

Blaine Tech Services, Inc.
Standard Operating Procedure

WATER LEVEL, SEPARATE PHASE LEVEL AND TOTAL WELL DEPTH MEASUREMENTS (GAUGING)

Routine Water Level Measurements

1. Establish that water or debris will not enter the well box upon removal of the cover.
2. Remove the cover using the appropriate tools.
3. Inspect the wellhead (see Wellhead Inspections).
4. Establish that water or debris will not enter the well upon removal of the well cap.
5. Unlock and remove the well cap lock (if applicable). If lock is not functional cut it off.
6. Loosen and remove the well cap. **CAUTION: DO NOT PLACE YOUR FACE OR HEAD DIRECTLY OVER WELLHEAD WHEN REMOVING THE WELL CAP. WELL CAP MAY BE UNDER PRESSURE AND/OR MAY RELEASE ACCUMULATED AND POTENTIALLY HARMFUL VAPORS.**
7. Verify and identify survey point as written on S.O.W.
 - TOC: If survey point is listed as Top of Casing (TOC), look for the exact survey point in the form of a notch or mark on the top of the casing. If no mark is present, use the north side of the casing as the measuring point.
 - TOB: If survey point is listed as Top of Box (TOB), the measuring point will be established manually. Place the inverted wellbox lid halfway across the wellbox opening and directly over the casing. The lower edge of the inverted cover directly over the casing will be the measuring point.
8. Put new Latex or Nitrile gloves on your hands.
9. Slowly lower the Water Level Meter probe into the well until it signals contact with water with a tone and/or flashing a light.
10. Gently raise the probe tip slightly above the water and hold it there. Wait momentarily to see if the meter emits a tone, signaling rising water in the casing. Gently lower the probe tip slightly below the water. Wait momentarily to see if the meter stops emitting a tone, signaling dropping water in the casing. Continue process until water level stabilizes indicating that the well has equilibrated.
11. While holding the probe at first contact with water and the tape against the measuring point, note depth. Repeat twice to verify accuracy. Write down measurement on Well Gauging Sheet under Depth to Water column.
12. Recover probe, replace and tighten well cap, replace lock (if applicable), replace well box cover and tighten hardware (if applicable)

Water Level and Separate Phase Thickness Measurements in Wells Suspected of Containing Separate Phase

1. Establish that water or debris will not enter the well box upon removal of the cover.
2. Remove the cover using the appropriate tools.
3. Inspect the wellhead (see Wellhead Inspections).
4. Establish that water or debris will not enter the well upon removal of the well cap.

5. Unlock and remove the well cap lock (if applicable). If lock is not functional cut it off.
6. Loosen and remove the well cap. CAUTION: DO NOT PLACE YOUR FACE OR HEAD DIRECTLY OVER WELLHEAD WHEN REMOVING THE WELL CAP. WELL CAP MAY BE UNDER PRESSURE AND/OR MAY RELEASE ACCUMULATED AND POTENTIALLY HARMFULL VAPORS.
7. Verify and identify survey point as written on S.O.W.
 - TOC: If survey point is listed as Top of Casing (TOC), look for the exact survey point in the form of a notch or mark on the top of the casing. If no mark is present, use the north side of the casing as the measuring point.
 - TOB: If survey point is listed as Top of Box (TOB), the measuring point will be established manually. Place the inverted well box lid halfway across the well box opening and directly over the casing. The lower edge of the inverted cover directly over the casing will be the measuring point.
8. Put new Nitrile gloves on your hands.
9. Slowly lower the tip of the Interface Probe into the well until it emits either a solid or broken tone.
 - BROKEN TONE: Separate phase layer is not present. Go to Step 8 of Routine Water Level Measurements shown above to complete gauging process using the Interface probe as you would a Water Level Meter.
 - SOLID TONE: Separate phase layer is present. Go to the next step.
10. Gently raise the probe tip slightly above the separate phase layer and hold it there. Wait momentarily to see if the meter emits a tone, signaling rising water in the casing. Gently lower the probe tip slightly below the separate phase layer. Wait momentarily to see if the meter stops emitting a tone, signaling dropping water in the casing. Continue process until water level stabilizes indicating that the well has equilibrated.
11. While holding the probe at first contact with the separate phase layer and the tape against the measuring point, note depth. Repeat twice to verify accuracy. Write down measurement on Well Gauging Sheet under Depth to Product column.
12. Gently lower the probe tip until it emits a broken tone signifying contact with water. While holding the probe at first contact with water and the tape against the measuring point, note depth. Repeat twice to verify accuracy. Write down measurement on Well Gauging Sheet under Depth to Water column.
13. Recover probe, replace and tighten well cap, replace lock (if applicable), replace well box cover and tighten hardware (if applicable).

Routine Total Well Depth Measurements

1. Lower the Water Level Meter probe into the well until it lightens in your hands, indicating that the probe is resting at the bottom of well.
2. Gently raise the tape until the weight of the probe increases, indicating that the probe has lifted off the well bottom.
3. While holding the probe at first contact with the well bottom and the tape against the well measuring point, note depth. Repeat twice to verify accuracy. Write down measurement on Well Gauging Sheet under Total Well Depth column.

4. Recover probe, replace and tighten well cap, replace lock (if applicable), replace well box cover and tighten hardware (if applicable).

Blaine Tech Services, Inc.
Standard Operating Procedure

WELL WATER EVACUATION (PURGING)

Purpose

Evacuation of a predetermined minimum volume of water from a well (purging) while *simultaneously* measuring water quality parameters is typically required prior to sampling. Purging a minimum volume guarantees that actual formation water is drawn into the well. Measuring water quality parameters either verifies that the water is stable and suitable for sampling or shows that the water remains unstable, indicating the need for continued purging. Both the minimum volume and the stable parameter qualifications need to be met prior to sampling. This assures that the subsequent sample will be representative of the formation water surrounding the well screen and not of the water standing in the well.

Defining Casing Volumes

The predetermined minimum quantity of water to be purged is based on the wells' casing volume. A casing volume is the volume of water presently standing within the casing of the well. This is calculated as follows:

$$\text{Casing Volume} = (\text{TD} - \text{DTW}) \text{ VCF}$$

1. Subtract the wells' depth to water (DTW) measurement from its total depth (TD) measurement. This is the height of the water column in feet.
2. Determine the well casings' volume conversion factor (VCF). The VCF is based on the diameter of the well casing and represents the volume, in gallons, that is contained in one (1) foot of a particular diameter of well casing. The common VCF's are listed on our Well Purge Data Sheets.
3. Multiply the VCF by the calculated height of the water column. This is the casing volume, the amount of water in gallons standing in the well.

Remove Three to Five Casing Volumes

Prior to sampling, an attempt will be made to purge all wells of a minimum of three casing volumes and a maximum of five casing volumes except where regulations mandate the minimum removal of four casing volumes.

Choose the Appropriate Evacuation Device Based on Efficiency

In the absence of instructions on the SOW to the contrary, selection of evacuation device will be based on efficiency.

Measure Water Quality Parameters at Each Casing Volume

At a minimum, water quality measurements include pH, temperature and electrical conductivity (EC). Measurements are made and recorded at least once every casing volume. They are considered stable when all parameters are within 10% of their previous measurement.

Note: The following instructions assume that well has already been properly located, accessed, inspected and gauged.

Prior to Purging a Well

1. Confirm that the well is to be purged and sampled per the SOW.
2. Confirm that the well is suitable based on the conditions set by the client relative to separate phase.
3. Calculate the wells' casing volume.
4. Put new Latex or Nitrile gloves on your hands.

Purging With a Bailer (Stainless Steel, Teflon or Disposable)

1. Attach bailer cord or string to bailer. Leave other end attached to spool.
2. Gently lower empty bailer into well until well bottom is reached.
3. Cut cord from spool. Tie end of cord to hand.
4. Gently raise full bailer out of well and clear of well head. Do not let the bailer or cord touch the ground.
5. Pour contents into graduated 5-gallon bucket or other graduated receptacle.
6. Repeat purging process.
7. Upon removal of first casing volume, fill clean parameter cup with purgewater, empty the remainder of the purgewater into the bucket, lower the bailer back into the well and secure the cord on the Sampling Vehicle.
8. Use the water in the cup to collect and record parameter measurements.
9. Continue purging until second casing volume is removed.
10. Collect parameter measurements.
11. Continue purging until third casing volume is removed.
12. Collect parameter measurements. If parameters are stable, stop purging. If parameters remain unstable, continue purging until stabilization occurs or the fifth casing volume is removed.

Purging With a Pneumatic Pump

1. Position Pneumatic pump hose reel over the top of the well.
2. Gently unreel and lower the pump into the well. Do not contact the well bottom.
3. Secure the hose reel.
4. Begin purging into graduated 5-gallon bucket or other graduated receptacle.
5. Adjust water recharge duration and air pulse duration for maximum efficiency.
6. Upon removal of first casing volume, fill clean parameter cup with water.
7. Use the water in the cup to collect and record parameter measurements.
8. Continue purging until second casing volume is removed.

9. Collect parameter measurements.
10. Continue purging until third casing volume is removed.
11. Collect parameter measurements. If parameters are stable, stop purging. If parameters remain unstable, continue purging until stabilization occurs or the fifth casing volume is removed.
12. Upon completion of purging, gently recover the pump and secure the reel.

Purging With a Fixed Speed Electric Submersible Pump

1. Position Electric Submersible hose reel over the top of the well.
2. Gently unreel and lower the pump to the well bottom.
3. Raise the pump 5 feet off the bottom.
4. Secure the hose reel.
5. Begin purging.
6. Verify pump rate with flow meter or graduated 5-gallon bucket
7. Upon removal of first casing volume, fill clean parameter cup with water.
8. Use the water in the cup to collect and record parameter measurements.
9. Continue purging until second casing volume is removed.
10. Collect parameter measurements.
11. Continue purging until third casing volume is removed.
12. Collect parameter measurements. If parameters are stable, stop purging. If parameters remain unstable, continue purging until stabilization occurs or the fifth casing volume is removed.
13. Upon completion of purging, gently recover the pump and secure the reel.

Blaine Tech Services, Inc.
Standard Operating Procedure

SAMPLE COLLECTION FROM GROUNDWATER WELLS USING BAILERS

Sampling with a Bailer (Stainless Steel, Teflon or Disposable)

1. Put new Latex or Nitrile gloves on your hands.
2. Determine required bottle set.
3. Fill out sample labels completely and attach to bottles.
4. Arrange bottles in filling order and loosen caps (see Determine Collection Order below).
5. Attach bailer cord or string to bailer. Leave other end attached to spool.
6. Gently lower empty bailer into well until water is reached.
7. As bailer fills, cut cord from spool and tie end of cord to hand.
8. Gently raise full bailer out of well and clear of well head. Do not let the bailer or cord touch the ground. If a set of parameter measurements is required, go to step 9. If no additional measurements are required, go to step 11.
9. Fill a clean parameter cup, empty the remainder contained in the bailer into the sink, lower the bailer back into the well and secure the cord on the Sampling Vehicle. Use the water in the cup to collect and record parameter measurements.
10. Fill bailer again and carefully remove it from the well.
11. Slowly fill and cap sample bottles. Fill and cap volatile compounds first, then semi-volatile, then inorganic. Return to the well as needed for additional sample material.

Fill 40-milliliter vials for volatile compounds as follows: Slowly pour water down the inside on the vial. Carefully pour the last drops creating a convex or positive meniscus on the surface. Gently screw the cap on eliminating any air space in the vial. Turn the vial over, tap several times and check for trapped bubbles. If bubbles are present, repeat process.

Fill 1 liter amber bottles for semi-volatile compounds as follows: Slowly pour water into the bottle. Leave approximately 1 inch of headspace in the bottle. Cap bottle.

Field filtering of inorganic samples using a stainless steel bailer is performed as follows: Attach filter connector to top of full stainless steel bailer. Attach 0.45 micron filter to connector. Flip bailer over and let water gravity feed through the filter and into the sample bottle. If high turbidity level of water clogs filter, repeat process with new filter until bottle is filled. Leave headspace in the bottle. Cap bottle.

Field filtering of inorganic samples using a disposable bailer is performed as follows: Attach 0.45 micron filter to connector plug. Attach connector plug to bottom of full disposable bailer. Water will gravity feed through the filter and into the sample bottle. If high turbidity level of water clogs filter, repeat process with new filter until bottle is filled. Leave headspace in the bottle. Cap bottle.

12. Bag samples and place in ice chest.
13. Note sample collection details on well data sheet and Chain of Custody.

Appendix B

Well Monitoring Data Sheets and Well Gauging Data,
Dated February 19, 2007
Blaine Tech Services, Inc.

SPH or Purge Water Drum Log

Client: Blymer Eng. @ Former Fiesta Beverage
 Site Address: 966 89th Ave, Oakland

STATUS OF DRUM(S) UPON ARRIVAL					
Date	6/29/05	6/02/06	6/05/06	6/22/06	2/19/07
Number of drum(s) empty:		0 BTS			
Number of drum(s) 1/4 full:					1
Number of drum(s) 1/2 full:	1				
Number of drum(s) 3/4 full:			1		
Number of drum(s) full:			1	30	30
Total drum(s) on site:	1	0 BTS	2 BTS	30	30
Are the drum(s) properly labeled?	N		Y	Y	Y
Drum ID & Contents:	-		Purge H ₂ O	H ₂ O?	Purge H ₂ O
If any drum(s) are partially or totally filled, what is the first use date:	-	-	6/09/06		

- If you add any SPH to an empty or partially filled drum, drum must have at least 20 gals. of Purgewater or DI Water.
- If drum contains SPH, the drum MUST be steel AND labeled with the appropriate label.
- All BTS drums MUST be labeled appropriately.

STATUS OF DRUM(S) UPON DEPARTURE					
Date	6/29/05	6/02/06	6/05/06	6/22/06	2/19/07
Number of drums empty:					
Number of drum(s) 1/4 full:					
Number of drum(s) 1/2 full:	2			1	
Number of drum(s) 3/4 full:		1			
Number of drum(s) full:		1	3		30
Total drum(s) on site:	2	2 BTS	3 BTS	30	30
Are the drum(s) properly labeled?	Y	Y	Y	Y	Y
Drum ID & Contents:	SP Purge Water	Purge H ₂ O	Purge H ₂ O	H ₂ O	H ₂ O

LOCATION OF DRUM(S)
 Describe location of drum(s): Near wall 3 against wall - see map / northeast corner
=> owner, 'Gary', intends to move to so. of Property (behind building)

FINAL STATUS					
Number of new drum(s) left on site this event	1	2	1	1	0
Date of inspection:	6/29/05	6/02/06	6/05/06	6/22/06	2/19/07
Drum(s) labelled properly:	Y	Y	Y	Y	Y
Logged by BTS Field Tech:	PC	we	sl	mm	SD
Office reviewed by:	ca	w	w	mt	W

WELLHEAD INSPECTION CHECKLIST

Date 2/19/07 Client Blymyer Engineers
 Site Address 966 89th Ave Oakland
 Job Number 070219-S4 Technician SL

Well ID	Well Inspected - No Corrective Action Required	Water Bailed From Wellbox	Wellbox Components Cleaned	Cap Replaced	Debris Removed From Wellbox	Lock Replaced	Other Action Taken (explain below)	Well Not Inspected (explain below)
MW-1R	X							
MW-2	X						X	
MW-3	X							
MW-4	X							
MW-5	X							
MW-6	X							
MW-7	X							
MW-8	X							
MW-9	X							

NOTES: MW-2 No bolts - underneath large
steel plate

WELL GAUGING DATA

Project # 070219-SL1 Date 2/19/07 Client Bymyer Engineers

Site 966 89th Ave Oakland

Well ID	Time	Well Size (in.)	Sheen / Odor	Depth to Immiscible Liquid (ft.)	Thickness of Immiscible Liquid (ft.)	Volume of Immiscibles Removed (ml)	Depth to water (ft.)	Depth to well bottom (ft.)	Survey Point: TOB or TOC	Notes
MW-1R	0831	2					7.94	21.55	↓	
MW-2	0840	2				8.12	23.96			
MW-3	0836	2	odor			8.12	24.91			
MW-4	0820	2				7.77	21.80			
MW-5	0845	2				8.61	19.75			
MW-6	0825	2				7.93	19.81			
MW-7	0805	2				7.85	21.77			
MW-8	0811	2				7.99	20.02			
MW-9	0814	2				8.08	22.07			

WELL MONITORING DATA SHEET

Project #: 070219-5L1	Client: Plymver Eng.
Sampler: SL	Date: 2/19/07
Well I.D.: MW-1R	Well Diameter: 2 3 4 6 8
Total Well Depth (TD): 2155	Depth to Water (DTW): 7.94
Depth to Free Product:	Thickness of Free Product (feet):
Referenced to: PVC Grade	D.O. Meter (if req'd): YSI HACH
DTW with 80% Recharge [(Height of Water Column x 0.20) + DTW]: 10.66	

Purge Method: Bailer Disposable Bailer Positive Air Displacement Electric Submersible Waterra Peristaltic Extraction Pump Other _____

Sampling Method: Bailer Disposable Bailer Extraction Port Dedicated Tubing Other: _____

2.2 (Gals.) X 3 = 6.6 Gals.	Well Diameter	Multiplier	Well Diameter	Multiplier
1 Case Volume	Specified Volumes	Calculated Volume	1"	0.04
			2"	0.16
			3"	0.37
			4"	0.65
			6"	1.47
			Other	radius ² * 0.163

Time	Temp (°F or °C)	pH	Cond. (mS or µS)	Turbidity (NTUs)	Gals. Removed	Observations
0955	12.2	6.95	677	>1000	2.2	Brown
1000	15.6	6.95	676	>1000	4.4	Slight Odor
1005	15.8	6.86	677	>1000	6.6	↓

Did well dewater? Yes No Gallons actually evacuated: 6.6

Sampling Date: 2/19/07 Sampling Time: 1010 Depth to Water: 10.10

Sample I.D.: MW-1R Laboratory: Kiff CalScience Other: McCampbell

Analyzed for: TPH-G BTEX MTBE TPH-D Oxygenates (5) Other:

EB I.D. (if applicable): @ Time Duplicate I.D. (if applicable):

Analyzed for: TPH-G BTEX MTBE TPH-D Oxygenates (5) Other:

D.O. (if req'd):	Pre-purge: 0.48 mg/L	Post-purge:	mg/L
O.R.P. (if req'd):	Pre-purge: 8 mV	Post-purge:	mV

WELL MONITORING DATA SHEET

Project #: <u>070219-SL1</u>	Client: <u>Blayne Eng.</u>
Sampler: <u>SL</u>	Date: <u>2/19/07</u>
Well I.D.: <u>MW-2</u>	Well Diameter: <u>2</u> 3 4 6 8
Total Well Depth (TD): <u>20.96</u>	Depth to Water (DTW): <u>9.12</u>
Depth to Free Product:	Thickness of Free Product (feet):
Referenced to: <u>PVC</u> Grade	D.O. Meter (if req'd): <u>YSI</u> HACH
DTW with 80% Recharge [(Height of Water Column x 0.20) + DTW]: <u>11.29</u>	

Purge Method: Bailer Disposable Bailer Positive Air Displacement Electric Submersible

Water: Peristaltic Extraction Pump Other _____

Sampling Method: Bailer Disposable Bailer Extraction Port Dedicated Tubing

Other: _____

<u>2.5</u> (Gals.) X <u>3</u> = <u>7.5</u> Gals.	Well Diameter	Multiplier	Well Diameter	Multiplier
1 Case Volume	Specified Volumes	Calculated Volume	1"	0.04
			2"	0.16
			3"	0.37
			4"	0.65
			6"	1.47
			Other	radius ² * 0.163

Time	Temp (°F or °C)	pH	Cond. (mS or µS)	Turbidity (NTUs)	Gals. Removed	Observations
1020	13.0	7.24	642	>1000	2.5	Brown
1025	16.1	7.13	636	>1000	5.0	↓
1030	16.3	7.06	629	>1000	7.5	

Did well dewater? Yes No Gallons actually evacuated: 7.5

Sampling Date: 2/19/07 Sampling Time: 1150 Depth to Water: 11.11

Sample I.D.: MW-2 Laboratory: Kiff CalScience Other: McCampbell

Analyzed for: TPH-G BTEX MTBE TPH-D Oxygenates (5) Other: _____

EB I.D. (if applicable): _____ Time _____ Duplicate I.D. (if applicable): _____

Analyzed for: TPH-G BTEX MTBE TPH-D Oxygenates (5) Other: _____

D.O. (if req'd):	<input checked="" type="checkbox"/> Pre-purge: <u>0.21</u> mg/L	<input type="checkbox"/> Post-purge: _____ mg/L
O.R.P. (if req'd):	<input checked="" type="checkbox"/> Pre-purge: <u>80</u> mV	<input type="checkbox"/> Post-purge: _____ mV

WELL MONITORING DATA SHEET

Project #: <u>070219-521</u>	Client: <u>Burver Eng.</u>
Sampler: <u>SL</u>	Date: <u>2/19/07</u>
Well I.D.: <u>MW-3</u>	Well Diameter: <u>2</u> 3 4 6 8
Total Well Depth (TD): <u>2491</u>	Depth to Water (DTW): <u>8.12</u>
Depth to Free Product:	Thickness of Free Product (feet):
Referenced to: <u>PVC</u> Grade	D.O. Meter (if req'd): <u>YSI</u> HACH
DTW with 80% Recharge [(Height of Water Column x 0.20) + DTW]: 2.7 <u>11.48</u>	

Purge Method: Bailer Disposable Bailer Positive Air Displacement Electric Submersible

Water: Peristaltic Extraction Pump Other _____

Sampling Method: Bailer Disposable Bailer Extraction Port Dedicated Tubing

Other: _____

<u>2.7</u> (Gals.) X <u>3</u> = <u>8.1</u> Gals.	Well Diameter	Multiplier	Well Diameter	Multiplier
1 Case Volume Specified Volumes Calculated Volume	1"	0.04	4"	0.65
	2"	0.16	6"	1.47
	3"	0.37	Other	radius ² * 0.163

Time	Temp (°F or °C)	pH	Cond. (mS or µS)	Turbidity (NTUs)	Gals. Removed	Observations
1050	13.7	7.10	638	>1000	2.7	Brown, Odor
1055	15.4	7.08	640	>1000	5.4	↓
1100	15.6	6.95	644	>1000	8.1	↓

Did well dewater? Yes No Gallons actually evacuated: 8.1

Sampling Date: 2/19/07 Sampling Time: 12:10 Depth to Water: 8.12

Sample I.D.: MW-3 Laboratory: Kiff CalScience Other: McStimbell

Analyzed for: TPH-G BTEX MTBE TPH-D Oxygenates (5) Other:

EB I.D. (if applicable): @ Time Duplicate I.D. (if applicable):

Analyzed for: TPH-G BTEX MTBE TPH-D Oxygenates (5) Other:

D.O. (if req'd):	<u>Pre-purge:</u> <u>0.08</u> mg/L	Post-purge:	mg/L
O.R.P. (if req'd):	<u>Pre-purge:</u> <u>81</u> mV	Post-purge:	mV

WELL MONITORING DATA SHEET

Project #: <u>07024-SL1</u>	Client: <u>Blymyer Eng.</u>
Sampler: <u>SL</u>	Date: <u>2/19/07</u>
Well I.D.: <u>MW-4</u>	Well Diameter: <u>2</u> 3 4 6 8
Total Well Depth (TD): <u>2190</u>	Depth to Water (DTW): <u>7.77</u>
Depth to Free Product:	Thickness of Free Product (feet):
Referenced to: <u>PVC</u> Grade	D.O. Meter (if req'd): <u>YSI</u> HACH
DTW with 80% Recharge [(Height of Water Column x 0.20) + DTW]: <u>10.58</u>	

Purge Method: Bailer Disposable Bailer Positive Air Displacement Electric Submersible Waterra Peristaltic Extraction Pump Other _____

Sampling Method: Bailer Disposable Bailer Extraction Port Dedicated Tubing Other _____

<u>22</u> (Gals.) X <u>3</u> = <u>6.6</u> Gals.	Well Diameter	Multiplier	Well Diameter	Multiplier
1 Case Volume	Specified Volumes	Calculated Volume	1"	0.04
			4"	0.65
			2"	0.16
			6"	1.47
			3"	0.37
			Other	radius ² * 0.163

Time	Temp (°F or °C)	pH	Cond. (mS or µS)	Turbidity (NTUs)	Gals. Removed	Observations
1125	13.7	7.14	675	>1000	2.2	Brown
1130	14.7	7.08	657	>1000	4.4	↓
1135	15.0	7.03	649	>1000	6.6	↓

Did well dewater? Yes No Gallons actually evacuated: 6.6

Sampling Date: 2/19/07 Sampling Time: 1140 Depth to Water: 8.03

Sample I.D.: MW-4 Laboratory: Kiff CalScience Other: McCampbell

Analyzed for: TPH-G BTEX MTBE TPH-D Oxygenates (5) Other: _____

EB I.D. (if applicable): _____ @ _____ Time Duplicate I.D. (if applicable): _____

Analyzed for: TPH-G BTEX MTBE TPH-D Oxygenates (5) Other: _____

D.O. (if req'd):	Pre-purge: <u>0.21</u> mg/L	Post-purge: _____ mg/L
O.R.P. (if req'd):	Pre-purge: <u>98</u> mV	Post-purge: _____ mV

WELL MONITORING DATA SHEET

Project #: 07024-921	Client: Blymyer Eng.
Sampler: SL	Date: 2/19/07
Well I.D.: MW-5	Well Diameter: 2 3 4 6 8
Total Well Depth (TD): 19.75	Depth to Water (DTW): 8.61
Depth to Free Product:	Thickness of Free Product (feet):
Referenced to: PVC Grade	D.O. Meter (if req'd): YSI HACH
DTW with 80% Recharge [(Height of Water Column x 0.20) + DTW]: 10.84	

Purge Method: Bailer Disposable Bailer Positive Air Displacement Electric Submersible

Water: Peristaltic Extraction Pump Other

Sampling Method: Bailer Disposable Bailer Extraction Port Dedicated Tubing

Other:

1.8 (Gals.) X 3 = 5.4 Gals.

1 Case Volume Specified Volumes Calculated Volume

Well Diameter	Multiplier	Well Diameter	Multiplier
1"	0.04	4"	0.65
2"	0.16	6"	1.47
3"	0.37	Other	radius ² * 0.163

Time	Temp (°F or °C)	pH	Cond (mS or µS)	Turbidity (NTUs)	Gals. Removed	Observations
0900	12.7	6.93	713	>1000	1.8	Brown
0905	13.9	6.82	679	>1000	3.6	↓
0910	14.1	6.73	670	>1000	5.4	

Did well dewater? Yes No Gallons actually evacuated: 5.4

Sampling Date: 2/19/07 Sampling Time: 0915 Depth to Water: 08.59

Sample I.D.: MW-5 Laboratory: Kiff CalScience Other: McCampbell

Analyzed for: TPH-G BTEX MTBE TPH-D Oxygenates (5) Other:

EB I.D. (if applicable): @ Time Duplicate I.D. (if applicable):

Analyzed for: TPH-G BTEX MTBE TPH-D Oxygenates (5) Other:

D.O. (if req'd):	Pre-purge: 1.98 mg/L	Post-purge:	mg/L
O.R.P. (if req'd):	Pre-purge: -114 mV	Post-purge:	mV

WELL MONITORING DATA SHEET

Project #: <u>070219 SL1</u>	Client: <u>Blymyer Eng.</u>
Sampler: <u>SL</u>	Date: <u>2/19/07</u>
Well I.D.: <u>MW-6</u>	Well Diameter: <u>2</u> 3 4 6 8
Total Well Depth (TD): <u>19.81</u>	Depth to Water (DTW): <u>7.93</u>
Depth to Free Product:	Thickness of Free Product (feet):
Referenced to: <u>PVC</u> Grade	D.O. Meter (if req'd): <u>YSI</u> HACH
DTW with 80% Recharge [(Height of Water Column x 0.20) + DTW]: <u>10.31</u>	

Purge Method: Bailer Disposable Bailer Positive Air Displacement Electric Submersible

Waterra Peristaltic Extraction Pump Other _____

Sampling Method: Bailer Disposable Bailer Extraction Port Dedicated Tubing

Other: _____

1.9 (Gals.) X 3 = 5.7 Gals.

1 Case Volume Specified Volumes Calculated Volume

Well Diameter	Multiplier	Well Diameter	Multiplier
1"	0.04	4"	0.65
2"	0.16	6"	1.47
3"	0.37	Other	radius ² * 0.163

Time	Temp (°F or °C)	pH	Cond. (mS or µS)	Turbidity (NTUs)	Gals. Removed	Observations
0930	14.6	6.58	699	>1000	1.9	Brown ↓
0935	14.9	6.73	737	>1000	3.8	
0940	15.6	6.74	740	>1000	5.7	

Did well dewater? Yes No Gallons actually evacuated: 5.7

Sampling Date: 2/19/07 Sampling Time: 1119 Depth to Water: 10.25

Sample I.D.: MW-6 Laboratory: Kiff CalScience Other McCampbell

Analyzed for: TPH-C BTEX MTBE TPH-D Oxygenates (5) Other:

EB I.D. (if applicable): @ _____ Time Duplicate I.D. (if applicable):

Analyzed for: TPH-G BTEX MTBE TPH-D Oxygenates (5) Other:

D.O. (if req'd):	<input checked="" type="checkbox"/> Pre-purge:	<u>0.21</u> mg/L	Post-purge:	mg/L
O.R.P. (if req'd):	Pre-purge:	<u>-30</u> mV	Post-purge:	mV

WELL MONITORING DATA SHEET

Project #: <u>070219-SL1</u>	Client: <u>Blymyer Eng.</u>
Sampler: <u>SL</u>	Date: <u>2/19/07</u>
Well I.D.: <u>MW-7</u>	Well Diameter: <u>2</u> 3 4 6 8
Total Well Depth (TD): <u>21.77</u>	Depth to Water (DTW): <u>7.85</u>
Depth to Free Product:	Thickness of Free Product (feet):
Referenced to: <u>PVC</u> Grade	D.O. Meter (if req'd): <u>YSI</u> HACH
DTW with 80% Recharge [(Height of Water Column x 0.20) + DTW]: <u>10.63</u>	

Purge Method: Bailer Disposable Bailer Positive Air Displacement Electric Submersible Waterra Peristaltic Extraction Pump Other _____

Sampling Method: Bailer Disposable Bailer Extraction Port Dedicated Tubing Other: _____

<u>2.2</u> (Gals.) X <u>3</u> = <u>6.6</u> Gals.	Well Diameter	Multiplier	Well Diameter	Multiplier
1 Case Volume	Specified Volumes	Calculated Volume	1"	0.04
			2"	0.16
			3"	0.37
			4"	0.65
			6"	1.47
			Other	radius ² * 0.163

Time	Temp (°F or °C)	pH	Cond. (mS or µS)	Turbidity (NTUs)	Gals. Removed	Observations
<u>1215</u>	<u>16.2</u>	<u>7.69</u>	<u>703</u>	<u>>1000</u>	<u>2.2</u>	<u>Brown</u>
<u>1220</u>	<u>17.1</u>	<u>7.36</u>	<u>702</u>	<u>>1000</u>	<u>4.4</u>	↓
<u>1225</u>	<u>17.2</u>	<u>7.21</u>	<u>703</u>	<u>>1000</u>	<u>6.6</u>	

Did well dewater? Yes No Gallons actually evacuated: 6.6

Sampling Date: 2/19/07 Sampling Time: 1230 Depth to Water: 8.16

Sample I.D.: MW-7 Laboratory: Kiff CalScience Other: McCampbell

Analyzed for: TPH-G BTEX MTBE TPH-D Oxygenates (5) Other:

EB I.D. (if applicable): @ _____ Time Duplicate I.D. (if applicable):

Analyzed for: TPH-G BTEX MTBE TPH-D Oxygenates (5) Other:

D.O. (if req'd):	Pre-purge: <u>0.10</u> mg/L	Post-purge:	mg/L
O.R.P. (if req'd):	Pre-purge: <u>110</u> mV	Post-purge:	mV

WELL MONITORING DATA SHEET

Project #: <u>070219-SL1</u>	Client: <u>Blumer Eng.</u>
Sampler: <u>SL</u>	Date: <u>2/19/07</u>
Well I.D.: <u>MW-8</u>	Well Diameter: <u>2</u> 3 4 6 8
Total Well Depth (TD): <u>20.02</u>	Depth to Water (DTW): <u>7.99</u>
Depth to Free Product:	Thickness of Free Product (feet):
Referenced to: <u>PVC</u> Grade	D.O. Meter (if req'd): <u>YSI</u> HACH
DTW with 80% Recharge [(Height of Water Column x 0.20) + DTW]: <u>10.40</u>	

Purge Method: Bailer Disposable Bailer Positive Air Displacement Electric Submersible Water Peristaltic Extraction Pump Other _____

Sampling Method: Bailer Disposable Bailer Extraction Port Dedicated Tubing Other: _____

1.9 (Gals.) X 3 = 5.7 Gals.
 1 Case Volume Specified Volumes Calculated Volume

Well Diameter	Multiplier	Well Diameter	Multiplier
1"	0.04	4"	0.65
2"	0.16	6"	1.47
3"	0.37	Other	radius ² * 0.163

Time	Temp (°F or °C)	pH	Cond (mS or µS)	Turbidity (NTUs)	Gals. Removed	Observations
<u>1235</u>	<u>15.2</u>	<u>7.23</u>	<u>702</u>	<u>>1000</u>	<u>1.9</u>	<u>Brown</u>
<u>1240</u>	<u>16.7</u>	<u>7.13</u>	<u>711</u>	<u>>1000</u>	<u>3.8</u>	<u>↓</u>
<u>1245</u>	<u>16.6</u>	<u>7.07</u>	<u>712</u>	<u>>1000</u>	<u>5.7</u>	

Did well dewater? Yes No Gallons actually evacuated: _____

Sampling Date: 2/19/07 Sampling Time: 1250 Depth to Water: 8.72

Sample I.D.: MW-8 Laboratory: Kiff CalScience Other McCampbell

Analyzed for: TPH-C BTEX MTBE TPH-D Oxygenates (5) Other: _____

EB I.D. (if applicable): _____ @ _____ Time Duplicate I.D. (if applicable): _____

Analyzed for: TPH-G BTEX MTBE TPH-D Oxygenates (5) Other: _____

D.O. (if req'd):	<u>Pre-purge:</u>	<u>0.11</u> mg/L	Post-purge:	_____ mg/L
O.R.P. (if req'd):	<u>Pre-purge:</u>	<u>102</u> mV	Post-purge:	_____ mV

WELL MONITORING DATA SHEET

Project #: <u>07029-SL1</u>	Client: <u>Bymyer Eng.</u>
Sampler: <u>SL</u>	Date: <u>2/19/07</u>
Well I.D.: <u>MW-9</u>	Well Diameter: <u>2</u> 3 4 6 8
Total Well Depth (TD): <u>72.07</u>	Depth to Water (DTW): <u>80.8</u>
Depth to Free Product:	Thickness of Free Product (feet):
Referenced to: <u>PVC</u> Grade	D.O. Meter (if req'd): <u>YSI</u> HACH
DTW with 80% Recharge [(Height of Water Column x 0.20) + DTW]: <u>10.88</u>	

Purge Method: Bailer Disposable Bailer Positive Air Displacement Electric Submersible Other _____

Waterra Peristaltic Extraction Pump Other _____

Sampling Method: Bailer Disposable Bailer Extraction Port Dedicated Tubing Other _____

2.2 (Gals.) X 3 = 6.6 Gals.
 1 Case Volume Specified Volumes Calculated Volume

Well Diameter	Multiplier	Well Diameter	Multiplier
1"	0.04	4"	0.65
2"	0.16	6"	1.47
3"	0.37	Other	radius ² * 0.163

Time	Temp (°F or °C)	pH	Cond. (mS or µS)	Turbidity (NTUs)	Gals. Removed	Observations
1315	15.8	7.56	646	>1000	2.2	Brown
1320	16.0	7.30	644	>1000	4.4	↓
1325	16.3	7.23	646	>1000	6.6	

Did well dewater? Yes No Gallons actually evacuated: 6.6

Sampling Date: 2/19/07 Sampling Time: 1330 Depth to Water: 9.32

Sample I.D.: MW-9 Laboratory: Kiff CalScience Other McCambell

Analyzed for: TPH-C BTEX MTBE TPH-D Oxygenates (5) Other:

EB I.D. (if applicable): @ Time Duplicate I.D. (if applicable):

Analyzed for: TPH-G BTEX MTBE TPH-D Oxygenates (5) Other:

D.O. (if req'd):	Pre-purge: <u>0.08</u> mg/L	Post-purge:	mg/L
O.R.P. (if req'd):	Pre-purge: <u>101</u> mV	Post-purge:	mV

Appendix C

Analytical Laboratory Report

Dated February 27, 2007

McC Campbell Analytical, Inc.



McC Campbell Analytical, Inc.

"When Quality Counts"

1534 Willow Pass Road, Pittsburg, CA 94565-1701
Web: www.mcccampbell.com E-mail: main@mcccampbell.com
Telephone: 877-252-9262 Fax: 925-252-9269

Blymyer Engineers, Inc. 1829 Clement Avenue Alameda, CA 94501-1395	Client Project ID: Former Fiesta Beverages	Date Sampled: 02/19/07
		Date Received: 02/21/07
	Client Contact: Mark Detterman	Date Reported: 02/27/07
	Client P.O.:	Date Completed: 02/27/07

WorkOrder: 0702471

February 27, 2007

Dear Mark:

Enclosed are:

- 1). the results of 9 analyzed samples from your **Former Fiesta Beverages project,**
- 2). a QC report for the above samples
- 3). a copy of the chain of custody, and
- 4). a bill for analytical services.

All analyses were completed satisfactorily and all QC samples were found to be within our control limits.

If you have any questions please contact me. McC Campbell Analytical Laboratories strives for excellence in quality, service and cost. Thank you for your business and I look forward to working with you again.

Best regards,

Angela Rydelius, Lab Manager

McC Campbell Analytical, Inc.



1534 Willow Pass Rd
 Pittsburg, CA 94565-1701
 (925) 252-9262

CHAIN-OF-CUSTODY RECORD

WorkOrder: 0702471

ClientID: BEIA

EDF

Fax

Email

HardCopy

ThirdParty

Report to:

Mark Detterman
 Blymyer Engineers, Inc.
 1829 Clement Avenue
 Alameda, CA 94501-1395

Email: MDetterman@blymyer.com
 TEL: (510) 521-3773 FAX: (510) 865-2594
 ProjectNo: Former Fiesta Beverages
 PO:

Bill to:

Accounts Payable
 Blymyer Engineers, Inc.
 1829 Clement Avenue
 Alameda, CA 94501-1395

Requested TAT: 5 days

Date Received: 02/21/2007

Date Printed: 02/22/2007

Sample ID	ClientSampID	Matrix	Collection Date	Hold	Requested Tests (See legend below)														
					1	2	3	4	5	6	7	8	9	10	11	12			
0702471-001	MW-1R	Water	2/19/07 10:10:00	<input type="checkbox"/>	A	A													
0702471-002	MW-2	Water	2/19/07 11:50:00	<input type="checkbox"/>	A														
0702471-003	MW-3	Water	2/19/07 12:10:00	<input type="checkbox"/>	A														
0702471-004	MW-4	Water	2/19/07 11:40:00	<input type="checkbox"/>	A														
0702471-005	MW-5	Water	2/19/07 9:15:00 AM	<input type="checkbox"/>	A														
0702471-006	MW-6	Water	2/19/07 11:15:00	<input type="checkbox"/>	A														
0702471-007	MW-7	Water	2/19/07 12:30:00	<input type="checkbox"/>	A														
0702471-008	MW-8	Water	2/19/07 12:50:00	<input type="checkbox"/>	A														
0702471-009	MW-9	Water	2/19/07 1:30:00 PM	<input type="checkbox"/>	A														

Test Legend:

1	G-MBTX_W	2	PREFD REPORT	3		4		5	
6		7		8		9		10	
11		12							

Prepared by: Melissa Valles

Comments: Run highest MTBE result by 8260 for all oxys/additives

NOTE: Samples are discarded 60 days after results are reported unless other arrangements are made. Hazardous samples will be returned to client or disposed of at client expense.



McC Campbell Analytical, Inc.

"When Quality Counts"

1534 Willow Pass Road, Pittsburg, CA 94565-1701
Web: www.mcccampbell.com E-mail: main@mcccampbell.com
Telephone: 877-252-9262 Fax: 925-252-9269

Blymyer Engineers, Inc. 1829 Clement Avenue Alameda, CA 94501-1395	Client Project ID: Former Fiesta Beverages	Date Sampled: 02/19/07
		Date Received: 02/21/07
	Client Contact: Mark Detterman	Date Extracted: 02/23/07-02/24/07
	Client P.O.:	Date Analyzed: 02/23/07-02/24/07

Gasoline Range (C6-C12) Volatile Hydrocarbons as Gasoline with BTEX and MTBE*

Extraction method: SW5030B

Analytical methods: SW8021B/8015Cm

Work Order: 0702471

Lab ID	Client ID	Matrix	TPH(g)	MTBE	Benzene	Toluene	Ethylbenzene	Xylenes	DF	% SS
001A	MW-1R	W	200,a,i	ND	8.0	0.80	12	8.7	1	101
002A	MW-2	W	51,a	7.1	8.0	1.6	1.0	2.8	1	91
003A	MW-3	W	280,a	ND	49	11	18	23	1	96
004A	MW-4	W	ND	ND	ND	ND	ND	ND	1	105
005A	MW-5	W	ND	5.6	ND	ND	ND	ND	1	111
006A	MW-6	W	ND	ND	ND	ND	ND	ND	1	106
007A	MW-7	W	ND	ND	ND	ND	ND	ND	1	115
008A	MW-8	W	ND,i	ND	ND	ND	ND	ND	1	115
009A	MW-9	W	ND	ND	ND	ND	ND	ND	1	109

Reporting Limit for DF =1; ND means not detected at or above the reporting limit	W	50	5.0	0.5	0.5	0.5	0.5	1	µg/L
	S	NA	NA	NA	NA	NA	NA	1	mg/Kg

* water and vapor samples and all TCLP & SPLP extracts are reported in ug/L, soil/sludge/solid samples in mg/kg, wipe samples in µg/wipe, product/oil/non-aqueous liquid samples in mg/L.

cluttered chromatogram; sample peak coelutes with surrogate peak.

+The following descriptions of the TPH chromatogram are cursory in nature and McC Campbell Analytical is not responsible for their interpretation: a) unmodified or weakly modified gasoline is significant; b) heavier gasoline range compounds are significant(aged gasoline?); c) lighter gasoline range compounds (the most mobile fraction) are significant; d) gasoline range compounds having broad chromatographic peaks are significant; biologically altered gasoline?; e) TPH pattern that does not appear to be derived from gasoline (stoddard solvent / mineral spirit?); f) one to a few isolated non-target peaks present; g) strongly aged gasoline or diesel range compounds are significant; h) lighter than water immiscible sheen/product is present; i) liquid sample that contains greater than ~1 vol. % sediment; j) reporting limit raised due to high MTBE content; k) TPH pattern that does not appear to be derived from gasoline (aviation gas). m) no recognizable pattern; n) TPH(g) range non-target isolated peaks subtracted out of the TPH(g) concentration at the client's request; p) see attached narrative.



QC SUMMARY REPORT FOR SW8021B/8015Cm

W.O. Sample Matrix: Water

QC Matrix: Water

WorkOrder: 0702471

Analyte	EPA Method SW8021B/8015Cm		Extraction SW5030B			BatchID: 26382			Spiked Sample ID: 0702466-016A			
	Sample	Spiked	MS	MSD	MS-MSD	LCS	LCSD	LCS-LCSD	Acceptance Criteria (%)			
	µg/L	µg/L	% Rec.	% Rec.	% RPD	% Rec.	% Rec.	% RPD	MS / MSD	RPD	LCS/LCSD	RPD
TPH(btex) [£]	ND	60	98.4	96	2.44	95.5	94.9	0.625	70 - 130	30	70 - 130	30
MTBE	ND	10	88.6	91	2.65	106	105	0.916	70 - 130	30	70 - 130	30
Benzene	ND	10	107	111	3.79	100	107	6.94	70 - 130	30	70 - 130	30
Toluene	ND	10	98.8	96.4	2.48	91.6	98.2	6.96	70 - 130	30	70 - 130	30
Ethylbenzene	ND	10	109	99.7	8.74	102	106	4.09	70 - 130	30	70 - 130	30
Xylenes	ND	30	107	96.7	9.84	100	100	0	70 - 130	30	70 - 130	30
%SS:	90	10	102	98	3.50	94	103	9.27	70 - 130	30	70 - 130	30

All target compounds in the Method Blank of this extraction batch were ND less than the method RL with the following exceptions:
NONE

BATCH 26382 SUMMARY

Sample ID	Date Sampled	Date Extracted	Date Analyzed	Sample ID	Date Sampled	Date Extracted	Date Analyzed
0702471-001	2/19/07 10:10 AM	2/24/07	2/24/07 12:28 AM	0702471-002	2/19/07 11:50 AM	2/24/07	2/24/07 5:56 PM
0702471-003	2/19/07 12:10 PM	2/24/07	2/24/07 2:05 AM				

MS = Matrix Spike; MSD = Matrix Spike Duplicate; LCS = Laboratory Control Sample; LCSD = Laboratory Control Sample Duplicate; RPD = Relative Percent Deviation.

% Recovery = 100 * (MS-Sample) / (Amount Spiked); RPD = 100 * (MS - MSD) / ((MS + MSD) / 2).

MS / MSD spike recoveries and / or %RPD may fall outside of laboratory acceptance criteria due to one or more of the following reasons: a) the sample is inhomogenous AND contains significant concentrations of analyte relative to the amount spiked, or b) the spiked sample's matrix interferes with the spike recovery.

£ TPH(btex) = sum of BTEX areas from the FID.

cluttered chromatogram; sample peak coelutes with surrogate peak.



QC SUMMARY REPORT FOR SW8021B/8015Cm

W.O. Sample Matrix: Water

QC Matrix: Water

WorkOrder: 0702471

EPA Method SW8021B/8015Cm		Extraction SW5030B			BatchID: 26386			Spiked Sample ID: 0702472-003A				
Analyte	Sample	Spiked	MS	MSD	MS-MSD	LCS	LCSD	LCS-LCSD	Acceptance Criteria (%)			
	µg/L	µg/L	% Rec.	% Rec.	% RPD	% Rec.	% Rec.	% RPD	MS / MSD	RPD	LCS/LCSD	RPD
TPH(btex) [£]	ND	60	96.9	95.6	1.40	103	95.8	7.18	70 - 130	30	70 - 130	30
MTBE	ND	10	92	93.4	1.44	83.1	104	22.4	70 - 130	30	70 - 130	30
Benzene	ND	10	102	106	3.14	110	99.5	9.72	70 - 130	30	70 - 130	30
Toluene	ND	10	94.4	96.8	2.52	102	92.6	10.1	70 - 130	30	70 - 130	30
Ethylbenzene	ND	10	95.3	100	4.79	105	104	1.45	70 - 130	30	70 - 130	30
Xylenes	ND	30	96	96.7	0.692	100	103	3.28	70 - 130	30	70 - 130	30
%SS:	106	10	95	99	4.16	104	96	8.13	70 - 130	30	70 - 130	30

All target compounds in the Method Blank of this extraction batch were ND less than the method RL with the following exceptions:
NONE

BATCH 26386 SUMMARY

Sample ID	Date Sampled	Date Extracted	Date Analyzed	Sample ID	Date Sampled	Date Extracted	Date Analyzed
0702471-004	2/19/07 11:40 AM	2/23/07	2/23/07 4:19 PM	0702471-005	2/19/07 9:15 AM	2/23/07	2/23/07 4:49 PM
0702471-006	2/19/07 11:15 AM	2/23/07	2/23/07 5:49 PM	0702471-007	2/19/07 12:30 PM	2/23/07	2/23/07 6:19 PM
0702471-008	2/19/07 12:50 PM	2/24/07	2/24/07 1:16 AM	0702471-009	2/19/07 1:30 PM	2/24/07	2/24/07 1:46 AM

MS = Matrix Spike; MSD = Matrix Spike Duplicate; LCS = Laboratory Control Sample; LCSD = Laboratory Control Sample Duplicate; RPD = Relative Percent Deviation.

% Recovery = 100 * (MS-Sample) / (Amount Spiked); RPD = 100 * (MS - MSD) / ((MS + MSD) / 2).

MS / MSD spike recoveries and / or %RPD may fall outside of laboratory acceptance criteria due to one or more of the following reasons: a) the sample is inhomogenous AND contains significant concentrations of analyte relative to the amount spiked, or b) the spiked sample's matrix interferes with the spike recovery.

£ TPH(btex) = sum of BTEX areas from the FID.

cluttered chromatogram; sample peak coelutes with surrogate peak.