Fiesta Beverages 7150 Island Queen Dr. Sparks, NV 89436



June 24, 2008

Ms. Barbara Jakub Alameda County Environmental Health Department 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 Ms. Jakub,

"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

RECEIVED

9:05 am, Jul 17, 2008

Alameda County
Environmental Health

Second Quarter 2008 Groundwater Monitoring Event

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

> June 10, 2008 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.

Mark E. Detterman, CEG

Sentor Geologist

Michael S. Lewis, REA

Vice President, Technical Services

Table of Contents

1.0 Introd	luction and Background	. 1
2.0 Groun	ndwater Sample Collection and Analytical Methods	.8
3.0 Groun	ndwater Flow Data and Groundwater Sample Analytical Results	.9
4.0 Intrin	sic Bioremediation Groundwater Sample Analytical Results	11
5.0 Conc	lusions and Recommendations	13
	Tables	
Table I:	Summary of Groundwater Elevation Measurements	
Table II:	Summary of Groundwater Sample Hydrocarbon Analytical Results	
Table III:	Summary of Groundwater Sample Fuel Oxygenate Analytical Results	
Table IV:	Summary of Groundwater Intrinsic Bioremediation Field Results	
Table V:	Summary of Groundwater Intrinsic Bioremediation Analytical Results	
	Figures	
	9	
Figure 1:	Site Location Map	
Figure 2:	Site Plan and Groundwater Gradient, May 29, 2008	
Figure 3:	Concentration of TPHG vs. Time in Well MW-3	
Figure 4:	Concentration of Benzene vs. Time in Well MW-3	
Figure 5:	Concentration of TPHG vs. Time in Well MW-1 and MW-1R	
Figure 6:	Concentration of Benzene vs. Time in Well MW-1 and MW-1R	
	Appendix	
* *	Standard Operating Procedures, Blaine Tech Services, Inc.	
Appendix B:	Well Monitoring Data Sheets and Well Gauging Data, Blaine Tech Services, Inday 29, 2008	c.,
Appendix C:		

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 crossgradient 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 bio-nutrient 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 was approved by the ACHCSA on May 4, 2007. A pre-remedial groundwater sampling event to determine pre-remedial bacterial populations in groundwater, in the event of a bacterial die-off related to remedial injections, occurred on April 27, 2007. Remedial activities began on May 22, 2007 with a volume test injection. The first injection of RegenOx occurred between June 4 and June 7, 2007, and the second event occurred on June 26 and 27, 2007. It was not possible to inject the entire volume of RegenOx specified by Regenesis due to resurfacing of the injected material. On August 9, 2007, an abbreviated interim round of sampling occurred on selected wells (MW-1R, MW-2, MW-3, and MW-5) to help determine the progress of the remedial actions at the site. Elevated concentrations

of hydrocarbons were detected in plume core wells MW-1R and MW-3. As a consequence, an additional round of RegenOx injection occurred on September 12 and 13, 2007. These events were reported under separate cover.

On August 28, 2007, twenty-three 55-gallon drums of soil and fifteen 55-gallon drums of purge water, development water, and groundwater were removed from the subject site. The drums were transported by NRC Environmental to Crosby and Overton in Long Beach, California. The drums of soil represented soil cuttings from the installation of all soil bores and wells since 1993. The drums of water had accumulated since the installation of wells MW-1R, and MW-4 through MW-9, as well as water return flow to the surface during remedial injection activities.

On March 28, 2008, Blymyer Engineers received word that a new case worker, Ms. Barbara Jakub, had been assigned to the project by the ACEHD.

2.0 Groundwater Sample Collection and Analytical Methods

This report documents the interim sampling of groundwater conducted for the Second Quarter 2008 groundwater monitoring event at the subject site (Figure 1). Quarterly groundwater samples were collected from monitoring wells MW-1R and MW-2 through MW-9 on May 29, 2008. 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 during the sampling event. 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), Oxygen-Reduction Potential (ORP), and Ferrous Iron were measured post-purge. 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 for each event 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 from all wells were analyzed for TPH as gasoline by Modified EPA Method 8015; and BTEX and MTBE by EPA Method 8021B. Tables II to V summarize current and previous analytical results for groundwater samples. The laboratory analytical report 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 8.39 to 9.06 feet below the top of the casings. Depth to groundwater in general has increased an average of 0.47 feet since the March 2008 sampling event; however, there were significant divergences from the average. In well MW-6 the depth to water decreased 1.4 feet, in well MW-1R it increased by 0.02 feet, and in well MW-2 it increased 1.35 feet. In the previous event, mounding in the vicinity of wells MW-2 and MW-3 suggested the infiltration of rainwater in the vicinity well MW-2. It had been observed that the integrity of pavement in the area of well MW-2 had been impacted as a result of the vigorousness of remedial chemical oxidation of hydrocarbons in the vicinity of the well. The current record dry spring appears to have returned the groundwater flow direction to the typical flow direction for the site. Flows to the west are generally consistent with historical data. Conversely, southern and eastern flow directions have also been observed previously. The average gradient was calculated at approximately 0.014 feet/foot for the current event.

Between the February and May sampling events, concentrations of TPH as gasoline and BTEX rose modestly in well MW-1R, but continued to decrease in wells MW-2 and MW-3. All compounds were non-detectable in well MW-2, while only benzene was detected in well MW-3. All other wells (upgradient, downgradient, and lateral) were non-detectable for TPH as gasoline and BTEX. TPH as gasoline and benzene in well MW-1R were over the generic non-drinking water ESL. During the previous quarter, all contaminants in all wells were below their respective non-drinking water ESL. Recent previous higher concentrations in wells immediately adjacent to the former tank basin are presumed to have been the result of the mobilization of hydrocarbons from soil to groundwater during the injection of RegenOx between June and September 2007. The September 2007 injection (third and last) was concentrated in the vicinity of wells MW-1R, MW-2, and upgradient of MW-3, and likely mobilized contaminants near these locations.

Concentrations of MTBE were not detected any of the wells sampled during the May 2008 sampling event. MTBE has not been detected in all wells for three consecutive quarters.

Data gathered prior to remedial injections have provided evidence at the site for recontamination of groundwater upon rise up into contaminated soil (MW-3; Figures 3 and 4) as well as drainage from

soil to groundwater as groundwater drops in elevation at the site (MW-1 / MW-1R; Figures 5 and 6). Data from the current quarter is consistent with these historical trends. This is a departure from the two previous monitoring events. Data from the two previous quarters is assumed to have been related to the injection of RegenOx fluids, and was presumed to indicate some progress in remedial treatments. Data from MW-1R over the past two quarters tended to suggest that granular backfill and the soil immediately adjacent to and beneath the 6-inch-diameter gas main located approximately 5 feet to the north of well MW-1R and former MW-1 might be providing a reservoir for hydrocarbons not easily and effectively reached. (For consistency all groundwater elevations in Figures 3 to 6 utilized the GeoTracker wellhead survey elevations to determine the groundwater elevation.)

Four quarters of groundwater samples have been analyzed for the fuel oxygenates di-isopropyl ether (DIPE), ethyl *tert*-butyl ether (ETBE), MTBE, *tert*-amyl methyl ether (TAME), and *tert*-butyl alcohol (TBA), by EPA Method 8260B. Ethanol and methanol have also been analyzed and were nondetectable. Due to the consistency of the data, fuel oxygenate analysis was eliminated. Only MTBE and TAME have been detected in groundwater (June 2003; Table III). 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 May sampling event; however, post-purge field parameters were collected. RNA parameters were collected near the bottom of the well, within the screened interval, to collect representative values of vicinity groundwater and to minimize the effect of standard purging on the parameters. 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 (Mn⁴⁺) to soluble manganese (Mn²⁺), insoluble ferric iron (Fe³⁺) to soluble ferrous iron (Fe²⁺), 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. DO was present in a moderately tight range of concentrations in post-purge groundwater (ranging from 0.72 milligrams per liter (mg/L) to 1.75 mg/L). The current concentrations are generally higher than previous recent data, although previous sampling events in June of a given year have generally yielded higher DO concentrations. This tends to suggest that the higher concentrations of DO are related to infiltration of rainwater into the subsurface as opposed to overly vigorous purging. As seen previously at the site, the lower concentrations of DO have been identified as a limiting factor retarding biological activity. While this may be less prevalent this quarter, it is likely that this is a continuing trend.

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. In general, plume core well MW-1R and downgradient wells MW-7, MW-8, and MW-9 contain lower ORP concentrations than the remaining wells. This suggests a two-fold process of more oxygenated rainwater infiltration in the vicinity of the tank basin that has not yet fully migrated across the entire area of study, and also suggests a higher demand for oxygen in the vicinity of well MW-1R. Higher ORP values have previously been located outside the plume core which suggests that the strongest demand for oxygen is located in the plume core. The lower ORP value in well MW-1R tends to suggest that the residual benefit from the injection of the RegenOx remedial product has dissipated.

Ferrous iron was also investigated during the current sampling event and was only detected in wells MW-1R and MW-3. During the previous monitoring event all wells were non-detectable for ferrous iron. This change also suggests that the supply of DO from the injection of the RegenOx remedial product has ceased. The presence of ferrous iron in well MW-1R indicates that Mn – Fe degrading microbial colonies near this well have resumed microbial degradation of the contaminants (at a significantly slower rate) due to the relative lack of DO in the vicinity of the well. It is likely that the concentration of DO in other wells has not allowed these colonies to reestablish sufficiently yet.

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 summary and conclusions were generated from the available data discussed above:

- Depth to groundwater in general has increased an average of 0.47 feet since the March 2008 sampling event; however, there were significant divergences from the average. This is likely due to the record dry spring and the lack of rainwater infiltration into the subsurface in the vicinity of wells MW-2 and MW-3 (mounding was documented during the previous monitoring and sampling event). For the current event, an average gradient was calculated at approximately 0.014 feet/foot.
- Between the February and May sampling events, concentrations of TPH as gasoline and BTEX
 rose modestly in well MW-1R, but continued to decrease in wells MW-2 and MW-3. Both
 represent predictable trends based on declining groundwater levels alone. All other wells were
 non-detectable for TPH as gasoline and BTEX.
- MTBE was not detected in any of the wells during the May 2008 sampling event. MTBE has
 not been detected in all wells for three consecutive quarters.
- Only TPH as gasoline in well MW-1R was over the generic non-drinking water ESL. During
 the previous quarter, all contaminants in all wells were below their respective non-drinking
 water ESL.
- Recent higher concentrations in the vicinity of the former tank basin are presumed to have been
 the result of a mobilization of hydrocarbons from soil to groundwater due to the injection of
 RegenOx fluids between June and September 2007. The last injection was focused near plume
 core wells and likely mobilized contaminants near the locations.
- Current concentrations of DO are generally higher than recent data; however, previous data from mid-year sampling events have generally yielded higher concentrations of DO. This line of evidence suggests the higher concentrations are related to rainwater infiltration. Lower concentrations of DO have typically been identified as a limiting factor at the site.
- ORP values in plume core well MW-1R and downgradient wells MW-7, MW-8, and MW-9
 contain lower ORP concentrations than, upgradient, and lateral wells. This is likely related to a

two-fold process of more oxygenated rainwater infiltrating in the vicinity of the tank basin that has not yet migrated to the downgradient wells, and also a higher oxygen demand in the plume core as represented by well MW-1R. Higher ORP values in wells outside the plume core have previously suggested the strongest demand for oxygen is appropriately located in the plume core. The lower ORP value in well MW-1R suggests that the benefits from the injection RegenOx has largely ceased.

- Ferrous iron was detected only in wells MW-1R and MW-3 this quarter, while all wells have been non-detectable previously for ferrous iron. The presence of ferrous iron in these two wells suggest that the supply of DO derived from the injected RegenOx has been largely used.
- The low DO and ferrous iron in well MW-1R indicate that Mn Fe degrading microbial colonies near this well have resumed microbial degradation of the contaminants at a significantly slower rate.
- Pre-remedial injection data have documented 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. Data from the current quarter is again consistent with these historical
 trends. This is a change from the two previous monitoring events which were affected by the
 injection of RegenOx fluids at the site.
- Data from MW-1R for the previous two quarters has suggested that granular backfill and the soil immediately adjacent to and beneath the 6-inch-diameter gas main located approximately 5 feet to the north of well MW-1R and former MW-1 might be providing a reservoir for hydrocarbons not easily and effectively reached.

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

- The next quarterly groundwater sampling event should occur in August 2008.
- Collection of limited RNA indicator data should be continued as a modest cost saving measure.
 The collection of DO, ORP, and ferrous iron field data may help contribute to the understanding of biodegradation beneath the site. Collection of additional laboratory RNA indicator data can be resumed if a need is documented.

- Interim corrective actions should be continued in accordance with the approved ICAP. Future quarterly monitoring events can be used to gauge the effectiveness of corrective actions.
- A copy of this report will be forwarded to:

Ms. Barbara Jakub Alameda County Environmental Health Department Environmental Protection Division 1131 Harbor Bay Parkway, Suite 250 Alameda, CA 94502-6577

Table I, Summary of Groundwater Elevation Measurements BEI Job No. 203004, Former Fiesta Beverage 966 89th Avenue, Oakland, California **TOC** Elevation Depth to Water Water Surface Elevation Well ID Date (feet) (feet) (feet) MW-1 18.72 8/6/1993 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 10.34 8.38 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 8.90 9/18/2003 9.82 12/15/2003 8.15 10.57 8.67 10.05 6/15/2004 12/15/2004 7.99 10.73 6/29/2005 7.88 10.84 21.70 5/8/2006 Destroyed Destroyed 2/19/2007 Destroyed Destroyed 6/21/2007 Destroyed Destroyed 11/8/2007 Destroyed Destroyed 2/28/2008 Destroyed Destroyed 5/29/2008 Destroyed Destroyed MW-1R 21.73 8.49 13.24 6/12/2006 7.94 13.79 2/19/2007 8.71 13.02 6/21/2007 8/9/2007 8.83 12.90 9.80 11/8/2007 11.93 2/28/2008 8.74 12.99 5/29/2008 8.76 12.97

Table I, Summary of Groundwater Elevation Measurements BEI Job No. 203004, Former Fiesta Beverage 966 89th Avenue, Oakland, California **TOC** Elevation Depth to Water Water Surface Elevation Well ID Date (feet) (feet) (feet) MW-2 18.44 8/6/1993 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 8.24 1/18/2001 10.20 4/25/2001 7.88 10.56 3/17/03* 7.08 11.36 8.90 9.54 6/23/2003 9.83 9/18/2003 8.61 7.97 12/15/2003 10.47 6/15/2004 8.42 10.02 12/15/2004 8.00 10.44 6/29/2005 9.51 8.93 21.45 6/12/2006 8.25 13.20 2/19/2007 8.12 13.33 6/21/2007 9.00 12.45 8/9/2007 8.62 12.83 11/8/2007 8.60 12.85 2/28/2008 7.20 14.25 5/29/2008 8.55 12.90

Table I, Summary of Groundwater Elevation Measurements BEI Job No. 203004, Former Fiesta Beverage 966 89th Avenue, Oakland, California **TOC** Elevation Depth to Water Water Surface Elevation Well ID Date (feet) (feet) (feet) MW-3 19.01 8/6/1993 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 9.05 9.96 6/23/2003 9.11 9.90 9/18/2003 12/15/2003 8.03 10.98 8.85 10.16 6/15/2004 12/15/2004 8.84 10.17 6/29/2005 9.00 10.01 22.02 6/12/2006 8.62 13.40 2/19/2007 8.12 13.90 6/21/2007 9.86 12.16 8/9/2007 9.60 12.42 11/8/2007 8.83 13.19 2/28/2008 7.99 14.03 5/29/2008 8.57 13.45 MW-4 21.34 8.37 12.97 6/12/2006 7.77 2/19/2007 13.57 6/21/2007 8.48 12.86 11/8/2007 8.61 12.73 2/28/2008 7.73 13.61 5/29/2008 8.39 12.95

Table I, Summary of Groundwater Elevation Measurements BEI Job No. 203004, Former Fiesta Beverage 966 89th Avenue, Oakland, California **TOC** Elevation Depth to Water Water Surface Elevation Well ID Date (feet) (feet) (feet) MW-5 22.53 8.75 13.78 6/12/2006 2/19/2007 8.61 13.92 6/21/2007 9.05 13.48 8/9/2007 9.17 13.36 11/8/2007 9.11 13.42 8.18 2/28/2008 14.35 5/29/2008 9.06 13.47 MW-6 21.97 6/12/2006 8.59 13.38 2/19/2007 7.93 14.04 6/21/2007 9.83 12.14 9.58 12.39 11/8/2007 9.90 2/28/2008 12.07 8.50 13.47 5/29/2008 MW-7 21.21 6/12/2006 8.31 12.90 2/19/2007 7.85 13.36 6/21/2007 8.51 12.70 11/8/2007 8.68 12.53 2/28/2008 7.81 13.40 5/29/2008 8.60 12.61 MW-8 20.97 6/12/2006 8.37 12.60 2/19/2007 7.99 12.98 6/21/2007 8.53 12.44 8.61 12.36 11/8/2007 7.79 13.18 2/28/2008 5/29/2008 8.61 12.36

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-9	6/12/2006		8.50	12.48					
	2/19/2007	20.98	8.08	12.90					
	6/21/2007		8.55	12.43					
	11/8/2007		8.67	12.31					
	2/28/2008		8.02	12.96					
	5/29/2008		8.51	12.47					

Notes: TOC = Top of Casing

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

NM = Not measured

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

Elevations in feet above mean sea level

	966 89th Avenue, Oakland, California										
Well ID	Sample Date	Modified EPA Method 8015 (μg/L)		EPA	Method 8020 α (μg/L)	or 8021B					
		TPH as Gasoline	Benzene	Toluene	Ethylbenzene	Total Xylenes	MTBE				
1	MCL	N/A	1	150	700	1,750	13				
Drinking V	Vater Source 1	100	1	40	30	20	5				
So	nking Water urce 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				
	6/21/2007	Destroyed	Destroyed	Destroyed	Destroyed	Destroyed	Destroyed				
	11/8/2007	Destroyed	Destroyed	Destroyed	Destroyed	Destroyed	Destroyed				
	2/28/2008	Destroyed	Destroyed	Destroyed	Destroyed	Destroyed	Destroyed				
	5/29/2008	Destroyed	Destroyed	Destroyed	Destroyed	Destroyed	Destroyed				

, , ,									
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		
1	MCL	N/A	1	150	700	1,750	13		
Drinking V	Vater Source 1	100	1	40	30	20	5		
	nking Water urce 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		
	6/21/2007	<50	< 0.5	< 0.5	<0.5	< 0.5	< 5.0		
	8/9/2007	870 ^a	140	6.30	23	22	<10		
	11/8/2007	3,800 ^a	330	22	140	130	<30		
	2/28/2008	150 ^a	5.5	< 0.5	3.9	2.2	<5.0		
	5/29/2008	690 ^a	44	2	35	7.8	<5.0		

	966 89th Avenue, Oakland, California										
Well ID	Sample Date	Modified EPA Method 8015 (μg/L)		EPA	Method 8020 α (μg/L)	or 8021B					
		TPH as Gasoline	Benzene	Toluene	Ethylbenzene	Total Xylenes	MTBE				
]	MCL	N/A	1	150	700	1,750	13				
Drinking V	Water Source 1	100	1	40	30	20	5				
	nking Water urce ²	500	46	130	290	100	1,800				
MW-2	8/6/1993	2,700	1.3	1.7	2.0	8.1	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	8.2	10	13	NA				
	12/15/1998	200	62	17	4.9	14	4.4 ^b				
	1/18/2001	300 ^a	74	26	7.3	21	7.3				
	4/25/2001	<50°	4.5	2.2	0.6	1.9	< 5.0				
	3/17/2003*	78 ^a	26	3.3	1.5	3.5	NA ^d				
	6/23/2003	160 ^a	51	1.6	1.2	1.8	NA				
	9/18/2003	<50	2.1	<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	1.8	1.2	<30				
	12/15/2004	<50	11	0.97	0.6	0.9	7.8				
	6/29/2005	130	29	2.000	3.3	3.4	6.7				
	6/13/2006	150 ^a	59	3.0	3.4	2.7	11				
	2/19/2007	51 ^a	8	1.6	1.0	2.8	7.1				
	6/21/2007	<50	< 0.5	< 0.5	<0.5	<0.5	<5.0				
	8/9/2007	<50	< 0.5	< 0.5	<0.5	<0.5	<5.0				
	11/8/2007	160 ^a	23	5.0	5.3	14	<10				
	2/28/2008	<50	1.3	< 0.5	<0.5	<0.5	<5.0				
	5/29/2008	<50	< 0.5	<0.5	<0.5	<0.5	<5.0				

	966 89th Avenue, Oakland, California										
Well ID	Sample Date	Modified EPA Method 8015 (µg/L)		EPA	Method 8020 α (μg/L)	or 8021B					
		TPH as Gasoline	Benzene	Toluene	Ethylbenzene	Total Xylenes	MTBE				
I	MCL	N/A	1	150	700	1,750	13				
Drinking V	Vater Source 1	100	1	40	30	20	5				
	nking Water urce 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				
	6/21/2007	1,500 ^a	120	64	62	250	<50				
	8/9/2007	2,400 ^a	140	19	100	110	<65				
	11/8/2007	440 ^a	7.2	3.3	8.6	26	<15				
	2/28/2008	320 ^a	10	5.8	9.6	32	<12				
	5/29/2008	<50	1.0	<0.5	<0.5	< 0.5	<5.0				

700 07th Avenue, Oakiana, Camorina									
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		
I	MCL	N/A	1	150	700	1,750	13		
Drinking V	Vater Source 1	100	1	40	30	20	5		
	nking Water urce ²	500	46	130	290	100	1,800		
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		
	6/21/2007	<50	< 0.5	< 0.5	<0.5	< 0.5	5.9		
	11/8/2007	<50	< 0.5	< 0.5	<0.5	< 0.5	< 5.0		
	2/28/2008	<50	< 0.5	< 0.5	<0.5	< 0.5	< 5.0		
	5/29/2008	<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		
	6/21/2007	<50	< 0.5	< 0.5	<0.5	< 0.5	5.4		
	11/8/2007	<50	< 0.5	< 0.5	<0.5	< 0.5	< 5.0		
	2/28/2008	<50	< 0.5	< 0.5	<0.5	< 0.5	< 5.0		
	5/29/2008	<50	< 0.5	< 0.5	<0.5	< 0.5	< 5.0		
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		
	6/21/2007	<50	< 0.5	< 0.5	< 0.5	<0.5	<5.0		
	11/8/2007	<50	< 0.5	< 0.5	< 0.5	<0.5	<5.0		
	2/28/2008	<50	< 0.5	< 0.5	<0.5	<0.5	<5.0		
	5/29/2008	<50	< 0.5	< 0.5	<0.5	<0.5	<5.0		

700 07th Avenue, Oakiana, Camorina									
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		
I	MCL	N/A	1	150	700	1,750	13		
Drinking V	Vater Source 1	100	1	40	30	20	5		
	nking Water urce ²	500	46	130	290	100	1,800		
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		
	6/21/2007	<50	< 0.5	< 0.5	<0.5	<0.5	< 5.0		
	11/8/2007	<50	< 0.5	< 0.5	<0.5	< 0.5	< 5.0		
	2/28/2008	<50	< 0.5	< 0.5	<0.5	< 0.5	< 5.0		
	5/29/2008	<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		
	6/21/2007	<50	< 0.5	< 0.5	<0.5	< 0.5	< 5.0		
	11/8/2007	<50	< 0.5	< 0.5	<0.5	< 0.5	< 5.0		
	2/28/2008	<50	< 0.5	< 0.5	<0.5	< 0.5	< 5.0		
	5/29/2008	<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		
	6/21/2007	<50	< 0.5	< 0.5	<0.5	<0.5	5.6		
	11/8/2007	<50	< 0.5	< 0.5	<0.5	<0.5	< 5.0		
	2/28/2008	<50	< 0.5	< 0.5	<0.5	<0.5	<5.0		
	5/29/2008	<50	< 0.5	< 0.5	<0.5	<0.5	<5.0		

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

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

 $\langle x \rangle$ = 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.

Note: Shaded cell indicates that detected concentration exceeds

Non-Drinking Water ESL

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

Well ID	Sample Date				EPA Me	thod 8260E	3 (ug/L)			
well ID	Sample Date	TAME	TBA	EBD	1,2-DCA	DIPE	Ethanol	ETBE	Methanol	MTBE
Drinking	Water Source 1	NV	12	0.05	0.5	NV	50,000	NV	NV	5
	rinking Water Source ²	NV	18,000	152	204	NV	50,000	NV	NV	1,800
	3/17/2003	8.3	<5.0	NA	NA	< 0.50	NA	< 0.50	NA	10.0
MW-1	6/23/2003	6.4	<25	NA	NA	<2.5	NA	<2.5	NA	8.0
IVI VV - I	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
	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
MW-2	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
	3/17/2003	4.3	8.6	NA	NA	< 0.50	NA	< 0.50	NA	10.0
MANU 2	6/23/2003	2.6	<5.0	NA	NA	< 0.50	NA	< 0.50	NA	5.6
MW-3	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
MW-4	6/12/2006	NA	NA	NA	NA	NA	NA	NA	NA	6.1

Table III, Summary of Groundwater Sample Fuel Oxygenate Analytical Results BEI Job No. 203004, Former Fiesta Beverage 966 89th Avenue, Oakland, California EPA Method 8260B (ug/L) Sample Date Well ID **TAME TBA EBD** 1,2-DCA **DIPE ETBE** Methanol **MTBE** Ethanol Drinking Water Source 1 NV 12 0.05 0.5 NV 50,000 NV NV 5 Non-Drinking Water NV 18,000 152 204 NV 50,000 NV NV 1,800 Source ²

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 = Methly tert-butyl ether

 $(\mu g/L)$ = 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
- 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 Field Meter Field Meter Field Meter Field Test Kit Field Meter Dissoved Oxidation Ferrous Iron Field Field pH Well ID Sample Date Oxygen Reduction **Temperature Potential** (mg/L)(Fe 2+)(oF/oC)pH units (mV) MW-1 3/17/2003 NA NA NA 60.4 / 60.0 * 7.1 / 7.30.4 6.9 / 6.96/23/2003 NA NA 61.0 / 61.0 * 9/18/2003 0.4 NA NA 65.1 / 62.9 * 7.1 / 6.9NA NA 6.8 / 6.712/15/2003 1.1 13.1 / 13.4 6/15/2004 0.1 NA NA 64.5 / 63.4 * 6.9 / 7.012/15/2004 NA NA NA 15.4 / 17.5 7.0 / 6.90.24 / 0.171.0 4.5 19.78 / 21.63 7.15 / 7.086/29/2005 5/8/2006 Destroyed Destroyed Destroyed Destroyed Destroyed 2/19/2007 Destroyed Destroyed Destroyed Destroyed Destroyed 6/21/2007 Destroyed Destroyed Destroyed Destroyed Destroyed 11/8/2007 Destroyed Destroyed Destroyed Destroyed Destroyed 2/28/2008 Destroyed Destroyed Destroyed Destroyed Destroyed 5/29/2008 Destroyed Destroyed Destroyed Destroyed Destroyed MW-1R 6.90 / 6.926/13/2006 0.87 / 0.37172.9 / 172.9 0 / 017.31 / 17.36 2/19/2007 0.48 8.0 NA 12.2 / 15.8 6.95 / 6.86 0.62 22.0 NA 19.6 7.1 6/21/2007 11/8/2007 0.3 -60 NA 64.4 6.9 2/28/2008 0.28 0.0 156 63.2 6.98 5/29/2008 0.72 97 0.6 17.3 7.12

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

	ı		,			
		Field Meter	Field Meter	Field Test Kit	Field Meter	Field Meter
W 11 ID	G 1 D	Dissoved	Oxidation	Ferrous Iron	Field	Field pH
Well ID	Sample Date	Oxygen	Reduction		Temperature	
			Potential			
		(mg/L)	(mV)	(Fe 2+)	(o F / o C)	pH units
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
	6/21/2007	0.18	46	NA	18.3	7.1
	11/8/2007	0.4	209	NA	64.0	7.07
	2/28/2008	0.29	191	0.0	63.1	6.98
	5/29/2008	1.53	212	0.0	17.8	7.18

Table IV, Summary of Groundwater Intrinsic Bioremediation Field Results BEI Job No. 203004, Former Fiesta Beverage 966 89th Avenue, Oakland, California Field Meter Field Meter Field Test Kit Field Meter Field Meter Dissoved Oxidation Ferrous Iron Field Field pH Well ID Sample Date Oxygen Reduction **Temperature Potential** (mg/L)(Fe 2+)(o F / o C)pH units (mV) MW-3 3/17/2003 NA NA NA 63.3 / 60.9 * 7.4 / 7.666.4 / 66.9 * 7.3 / 7.26/23/2003 0.7 NA NA 9/18/2003 0.4 NA NA 63.7 / 62.6 * 7.1 / 7.1NA NA 14.7 / 15.1 6.5 / 6.412/15/2003 1.6 6/15/2004 0.0 NA NA 63.1 / 62.3 * 7.5 / 7.112/15/2004 NA NA NA 15.4 / 16.7 7.2 / 7.00.72 / 0.78141.7 / -67.6 0.9 17.65 / 18.79 6.94 / 7.026/29/2005 7.02 / 6.981.01 / 0.41170.0 / 168.5 0 / 06/13/2006 17.30 / 17.15 0.08 13.7 / 15.6 7.10 / 6.952/19/2007 81 NA 6/21/2007 0.10 39 NA 18.1 7.2 11/8/2007 0.30 -30 NA 62.5 7.04 2/28/2008 0.32 132 0.0 61.2 5.45 0.77 7.19 5/29/2008 186 0.6 16.3 MW-4 6/12/2006 0.67 / 0.33164.3 / 161.0 0.5 / 016.90 / 16.79 6.82 / 6.7998 2/19/2007 0.21 NA 13.7 / 15.0 7.14 / 7.030.31 NA 16.4 7.0 6/21/2007 118 11/8/2007 0.30 222 NA 62.7 6.96 2/28/2008 0.28 0.0 7.01 173 61.6 5/29/2008 1.07 228 0.0 16.2 6.81

Table IV, Summary of Groundwater Intrinsic Bioremediation Field Results BEI Job No. 203004, Former Fiesta Beverage 966 89th Avenue, Oakland, California Field Meter Field Meter Field Test Kit Field Meter Field Meter Dissoved Oxidation Ferrous Iron Field Field pH Well ID Sample Date Oxygen Reduction **Temperature** Potential (mg/L)(Fe 2+)(o F / o C)pH units (mV) MW-5 6/12/2006 0.61 / 0.31175.2 / 169.0 0 / 018.40 / 18.01 7.01 / 6.94 12.7 / 14.1 2/19/2007 1.98 -114 NA 6.93 / 6.736/21/2007 1.23 99 NA 16.8 7.1 11/8/2007 0.30 NA 63.9 6.85 211 2/28/2008 0.26 213 0.0 62.6 7.14 5/29/2008 0.80 249 0.0 16.5 7.18 MW-6 181.2 / 174.8 0 / 017.25 / 17.32 6/13/2006 3.10 / 0.816.94 / 6.830.21 -30 NA 6.58 / 6.742/19/2007 14.6 / 15.6 0.26 NA 16.2 7.1 6/21/2007 102 11/8/2007 0.60 -8 NA 63.5 6.99 2/28/2008 0.37 212 0.0 60.8 6.93 5/29/2008 1.75 194 0.0 16.3 7.22 MW-7 0.59 / 0.27172.5 / 171.8 0.5 / 0.218.14 / 18.00 6/12/2006 6.90 / 6.872/19/2007 0.10 110 NA 16.2 / 17.2 7.69 / 7.217.0 6/21/2007 0.14 123 NA 17.3 0.30 227 NA 64.5 6.90 11/8/2007

142

83

0.0

0.0

64.2

17.8

7.00

7.17

2/28/2008

5/29/2008

0.27

1.47

Table IV, Summary of Groundwater Intrinsic Bioremediation Field Results BEI Job No. 203004, Former Fiesta Beverage 966 89th Avenue, Oakland, California Field Meter Field Meter Field Meter Field Test Kit Field Meter Dissoved Oxidation Ferrous Iron Field Field pH Well ID Sample Date Reduction Oxygen **Temperature** Potential (mg/L)(Fe 2+)(o F / o C)pH units (mV) MW-8 6/12/2006 0.37 / 0.33186.1 / 180.4 0 / 018.55 / 18.39 6.85 / 6.857.23 / 7.07 2/19/2007 0.11 102 NA 15.2 / 16.6 6/21/2007 0.12 NA 17.2 7.1 111 0.30 232 NA 64.3 7.01 11/8/2007 2/28/2008 0.26 206 0.0 63.1 7.08 5/29/2008 1.23 72 0.0 17.5 7.22 MW-9 2.01 / 1.87 206.0 / 191.0 0 / 016.88 / 16.91 6.63 / 6.666/12/2006 0.08 NA 15.8 / 16.3 7.56 / 7.232/19/2007 101 0.12 7.1 6/21/2007 112 NA 16.5 11/8/2007 0.40 230 NA 65.1 6.94

208

94

0.0

0.0

62.1

17.1

7.01

7.33

Notes: mV = Millivolts

mg/L = Milligrams per liter

° F / ° C = degrees Fahrenheit / degrees Centigrade

0.26

1.44

* = degrees Fahrenheit

2/28/2008

5/29/2008

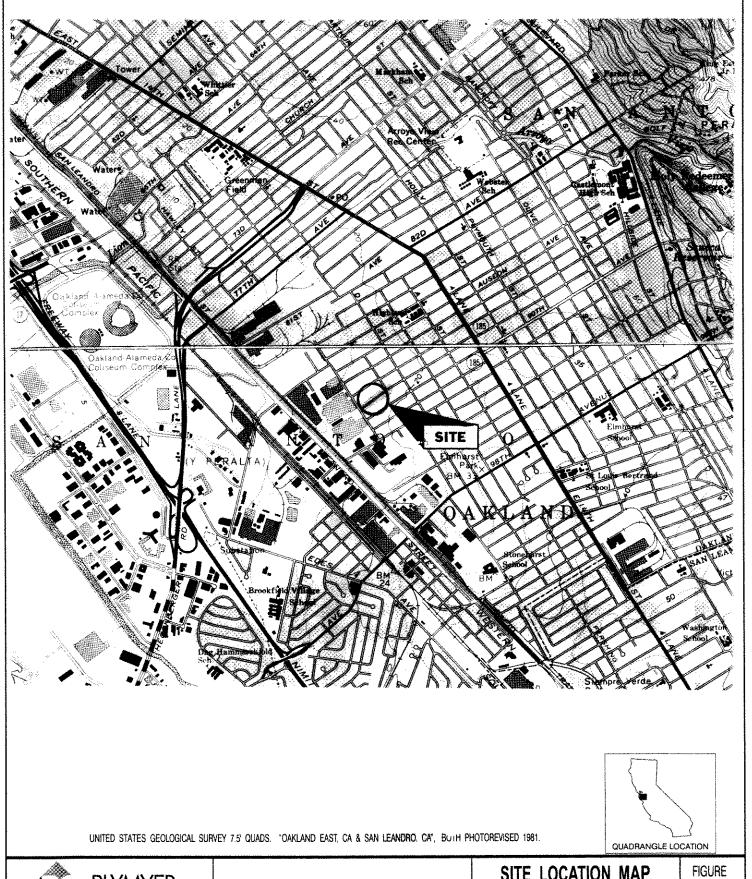
2.6 / 2.2 = Initial reading (pre-purge) / Final reading (post-purge)

NA = Not analyzed

Table V, S	Table V, Summary of Groundwater Intrinsic Bioremediation Analytical Results BEI Job No. 203004, Former Fiesta Beverage 966 89th Avenue, Oakland, California											
		Method SM 5310B	Method	E300.1	Method RSK 174							
Well ID	Sample Date	CO_2	Nitrate (as N)	Methane								
			μg/L									
MW-1	6/29/2005	490	490 <0.1 5									
	5/8/2006	Destroyed	Destroyed Destroyed									
MW-1R	6/13/2006	290	4.3	46	24							
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-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	51	<0.5								
MW-8	6/12/2006	330	<0.5									
MW-9	6/12/2006	240	240 8.3 44									

Notes: SM = Standard Method

 $mg/L = Milligrams per liter \ \mu g/L = Micrograms per liter \ CO_2 = Carbon Dioxide$



BEI JOB NO.

3-19-03

203004

SCALE IN FEET

1000

2000



SITE LOCATION MAP

FORMER FIESTA BEVERAGE 966 89TH AVE. OAKLAND, CA

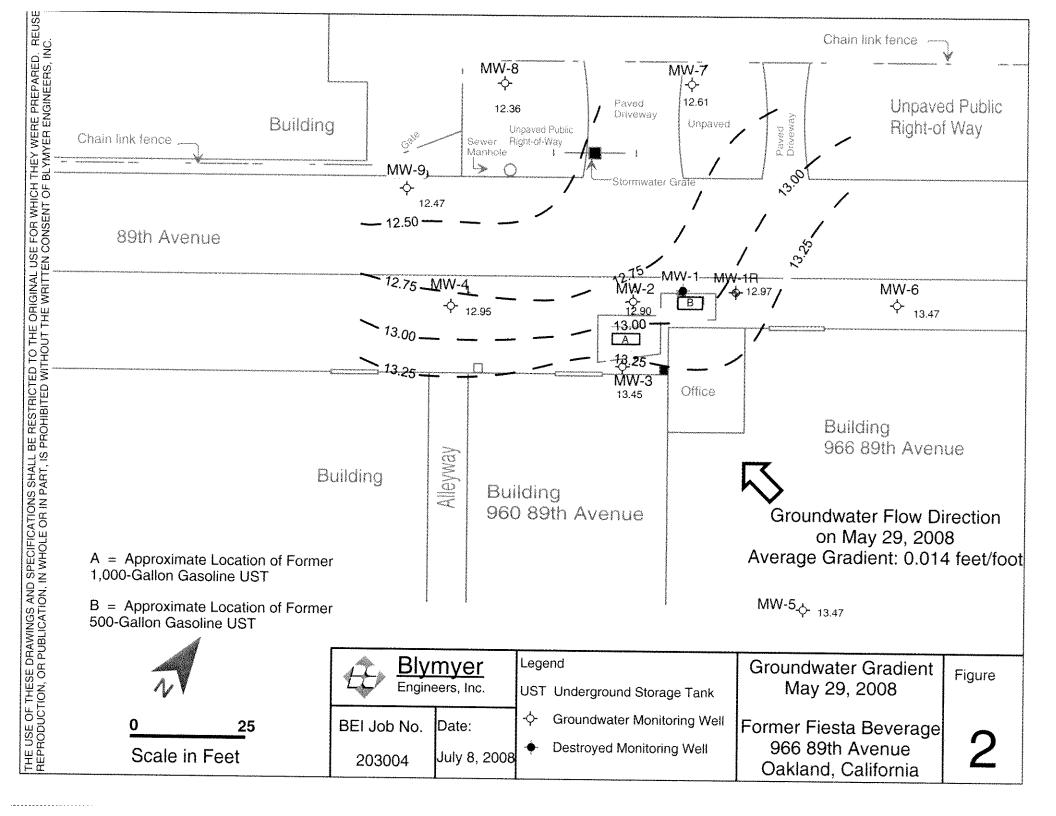


Figure 3: Concentration TPHG vs. Time in MW-3

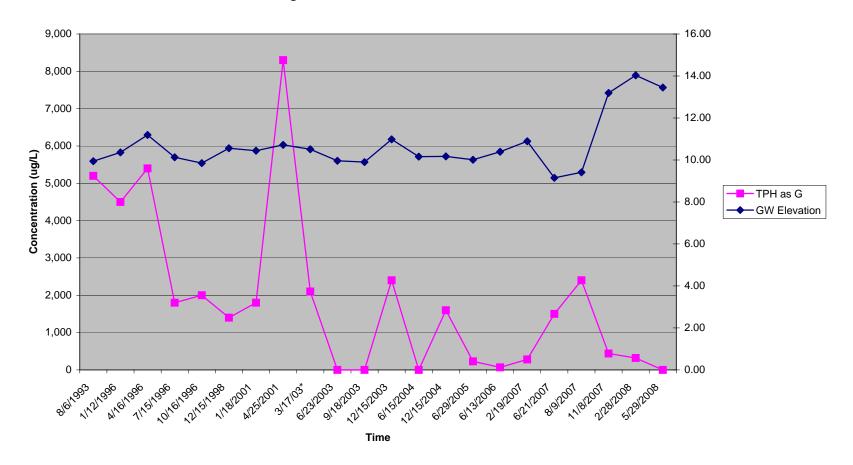


Figure 4: Concentration Benzene vs. Time in MW-3

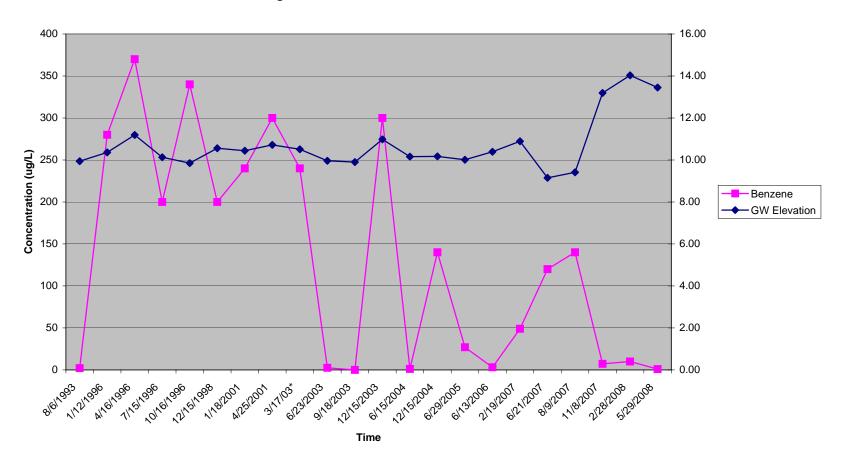


Figure 5: Concentration TPHG vs. TIme in MW-1 / MW-1R

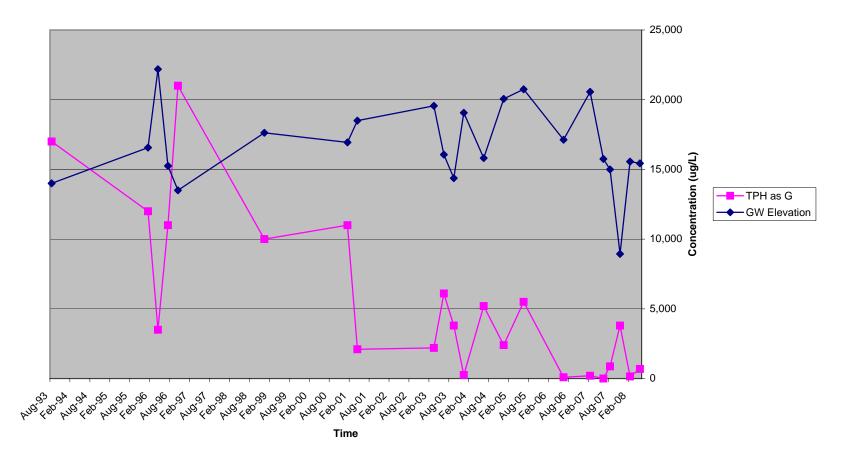
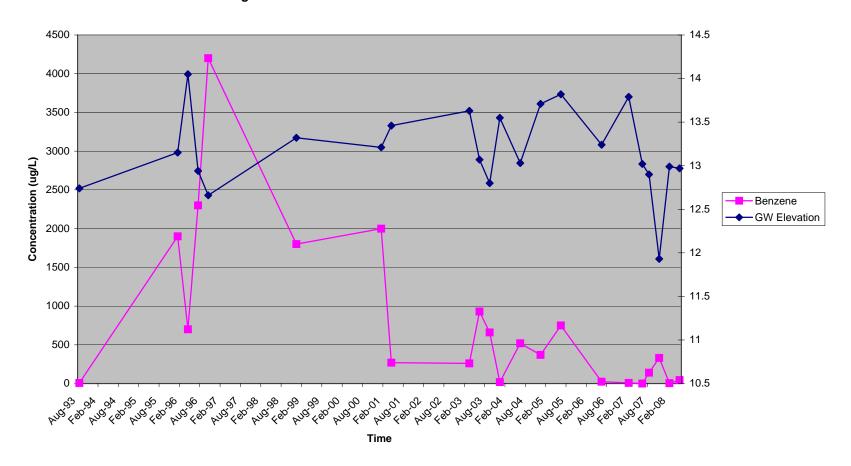


Figure 6: Concentation Benzene vs. Time in Well MW-1 / MW-1R



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 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 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.

GAUGING SOP Page 2 of 3

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.

GAUGING SOP Page 3 of 3

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

PURGING SOP Page 1 of 3

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:

Casing Volume = (TD - DTW) 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.

Sampling SOP

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.
- 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.
- 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.
- 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.

BLAINE TECH SERVICES, INC

Page 1 of 1

Appendix B

Well Monitoring Data Sheets and Well Gauging Data,

Blaine Tech Services, Inc. May 29, 2008

BLA	NE	SAN	JOSE C	1680 RO	GERS AVEN NIA 95112-11	UE OF		co	NDUC	T ANA	LYSIS	TO DI	ETECT	-	LAB			McCampbel	i	DHS#
TECH SEI				FAX	(408) <mark>573-77</mark>	71									ALL A	NALYSES N	AUST	MEET SPECIF DRNIA DHS AN	FICATIONS AN	D DETECTION
		INC.		PHONE	(408) 573-05	55										☐ EPA	ALII C		ID □ RWQCB RE	GION
CHAIN OF CL	JSTODY	BTS ;	# 080	52	9-4Wi	3S										LIA OTHER	₹			
CLIENT	Blymy	er Engin	eers, Ir	ic.		CONTAINERS		B)							SPEC	IAL INSTRU	JCTIO	NS		
SITE	Former	Fiesta I	Beverag	ge		TNC 7		(8021B)							Invo	ice and D			ъ.	
	966 89	th Aven	ue			ALL C	Q								}				nyer Engine	ers, Inc.
	Oakland	d, CA				TEA	(8015M)	& MTBE							ı	: Mark D				
			MATRIX	x COI	NTAINERS	COMPOSITE	08)	\ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \							EDF	Format I	Requ	ured.		
	1		S= SOIL W=H ₂ 0			OO	TPH-G	BTEX									. 1			
SAMPLE I.D.	DATE	TIME		TOTAL		0		BI	ļ						ADD'L	INFORMAT	ION	STATUS	CONDITION	LAB SAMPLE#
MW-1R	05/29/05	-	W	3	HCL Voa		Х	X												
MW-2		1117	W	3	HCL Voa		Х	Х												
MW-3		1218	W	3	HCL Voa		Х	х												
MW-4		0956	W	3	HCL Voa		Х	Х												
MW-5		1027	W	3	HCL Voa		Х	Х												
MW-6		1650	W	3	HCL Voa		Х	Х												
MW-7		1305	W	3	HCL Voa		Х	Х												
MW-8		1330	W	3	HCL Voa		Х	Х										7. 1		
MW-9	\underline{V}	1355	W	3	HCL Voa		х	Х									\dashv			
																	\dashv			
SAMPLING COMPLETED	DATE US 29 H		SAMPLIN PERFOR	NG RMED BY	WILL	14	n i	UON	16			<u> </u>		<u> </u>		TS NEEDED		s contracted		· · · · · · · · · · · · · · · · · · ·
RELEASED BY	th					DATE	9/0		TIME	12		RECE	IVED I	3Y_	J 5/	MPLE INSTOB			DATE	TIME
RELEASED BY	perplo	Cust	حیا ا نب)	1	DATE	a/576		TIME	- ১৯		REGE	IVED E	BY	1/2/1	1 de	141		DATE = /30/5	8 1612 TIME 8 1308
RELEASED BY				/		DATE			TIME			RECE	IVED E	BY	<u> </u>	n -	74		DATE	TIME

TIME DATE TIME SHIPPED VIA DATE SENT TIME SENT COOLER#

SPH or Purge Water Drum Log

Client:	JM		
Site Address: 966	89 th Mr. Oaliland	CA	

STATUS OF DRUM(S) UPON ARRIVAL										
Date	6/21/07	8/09/07	11/8/07	2/28/08	05/29/20					
Number of drum(s) empty:										
Number of drum(s) 1/4 full:										
Number of drum(s) 1/2 full:										
Number of drum(s) 3/4 full:										
Number of drum(s) full:	4 (Ø5 4	4	6					
Total drum(s) on site:	4(100)	4	5	Ś	67					
Are the drum(s) properly labeled?	γ	Y	Y	Y	4 NON BTS 2 BTS YEA					
Drum ID & Contents:	MANNAMO	purey H20	perze Mro	Pugett20	Tr are the					
If any drum(s) are partially or totally filled, what is the first use date:	mjecton	06/02/06		100 DATE	WA					

- 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	6/21/07	08/09/07	11/8/57	2/28/08	05/29/50					
Number of drums empty:										
Number of drum(s) 1/4 full:										
Number of drum(s) 1/2 full:					0					
Number of drum(s) 3/4 full:										
Number of drum(s) full:	5 1075 4 Nen	27.55.5	5 4004	5	B					
Total drum(s) on site:	6	6	6	6	4 MIN BCS					
Are the drum(s) properly labeled?	4	Y	4	X	Y					
Drum ID & Contents:	Perge 1/20	Purgetho	parxwater	Purple H20	purge the.					

LOCATION OF DRUM(S)

Describe location of drum(s): Biland building through All up dec Don't Let SLIDING DOOR DOOR OUT

FINAL STATUS		_				
Number of new drum(s) left on site this event	2	3/10/07	5	λ.	O	
Date of inspection:	6/2/ler	8/19/07	11/8/07	2/28/08	05/29/00	
Drum(s) labelled properly:	Ÿ	NIA	Y	Y	Y	
Logged by BTS Field Tech:	DR	MM	K5/	IW	WV	
Office reviewed by:	W	N	NOW	B	pow	

TEST EQUIPMENT CALIBRATION LOG

PROJECT NAM	IE FORMER 89	FIESTA BEV	ERHGE AND, OF	PROJECT NUMBER 086529-WW1							
NAME	EQUIPMENT NUMBER	DATE/TIME OF TEST	STANDARDS USED	EQUIPMENT READING	CALIBRATED TO: OR WITHIN 10%:	TEMP.	INITIALS				
HYRON L ULTRAMETER	6211286	05/29/8	pt: 4,7,10 Cond: 3500	pH: 7.00, 4.00 10:00 cond: 39:00	Yes	21°C	w				
(1	('	11	ORP: 236	DPP: 23.6	Yes	2100	un				
ZIOO P TURBIDIMETEA	· · · · · · · · · · · · · · · · · · ·	0720	NW, CO.1,72		Yes	NA	wn				
YSI 550A DO METER	OBENZY	05/29/08	D0%: 100 @10 mg/L	D090:99.8 09.52 mg/L	Yes	17.9°C	WW				
			-	٠.							

WELLHEAD INSPECTION CHECKLIST

Page (of /

	Date <u>05</u>	/20	1/06	C	Client	Bl	41	HER						
	Date <u>05</u> Site Addres	is and	-966 g	39+	h A	VE, C	AL	LANI	, CA					
	Job Numbe	r <u>50</u>	30529	- M	~				chnician					
	Mall ID		Well Inspected - No Corrective		er Bailed From	Compon	ents	Cap Replaced	Debris Removed From	Lock Replaced	Other Action Taken (explain		Well Not Inspected (explain	
	Well ID MW-11	2	Action Required		/ellbox	Cleane	ea		Wellbox	,	below)		below)	-
~		~ ~~~~~~~~~~~~		2/	2 B0	C75	M	SSING	\times	COVER	ED BY	4-	7FE	-
٠. ٠.	MW-3		\mathcal{D}		······································			77-74-14-14-14-14-14-14-1-1-1-1-1-1-1-1-						
K	Mw-	İ	X	100	TE: (NDE	z n	MET	ti pla	TE (0.5				*
	Mw-S		_	91	1ALI	HULE		ANNUL H	SEA	2.0)	(1)			
	MW-6		X											
	MW-													
	Mw-		(600)	H	NLE	= 11	J	ANNU	LAR S	EAL	(0,51			
	MW-	1		14	OLE	= 11	7	ANNV	LAR	SEHC	(0-9		<u> </u>	
	2.4													
	. s ^{e .}												****	
	*	*. 1												
				4							✓.	9.	ŧ	
go.	÷.							·						
٥														
lgen.	NOTES:													
2		***************************************				·* · · · · · · · · · · · · · · · · · ·		7		ne .				-
							. 100							-
														- <i>B</i>
-										77-140-3-11			\$ 14. J	
													i de la companya de	Salt a

WELL GAUGING DATA

Project # 080579-WW | Date 05/29/08 Client BLYMER

Site 966 89th AVE, OAKLAND, CA

						Thickness				Survey	
1			Well	C1 /	Depth to	of	Immiscibles		D 41 . 11	Point:	
	Well ID	Time	Size	Sheen / Odor	1	Immiscible	l .	Depth to water	_	TOBIOF	NI -4
ŀ	well ID '	1 line	(in.)	Odor	Liquid (ft.)	Liquia (II.)	(ml)	(ft.)	bottom (ft.)	100	Notes
	MW-IR	B53	2					8.76	21.50	1	
	MW-2	0933	2					8.55	23.75	TA CONTINUE OF THE PARTY.	PLATE
	mw-3	0908	2					8.57	23.90	-	
	Mw-4	0942	2					8.39	21.77		PLATE
	mw-5	0916	2					9.06	19.70	C. Transpagning to the Control of th	
	MW-6	0914	2					8,50	19.75		
	MW-7	6858	2					8.60	2/068	Anna de Constantes de Constant	¥ .
	mw-8 mw-9							8-61	19.64		
	m-9	0902	2			er		8,51	22.04	V	
										*	The state of the s
				更高	**		·		1		

							V-10				

X

W.LL MONITORING DATA SHE

	0,221,000,000							
Project #: 080529-WWI	Client: BLYMER ENG	-						
Sampler: WW	Date: 05/29/08							
Well I.D.: MW-R	Well Diameter: (2) 3 4	6 8						
Total Well Depth (TD): 71.50	Depth to Water (DTW): 8	.76						
Depth to Free Product:	Thickness of Free Product (feet):							
Referenced to: Grade	D.O. Meter (if req'd): ASI HACH							
DTW with 80% Recharge [(Height of Water	: Column x 0.20) + DTW]:	1.31						
Purge Method: Bailer Disposable Bailer Positive Air Displacement Electric Submersible Other Control Contr	Waterra Sampling Method: Peristaltic ction Pump Other: Well Diameter Multiplier Well I	Extraction Port Dedicated Tubing Diameter Multiplier 0.65 1.47						
1 Case Volume Specified Volumes Calculated V	olume							
Temp Cond. (°F or C) pH (mS or (s)) 1236 17.9 7.18 322	Turbidity (NTUs) Gals. Removed >1000 7	Observations						
1240 17.3 7.12 833	>1000	1/						
Did well dewater? Yes (NO)	Gallons actually evacuated:	Fe ²⁺ ; 0.6						
Sampling Date: 05/29/68 Sampling Tim	ne: 1245 Depth to Wate	er: 9.0 0						
Sample I.D.: MW-1R	Laboratory: Kiff CalScience	Other Mc CAMPBELL						
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	O Oxygenates (5) Other:							
D.O. (if req'd): Pre-purge:	mg/L Post-purge: 0,72 mg/L							
	mV = Post purgo:	α 7 mV						

W_LL MONITORING DATA SHE I

Project #:	08052	9-W	WI	Client:	BLYA	TER EN	JG				
Sampler:	WW			Date:	05/	29/08					
Well I.D.:		2			iameter	:(2) 3	4	6 8			
Total Well			3.75	Depth	to Water	r (DTW):	8.5	55			
Depth to Fr				Thickness of Free Product (feet):							
Referenced		PVC	Grade	D.O. Meter (if req'd): HACH							
DTW with	80% Rech	arge [(H	leight of Water	Columi	1×0.20) + DTW]:		11.59			
7 4	Disposable B Positive Air I Electric Subn Gals.) X	Displaceme	Other		Well Diamete 1" 2" 3"		Other:	Bailer Disposable Bailer Extraction Port Dedicated Tubing iameter Multiplier 0.65 1.47 radius² * 0.163			
Time	Temp (°F or C)	рН 7.33	Cond. (mS or (S)	1	oidity FUs)	Gals. Remo	oved	Observations			
1110	17.9	7.20	215	L(I	7	4.2					
1112	17.8	7.18	813	5	22	6.3	>				
anst pur	(h) (n) a	4	La Contraction	D+7	0(Fe 2+; 0			
Post	1	dings	taken (w +	Callen	o o otvoll	I avio anata	l	/ ?			
Did well de		Yes (9/08	No. Sampling Tim		17	y evacuated Depth to V		::11.09			
Sampling D		7/00	Damping 1 iii					M (1,00)			
Sample I.D.	: MW-			Labora			cience	Other IC CAPITORS			
Analyzed for	or: CTPH-G	BTEX	MTBE TPH-D	Oxygen	ates (5)	Other:					
EB I.D. (if	applicable)):	@ Time	Duplic	ate I.D.	(if applicab	ole):				
Analyzed for	or: TPH-G	BTEX	MTBE TPH-D								
D.O. (if req	'd): Pi	re-purge:		mg/ _L	Р	ost-purge:		(.53 ^{mg} / _L			
O.R.P. (if re	ea'd): Pi	re-purge:		mV	P	ost-purge:		212 mV			

W.LL MONITORING DATA SHE T

Project #: 0805	29-W	WI	Client:	BLYA	1ER ENG				
Sampler: WW			Date:	05/	29/08				
Well I.D.: MW	-3		Well D			6 8			
Total Well Depth (T		5.90	Depth t	o Water	C.8 :(WTD):	7			
Depth to Free Produ			Thickness of Free Product (feet):						
Referenced to:	AVO	Grade	D.O. M	eter (if	req'd):	ASI HACH			
DTW with 80% Rec	harge [(H	eight of Water	Column	$\times 0.20$) + DTW]:	11.64			
Purge Method: Bailer Disposable Positive Ai Electric Su (Gals.) X	r Displaceme		_	Well <u>Diamete</u> 1" 2"	0.04 4" 0.16 6"	Bailer Disposable Bailer Extraction Port Dedicated Tubing Diameter Multiplier 0.65 1.47			
	ecified Volum	es Calculated Vo		3"	0.37 Other	radius ² * 0.163			
Temp Time (°F or C	p _H 7.38	Cond. (mS or µS)	Turb (NT	idity 'Us)	Gals. Removed	Observations			
1209 15-8	7.2	939	78		5	((
211 16.5	1/- ((811	>10		7.5	<i>(</i> (
70st purpe po	iramete 19	s @ 20'				Fe ^{2t} : 0.6			
Did well dewater?	Yes	(NO)	Gallons	actuall	y evacuated:	7.5			
Sampling Date: 05/	29/08	Sampling Time	e: 12	18	Depth to Water	r: 11,64			
Sample I.D.: MW	1-3		Labora	tory:	Kiff CalScience	Other Mc CAMPBEL			
Analyzed for: TPH-	G BTEX	MTBE TPH-D	Oxygena	ites (5)	Other:				
EB I.D. (if applicab	le):	@ Time	Duplica	ate I.D.	(if applicable):				
Analyzed for: ТРН-	G BTEX	MTBE TPH-D	Oxygena	ites (5)	Other:				
D.O. (if req'd):	Pre-purge:		mg/L	F	Post-purge:	D.77 mg/L			
O.R.P. (if req'd):	Pre-purge:		mV	F	Post-purge:	196 mV			

WELL MONII	URING DATA	SHEEL						
Project #: 080529-WWI	Client: BLYMER ENG							
Sampler: WW	Date: 05/29/08							
Well I.D.: MW-4	Well Diameter: (2) 3 4 6 8							
Total Well Depth (TD): 2(77	Depth to Water (DTW): 8.39							
Depth to Free Product:	Thickness of Free Product (feet):							
Referenced to: Grade	D.O. Meter (if	req'd):	ASI HACH					
DTW with 80% Recharge [(Height of Water	Column x 0.20) + DTW]:	1.07					
Purge Method: Bailer Disposable Bailer Positive Air Displacement Extrace Electric Submersible Other	Waterra Peristaltic ction Pump Well Diamete	Sampling Method: Other:	Extraction Port Dedicated Tubing					
$\frac{2 - \hat{I}_{\text{Case Volume}}}{1 \text{ Case Volume}} = \frac{6.3}{\text{Calculated Volumes}}$	Gals. 1" 2" 3"	0.04 4" 0.16 6" 0.37 Other	0.65 1.47 radius ² * 0.163					
Temp Cond. Time (°F or C) pH (mS or μS) 0940 16.7 6.78 719	Turbidity (NTUs)	Gals. Removed	Observations					
0950 16.6 6.82 674	959	4.2						
0951 16.2 6.81 6.80649	>1000	6.3						
post purge readings taken	@ 181		Fe2+: 0					
Did well dewater? Yes No	Gallons actuall	ly evacuated: (6.3					
Sampling Date: 05/29/68 Sampling Tim	e: 0956	Depth to Water	r: 9,70					
Sample I.D.: Mw-4	Laboratory:	Kiff CalScience	Other Mc CAMPBEL					
Analyzed for: TPH-G BTEX MTBE TPH-D	Oxygenates (5)	Other:						
EB I.D. (if applicable):	Duplicate I.D.	(if applicable):						
Analyzed for: TPH-G BTEX MTBE TPH-D	Oxygenates (5)	Other:						
D.O. (if req'd): Pre-purge:	mg/L P	ost-purge:	(.07 mg/L					
O.R.P. (if req'd): Pre-purge:	mV P	ost-purge:	728 mV					

	A A		ORGING DIE	ALE CEERLE						
Project #: 080	529-W	WI	Client: BLYMER ENG							
Sampler: WW			Date: 05	129/08						
Well I.D.: M	W-5		Well Diameter: (2) 3 4 6 8							
Total Well Dept		1.70	Depth to Water (DTW): 9.06							
Depth to Free Pr	roduct:	1000	Thickness of Free Product (feet):							
Referenced to:	ATO .	Grade	D.O. Meter	(if req'd):	(TSI) HACH					
DTW with 80%	Recharge [(H	leight of Water	Column x 0.	20) + DTW]: /	0.64					
Purge Method: Baile Dispo			Waterra Peristaltic tion Pump	Sampling Method Other ameter Multiplier Well	Extraction Port Dedicated Tubing : Diameter Multiplier					
(Gals.) 2	X 3 Specified Volum	$\frac{1}{10000000000000000000000000000000000$	Gals.	0.04 4" 0.16 6" 0.37 Othe	0.65 1.47 r radius ² * 0.163					
1	emp or C) pH	Cond. (mS or (µS)	Turbidity (NTUs)	Gals. Removed	Observations					
1020 16	57.16	716	>1000	3.4						
1022 16	,5 7.18	698	>100	5.1						
Post purey	paramete	rs @ 16			Fezt: 0					
Did well dewate	8	6 0	Gallons actu	ually evacuated:	5-1					
Sampling Date:	05/29/08	Sampling Time	e: 1027	Depth to Wate	er: 9-12					
Sample I.D.:	1W-5		Laboratory:	Kiff CalScienc	e Other Mc CAMPBEL					
Analyzed for: C	ТРН-G ВТЕХ	MTBE TPH-D	Oxygenates (5	5) Other:						
EB I.D. (if appli	cable):	@ Time	Duplicate I.	D. (if applicable):						
Analyzed for:	TPH-G BTEX	MTBE TPH-D	Oxygenates (5	5) Other:						
D.O. (if req'd):	Pre-purge:		mg/L	Post-purge:	0.80 mg/L					
O.R.P. (if req'd)	: Pre-purge:		mV -	Post-purge:	249 mV					

W_LL MONITORING DATA SHL_ [

Project #: O	8052	9-W	WI	Client:	BLYM	IER ENG				
Sampler: 1	IW			Date:	05/	29/08				
Well I.D.:	MW-	6		Well Diameter: (2) 3 4 6 8						
Total Well D	Depth (TD): (9.75	Depth to Water (DTW): 8.50						
Depth to Fre	e Product			Thickness of Free Product (feet):						
Referenced t	:o:	AVO	Grade	D.O. M	eter (if	req'd):	YSI HACH			
DTW with 8	0% Recha	arge [(H	eight of Water	Column	x 0.20)	+ DTW]:	10.75			
Purge Method:	Bailer Disposable Ba Positive Air I Electric Subm	ailer Displacemer		Waterra Peristaltic tion Pump	Well Diameter	Sampling Method: Other:	Bailer Disposable Bailer Extraction Port Dedicated Tubing Multiplier 0.65			
1 Case Volume	als.) X	3 fied Volum	$= \frac{5 - 4}{\text{Calculated Vo}}$	_Gals. lume	2" 3"	0.16 6" 0.37 Other	1.47 radius ² * 0.163			
Time	Temp	Нq	Cond. (mS or μS)	Turb (NT	-	Gals. Removed	Observations			
1041	16.4	7-31	772	>/0/	J O	1.8	odor			
1043	16.8	7.17	771	>10		3.6	ع په			
1045	16.3	7.22	764	>10	10	54	(1			
		,								
post pur	al pan	omete	18 taken 6	16			Fezt: 0			
Did well dev	vater?	Yes /	Ŋ	Gallons	actuall	y evacuated:	54			
Sampling Da	ate: 05/2	9/08	Sampling Time	e: 10	50	Depth to Water	r: /0 - 6 6			
Sample I.D.:	Mw.	-6		Labora	tory:	Kiff CalScience	Other Mc CAMPBEL			
Analyzed for	r: TPH-G	BTEX	MTRE TPH-D	Oxygena	ites (5)	Other:				
EB I.D. (if a	pplicable)):	@ Time	Duplica	ate I.D.	(if applicable):				
Analyzed for		BTEX	MTBE TPH-D	Oxygena		Other:	-			
D.O. (if req'	d): P	re-purge:		mg/ _L	P	ost-purge:	1.75 mg/L			
O.D.D. Gfro	a'd). D			mV	P	ost-purge:	MV			

W_LL MONIT	ORING DATA	SHELL					
Project #: 080529-WWI	Client: BLY	MER ENG					
Sampler: WW	Date: 05/29/08						
Well I.D.: MW-7	Well Diameter		6 8				
Total Well Depth (TD): 21.68	Depth to Water (DTW): 8.60						
Depth to Free Product:	Thickness of F	ree Product (fee	et):				
Referenced to: Grade	D.O. Meter (if	req'd):	ASI HACH				
DTW with 80% Recharge [(Height of Water	Column x 0.20)) + DTW]: /	1.22				
Purge Method: Bailer Disposable Bailer	Waterra Peristaltic ction Pump	Sampling Method: Other:	Bailer Disposable Bailer Extraction Port Dedicated Tubing				
$\frac{2 \cdot (Gals.) \times 3}{1 \cdot Case \ Volume} = \frac{6 \cdot 3}{Specified \ Volumes} = \frac{1}{Calculated \ Volumes}$	Gals. Well Diamete	n Multiplier Well II 0.04 4" 0.16 6" 0.37 Other	Diameter Multiplier 0.65 1.47 radius ² * 0.163				
Temp Cond. Time (°F or C) pH (mS or μS)	Turbidity (NTUs)	Gals. Removed	Observations				
1304 18.5 7.36 743	21000	21	oder				
1207 183 7.16 745	>1000	4.2	((
1310 17.9 7.17 742	2/000	6-3	`(
post pure readings @ 17	1		Fezt: O				
Did well dewater? Yes No	Gallons actuall	y evacuated:	6.3				
Sampling Date: 05/29/08 Sampling Tim	e: 1305	Depth to Water	r: 8-71				
Sample I.D.: MW-7	Laboratory:	Kiff CalScience	Other Mc CAMPBELL				
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:	garage and the contract of the				
D.O. (if req'd): Pre-purge:	mg/ _L F	ost-purge:	1-47 mg/L				

O.R.P. (if req'd):

Pre-purge:

-mV

Post-purge:

WLLL MONITORING DATA SHE. I

Project #: 080529-WWI	Client: BLYMER ENG						
Sampler: WW	Date: 05/29/08						
Well I.D.: MW-8	Well Diameter: (2) 3 4 6 8						
Total Well Depth (TD): 19.64	Depth to Water (DTW): 8.6						
Depth to Free Product:	Thickness of Free Product (feet):						
Referenced to: Grade	D.O. Meter (if req'd):						
DTW with 80% Recharge [(Height of Wat	er Column x 0.20) + DTW]: 0 82						
Purge Method: Bailer Disposable Bailer Positive Air Displacement Ext	Waterra Sampling Method: Bailer Peristaltic Disposable Bailer Extraction Pump Other:						
$\frac{1}{1} \frac{S}{\text{Case Volume}} \text{(Gals.) X } \frac{3}{\text{Specified Volumes}} = \frac{S L}{\text{Calculated}}$	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						
Temp Cond. Time (°F or C) pH (mS or us)	Turbidity (NTUs) Gals. Removed Observations						
1320 11.1 1.10 / 1.	7/000 3.6						
1325 17.5 7.22 188	>1000 5.4						
post pure readings (a) 16'	Fe ^{2t} : O						
Did well dewater? Yes No	Gallons actually evacuated: S. 4						
Sampling Date: 05/29/08 Sampling Ti	ime: 1330 Depth to Water: 8.73						
Sample I.D.: MW-2	Laboratory: Kiff CalScience Other Mc CAMPBEL						
Analyzed for: TPH-G BTEX MTBE TPH-L	Oxygenates (5) Other:						
EB I.D. (if applicable):	Duplicate I.D. (if applicable):						
Analyzed for: TPH-G BTEX MTBE TPH-I							
D.O. (if req'd): Pre-purge:	mg/L Post-purge: 1.23 mg/L						
OPP (ifred'd): Pre-purge:	mV Post-purge: -12 mV						

V LL MONITORING DATA SHL T

Project #:	9057	19-n	WI	Client: BLYMER ENG.						
Sampler:	WW			Date: 05-	29-08					
Well I.D.:	HW-9			Well Diameter: (2) 3 4 6 8						
Total Well	Depth (TD)): Z [20 Y	Depth to Water (DTW): 3.51						
Depth to Fr	ee Produc	t:		Thickness of F	ree Product (fe	et):				
Referenced	to:	ANC?	Grade	D.O. Meter (if req'd): YSI HACH						
DTW with	80% Rech	arge [(H	eight of Water	Column x 0.20)) + DTW]: (1.22				
	Bailer Disposable B Positive Air I Electric Subn	Displaceme		Waterra Peristaltic tion Pump	Sampling Method:	Disposable Bailer Extraction Port Dedicated Tubing				
2.2 (() 1 Case Volume		3 Ified Volum	$\frac{1}{1000} = \frac{6 - 6}{1000}$ Calculated Vo	Well Diamete 1" 2" 3"	r Multiplier Well I 0.04 4" 0.16 6" 0.37 Other	Diameter Multiplier 0.65 1.47 radius ² * 0.163				
Time	Temp	рН	Cond. (mS or as)	Turbidity (NTUs)	Gals. Removed	Observations				
1344	(1.)	1.30	612	552	2.1	odur				
1347	16.9	076	674	4//	4-4	(r				
1350	17.1	7.33	674	713	6.6	((
	1 0	eading		(11 11		Fezt: 0				
Did well de	j		No.	Gallons actually S		0.0				
Sampling D		29/08	Sampling Time	e: 1523	Depth to Wate					
Sample I.D.	: MW	$\frac{9}{2}$		Laboratory:	Kiff CalScience	Other Mc CAMPBELL				
Analyzed fo	or: TPH-G	BTEX	MTBE TPH-D	Oxygenates (5)	Other:					
EB I.D. (if a	applicable)):	@ Time	Duplicate I.D.	(if applicable):					
Analyzed fo	r: TPH-G	BTEX	MTBE TPH-D	Oxygenates (5)	Other:	1./.				
D.O. (if req'	d): Pr	re-purge:		mg/ _L P	ost-purge:	1.4L/ mg/L				
O.R.P. (if re	ea'd). Pr	e-mirge		mV P	ost-purge:	94 mV				

Appendix C
Analytical Laboratory Report

McCampbell Analytical, Inc. June 5, 2008

McCampbell Analytical, Inc.

1534 Willow Pass Road, Pittsburg, CA 94565-1701 Telephone: 877-252-9262 Fax: 925-252-9269

Blymyer Engineers, Inc.	Client Project ID: #080529-WW1; Former	Date Sampled: 05/29/08
1829 Clement Avenue	Fiesta Beverage, Oakland	Date Received: 05/30/08
Alameda, CA 94501-1395	Client Contact: Mark Detterman	Date Reported: 06/05/08
Thansea, 611 7 1301 1373	Client P.O.:	Date Completed: 06/05/08

WorkOrder: 0805790

June 05, 2008

Dear	N/	ar	·

Enclosed within are:

- 1) The results of the 9 analyzed samples from your project: #080529-WW1; Former Fiesta Beve
- 2) A QC report for the above samples,
- 3) A copy of the chain of custody, and
- 4) An invoice 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 or concerns, please feel free to give me a call. Thank you for choosing

McCampbell Analytical Laboratories for your analytical needs.

Best regards,

Angela Rydelius Laboratory Manager

McCampbell Analytical, Inc.

			10	880 POG	ERS AVENU	E		COND	LICT	ANIALN	/CIC T	ODE	TECT		LAB McCampbell phs #
BLA	NE	SAN J		LIFORN	IIA 95112-110)5	_	COND	001	ANAL	313 1	O DE	ECI		ALL ANALYSES MUST MEET SPECIFICATIONS AND DETECTION
TECH SER					(408) 573-777 (408) 573-055										LIMITS SET BY CALIFORNIA DHS AND PRINCE PA REGION
					10 M										LIA
CHAIN OF CUS	TODY	BTS#	080	520	7-hm1	S									OTHER
CLIENT	Blymyer		1.0			CONTAINERS		B							SPECIAL INSTRUCTIONS
SITE						Ā		(8021B)							Invoice and Report to: Blymyer Engineers, Inc.
	Former Fiesta Beverage							(8)							500 State 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1
966 89th Avenue						ALL	5M)	MTBE							Attn: Mark Detterman
	Oakland,	CA				SITE	(801	IM							EDF Format Required.
			MATRIX	CON	NTAINERS	COMPOSI		8							
	1 1		S= SOIL W=H ₂ 0		í	8	TPH-G	BTEX							
SAMPLE I.D.	DATE	TIME	S=X	TOTAL		0	TP	BT							ADD'L INFORMATION STATUS CONDITION LAB SAMPLE #
MW-1R	05/29/08	1245	W	3	HCL Voa		х	х							
MW-2		1117	W	3	HCL Voa		х	х							
MW-3		1218	w	3	HCL Voa		х	х							
MW-4		0956	w	3	HCL Voa		х	х							
MW-5	_	1027	w	3	HCL Voa		х	х						01	
MW-6	_	1050	w	3	HCL Voa		X	X					CE /t	CONE	DITION APPROPRIATE CONTAINERS
MW-7	_	1305	W	3	HCL Voa		X	x					MEAD	CDAC	NATED IN LAB PRESERVED IN LAB
MW-8	- +	1330	W		HCL Voa		X	x						ERVAT	VOAS 10 8 G ME INSTITUTE
MW-9	1	1355	W	3				x							
10100-9		132	VV	3	HCL Voa		Х	^							
SAMPLING COMPLETED	DATE OS 29 VA	TIME	SAMPL PERFO	ING RMED B	y WIL	14	M	WON	6	-					RESULTS NEEDED NO LATER THAN As contracted
RELEASED BY	1h	<u> </u>				DAT	E 29/		IME 16	12		REC	EIVED	ВУ	2 SAMPLE DATE US/29/08 1612
RELEASED BY	Jaglo	Cust	-l,:	.)		DAT	E 30/0		TIME	308			IVED	BY	b h to DATE TIME 130 X
RELEASED BY	1/	1		-)		DAT	Ę .	1	TIME			RECE	IVE	BY	DATE 5/30/88 TIME
C IOUIDDED VII	takla	nt				-	20/0			45	_	1000	(1	-a-1-8 0/30/-8
SHIPPED VIA					1	DAT	E SEN	11		SENT		600	LER#		

McCampbell Analytical, Inc.

CHAIN-OF-CUSTODY RECORD

Page 1 of 1

	g, CA 94565-1701					Work	Order	: 08057	790	(ClientC	Code: E	BEIA				
(, , , ,			WriteOr	n ☑ EDF		Excel		Fax	[✓ Email		Hard	dCopy	Thir	dParty	☐ J -1	flag
Report to: Mark Detterman Blymyer Engineers, Inc. 1829 Clement Avenue Alameda, CA 94501-1395 (510) 521-3773 FAX (510) 865-2594		Email: cc: PO: ProjectNo:	MDetterman@blymyer.com : #080529-WW1; Former Fiesta Beverage, Oakland				Bl <u>y</u> 18	counts /myer E 29 Clen ameda,	nginee nent Av	rs, Inc. renue	95	Requeste Date Rec Date Pri			eived: 05/30/20		
									Ren	uested	Tosts	(See le	gend h	elow)			
Lab ID	Client ID		Matrix	Collection Date	Hold	1	2	3	4	5	6	7	8	9	10	11	12
0805790-001	MW-1R		Water	5/29/2008 12:45		Α	Α		-			-				1	
0805790-002	MW-2		Water	5/29/2008 11:17		Α										1	
0805790-003	MW-3		Water	5/29/2008 12:18		Α										1	
0805790-004	MW-4		Water	5/29/2008 9:56		Α										1	
0805790-005	MW-5		Water	5/29/2008 10:27		Α										1	
0805790-006	MW-6		Water	5/29/2008 10:50		Α										1	
0805790-007	MW-7		Water	5/29/2008 13:05		Α										1	
0805790-008	MW-8		Water	5/29/2008 13:30		Α										1	
0805790-009	MW-9		Water	5/29/2008 13:55		Α										1	
Test Legend:								Γ-					Г	_ 1			
1 G-MB1		PREDF RI	EPORT	3				4					=	5			
6	7			8				9					Ŀ	10			
Community:	12												Prepa	ared by	: Ana V	/enegas	<u> </u>

Comments:

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

Sample Receipt Checklist

Client Name:	Blymyer Eng	gineers, Inc.				Date	and Time Receive	d: 5/30/2008	7:19:10 PM
Project Name:	#080529-WW	/1; Former Fie	esta Bevera	age,	Oaklaı	nd Che	cklist completed a	nd reviewed by:	Ana Venegas
WorkOrder N°:	0805790	Matrix <u>W</u>	<u>/ater</u>			Carr	ier: <u>Derik Carta</u>	n (MAI Courier)	
			Chain o	f Cu	stody (C	COC) Inforn	nation		
Chain of custod	y present?		,	Yes	V	No 🗆			
Chain of custody	y signed when re	elinquished and r	eceived?	⁄es	V	No 🗆			
Chain of custod	y agrees with sa	mple labels?	,	⁄es	✓	No 🗌			
Sample IDs note	d by Client on CC	OC?	,	Yes	V	No 🗆			
Date and Time o	of collection noted	by Client on CO	C? \	Yes	✓	No 🗆			
Sampler's name	noted on COC?		`	⁄es	V	No 🗆			
			San	nple	Receipt	t Informatio	on_		
Custody seals ir	ntact on shipping	container/cooler	?	⁄es		No 🗆		NA 🔽	
Shipping contain	ner/cooler in good	d condition?	,	⁄es	V	No 🗆			
Samples in prop	per containers/bo	ttles?	,	Yes	✓	No 🗆			
Sample containe	ers intact?		,	Yes	✓	No 🗆			
Sufficient sampl	e volume for indi	cated test?	,	⁄es	✓	No 🗌			
		<u>Sam</u>	ple Preserva	atior	n and Ho	old Time (H	T) Information		
All samples rece	eived within holdi	ng time?	,	⁄es	✓	No 🗌			
Container/Temp	Blank temperatur	re	C	Coole	er Temp:	8.6°C		NA \square	
Water - VOA via	als have zero hea	adspace / no bub	bles?	Yes	V	No 🗆	No VOA vials su	ubmitted	
Sample labels c	checked for corre	ct preservation?	,	Yes	✓	No 🗌			
TTLC Metal - pH	l acceptable upor	n receipt (pH<2)?	`	⁄es		No 🗆		NA 🗹	
* NOTE: If the "	'No" box is check	ked, see commer	nts below.						
	=====	=====	====	==	===				
Client contacted	:	D	ate contacted	i:			Contac	eted by:	
Comments:									

"When Ouality Counts"

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

Blymyer Engineers, Inc.	Client Project ID: #080529-WW1; Former Fiesta	Date Sampled: 05/29/08							
1829 Clement Avenue	Beverage, Oakland	Date Received: 05/30/08							
Alameda, CA 94501-1395	Client Contact: Mark Detterman	Date Extracted: 06/01/08-06/	/03/08						
114111444, 6117 1001 1070	Client P.O.:	Date Analyzed 06/01/08-06/	/03/08						
Gasoline Range (C6-C12) Volatile Hydrocarbons as Gasoline with BTEX and MTBE*									

Extracti	on method SW5030B		Anal	Work Order	: 0805	790				
Lab ID	Client ID	Matrix	TPH(g)	MTBE	Benzene	Toluene	Ethylbenzene	Xylenes	DF	% SS
001A	MW-1R	W	690,a	ND	44	1.9	35	7.8	1	116
002A	MW-2	W	ND	ND	ND	ND	ND	ND	1	98
003A	MW-3	W	ND	ND	1.0	ND	ND	ND	1	93
004A	MW-4	W	ND	ND	ND	ND	ND	ND	1	96
005A	MW-5	W	ND	ND	ND	ND	ND	ND	1	94
006A	MW-6	W	ND	ND	ND	ND	ND	ND	1	98
007A	MW-7	W	ND	ND	ND	ND	ND	ND	1	102
008A	MW-8	W	ND	ND	ND	ND	ND	ND	1	118
009A	MW-9	W	ND	ND	ND	ND	ND	ND	1	93
Rep	porting Limit for DF =1;	W	50	5.0	0.5	0.5	0.5	0.5	1	μg/L
	ND means not detected at or above the reporting limit		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 McCampbell Analytical is not responsible for their interpretation: a) unmodified or weakly modified gasoline is significant; b) heavier gasoline range compounds are significant(aged gasoline?); c) lighter gasoline range compounds (the most mobile fraction) are significant; d) gasoline range compounds having broad chromatographic peaks are significant; biologically altered gasoline?; e) TPH pattern that does not appear to be derived from gasoline (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.



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

QC SUMMARY REPORT FOR SW8021B/8015Cm

W.O. Sample Matrix: Water QC Matrix: Water WorkOrder: 0805790

EPA Method SW8021B/8015Cm	Extraction SW5030B				BatchID: 35964			Sp	Spiked Sample ID: 0805735-001A			
Analyte	Sample	Spiked	MS	MSD	MS-MSD	LCS	LCSD	LCS-LCSD	Acce	eptance	Criteria (%)	
7 thaty to	μg/L	μg/L	% Rec.	% Rec.	% RPD	% Rec.	% Rec.	% RPD	MS / MSD	RPD	LCS/LCSD	RPD
TPH(btex [£])	ND	60	81.5	97.1	17.5	77.6	85.8	10.0	70 - 130	20	70 - 130	20
MTBE	ND	10	77.6	94	19.2	74.2	80.8	8.47	70 - 130	20	70 - 130	20
Benzene	ND	10	76.6	89.2	15.1	77.9	82	5.18	70 - 130	20	70 - 130	20
Toluene	ND	10	75.7	91.2	18.6	76.8	80.5	4.80	70 - 130	20	70 - 130	20
Ethylbenzene	ND	10	76.2	88.3	14.8	78.6	82.6	5.01	70 - 130	20	70 - 130	20
Xylenes	ND	30	76.4	85.3	11.1	72.8	77.3	6.03	70 - 130	20	70 - 130	20
%SS:	102	10	101	102	1.55	101	101	0	70 - 130	20	70 - 130	20

All target compounds in the Method Blank of this extraction batch were ND less than the method RL with the following exceptions:

NONE

BATCH 35964 SUMMARY

Lab ID	Date Sampled	Date Extracted	Date Analyzed	Lab ID	Date Sampled	Date Extracted	Date Analyzed
0805790-001A	05/29/08 12:45 PM	06/03/08	06/03/08 3:35 AM	0805790-002A	05/29/08 11:17 AM	06/01/08	06/01/08 1:17 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 = <math>100 * (MS - MSD) / ((MS + MSD) / 2).

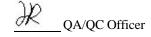
MS / MSD spike recoveries and / or %RPD may fall outside of laboratory acceptance criteria due to one or more of the following reasons: a) the sample is inhomogenous AND contains significant concentrations of analyte relative to the amount spiked, or b) the spiked sample's matrix interferes with the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

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

cluttered chromatogram; sample peak coelutes with surrogate peak.

N/A = not enough sample to perform matrix spike and matrix spike duplicate.

NR = matrix interference and/or analyte concentration in sample exceeds spike amount for soil matrix or exceeds 2x spike amount for water matrix or sample diluted due to high matrix or analyte content, or inconsistency in sample containers.



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

QC SUMMARY REPORT FOR SW8021B/8015Cm

W.O. Sample Matrix: Water QC Matrix: Water WorkOrder: 0805790

EPA Method SW8021B/8015Cm	Extraction SW5030B				BatchID: 36010 Sp				piked Sample ID: 0805790-008A			
Analyte	Sample	Spiked	MS	MSD	MS-MSD	LCS	LCSD	LCS-LCSD	Acce	eptance	Criteria (%)	
7 tildiyte	μg/L	μg/L	% Rec.	% Rec.	% RPD	% Rec.	% Rec.	% RPD	MS / MSD	RPD	LCS/LCSD	RPD
TPH(btex)	ND	60	92.9	91.4	1.69	82.8	81.1	2.09	70 - 130	20	70 - 130	20
MTBE	ND	10	103	103	0	89.3	90.2	1.00	70 - 130	20	70 - 130	20
Benzene	ND	10	87.2	88.8	1.78	80.5	74.8	7.35	70 - 130	20	70 - 130	20
Toluene	ND	10	96.3	98.6	2.34	89.3	86.5	3.24	70 - 130	20	70 - 130	20
Ethylbenzene	ND	10	94.1	96.8	2.90	86.8	82.4	5.18	70 - 130	20	70 - 130	20
Xylenes	ND	30	104	107	2.56	96.4	94.7	1.74	70 - 130	20	70 - 130	20
%SS:	118	10	92	93	0.630	94	93	0.392	70 - 130	20	70 - 130	20

All target compounds in the Method Blank of this extraction batch were ND less than the method RL with the following exceptions:

NONE

BATCH 36010 SUMMARY

Lab ID	Date Sampled	Date Extracted	Date Analyzed	Lab ID	Date Sampled	Date Extracted	Date Analyzed
0805790-003A	05/29/08 12:18 PM	06/01/08	06/01/08 1:49 AM	0805790-004A	05/29/08 9:56 AM	06/01/08	06/01/08 2:21 AM
0805790-005A	05/29/08 10:27 AM	06/01/08	06/01/08 2:53 AM	0805790-006A	05/29/08 10:50 AM	06/01/08	06/01/08 3:25 AM
0805790-007A	05/29/08 1:05 PM	06/01/08	06/01/08 3:56 AM	0805790-008A	05/29/08 1:30 PM	1 06/01/08	06/01/08 4:28 AM
0805790-009A	05/29/08 1:55 PM	06/01/08	06/01/08 5:00 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.

N/A = not enough sample to perform matrix spike and matrix spike duplicate.

NR = matrix interference and/or analyte concentration in sample exceeds spike amount for soil matrix or exceeds 2x spike amount for water matrix or sample diluted due to high matrix or analyte content, or inconsistency in sample containers.

