Fiesta Beverages 7150 Island Queen Dr. Sparks, NV 89436

RECEIVED

2:02 pm, Oct 17, 2008

Alameda County Environmental Health

Oct 1 , 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

Calley

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

Third Quarter 2008 Groundwater Monitoring Event

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

> September 18, 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.

CERTIFIED

Mark E. Detterman, CEG Senior Geologist

And: Mussile

Michael S. Lewis, REA Vice President, Technical Services

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1.0 Introduction and Background

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

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

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

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

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

On January 16, 2003, a new case manager, Mr. Amir Gholami, was appointed by the ACHCSA. On September 17, 2003, a workplan for a Geoprobe⁷ investigation of the site was submitted to the ACHCSA. The intent was to attempt to determine the lateral and vertical extent of impacted soil and groundwater in order to better target the residual contamination in future remedial actions to be determined. Due to the lack of a response from the ACHCSA, on February 17, 2004, Blymyer Engineers issued a *Letter of Intent to Proceed: Geoprobe⁷ Investigation*.

The Fourth Quarter 2003 Groundwater Monitoring Event report, dated January 6, 2004, recommended that analysis for fuel oxygenates by EPA Method 8260B be eliminated from the analytical program. It was reasoned that the data generated to date had been very consistent, and further quantification would not significantly add to the level of understanding at the site.

Additionally, the concentration of methyl *tert*-butyl ether (MTBE) can be monitored using EPA Method 8021B for no additional cost, and the resultant concentration of MTBE can be used as a proxy for the approximate concentration of the remaining fuel oxygenates. Based on the lack of response from the ACHCSA, it has been presumed that this was found reasonable and acceptable.

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

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

On July 6, 2005, the new case manager for the ACHCSA, Mr. Barney Chan, issued the letter *Fuel Leak Case RO0000314* commenting on the December 14, 2004 report. The ACHCSA determined that the collection of additional data is needed to progress the site towards closure. The letter requested a workplan to clear well MW-1 of several feet of sediment due to the potential for groundwater gradient biasing, requested further definition of the groundwater and soil plumes through the installation of additional wells and soil bores, requested a conduit study, and requested a Feasibility Study and Remedial Action Plan.

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

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

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

The RI/FS Report documented the destruction of well MW-1, the installation of replacement well MW-1R, and the installation of wells MW-4 through MW-9. The soil and groundwater data collected in the effort achieved vertical delineation, as well as upgradient, lateral, and downgradient delineation of all hydrocarbon compounds in soil and groundwater, with the exception of MTBE in groundwater. MTBE was delineated to below the MCL and the *non-drinking water* ESL goal for the compound, but was slightly above the *drinking water* goal. Because the site is in an area that is not known to extensively use groundwater as a drinking water source, the numeric remedial goals were predominantly compared to the *non-drinking water* ESL goals; however, the ACHCSA may ultimately apply *drinking water* ESL goals to remedial efforts at the site.

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

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

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

Six potential remedial options were evaluated for appropriateness at the site; monitored natural attenuation (MNA), groundwater pump and treat, enhanced insitu bioremediation (EIB), air sparging-vapor recovery (ASVR), dual phase extraction, and insitu chemical oxidation (ISCO). A combination of EIB and ISCO was selected as the most appropriate remedial technology for the site due to multiple factors. ISCO was selected for the vicinity of the former tank excavation and would consist of the injection of the commercial oxidation product RegenOx. Chemical oxidation of

residual source soil and groundwater containing higher hydrocarbon concentrations is anticipated to eliminate potential residual free-phase hydrocarbons in the tank vicinity. EIB using Oxygen Releasing Compound Advanced (ORC Advanced) was selected for the larger area around and downgradient of the former tank location. Petroleum hydrocarbon compounds are recognized to degrade favorably and rapidly under aerobic (oxygen rich) conditions. To stimulate aerobic bacterial activity and increase the rate of biodegradation within the hydrocarbon plume, non-toxic inorganic chemicals (bionutrients) can be added to the groundwater that release oxygen, nitrogen ORC and phosphate, such as 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 "bioneeds," 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 was notified that a new case worker, Ms. Barbara Jakub, had been assigned to the project by the ACHCSA.

2.0 Groundwater Sample Collection and Analytical Methods

This report documents the interim sampling of groundwater conducted for the Third 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 August 27, 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 8.72 to 9.60 feet below the top of the casings. Depth to groundwater in general has increased an average of 0.41 feet since the May 2008 sampling event; however, there were significant divergences from the average. In well MW-3 and MW-6 the depth to water increased by 1.03 and 1.02 feet, respectively, and in wells MW-7 and MW-8 it increased by only 0.12 and 0.15 feet, respectively. In the First Quarter 2008 groundwater event, mounding in the vicinity of wells MW-2 and MW-3 suggested the infiltration of rainwater in the vicinity well MW-2. The integrity of the pavement in the area of well MW-2 appears to have been compromised by the vigorous remedial chemical oxidation reactions. During the previous quarterly event, the dry spring appears to have returned the groundwater flow direction towards the west, generally the typical flow direction for the site. During the current event, groundwater flow remains towards the west; however, a slight mound is present around wells MW-1R and MW-2. Surface water infiltration may be responsible for the slight mound as ponded surface water can be present near these two wells during the dry season. Similar to the previous quarterly event, as well as the predominance of historic groundwater flow directions, groundwater flow is towards the west. The average gradient across the full area of investigation was calculated at approximately 0.007 feet/foot for the current event.

Between the May and August quarterly sampling events all concentrations of TPH as gasoline and BTEX decreased markedly in well MW-1R. The concentrations of TPH as gasoline and benzene in well MW-1R (190 and 14 ug/L, respectively) were below the generic non-drinking water ESLs but over generic drinking water ESLs. All other concentrations in well MW-1R were below all ESL values. In wells MW-2 and MW-3, except for benzene, which increased very slightly, all hydrocarbon concentrations remained nondetectable. The concentration of benzene in wells MW-2 and MW-3 (1.1 and 1.3 ug/L, respectively) are significantly below the generic non-drinking water ESL of 46 ug/L, and slightly above the drinking water ESL of 1.0 ug/L. All other wells (upgradient, downgradient, and lateral) were non-detectable for TPH as gasoline and BTEX.

During the First Quarter 2008, all contaminants in all wells were below their respective non-drinking water ESL; however, during the Second Quarter 2008 the concentration of TPH as gasoline rose

above the non-drinking water ESL in well MW-1R. Recent 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 is assumed to have mobilized contaminants near these locations.

Concentrations of MTBE were not detected any of the wells sampled during the current sampling event. MTBE has not been detected in all wells for four 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 probable drainage from soil to groundwater as groundwater drops in elevation at the site (MW-1 / MW-1R; Figures 5 and 6). Data from well MW-3 for the current quarter is generally consistent with the historical trend (this quarter, decreasing groundwater and decreasing concentrations) at the well; however, data from well MW-1 / MW-1R is not consistent with historical trends (this quarter, decreasing concentrations with decreasing groundwater levels). This might represent a change, a decrease in availability in the residual soil contamination presumed to be the source of groundwater impacts in well MW-1R. Data from the recent quarters has been somewhat atypical, and has been assumed to be related to the injection of RegenOx fluids. This has generally been presumed to indicate progress in remedial treatments. Specifically, 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-inchdiameter 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. This appears to have ceased during this quarter. (For consistency all groundwater elevations in Figures 3 to 6 utilized the GeoTracker wellhead survey elevations to determine the groundwater elevation.)

Groundwater samples have been analyzed for four quarters for 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 a listed MCL of 13 Fg/L.

4.0 Intrinsic Bioremediation Groundwater Sample Analytical Results

Intrinsic bioremediation laboratory analytical parameters were not collected during the current 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 (Mn²⁺) 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 very tight range of concentrations in post-purge groundwater, ranging from 0.11 milligrams per liter (mg/L) to 0.28 mg/L. The current concentrations are markedly lower than recent quarterly data. The data trend continues to indicate that recent higher concentrations of DO were related to RegenOx injection or 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.

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 wells MW-1R and MW-3 contain lower ORP concentrations than the remaining wells. ORP values in these two wells are 65 and 128 mV, respectively, whereas all other wells are at or above 200 mV. Except for a slight decline in wells MW-2 and MW-4, the ORP reading in all other wells have increased since the previous sampling event. This generally supports previous observations that more oxygenated rainwater infiltration in the vicinity of the tank basin was in the process of migrating across the area of study. It also suggests a higher demand for oxygen in the vicinity of well MW-1R and MW-3. Higher ORP values have generally been located outside the plume core, suggesting that the strongest demand for oxygen is located in the plume core. The continued decrease in 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 not detected in any well this quarter. During the previous monitoring event detectable ferrous iron returned to wells MW-1R and MW-3 for the first time since the remedial injections. While less clear this quarter, the detection of ferrous iron during the previous quarter suggests that the supply of DO from the injection of the RegenOx remedial product and rainwater infiltration has ceased. In general, the presence of ferrous iron in wells indicates that Mn – Fe degrading microbial colonies near the wells 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.

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 0.41 feet since the May 2008 sampling event; however, there were significant divergences from the average. Depth to groundwater in several wells increased by slightly over 1 foot, whereas in several wells it increased by only approximately 0.12 feet. A slight mound at wells MW-1R and MW-2 may be the result of the infiltration of ponded surface water. For the current event, groundwater flows towards the northwest at an average gradient calculated at approximately 0.007 feet/foot.
- Between the May and August quarterly sampling events, all concentrations of TPH as gasoline and BTEX decreased markedly in well MW-1R. The concentrations of TPH as gasoline and benzene in well MW-1R were below the generic non-drinking water ESLs. All other concentrations in well MW-1R were below all ESL values.
- In wells MW-2 and MW-3, except for benzene, which increased very slightly, all hydrocarbon concentrations remained nondetectable. The benzene concentrations were significantly below the generic non-drinking water ESL and slightly above the ESL for a drinking water source.
- All other wells (upgradient, downgradient, and lateral) were non-detectable for TPH as gasoline and BTEX.
- MTBE was not detected in any of the wells during the current sampling event. MTBE has not been detected in all wells for four consecutive quarters.
- The current concentrations of DO are markedly lower than recent quarterly data. The data trend
 continues to indicate that recent higher concentrations of DO were related to RegenOx injection
 or infiltration of rainwater. Lower concentrations of DO have typically been identified as a
 limiting factor at the site.
- Plume core wells MW-1R and MW-3 contain lower ORP concentrations than the remaining wells. Except for a slight decline in wells MW-2 and MW-4, the ORP reading in all other wells has increased since the previous sampling event. This generally supports previous observations that more oxygenated rainwater infiltration in the vicinity of the tank basin was in the process of

migrating across the area of study. It also suggests a higher demand for oxygen in the vicinity of wells MW-1R and MW-3. The continued decrease in 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 not detected any well this quarter. During the previous monitoring event detectable ferrous iron returned to wells MW-1R and MW-3 for the first time since the remedial injections. When present ferrous iron indicates that Mn Fe degrading microbial colonies near the wells 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.
- 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 well MW-3 for the current quarter is generally consistent with the historical trend (this quarter, decreasing groundwater and decreasing concentrations); however, data from well MW-1 / MW-1R is not consistent with historical trends (this quarter, decreasing concentrations with decreasing groundwater levels). This might represent a decrease in availability in the residual soil contamination presumed to be the source of groundwater impacts in well MW-1R.
- Data from MW-1R for recent 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 November 2008.
- Collection of limited RNA indicator data can be discontinued due to the lack of residual oxygen
 from the remedial injections, and the return of more anaerobic conditions. Collection of
 additional laboratory RNA indicator data can be resumed if a need is documented.

- If hydrocarbon concentrations continue to decrease in the next monitoring event, the site should be recommended for site closure.
- 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

W II ID	Б.	TOC Elevation	Depth to Water	Water Surface Elevation
Well ID	Date	(feet)	(feet)	(feet)
MW-1	8/6/1993	18.72	8.96	9.76
	1/12/1996		8.55	10.17
	4/16/1996		7.65	11.07
	7/15/1996		8.76	9.96
	10/16/1996		9.04	9.68
	12/15/1998		8.38	10.34
	1/18/2001		8.49	10.23
	4/25/2001		8.24	10.48
	3/17/03*		8.08	10.64
	6/23/2003		8.63	10.09
	9/18/2003		8.90	9.82
	12/15/2003		8.15	10.57
	6/15/2004		8.67	10.05
	12/15/2004		7.99	10.73
	6/29/2005		7.88	10.84
	5/8/2006	21.70	Destroyed	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
	8/27/2008		Destroyed	Destroyed
MW-1R	6/12/2006	21.73	8.49	13.24
	2/19/2007		7.94	13.79
	6/21/2007		8.71	13.02
	8/9/2007		8.83	12.90
	11/8/2007		9.80	11.93
	2/28/2008		8.74	12.99
	5/29/2008		8.76	12.97
	8/27/2008		9.02	12.71

Well ID	Date	TOC Elevation (feet)	Depth to Water (feet)	Water Surface Elevation (feet)
MW-2	8/6/1993	18.44	8.68	9.76
	1/12/1996		8.24	10.20
	4/16/1996		7.41	11.03
	7/15/1996		8.45	9.99
	10/16/1996		8.73	9.71
	12/15/1998		8.05	10.39
	1/18/2001		8.24	10.20
	4/25/2001		7.88	10.56
	3/17/03*		7.08	11.36
	6/23/2003		8.90	9.54
	9/18/2003		8.61	9.83
	12/15/2003		7.97	10.47
	6/15/2004		8.42	10.02
	12/15/2004		8.00	10.44
	6/29/2005		9.51	8.93
	6/12/2006	21.45	8.25	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
	8/27/2008		8.76	12.69

W-11 ID	Dete	TOC Elevation	Depth to Water	Water Surface Elevation
Well ID	Date	(feet)	(feet)	(feet)
MW-3	8/6/1993	19.01	9.07	9.94
	1/12/1996]	8.65	10.36
	4/16/1996]	7.82	11.19
	7/15/1996		8.88	10.13
	10/16/1996		9.16	9.85
	12/15/1998		8.45	10.56
	1/18/2001		8.57	10.44
	4/25/2001		8.29	10.72
	3/17/03*		8.50	10.51
	6/23/2003		9.05	9.96
	9/18/2003		9.11	9.90
	12/15/2003		8.03	10.98
	6/15/2004		8.85	10.16
	12/15/2004		8.84	10.17
	6/29/2005		9.00	10.01
	6/12/2006	22.02	8.62	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
	8/27/2008		9.60	12.42
MW-4	6/12/2006	21.34	8.37	12.97
	2/19/2007		7.77	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
	8/27/2008		8.76	12.58

W II ID	Б.,	TOC Elevation	Depth to Water	Water Surface Elevation
Well ID	Date	(feet)	(feet)	(feet)
MW-5	6/12/2006	22.53	8.75	13.78
	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
	2/28/2008		8.18	14.35
	5/29/2008		9.06	13.47
	8/27/2008		9.31	13.22
MW-6	6/12/2006	21.97	8.59	13.38
	2/19/2007		7.93	14.04
	6/21/2007		9.83	12.14
	11/8/2007		9.58	12.39
	2/28/2008		9.90	12.07
	5/29/2008		8.50	13.47
	8/27/2008		9.52	12.45
MW-7	6/12/2006	21.21	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
	8/27/2008		8.72	12.49
MW-8	6/12/2006	20.97	8.37	12.60
	2/19/2007		7.99	12.98
	6/21/2007		8.53	12.44
	11/8/2007		8.61	12.36
	2/28/2008		7.79	13.18
	5/29/2008		8.61	12.36
	8/27/2008		8.76	12.21

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		8.08	12.90					
	6/21/2007		8.55	12.43					
	11/8/2007	20.98	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

8/27/2008

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

8.81

12.17

Elevations in feet above mean sea level

Table II, Summary of Groundwater Sample Hydrocarbon Analytical Results BEI Job No. 203004, Former Fiesta Beverage 966 89th Avenue, Oakland, California Modified EPA EPA Method 8020 or 8021B Method 8015 $(\mu g/L)$ Well ID Sample Date $(\mu g/L)$ TPH as Gasoline Benzene Toluene Ethylbenzene Total Xylenes **MTBE MCL** N/A 1 150 700 1,750 13 Drinking Water Source 1 100 1 40 30 20 5 Non-Drinking Water 500 46 130 290 100 1,800 Source ² MW-1 8/6/1993 17,000 7.1 9.2 53 NA 8.4 1/12/1996 12,000 1,900 840 370 1,100 NA 4/16/1996 NA 3,500 **700** 55 100 180 7/15/1996 2,300 450 350 910 NA 11,000 21,000 4,200 2,200 650 10/16/1996 2,600 NA 12/15/1998 10,000 1,800 520 270 1,100 <350 1/18/2001 320 320 <120 11,000^a 2,000 1,100 4/25/2001 2.100 a, c 270 46 **59** 130 < 5.0 NA^{d} **36** 3/17/2003* 2,200^a 260 19 54 6/23/2003 6,100^a 930 53 99 200 NA 9/18/2003 13 24 **34** NA 3.800^a 660 260 a 19 1.1 < 0.5 1.5 12/15/2003 NA 6/15/2004 5,200°a **520** 13 **38** 39 < 50 12/15/2004 <15 2,400°a 370 8.2 13 14 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

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6/21/2007

11/8/2007

2/28/2008

5/29/2008

8/27/2008

Destroyed

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

		700 07 tii 1	ivenue, Oan	inaria, carr	OT III G		
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	Water 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
	8/27/2008	190 ^a	14	< 0.5	8.1	1.5	< 5.0

Table II, Summary of Groundwater Sample Hydrocarbon Analytical Results BEI Job No. 203004, Former Fiesta Beverage 966 89th Avenue, Oakland, California Modified EPA EPA Method 8020 or 8021B Method 8015 $(\mu g/L)$ Well ID Sample Date $(\mu g/L)$ TPH as Gasoline Benzene Toluene Ethylbenzene **Total Xylenes MTBE MCL** N/A 1 150 700 1,750 13 Drinking Water Source 1 100 1 40 30 20 5 Non-Drinking Water 500 46 130 290 100 1,800 Source ² MW-2 8/6/1993 2,700 1.3 1.7 2.0 NA 8.1 94 1/12/1996 2,700 600 310 220 NA 4/16/1996 190 **39** 10 NA 11 14 700 NA 7/15/1996 160 33 34 48 10/16/1996 190 48 8.2 10 13 NA 4.4 b 12/15/1998 200 **62 17** 4.9 14 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 $NA^{\,d}$ 78^a 26 1.5 3.5 3/17/2003* 3.3 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 12/15/2003 < 50 < 0.5 < 0.5 < 0.5 NA 95 ^a 6/15/2004 15 1.3 1.8 1.2 < 30 12/15/2004 < 50 0.97 0.9 **7.8** 11 0.6 6/29/2005 130 29 2.000 3.3 3.4 **6.7** 150 a **59** 3.0 3.4 2.7 11 6/13/2006 51 ^a 2/19/2007 8 1.0 2.8 7.1 1.6 < 50 < 0.5 < 5.0 6/21/2007 < 0.5 < 0.5 < 0.5 8/9/2007 < 50 < 0.5 < 0.5 < 0.5 < 0.5 < 5.0 5.3 11/8/2007 160 a 23 5.0 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

1.1

< 0.5

< 0.5

< 0.5

< 5.0

8/27/2008

< 50

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

Well ID	Sample Date	Modified EPA Method 8015 (µg/L)		EPA	Method 8020 (μg/L)	or 8021B	
		TPH as Gasoline	Benzene Toluene Ethylbenzene Total Xylenes March 1 150 700 1,750 1 40 30 20 46 130 290 100 2.1 2.9 3.6 17 280 180 120 470 370 340 160 580 200 220 66 250 340 140 100 300 200 39 72 150 240 41 86 120 300 330 200 1,100 240 78 10 280 2.5 0.6 0.69 1.4 <0.5	MTBE			
I	MCL	N/A	1	150	700	1,750	13
Drinking V	Vater Source 1	100	1	40	30	20	5
Non-Drinking Water Source 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°	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
	8/27/2008	<50	1.3	< 0.5	< 0.5	< 0.5	< 5.0

Table II, Summary of Groundwater Sample Hydrocarbon Analytical Results BEI Job No. 203004, Former Fiesta Beverage 966 89th Avenue, Oakland, California Modified EPA EPA Method 8020 or 8021B Method 8015 $(\mu g/L)$ Sample Date $(\mu g/L)$ TPH as Gasoline Benzene Toluene Ethylbenzene **Total Xylenes MTBE MCL** N/A 1 150 700 1,750 13 Drinking Water Source 1 100 1 40 30 20 5 Non-Drinking Water 500 46 130 290 100 1,800 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 < 0.5 11/8/2007 < 50 < 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 8/27/2008 < 50 < 0.5 < 0.5 < 0.5 < 0.5 < 5.0

Well ID

Table II, Summary of Groundwater Sample Hydrocarbon Analytical Results BEI Job No. 203004, Former Fiesta Beverage 966 89th Avenue, Oakland, California Modified EPA Method 8015 EPA Method 8020 or 8021B

Well ID	Sample Date	Modified EPA Method 8015 (µg/L)	hod 8015 EPA Method 8020 or 8021B (u.g/L.)						
		TPH as Gasoline	Benzene	Toluene	Ethylbenzene	Total Xylenes	MTBE		
1	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		
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		
	8/27/2008	<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		
	8/27/2008	<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		
	8/27/2008	<50	< 0.5	< 0.5	< 0.5	<0.5	< 5.0		

	Table II, Summary of Groundwater Sample Hydrocarbon Analytical Results BEI Job No. 203004, Former Fiesta Beverage 966 89th Avenue, Oakland, California									
Well ID	Sample Date	Modified EPA Method 8015 (µg/L)	EPA Method 8020 or 8021B (µg/L)							
		TPH as Gasoline	Benzene	Toluene	Ethylbenzene	Total Xylenes	MTBE			
1	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.

Bold results indicate detectable analyte concentrations.

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

^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.

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

Well ID	Cample Date				EPA Me	thod 8260E	(ug/L)			
Well ID	Sample Date	TAME	TBA	EBD	1,2-DCA	EPA Method 8260B (ug/L) C2-DCA DIPE Ethanol ETBE Methanol MTBE	MTBE			
Drinking	Water Source 1	NV	12	0.05	0.5	NV	50,000	NV	NV	5
Non-Drinking 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
	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
MW-3	6/23/2003	2.6	< 5.0	NA	NA	< 0.50	NA	< 0.50	NA	5.6
1V1 VV - 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

Well ID	Sample Date				EPA Me	thod 8260B	(ug/L)			
	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
Non-Drinking Water Source ²		NV	18,000	152	204	NV	50,000	NV	NV	1,800

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

Bold results indicate detectable analyte concentrations.

Note: Shaded cell indicates that detected concentration exceeds ESL

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

		Field Meter	Field Meter	Field Test Kit	Field Meter	Field Meter
Wall ID	Comple Date	Dissoved	Oxidation	Ferrous Iron	Field	Field pH
Well ID	Sample Date	Oxygen	Reduction Potential		Temperature	
		(mg/L)	(mV)	(Fe 2+)	(o F / o C)	pH units
MW-1	3/17/2003	NA	NA	NA	60.4 / 60.0 *	7.1 / 7.3
	6/23/2003	0.4	NA	NA	61.0 / 61.0 *	6.9 / 6.9
	9/18/2003	0.4	NA	NA	65.1 / 62.9 *	7.1 / 6.9
	12/15/2003	1.1	NA	NA	13.1 / 13.4	6.8 / 6.7
	6/15/2004	0.1	NA	NA	64.5 / 63.4 *	6.9 / 7.0
	12/15/2004	NA	NA	NA	15.4 / 17.5	7.0 / 6.9
	6/29/2005	0.24 / 0.17	1.0	4.5	19.78 / 21.63	7.15 / 7.08
	5/8/2006	Destroyed	Destroyed	Destroyed	Destroyed	Destroyed
	2/19/2007	Destroyed	Destroyed	Destroyed	Destroyed	Destroyed
	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
	8/27/2008	Destroyed	Destroyed	Destroyed	Destroyed	Destroyed
MW-1R	6/13/2006	0.87 / 0.37	172.9 / 172.9	0/0	17.31 / 17.36	6.90 / 6.92
	2/19/2007	0.48	8.0	NA	12.2 / 15.8	6.95 / 6.86
	6/21/2007	0.62	22.0	NA	19.6	7.1
	11/8/2007	0.3	-60	NA	64.4	6.9
	2/28/2008	0.28	156	0.0	63.2	6.98
	5/29/2008	0.72	97	0.6	17.3	7.12
	8/27/2008	0.18	65	0.0	66.2	6.8

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

900 87th Avenue, Oakianu, Cambi ma										
		Field Meter	Field Meter	Field Test Kit	Field Meter	Field Meter				
Well ID	Sample Date	Dissoved	Oxidation	Ferrous Iron	Field	Field pH				
		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				
	8/27/2008	0.14	202	0.0	72.1	6.56				

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

		Field Motor	Field Motor	Field Test Vit	Field Motor	Field Motor
		Field Meter	Field Meter	Field Test Kit		Field Meter
Well ID	Sample Date	Dissoved Oxygen	Oxidation Reduction	Ferrous Iron	Field Temperature	Field pH
	2 3333- F 33 2 333	Oxygen	Potential		Temperature	
		(mg/L)	(mV)	(Fe 2+)	(o F / o C)	pH units
MW-3	3/17/2003	NA	NA	NA	63.3 / 60.9 *	7.4 / 7.6
	6/23/2003	0.7	NA	NA	66.4 / 66.9 *	7.3 / 7.2
	9/18/2003	0.4	NA	NA	63.7 / 62.6 *	7.1 / 7.1
	12/15/2003	1.6	NA	NA	14.7 / 15.1	6.5 / 6.4
	6/15/2004	0.0	NA	NA	63.1 / 62.3 *	7.5 / 7.1
	12/15/2004	NA	NA	NA	15.4 / 16.7	7.2 / 7.0
	6/29/2005	0.72 / 0.78	141.7 / -67.6	0.9	17.65 / 18.79	6.94 / 7.02
	6/13/2006	1.01 / 0.41	170.0 / 168.5	0/0	17.30 / 17.15	7.02 / 6.98
	2/19/2007	0.08	81	NA	13.7 / 15.6	7.10 / 6.95
	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
	5/29/2008	0.77	186	0.6	16.3	7.19
	8/27/2008	0.15	128	0.0	65.7	7.08
MW-4	6/12/2006	0.67 / 0.33	164.3 / 161.0	0.5 / 0	16.90 / 16.79	6.82 / 6.79
	2/19/2007	0.21	98	NA	13.7 / 15.0	7.14 / 7.03
	6/21/2007	0.31	118	NA	16.4	7.0
	11/8/2007	0.30	222	NA	62.7	6.96
	2/28/2008	0.28	173	0.0	61.6	7.01
	5/29/2008	1.07	228	0.0	16.2	6.81
	8/27/2008	0.20	217	0.0	72.7	6.83

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
Well ID	Sample Date	Dissoved Oxygen	Oxidation Reduction Potential	Ferrous Iron	Field Temperature	Field pH
		(mg/L)	(mV)	(Fe 2+)	(o F / o C)	pH units
MW-5	6/12/2006	0.61 / 0.31	175.2 / 169.0	0/0	18.40 / 18.01	7.01 / 6.94
	2/19/2007	1.98	-114	NA	12.7 / 14.1	6.93 / 6.73
	6/21/2007	1.23	99	NA	16.8	7.1
	11/8/2007	0.30	211	NA	63.9	6.85
	2/28/2008	0.26	213	0.0	62.6	7.14
	5/29/2008	0.80	249	0.0	16.5	7.18
	8/27/2008	0.11	265	0.0	64.7	6.46
MW-6	6/13/2006	3.10 / 0.81	181.2 / 174.8	0 / 0	17.25 / 17.32	6.94 / 6.83
	2/19/2007	0.21	-30	NA	14.6 / 15.6	6.58 / 6.74
	6/21/2007	0.26	102	NA	16.2	7.1
	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
	8/27/2008	0.14	241	0.0	65.0	6.83
MW-7	6/12/2006	0.59 / 0.27	172.5 / 171.8	0.5 / 0.2	18.14 / 18.00	6.90 / 6.87
	2/19/2007	0.10	110	NA	16.2 / 17.2	7.69 / 7.21
	6/21/2007	0.14	123	NA	17.3	7.0
	11/8/2007	0.30	227	NA	64.5	6.90
	2/28/2008	0.27	142	0.0	64.2	7.00
	5/29/2008	1.47	83	0.0	17.8	7.17
	8/27/2008	0.21	196	0.0	76.1	6.83

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 Meter Field Test Kit Dissoved Oxidation Ferrous Iron Field Field pH Well ID Sample Date Reduction Temperature Oxygen Potential (mg/L)(Fe 2+) (o F / o C)pH units (mV) MW-8 0.37 / 0.33186.1 / 180.4 0 / 06.85 / 6.856/12/2006 18.55 / 18.39 NA 15.2 / 16.6 2/19/2007 0.11 102 7.23 / 7.07 6/21/2007 0.12 111 NA 17.2 7.1 11/8/2007 0.30 232 NA 64.3 7.01 7.08 2/28/2008 0.26 206 0.0 63.1 72 1.23 0.017.5 7.22 5/29/2008 0.26 190 74.8 8/27/2008 0.0 6.29 MW-9 2.01 / 1.87 206.0 / 191.0 0 / 06/12/2006 16.88 / 16.91 6.63 / 6.66 2/19/2007 0.08 101 NA 15.8 / 16.3 7.56 / 7.236/21/2007 0.12 112 NA 7.1 16.5 0.40 230 NA 65.1 6.94 11/8/2007 2/28/2008 0.26 208 0.0 62.1 7.01 1.44 94 0.0 17.1 7.33 5/29/2008

203

72.2

7.69

0.0

Notes: mV = Millivolts

mg/L = Milligrams per liter

° F / ° C = degrees Fahrenheit / degrees Centigrade

0.28

* = degrees Fahrenheit

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

NA = Not analyzed

8/27/2008

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)	Sulfate	Methane						
			μg/L								
MW-1	6/29/2005	490	<0.1	5	5,900						
	5/8/2006	Destroyed	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	6	51	< 0.5						
MW-8	6/12/2006	330	7.3	46	< 0.5						
MW-9	6/12/2006	240	8.3	44	1.1						

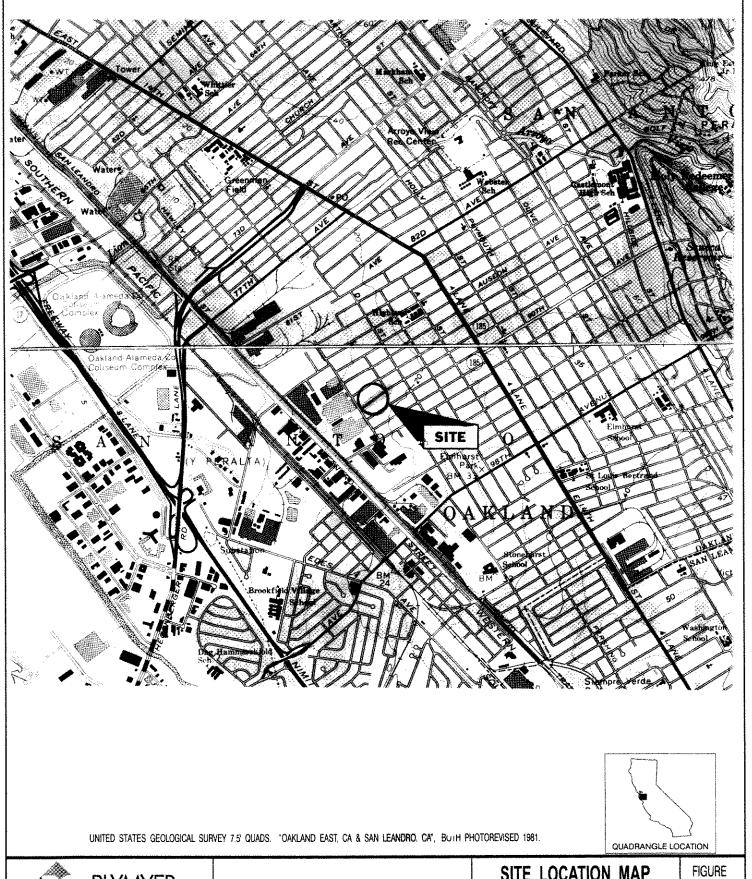
Notes: SM = Standard Method

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

Table V	Table VI, Summary of Groundwater Bacteria Enumeration Analytical Results BEI Job No. 203004, Former Fiesta Beverage 966 89th Avenue, Oakland, California										
		Aerobic Bacteria									
		Method 921	Method 9215A (HPC) / SM 9215 B Modified								
Well ID San	Sample Date	Hydrocarbon Degraders	Total Heterotrophs	Target Hydrocarbons Tested							
		cfu/ml									
MW-1R	4/27/2007	1,000	1,000	Gasoline/Diesel							
	8/9/2007	2,000	10,000	Gasoline/Diesel							
MW-2	4/27/2007	1,000	3,000	Gasoline/Diesel							
MW-5	8/9/2007	300	3,000	Gasoline/Diesel							
MW-6	4/27/2007	600	1,000	Gasoline/Diesel							
MW-9	4/27/2007	200	300	Gasoline/Diesel							

Notes: SM = Standard Method

cfu/ml = Colony forming units per milliliter



BEI JOB NO.

3-19-03

203004

SCALE IN FEET

1000

2000



SITE LOCATION MAP

FORMER FIESTA BEVERAGE 966 89TH AVE. OAKLAND, CA

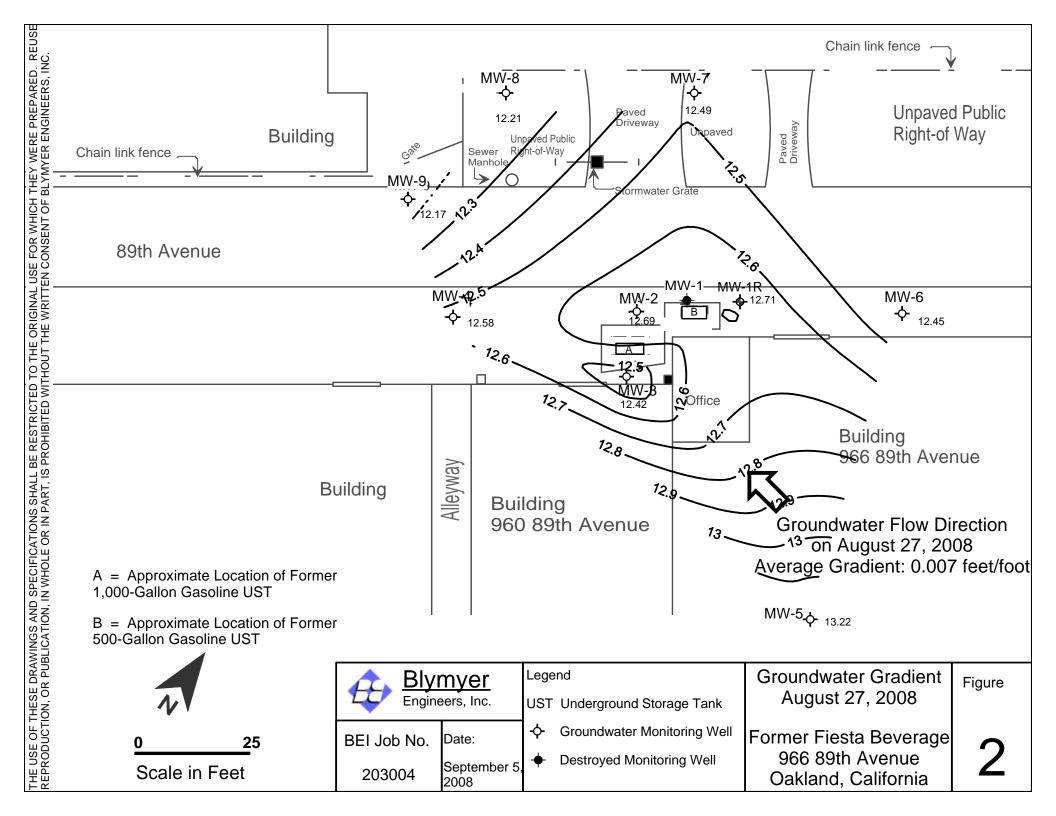


Figure 3: Concentration of TPHG vs. Time in Well MW-3

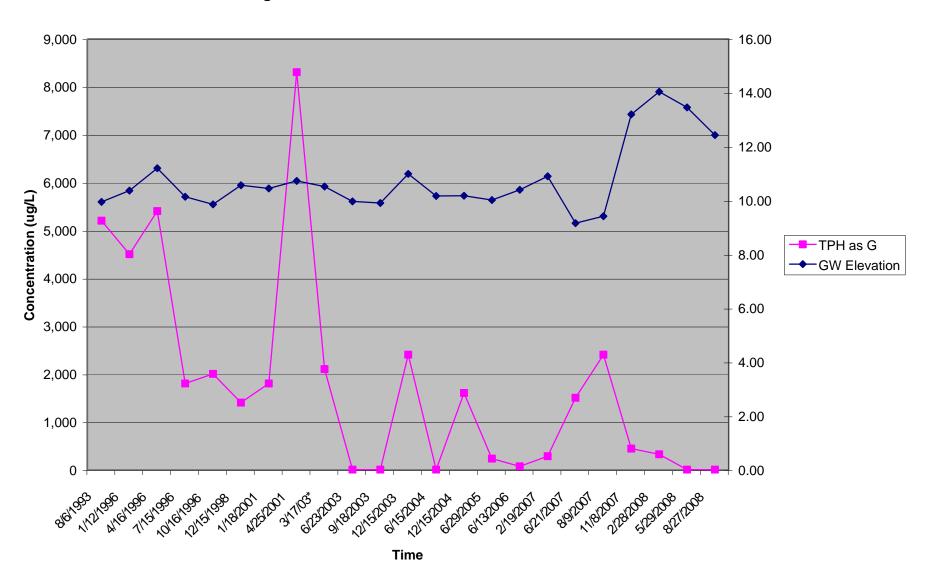


Figure 4: Concentration of Benzene vs. Time in Well MW-3

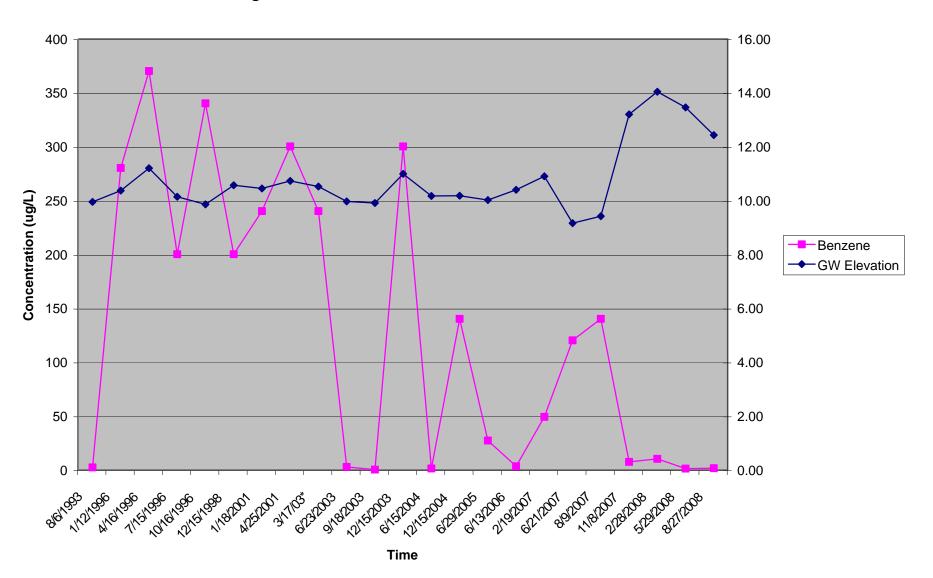


Figure 5: Concentration of TPHG vs. Tlme in Well MW-1 & MW-1R

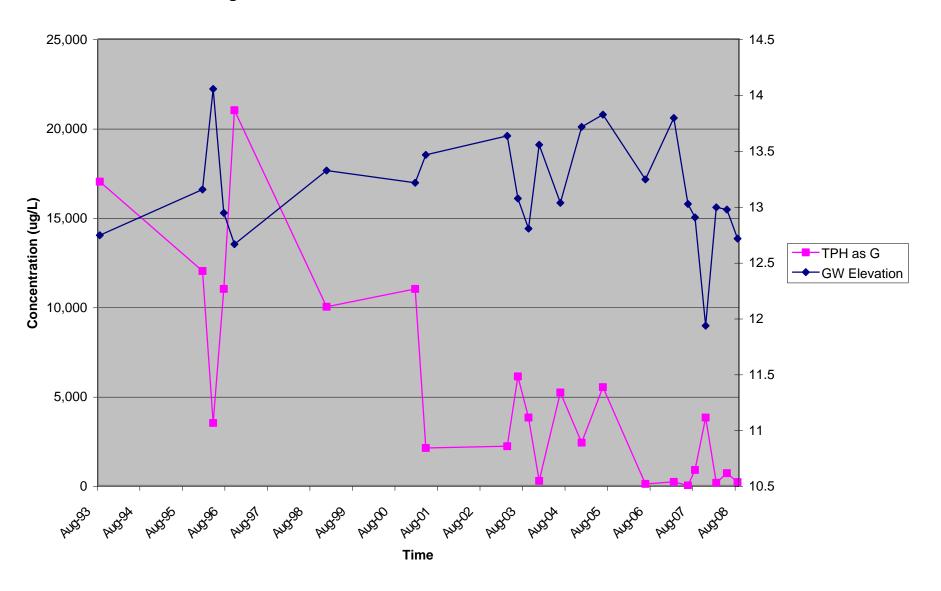
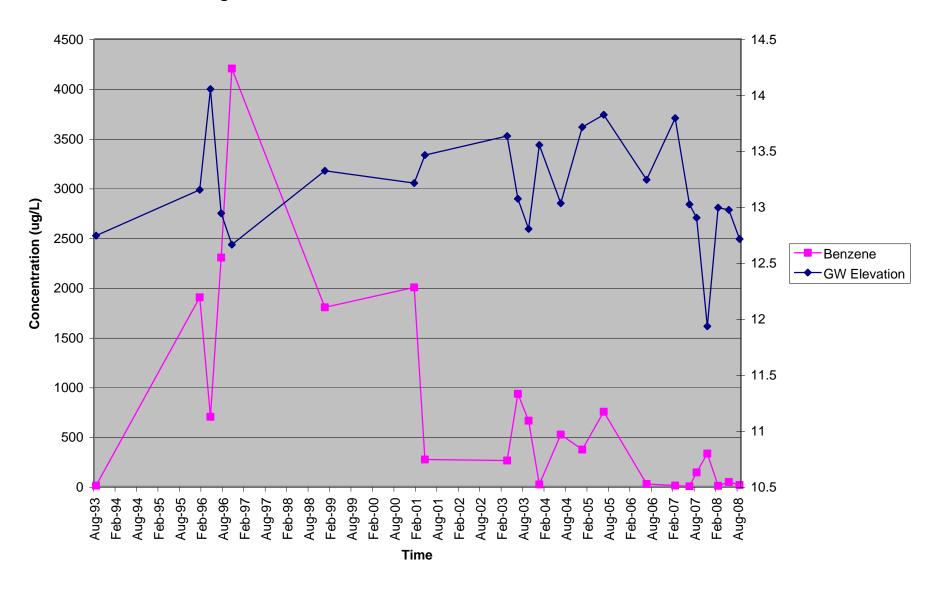


Figure 6: Concentration of Benzene vs. Tlme in Well MW-1 & MW-1R



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

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Appendix B

Well Monitoring Data Sheets and Well Gauging Data,

Blaine Tech Services, Inc. August 27, 2008

	S	rH or Pu	rge Water	Drum L	0g		
Client: Site Address:	Blyms 966 84 th	Av. C	alilund C	<u> </u>			
STATUS OF DR	UM(S): UPON	I ARRIVAI				Every and the second of the se	
Agent Colonia (Colonia) (Colonia) (Company) (Colonia) (C	Date	6/21/07	8/09/07	11/8/07	2/28/08	05/29/32	8/27/08
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:						

purge Mro

NO DATE

NA

0

05/29/08

WW

NW

8/27/08

IW

1

2/28/08

IW

11/8/07

65

NOW

6

PURGE H20

STAG ON

- If you add any SPH to an empty or partially filled drum, drum must have at least 20 gals. of Purgewater or DI Water.

06/02/06

- -If drum contains SPH, the drum MUST be steel AND labeled with the appropriate label.
- -All BTS drums MUST be labeled appropriately.

Are the drum(s) properly labeled?

If any drum(s) are partially or totally

filled, what is the first use date:

Number of drum(s) full: Total drum(s) on site:

Drum ID & Contents:

FINAL STATUS

Date of inspection:

Office reviewed by:

Drum(s) labelled properly:

Logged by BTS Field Tech:

this event

Number of new drum(s) left on site

STATUS OF DRUM(S) UPON	DEPARTI	JRE				
Date	6/21/07	08/09/07	11/8/07	2/28/08	05/29/50	8/27/08
Number of drums empty:						
Number of drum(s) 1/4 full:					0	
Number of drum(s) 1/2 full:					0	
Number of drum(s) 3/4 full:						
Number of drum(s) full:	5 4 Nen	5 18TS2	5 4non	5	b	le le
Total drum(s) on site:	6	6	6	6	4 WIN 1875	ヌ
Are the drum(s) properly labeled?	4	ΙY	7	Y	Y	S BTS 2NO
Drum ID & Contents:	Perge 1t20	Purgetho	parknata	Purge H20	punge the	PURGEHZE
LOCATION OF DRUM(S)						
Describe location of drum(s): ງູ່	chard bu	ilding th	lough Asi	Hup der	Don't L	er Out

NICH

WW

2

6/2160

Page ____ of ____

WELLHEAD INSPECTION CHECKLIST

Date <u>8/2</u>	7/08	Client	BLYMY	ier e	NG/NEE	es		
Site Address	966 89	the Ave	WE,	OAKLA	ND			
	080827-1u					14N 4	VILLIAM	5
Well ID	Well Inspected - No Corrective Action Required	Water Bailed From Wellbox	Wellbox Components Cleaned	Cap Replaced	Debris Removed From Wellbox	Lock Replaced	Other Action Taken (explain below)	Well Not Inspected (explain below)
MW-IR	X							
MW-2		D OF	2 801	TS PR	esent,	well B	oxforc	FDIRT
MW-3		loF	2 3	4" BE	LTS S	TRIPPE	D	PDIRT
MW-4	X	UN'	ser l	arge	META	LPL	ATE	
MW-5	X							
MW-4	X							
MW-7	X							
MW-8	X							
MW-9	X							
	A STATE OF THE STA							
NOTES:							V	
								,
			· · · · · · · · · · · · · · · · · · ·					

WELL GAUGING DATA

Project # 080827-IW-I Date 8/27/08 Client BLYMYER ENGINEERS

Site FORMER FIESTA BEVERAGE, 966 89th AVENUE, DAKLAND

Well ID	Time	Well Size (in.)	Sheen / Odor		Thickness of Immiscible Liquid (ft.)	Immiscibles Removed	1	Depth to well bottom (ft.)	Survey Point: TOB or TOC	Notes	
MW-IR	0902	2					9.02	21.53			
MW-2	0832	2					8.76	23.90			
MW-3	0905	2			-		9.60	24.90			
MW-4	1341	2	COVER	ed an	D PA	rked Togavse	8.76	21.80		PANEL	BI
MW-5	0918	2_					9.31	19.75		IN SHOP	
MW-6	0910	2					9.52	19.81			
MW-7	0850	2					8.72	21.75			
MW-8	0856	2					8.76	20.00			
MW-9	0847	2					18.8	22.05			
						-					
											1
											,

			LWUNII	ORING DAT	ASH. I'			
Project #:	080827.	-IW-1		Client: Blym	YER ENGINE	ERS		
Sampler:	IW			Date: 8/27/08				
Well I.D.:	MW - 1	.R		Well Diameter: 2 3 4 6 8				
Total Well	Depth (TI)): 21	1.53	Depth to Water (DTW): 9.02				
Depth to Fr	ee Produc	t:		Thickness of Free Product (feet):				
Referenced	to:	(ÉVC)) Grade	D.O. Meter (i		WSD HACH		
DTW with	80% Rech	arge [(F	Height of Water	Column x 0.20	0) + DTW]: ((.52		
2.0	Disposable B Positive Air I Electric Subn Gals.) X	Displaceme mersible	ent Extrac Other = 6.0	Waterra Peristaltie tion Pump Gals. Well Diame " 2" 3"	0.04 4" 0.16 6"	Disposable Bailer Extraction Port Dedicated Tubing Multiplier 0.65 1.47		
1 Case Volume	Speci	ified Volum	nes Calculated Vo	olume 3"	0.37 Other	radius ² * 0.163		
Time	Temp For °C)	pH	Cond. (mS or (S)	Turbidity (NTUs)	Gals. Removed	Observations		
1038	66.2			641	2.0	ODOR		
1032	66.1	6.84		7/000	4.0))		
1045	66.2	6.80	872	>1000	6.0	18		
POST PUR	pe par	AMETER	es takene	18' bg5		Fe2+ = 0.0		
Did well dev	water?	Yes (NO NO	Gallons actual	ly evacuated:	6.0		
Sampling Da	ate: 8/27	1/08	Sampling Time	: 1050	Depth to Water	r: 10.83		
Sample I.D.:	: MW-1	R		Laboratory:	Kiff CalScience	Other McCambell		
Analyzed for	r: TPH-G	BTEX	МТВ Р ТРН-D	Oxygenates (5)	Other:			
EB I.D. (if a	pplicable)	•	@ Time	Duplicate I.D.	(if applicable):	~		
Analyzed for	r: TPH-G	BTEX		Oxygenates (5)	Other:			
D.O. (if req'o	d): Pr	e-purge:		mg/L	ost-purgę:	o.18 mg/L		

mV

Post-purge:

65

mV

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

Pre-purge:

V LL MONITORING DATA SH. T

	·			ORINGDALA	1 211 1			
Project #:	080827.	-1W-1	90° 90° 4644 - 10 - 14	Client: BLYM)	TER ENGINE	ERS		
Sampler:	IW			Date: 8/27/	_			
Well I.D.:	MW-2			Well Diameter: 2 3 4 6 8				
Total Well	Depth (TI)): 2	23.90	Depth to Water	er (DTW): 8. 7	76		
Depth to Fr	ee Produc	t:		Thickness of F	Free Product (fe	et):		
Referenced	to:	(PVC)) Grade	D.O. Meter (if	req'd):	YSD HACH		
DTW with	80% Rech	arge [(H	leight of Water	Column x 0.20		1.79		
Purge Method: Bailer Disposable Bailer Positive Air Displacement Electric Submersible Other Waterra Positive Air Displacement Evtraction Pump Other Other: Well Diameter Multiplier Well Diameter Multiplier 1" 0.04 4" 0.65								
2.4 (Gals.) X	3	= 7.2	Gals.	0.04 4" 0.16 6"	1.47		
l Case Volume		ified Volum		- 11 20	0.37 Other	radius ² * 0.163		
Time	Temp For °C)	1	Cond. (mS or AS)	Turbidity (NTUs)	Gals. Removed	Observations		
1358	71.6	661	882	339	2.4	ODOR		
1403	72.3	6.58	871	370	4.8	q f		
1408	72.1	6.56	876	564	7.2	0.0		
POST PUR	be par	AMETER	es takeve	21'459		Fe2+ = 0.0		
Did well dev	water?	Yes 2	N 0	Gallons actually	y evacuated:	7.2		
Sampling Da	ate: 8/27	1/08	Sampling Time	e: 14le	Depth to Water	r: 9.90		
Sample I.D.:	: MW-	2		Laboratory:	Kiff CalScience	Other McCAMBELL		
Analyzed for	r: (TPH-G)	BTEXX	MTBB TPH-D	Oxygenates (5)	Other:			
EB I.D. (if a	pplicable)	1	Time	Duplicate I.D. ((if applicable):			
Analyzed for	r: TPH-G	BTEX 1		Oxygenates (5)	Other:			
D.O. (if req'o	d): Pr	e-purge:		mg/L PC	ost-purge:	0-14 mg/L		

mV

Post-purge:

202 mV

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

Pre-purge:

			TIVIOIATI	ORINGDALA	A OIL I			
Project #:	080827.	-1W-1		Client: BLYM	YER ENGINE	ERS		
Sampler:	IW			Date: %/27 /				
Well I.D.:	MW-3	3		Well Diameter	r: ② 3 4	6 8		
Total Well	Depth (TI)):	24.90	Depth to Wate	er (DTW): 9.	.60		
Depth to Fr	ee Produc	t:		Thickness of Free Product (feet):				
Referenced	to:	(PVC)	Grade	D.O. Meter (if	req'd):	WSD HACH		
DTW with	80% Rech	arge [(H	leight of Water	Column x 0.20)) + DTW]: 12	2.66		
Purge Method:	Bailer XDisposable B Positive Air I Electric Subn	Displaceme:		Well Diamet		Disposable Bailer Extraction Port Dedicated Tubing : Diameter Multiplier		
2.4 (C) 1 Case Volume		3 ified Volum	$\frac{1}{\text{nes}} = \frac{7.2}{\text{Calculated Vo}}$	- 11 3"	0.04 4" 0.16 6" 0.37 Other	0.65 1.47 radius ² * 0.163		
Time	Temp For °C)	1 1	Cond. (mS or aS)	Turbidity (NTUs)	Gals. Removed	Observations		
1120	65.4	T	961	531	2.4	STRONG ODOR		
1126	65.5	7.11	971	217	4.8	11		
1132	65.7	7.08	979	142	7.2	1,		
POST PUR	ue par	AMETER	es takeve	Z1' bs5		Fe2+ = 0.0 mg/		
Did well dev	water?	Yes	No	Gallons actuall	y evacuated:	J		
Sampling D	ate: 8/27	1/08	Sampling Time	: 1138	Depth to Wate	r:		
Sample I.D.	: MW-3	3		Laboratory:	Kiff CalScience	Other Mclambell		
Analyzed fo	r: TPH-G	BTEX	MTBB TPH-D	Oxygenates (5)	Other:			
EB I.D. (if a	pplicable)		@ Time	Duplicate I.D.	(if applicable):			
Analyzed fo	r: TPH-G	BTEX	MTBE TPH-D	Oxygenates (5)	Other:			
D.O. (if req'	d): Pr	e-purge:	And the control of th	mg/L (P	ost-purge:	0.15 mg/L		
O.R.P. (if re	<u>q'd):</u> Pr	e-purge:		mV P	ost-purge;	128 mV		

mV

Post-purge:

128

mV

	***************************************	,		URING DATA	A SH. I				
Project #:	080827.	-1W-1		Client: BLYM	IER ENGINE	ERS			
Sampler:	IW			Date: 8/27/08					
Well I.D.:	Mw	- 4		Well Diameter: (2) 3 4 6 8					
Total Well	Depth (TI)): 2	1.80	Depth to Wate	r (DTW): 8.	76			
Depth to Fr	ee Produc	t:		Thickness of Free Product (feet):					
Referenced	to:	(PVC)	Grade	D.O. Meter (if		WSD HACH			
DTW with	80% Rech	arge [(H	Ieight of Water	Column x 0.20) + DTW]:	1.37			
·	Disposable E Positive Air I Electric Subr	Displaceme nersible	ent Extrac Other	Waterra - Peristaltie J w ction Pump ∴	Sampling Method	✓ Disposable Bailer Extraction Port Dedicated Tubing			
2.1 (1) 1 Case Volume	Gals.) X	S fied Volum	= 6.3	Gals. Solume Well Diameter 1" 2" 3"	er Multiplier Well 0.04 4" 0.16 6" 0.37 Othe	Diameter Multiplier 0.65 1.47 r radius ² * 0.163			
Time	Temp For °C)	рН	Cond. (mS or 瓜	Turbidity (NTUs)	Gals. Removed	Observations			
1536	72.1	6.62	812	124	2.1				
1539	72.6	6.81	857	189	4.2				
1543	72.7	6.83	842	310	6.5				
Post Pur.			es takevo			Fe ²⁺ =0.0			
			No)	Gallons actuall		6.3			
Sampling D			Sampling Time	e: 1548	Depth to Wate	T: [0.13]			
Sample I.D.	: MW-	4		Laboratory:	Kiff CalScience	Other M. CLAMBELL			
Analyzed fo	r: (TPH-G	BTEX		Oxygenates (5)	Other:				
EB I.D. (if a	pplicable)	•	@ Time	Duplicate I.D.	(if applicable):				
Analyzed fo	r: TPH-G	BTEX	MTBE TPH-D	Oxygenates (5)	Other:	- Marine consistent which is the strong control of the strong cont			
D.O. (if req'	d): Pr	e-purge:		mg/L (P	ost-purge:	0.20 mg/L			
O.R.P. (if re	q'd): Pr	e-purge:		mV EP	ost-purge:	217 mV			

Post-purge:

217

mV

LL MONITORING DATA SH. T

Project #:	080827.	-1W-1		Client: BLYM	YER ENGINE	ERS
Sampler:	lW			Date: 8/27		
Well I.D.:	MW-5)		Well Diamete	er: (2) 3 4	6 8
Total Well	Depth (TI	D): [4	1.75	Depth to Wate	er (DTW):	1.31
Depth to Fi	ee Produc			Thickness of	Free Product (fe	
Referenced	to:	(PVC)) Grade	D.O. Meter (i		WSD HACH
DTW with	80% Rech	arge [(H	Height of Water	Column x 0.20	0) + DTW]: \(40
Purge Method:	Bailer Disposable E Positive Air Electric Subr	Bailer Displaceme	ent Extrac Other	Waterra Peristaltic ction Pump Well Diame 1" 2" Gals.	Sampling Method	Bailer Disposable Bailer Extraction Port Dedicated Tubing
Time	Temp For °C)	pН	Cond. (mS or 4 \$)	Turbidity (NTUs)	Gals. Removed	Observations
0926	65.7	6.60	739	20	1.6	CLEAR
0934	64.9	6.47	735	9	3.2	4
0942	64.7	6.46	736	6	4.8	1,
POST PUR	be par	AMETER	es takene	17' bgs		Fe2+ = 0.0 m/2
Did well dev	water?	Yes	M9	Gallons actual	ly evacuated:	4.8
Sampling D	ate: 8/27	/08	Sampling Time	: 0948	Depth to Wate	r: 9.32
Sample I.D.	MW-	5		Laboratory:	Kiff CalScience	Other Mclambell
Analyzed fo	r: TPH-G	BTEX	MTBB TPH-D	Oxygenates (5)	Other:	
EB I.D. (if a	pplicable)	•	(a) Time	Duplicate I.D.	(if applicable):	
Analyzed for	r: TPH-G	BTEX	MTBE TPH-D	Oxygenates (5)	Other:	
O.O. (if req'o	d): Pro	e-purge:		mg/L P	ost-purge:	0.11 mg/L
R.P. (if re	q'd): Pro	e-purge:		mV e	ost-purge:	265 mV

LL MONITORING DATA SH. 1

Project #:	<u> </u>	-IW-1	*****	Client: BLYMYER ENGINEERS								
Sampler:	IW			Date: 8/27/08								
Well I.D.:	MW-6		·	Well Diameter: 2 3 4 6 8								
Total Well	Depth (TI)):	19.81	Depth to Water (DTW): 9.52								
Depth to Fr	ee Produc	t:		Thickness of Free Product (feet):								
Referenced	to:	(PVC)	Grade	D.O. Meter (if	req'd):	WSD HACH						
DTW with	80% Rech	arge [(H	Height of Water	Column x 0.20) + DTW]: \	.58						
1	Bailer Disposable B Positive Air l Electric Subr	Displaceme		Waterra Peristaltice to the cition Pump Well Diamete 1" 2"	Other: Other: Well I	✓Disposable Bailer Extraction Port Dedicated Tubing						
1 Case Volume	Gals.) X Speci	ified Volun		_ Gals. -	0.37 Other							
Time	Temp For °C)	pH 7.01	Cond. (mS or us)	Turbidity (NTUs)	Gals. Removed	Observations						
1012	65.2		838	408	3.2							
1014	45.0	6.83	834	612	4.8							
POST PUR	ce par	AM ete i	es takene	17' 695		Fe2+ = 0.0 mg/						
Did well der	water?	Yes	No	Gallons actuall	y evacuated:	H.20 W 4.8						
Sampling D	ate: 8/27	1/08	Sampling Time	e: 1019	Depth to Wate	r: 11.20						
Sample I.D.	: MW-(<u></u>		Laboratory:	Kiff CalScience	Other McCambell						
Analyzed fo	r: (TPH-G	BTEX	MTBB TPH-D	Oxygenates (5)	Other:							
EB I.D. (if a	pplicable)):	@ Time	Duplicate I.D.	(if applicable):							
Analyzed fo	r: TPH-G	BTEX	MTBE TPH-D	Oxygenates (5)	Other:	·						
D.O. (if req'	d): Pr	e-purge:		mg/L CP	ost-purge:	0.14 mg/L						
O.R.P. (if re	q'd): Pr	e-purge:		mV P	ost-purge:	241 mV						

	And self-field and an analysis of the self-self-self-self-self-self-self-self-	*	TINIONNI TI	UNING DA.	LA SOL L	
Project #:	080827.	-IW-I		Client: BLYA	AYER ENGINE	ERS
Sampler:	IW			Date: %/2 :	7/08	
Well I.D.:	MW-7			Well Diamet	ter: (2) 3 4	6 8
Total Well	Depth (TI)): 2	1.75	Depth to Wa	ter (DTW): 8	.72
Depth to Fr	ee Produc	t:		Thickness of	Free Product (fe	
Referenced	to:	QÛ)	Grade	D.O. Meter ((if req'd):	WSD HACH
DTW with	80% Rech	arge [(H	eight of Water	Column x 0.2	20) + DTW]: \[\big	.33
Purge Method:	Bailer Disposable B Positive Air I Electric Subr	Displaceme		Waterra Peristalfic	Sampling Method	✓ Disposable Bailer Extraction Port Dedicated Tubing
2.0 1 Case Volume	Jais.) X	3 ified Volum	es Calculated Vo	Gals. Slume Well Diar	meter Multiplier Well 0.04 4" 0.16 6" 0.37 Othe	Diameter Multiplier 0.65 1.47 r radius² * 0.163
Time	Temp (F) or °C)	pН	Cond. (mS or µS)	Turbidity (NTUs)	Gals. Removed	Observations
1510	74.9	6.66	767	102	2.0	
1515	75.8	6.81	788	211	4.0	
1520	76.1	6.83	771	315	6.0	
POST PUR	ce par	Ameter	es takene	18' Wg		Fe ²⁺ = 0.0
Did well de	water?	Yes	<i>9</i>	Gallons actua	ally evacuated:	6.0
Sampling D	ate: 8/27	1/08	Sampling Time	: 1526	Depth to Wate	er: 11.01
Sample I.D.	: WM	7		Laboratory:	Kiff CalScience	e Other McCAMBELL
Analyzed fo	or: TPH-G	BTEX	MTBB TPH-D	Oxygenates (5)	Other:	
EB I.D. (if a	pplicable)):	② Time	Duplicate I.D	O. (if applicable):	
Analyzed fo	or: TPH-G	BTEX	MTBE TPH-D	Oxygenates (5)	Other:	
D.O. (if req'	d): Pr	e-purge:		mg/ _L C	Post-purge:	0.21 mg/L
O.R.P. (if re	eq'd): Pr	e-purge:		mV .	Post-purge:	196 mV

		,		URING DATA		
Project #:	080827 -	-1W-1		Client: BLYM	ER ENGINE	ERS
Sampler:	IW			1		
Well I.D.:	MW-8	}		Well Diameter	·: (2) 3 4	6 8
Total Well	Depth (TE): 2 <i>e</i>	0.00	Depth to Wate	r (DTW): §. -	76
Depth to Fr	ee Product	-•		Thickness of F	ree Product (fee	et):
Referenced	to:	(PVC)	Grade	D.O. Meter (if	req'd):	YSD HACH
DTW with	80% Rech	arge [(H	leight of Water	Column x 0.20) + DTW]: ([.	.01
Purge Method:	Positive Air I	Displaceme	nt Extrac	etion Pump	Other:	✓ Disposable Bailer Extraction Port Dedicated Tubing
l Case Volume	Date: \$/27/08 Well I.D.: MW/-8 Well Diameter: 2 3 4 6 8					
	€For °C)		(mS or µ(S))	(NTUs)	Gals. Removed	Observations
	72.8			71000	1.8	SILTY
1433	74.5	6.32	820	>1000	3-6	17
1436	74.8	6.29	827	2/000	5.4	10
Date: \$ 27 68						
POST PUR	oe par	AMETE	es taken e	17' bs		Fe ²⁺ = 0.0
Did well de	Client: BLYMYON ENGINEERS ampler: IM Date: \$/27/68 Well LD.: MW/-8 Well Depth (TD): 20.00 Depth to Water (DTW): \$.76 Depth to Free Product: Thickness of Free Product (feet): eferenced to: **PCO** Grade** D.O. Meter (if req'd): **Positive Air Displacement Electric Submersible** **Positive Air Displacement Extraction Pump Other: **Positive Air Displacement Multidement Well Dementer Well Dementer					
Sampling D	Client: SLYMYOR ENGINEERS Impler: IM Date: \$1/27/68 Well Diameter: 2 3 4 6 8 Depth to Free Product: Thickness of Free Product (feet): Seferenced to: Septimon of Free Product (feet): Septimon					
Sample I.D.	: MW-	8		Laboratory:	Kiff CalScience	Other McCAMBELL
Analyzed fo	or: (TPH-G	BTEX	МТВВ ТРН-D	Oxygenates (5)	Other:	
EB I.D. (if a	applicable)	:		Duplicate I.D.	(if applicable):	
Analyzed fo	or: TPH-G	BTEX	MTBE TPH-D		Other:	
D.O. (if req'	(d): Pr	e-purge:	anny ang taon ang taon di basa ang alak kanalak kanalak kanalak an dianan anang perananan ang kanal Agang	mg/L (F	ost-purge:	0.26 mg/L

mV

Post-purge:

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

Pre-purge:

190

mV

LL MONITORING DATA SH. T

		*		OMING	DAIA	OII I	
Project #: ()80827 -	-1W-1		Client:	BLY MY	er engine	ERS
Sampler:	IW			Date:	8/27/	8	
Well I.D.:	MW-	7		Well Dia	meter	: ② 3 4	6 8
Total Well	Depth (TD)): 2	2.05	Depth to	Water	(DTW): 8.	81
Depth to Fr	ee Product	:		Thicknes	ss of F	ree Product (fe	et):
Referenced	to:	(PVC)	Grade	D.O. Me	ter (if	req'd):	WSD HACH
DTW with	80% Rech	arge [(H	leight of Water	Column	x 0.20)	+ DTW]:	.46
Purge Method:	Disposable B Positive Air I	Displaceme			ell Diamete	r Multiplier Well	✓ Disposable Bailer Extraction Port Dedicated Tubing : Diameter Multiplier
2. Case Volume		3 fied Volun	$\frac{1}{1000} = \frac{6.3}{\text{Calculated Vo}}$	_ Gals.	1" 2" 3"	0.16 6"	1.47
Time	Temp	рН	Cond. (mS or ω S)	(NTU	Js)	Gals. Removed	Observations
				317			
1335	72.2	7.69	716	515	,	6.3	
POST PUR	Positive Air Displacement Extraction Pump Other Extraction Pump Dedicated Tubing Other						
Did well der	water?	Yes	No	Gallons a	actuall	y evacuated:	6.3
Sampling D	ate: 8/27	1/08	Sampling Time	e: 134		Depth to Wate	er: 9.36
Sample I.D.	: mw-9	1		Laborato	ry:	Kiff CalScience	e Other Mclambell
Analyzed fo	r: TPH-G	BTEX	MTBB TPH-D	Oxygenate	es (5)	Other:	
EB I.D. (if a	pplicable)	:	@ Time	Duplicate	e I.D. ((if applicable):	
Analyzed fo	or: TPH-G	BTEX	MTBE TPH-D	Oxygenate	es (5)	Other:	
D.O. (if req'	d): Pr	e-purge:		mg/L	P	ost-purge:	0.28 mg/L

mV

Post-purge:

203

mV

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

Pre-purge:

Appendix C
Analytical Laboratory Report

McCampbell Analytical, Inc. September 4, 2008

McCampbell Analytical, Inc.

"When Quality 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: #080827-1W-1; Former	Date Sampled: 08/27/08
1829 Clement Avenue	Fiesta Beverage, 966 89th Ave	Date Received: 08/28/08
Alameda, CA 94501-1395	Client Contact: Mark Detterman	Date Reported: 09/04/08
7 Harroad, C11 7 7501 1575	Client P.O.:	Date Completed: 09/04/08

WorkOrder: 0808811

September 04, 2008

_			
Dear	N/	വഴ	7
17541	10	ш	N.

Enclosed within are:

- 1) The results of the 9 analyzed samples from your project: #080827-1W-1; Former Fiesta Bever
- 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.

	31 AI	IN LET				ERS AVENU			CON	IDUCT ANAL	YSIS TO	DETECT		LAB McCampbell DHS#
	BLAI ECH SER	VICES, INC.	3300000		FAX	IIA 95112-110 (408) 573-777 (408) 573-055	1							ALL ANALYSES MUST MEET SPECIFICATIONS AND DETECTION LIMITS SET BY CALIFORNIA DHS AND EPA RWQCB REGION
C	HAIN OF CUS	STODY	BTS#	080	827	- IW - /	S							LIA OTHER
C	LIENT	Blymyer	Engine	ers, In	c.		CONTAINERS		1B)					SPECIAL INSTRUCTIONS
S	TE	Former F	iesta B	everag	e		CONT		(8021B)					Invoice and Report to: Blymyer Engineers, Inc.
		966 89th	Avenu	ie			ALL	5M)	MTBE					Attn: Mark Detterman
		Oakland,	CA				SITE	(801;	M					EDF Format Required.
SA	AMPLE I.D.	DATE	TIME	MATRIX N=H ² 0	TOTAL	NTAINERS	C = COMPOSITE	TPH-G (BTEX &					Global ID = T0600101573 ADD'L INFORMATION STATUS CONDITION LAB SAMPLE #
1	W-1R	8/27/68		W	3	HCL Voa	Ŭ	x	x					ABBERRICH STATES SCHEMEN BUSINESS
	W-2		1410	w	3	HCL Voa		х	х					
, -	W-3	-	1138	w	3	HCL Voa		х	х					
1	W-4		1548	w	3	HCL Voa		х	х			IC	E/tº	ONDITION APPROPRIATE
-	W-5		0948	w	3	HCL Voa		х	х			HE	EAD SE	PACE ABSENT CONTAINERS
	W-6		1019	w	3	HCL Voa		х	х					VATIONPRESERVED IN LAB
√M	W-7		1526	W	3	HCL Voa		х	х					
HM	W-8		1441	W	3	HCL Voa		х	х					
+M	W-9		1341	W	3	HCL Voa		Х	Х					
C	AMPLING OMPLETED	DATE 8/27/08	TIME 1548	SAMPLI	ING RMED B	BY JAN	WIL	LIA	MS					RESULTS NEEDED NO LATER THAN As contracted
	LEASED BY	////	2_	-			BAT DAT	27/	08	/655				Dun (SAMPLECUSTODIA) 8/27/08 1655
IK	ELEASED BY	23	6	ang (Ch	(I		286	58	1310		RECEIVED	E	at 8-28-08 13:00
	ELEASED BY	Denk	Cant-				DAT	_		TIME 1645		RECEIVED	BY	DATE TIME 8.28 08
SH	HIPPED VIA						DAT	E SEN	IT	TIME SENT	C	OOLER#		

McCampbell Analytical, Inc.

CHAIN-OF-CUSTODY RECORD

Page 1 of 1

- 12 K	g, CA 94565-1701 52-9262					Work	Order	: 0808	811	(ClientC	Code: B	EIA				
			WriteOn	✓ EDF		Excel		Fax		Z Email		Hard	Сору	Thir	dParty	J-	flag
Report to: Mark Dettern Blymyer Eng 1829 Cleme Alameda, C. (510) 521-377	gineers, Inc. nt Avenue A 94501-1395	cc: PO: ProjectNo:	MDetterman@ #080827-1W- Beverage, 966	1; Former Fiesta			Bl ₂	counts ymyer E 29 Clen ameda,	nginee nent Av	rs, Inc. enue	95		Date	e Rece	ived:		
									Requ	uested	Tests	(See leg	gend be	elow)			
Lab ID 0808811-001	Client ID		Matrix	Collection Date	H <u>ol</u> d	1	2	3	4	5	6	7	8	9	10	11	12
	MW-1R		Water	8/27/2008 10:50	Н	A	Α					+	1				-
0808811-002	MW-2		Water	8/27/2008 14:16	H	A						+					
0808811-003	MW-3		Water	8/27/2008 11:38	H	A											
0808811-004	MW-4		Water	8/27/2008 15:48		A											
0808811-005	MW-5		Water	8/27/2008 9:48	H	A						+					
0808811-006	MW-6		Water	8/27/2008 10:19	H	A						+					
0808811-007	MW-7		Water	8/27/2008 15:26	Н	A											
0808811-008	MW-8		Water	8/27/2008 14:41	Н	A											
0808811-009	MW-9		Water	8/27/2008 13:41	Ш	Α											<u> </u>
	ΓΕΧ_W 2	PREDF RE	PORT	3				4						5			
6	7			8				9					[10			
11	12												Prepa	ared by	: Ana V	⁷ enegas	<u>s</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	ineers, Inc.			Date	and Time Received:	8/28/2008	6:56:29 PM
Project Name:	#080827-1W-	1; Former Fies	sta Bevera	ge, 966 8	9th Chec	klist completed and	reviewed by:	Ana Venegas
WorkOrder N°:	0808811	Matrix Wa	<u>ter</u>		Carrie	er: <u>Derik Cartan (</u>	(MAI Courier)	
			Chain of	Custody (COC) Inform	ation		
Chain of custody	y present?		Υe	es 🗸	No 🗆			
Chain of custody	y signed when rel	inquished and red	ceived? Ye	es 🗸	No 🗆			
Chain of custody	y agrees with san	ple labels?	Υe	es 🗸	No 🗌			
Sample IDs noted	d by Client on CO	D?	Υe	es 🗸	No 🗆			
Date and Time o	of collection noted l	by Client on COC?	Ye	es 🗸	No 🗆			
Sampler's name	noted on COC?		Υe	es 🗸	No 🗆			
			Samp	ole Receip	ot Information	<u>n</u>		
Custody seals in	ntact on shipping	container/cooler?	Ye		No 🗆		NA 🗹	
Shipping contain	ner/cooler in good	condition?	Υe	es 🗸	No 🗆			
Samples in prop	er containers/bot	les?	Υe	es 🗸	No 🗆			
Sample containe	ers intact?		Ye	es 🗸	No 🗆			
Sufficient sample	e volume for indic	ated test?	Υe	es 🗸	No 🗌			
		Sampl	e Preservat	ion and H	lold Time (HT	<u>) Information</u>		
All samples rece	eived within holdin	g time?	Υe	es 🗸	No 🗌			
Container/Temp	Blank temperature)	Co	oler Temp:	1.8°C		NA 🗆	
Water - VOA via	als have zero hea	dspace / no bubb	les? Ye	es 🔽	No 🗆	No VOA vials subr	nitted \square	
Sample labels cl	hecked for correc	t preservation?	Υe	es 🗸	No 🗌			
TTLC Metal - pH	l acceptable upon	receipt (pH<2)?	Υe	es \square	No 🗆		NA 🔽	
Samples Receive	red on Ice?		Υe		No 🗆			
			(Ice Type: \	WET ICE)			
* NOTE: If the "I	No" box is checke	ed, see comments	s below.					
	=====	=====	====		====	=====	=====	======
Client contacted:	:	Dat	e contacted:			Contacted	d by:	
Comments:								



"When Ouality Counts'

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

Blymyer Engineers, Inc.	Client Project ID: #080827-1W-1; Former Fiesta Beverage, 966 89th Ave	Date Sampled: 08/27/08
1829 Clement Avenue	Former Flesta Deverage, 900 89th Ave	Date Received: 08/28/08
	Client Contact: Mark Detterman	Date Extracted: 08/30/08-09/02/08
Alameda, CA 94501-1395	Client P.O.:	Date Analyzed 08/30/08-09/02/08

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

Analytical methods SW8021B/8015Cm Extraction method SW5030B Lab ID Client ID Matrix TPH(g) MTBE Benzene Toluene Ethylbenzene Xylenes DF % SS 001A MW-1R W 190,d1 ND 14 ND 8.1 117 002A W ND MW-2ND ND 1.1 ND ND 1 99 003A W MW-3 ND ND 1.3 ND ND ND 1 98 004A MW-4 W ND ND ND ND ND ND 1 103 005A MW-5 W ND ND ND ND ND ND 1 96 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,b1 ND ND ND ND ND 1 124 009A MW-9 W ND ND ND ND ND ND 95 Reporting Limit for DF = 1; W 50 5.0 0.5 0.5 0.5 0.5 $\mu g/L$ ND means not detected at or 1.0 0.05 0.005 0.005 0.005 0.005 mg/Kg above the reporting limit

- b1) aqueous sample that contains greater than ~1 vol. % sediment
- d1) weakly modified or unmodified gasoline is significant



^{*} 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:

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QC SUMMARY REPORT FOR SW8021B/8015Cm

W.O. Sample Matrix: Water QC Matrix: Water BatchID: 37876 WorkOrder 0808811

EPA Method SW8021B/8015Cm Extraction SW5030B Spiked Sample ID: 0808801-002												002	
Analyte	Sample	Spiked	MS	MSD	MS-MSD LCS LCSD LCS-LCSD					Acceptance Criteria (%)			
Analyto	μg/L	μg/L	% Rec.	% Rec.	% RPD	% Rec.	% Rec.	% RPD	MS / MSD	RPD	LCS/LCSD	RPD	
TPH(btexf)	ND	60	111	107	3.90	102	98.3	3.36	70 - 130	20	70 - 130	20	
MTBE	ND	10	99.7	99.3	0.399	99.4	104	4.84	70 - 130	20	70 - 130	20	
Benzene	ND	10	90.7	89.7	1.16	91.8	92.9	1.22	70 - 130	20	70 - 130	20	
Toluene	ND	10	91.1	87.3	4.30	91.7	92.8	1.17	70 - 130	20	70 - 130	20	
Ethylbenzene	ND	10	85.4	87.2	2.04	95.7	96	0.320	70 - 130	20	70 - 130	20	
Xylenes	ND	30	79.4	81	1.93	107	108	0.853	70 - 130	20	70 - 130	20	
%SS:	97	10	107	102	4.02	95	96	1.48	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 37876 SUMMARY

Lab ID	Date Sampled	Date Extracted	Date Analyzed	Lab ID	Date Sampled	Date Extracted	Date Analyzed
0808811-001A	08/27/08 10:50 AM	08/30/08	08/30/08 2:25 AM	0808811-002A	08/27/08 2:16 PM	09/02/08	09/02/08 11:02 PM
0808811-003A	08/27/08 11:38 AM	09/02/08	09/02/08 11:33 PM	0808811-004A	08/27/08 3:48 PM	08/30/08	08/30/08 8:29 AM
0808811-005A	08/27/08 9:48 AM	08/30/08	08/30/08 6:32 AM	0808811-006A	08/27/08 10:19 AM	08/30/08	08/30/08 7:06 AM
0808811-007A	08/27/08 3:26 PM	08/30/08	08/30/08 7:39 AM	0808811-008A	08/27/08 2:41 PM	08/30/08	08/30/08 8:13 AM
0808811-009A	08/27/08 1:41 PM	08/30/08	08/30/08 8:46 AM				

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

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

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

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

cluttered chromatogram; sample peak coelutes with surrogate peak.

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.

