



Bioremediation Work Plan In Situ Enhanced Natural Attenuation of Petroleum Ken's Auto Wash Ellensburg, Washington

Prepared for Ken's Auto

October 14, 2013 7168-09





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FINAL 2011 BIOREMEDIATION WORK PLAN IN SITU ENHANCED NATURAL ATTENUATION OF PETROLEUM KEN'S AUTO WASH ELLENSBURG, WASHINGTON

1.0 INTRODUCTION

This Work Plan describes the *in situ* bioremediation injection program and the sampling and analysis procedures to conduct groundwater sampling and analysis at the Ken's Auto Wash site, located at 1013 East University Way in Ellensburg, Washington (Figure 1). A draft final version of this Work Plan was originally issued on January 28, 2011. At the request of the Washington State Department of Ecology, we revised and finalized the plan to include a schedule for confirmational groundwater monitoring. The goal of this work is to accelerate biological degradation of residual gasoline-range hydrocarbon (TPH-G) in the former source area and reduce the likelihood of future groundwater concentration exceedances above the Washington State Model Toxics Control Act (MTCA) Method A cleanup levels (Chapter 173-340 WAC).

Following this introductory section that summarizes previous remedial actions and groundwater analytical results, we provide a description of Remedial Program Implementation, Groundwater Sampling and Analysis, Chemical Data Quality Review and Data Quality Assurance, and Evaluation and Reporting.

1.1 Site History and Background

The site is affected by a petroleum hydrocarbon release discovered during UST tightness testing in 1996 (Figure 2). Corrective actions were taken at that time, and the site USTs were subsequently removed in April 2005, as documented in the June 7, 2005, Gasoline UST Closure Report. Petroleum-impacted soils were removed downgradient of the UST area in October and November 2000, but a small volume of affected soil remained because of utilities at the site, as shown on Figure 2.

During the soil removal, oxygen-releasing compound (ORC) was added to the excavation backfill to promote biodegradation of residual petroleum hydrocarbons. ORC was also injected in the downgradient area of petroleum hydrocarbon-affected groundwater in February 2005, as documented in the April 6, 2005, Supplemental Strataprobe Exploration Report. Although concentrations of TPH in groundwater continued to decrease following UST removal, soil removal, and ORC injection, low-level TPH-G concentrations remain in groundwater downgradient of the residual source area.

Remedial alternatives were presented and evaluated in a Remedial Investigation and Feasibility Study (RI/FS) dated November 14, 2006. The RI/FS addressed requirements of an Agreed Order issued by the Washington State Department of Ecology (Ecology) for site cleanup assessment following an MTCA site hazard ranking of 2. Remedial technologies evaluated in the RI/FS were based on results of site investigation, soil cleanup, and monitoring efforts through 2006.

Monitored natural attenuation with free product removal was selected as the preferred remedial alternative following Ecology's review of the RI/FS. Monitored natural attenuation is a process where hydrocarbon-degrading microbes that occur naturally in subsurface soils degrade petroleum hydrocarbons. Appreciable free product has not been identified at the site since 2004, so current remedial actions have not incorporated sorbent socks to remove free product. Site monitoring continues in accordance with the selected RI/FS alternative. Ecology has not required any additional actions.

1.2 Site Groundwater Conditions

Petroleum-impacted soils remain downgradient of the former USTs beneath the adjacent sidewalk and portions of East University Way (Figure 2). This remaining source material is likely contributing to periodic exceedances of MTCA Method A cleanup criteria for TPH-G and benzene in groundwater near wells MW-14 and MW-6 (Table 1). The fluctuations in TPH-G and benzene concentrations appear to be related to seasonal groundwater level fluctuation.

1.2.1 TPH-G Testing Results Summary

The three most recent groundwater sampling and testing events were conducted in June 2008, October 2008, and October 2009. Historical data is presented in Table 1 and a brief summary of areas of interest are presented below.

- The October 2008 and 2009 groundwater samples from well MW-14 contained TPH-G concentrations of 2.04 and 2.03 milligrams per liter (mg/L), respectively. Although TPH-G was non-detect at a detection limit of 0.05 mg/L during June 2008, this was atypical for groundwater sampling results over the last five years, where TPH-G concentrations as high as 8.10 mg/L were detected.
- The June 2008, October 2008, and October 2009 samples from well MW-6 contained TPH-G at concentrations of 1.55, 0.855, and 0.501 mg/L, respectively.

For comparison, the MTCA Method A cleanup level for TPH-G in groundwater is 0.800 mg/L when benzene is present, and 1.00 mg/L when benzene is not present. Therefore, TPH-G concentrations in samples from well MW-14 exceeded the applicable MTCA Method A criteria in October 2008 and October 2009, as well as historically. The TPH-G concentration in the June 2008 sample from well MW-6 exceeded the MTCA criteria, but relatively low to non-detect benzene concentrations in October 2008 and October 2009 precluded TPH-G exceedances during these sampling events. TPH-G exceedances in MW-6 groundwater samples were common during previous sampling events.

1.2.2 Benzene Testing Results Summary

- Benzene was detected at a concentration of 4.76 micrograms per liter (ug/L) in the groundwater sample from well MW-14 in October 2008. Benzene was not detected above the 12.2 ug/L detection limit in October 2009. This reporting limit was unavoidably elevated above the 5 ug/L MTCA Method A cleanup level. Benzene was also was not detected at a detection limit of 0.5 ug/L during the June 2008 monitoring event.
- In well MW-6, benzene was detected at 1.93 ug/L in June 2008 and 3.1 ug/L in October 2008. The October 2009 benzene concentration in the MW-6 groundwater sample was non-detect at an unavoidably elevated detection limit of 7.59 ug/L.

1.3 Monitored Natural Attenuation Assessment

Natural attenuation appears to be progressing at the site within the relatively long-term time frame expected. During natural attenuation, hydrocarbondegrading microbes oxidize and metabolize petroleum hydrocarbons using electron acceptors such as dissolved oxygen, nitrate, ferric iron, manganese, sulfate, and carbon dioxide. Groundwater monitoring data indicate that dissolved oxygen, nitrate, ferric iron, and sulfate are currently being used as electron acceptors. However, natural attenuation in the source area appears to be limited by the availability of these acceptors.

Figures 3 and 4 plot nitrate and sulfate groundwater data along the plume axis (MW-4R, MW-14, and MW-6). The plot demonstrates that nitrate and sulfate electron acceptor concentrations decline as groundwater moves through the plume. MW-4R has higher sulfate concentrations than the downgradient wells MW-14 and MW-6. MW-4R and MW-14 have higher nitrate concentrations than the downgradient well MW-6. This indicates that sulfate and nitrate are being consumed for oxidative purposes. As the gradient across the site increases because of seasonal precipitation, relative concentrations of nitrate

and sulfate increase because of the influx of upgradient, naturally oxidative groundwater onto the site. As groundwater flux decrease and gradients flatten during the dry season, relative nitrate and sulfate concentration decline. This repeating trend is consistent with rate-limited desorption of TPH-G from the soil matrix and limited biological activity.

Groundwater data also suggests that much of the residual mass is located in the seasonal groundwater smear zone. Figure 5 shows the relationship between dissolved TPH-G concentrations and nitrate in MW-14, downgradient well MW-6, and groundwater elevations. Groundwater flux and the mass of incoming nitrate increases during the rainy season, as indicated in MW-14 data, but is consumed by the time groundwater reaches MW-6. At MW-6, seasonal high groundwater elevations, instead, result in higher dissolved TPH-G concentrations, an inverse relationship to that observed in the source area well (MW-14). This suggests that seasonal increases in oxidant loading through the source area are insufficient to quickly degrade the mobilized TPH-G.

As presented on Figure 6 and in Table 1, oxidative-reductive potential (ORP) values are predictably depressed around areas of residual contamination. ORP values less than -110 millivolts (mV) in MW-14 and MW-6 are consistent with sulfate reduction and methane production. These consistent ORP values in MW-14 and MW-6 area also suggest that native, bioavailable manganese and iron within the soil matrix has been converted to a reduced form. These reduced metals create additional oxidant demand in the form of reduced metal complexes such as ferrous sulfide.

Downgradient of MW-6, ORP values quickly rebound by the time groundwater reaches MW-13 and range from approximately +150 mV at the end of the rainy season to +250 mV at the end of the dry season. This abrupt shift in ORP is likely due to the seasonal infiltration of rainwater containing dissolved oxygen through the unpaved fairgrounds parking lot. This helps maintain an overall oxidized geochemistry in the vicinity of MW-13. Despite the seasonal mobilization of residual TPH-G from the source area, the contamination does not appear to represent a major threat to off-site receptors because rapid oxidation results in a highly truncated TPH-G plume.

2.0 REMEDIAL PROGRAM IMPLEMENTATION

The remedial program was formulated based on the nature of the contaminant (TPH-G), the target soil matrix (silty sand to sandy gravel with areas of gravel backfill), distribution (localized to shallow source area), and the availability of existing infrastructure (monitoring wells and air sparging pipe). The remedial

program will focus on using aqeous-phase transport of supplemental electron acceptors to enhance the natural attenuation already occurring at the site.

To enhance performance of the remedial program, hydrocarbon-degrading bacteria will be introduced during the first injection to reduce the acclimation time and maximize amendment usage. Conservative tracers will be introduced at two locations during this first injection to track groundwater movement, flux, and amendment use. Electron acceptor and surfactant amendments will be introduced quarterly to sustain degradation activity. A detailed description of the amendments and injection protocols are presented below.

2.1 Amendment Descriptions

A total of five amendments will be introduced into the groundwater as part of the remedial approach: (1) sodium bromide groundwater tracer; (2) sodium chloride groundwater tracer; (3) PetroBac[™], which is a combination of petroleum-degrading bacteria and surface-active agents designed to improve oxidant usage; (4) OxEA-aq[™], which is a blend of natural microbial oxidants with macro- and micronutrients to enhance petroleum destruction; and (5) Ivey-sol[®] 103, which is a nonionic surfactant designed to improve bioremediation of TPH-G. Material Safety Data Sheets (MSDS) for these products are included in Appendix B. The five amendments are summarized below.

Tracers. Conservative groundwater tracers will be used to track groundwater flow, velocity, and effective amendment distribution. The tracers used will include sodium bromide and sodium chloride salts.

PetroBac. ETEC, LLC of Portland, Oregon, manufactures and supplies the PetroBac amendment. PetroBac is a liquid containing multiple strains of proven hydrocarbon-degrading bacteria and a biodegradable surface-active agent. PetroBac will be freshly batched by ETEC with a guaranteed active plate count of 10⁸ colony-forming units per milliliter prior to injection. Fresh batching and plate count verification allows optimal activity. The biodegradable surface-active agent in PetroBac encourages the slow desorption of residual TPH-G from the soil matrix to improve petroleum degradation rates and overall oxidant consumption.

OxEA-aq. Bioremediation Specialists of Portland, Oregon, supplies the OxEA-aq amendment. OxEA-aq is a powder consisting of a highly soluble blend of nitrogen- and sulfur-based oxidants designed to enhance natural attenuation of petroleum by providing the same electron acceptors that existing site microbes are accustomed to using. The amendment also provides a diverse blend of both

macro- and micronutrients to support the rapid development of these native bacteria to further enhance hydrocarbon destruction.

Ivey-sol 103. Ivey International Inc. manufacturers Ivey-sol 103 and makes it available through EnviroSupply & Service of Irvine, California. Ivey-sol 103 is a liquid consisting of a patented, biodegradable, nonionic surfactant blend that selectively desorbs gasoline-range petroleum hydrocarbons to improve bioavailability and overall oxidant consumption.

2.2 Oxidative Treatment Calculations

Recent groundwater data was used to estimate the mass of OxEA-aq required to oxide residual petroleum contamination and restore natural oxidative geochemistry at the site. The result of these calculations is presented in Table 2. Results suggest that 2,000 pounds of OxEA-aq should be sufficient to completely destroy both dissolved and adsorbed petroleum hydrocarbons as well as return the site's geochemistry from sulfate-reducing/methanogenic back to nitrate-reducing/aerobic conditions. Because of differences in contaminant distribution or amendment contact, these values only provide an estimate and actual performance will be verified with groundwater quality data and may be adjusted accordingly.

The calculation assumptions presented in Table 2 represent best professional judgment for this site given the age of the plume, remedial actions performed to date, and recent analytical data. These assumptions include:

- Generalizing a 1.5 mg/L TPH-G plume with a very low fraction of aromatics, based on available groundwater monitoring data;
- Incorporating a small concentration value for heavier diesel (0.005 mg/L) hydrocarbons because of the low percentage of aromatic TPH fractions. This captures the deviation from the normal heptane-range average for gasoline and accounts for bias toward heavier fractions commonly observed in weathered gasoline.
- Assuming dissolved hydrogen or equivalent organic acid concentrations sufficient to generate 5 mg/L of methane will be oxidized, based on ORP values; and
- Assuming that an average groundwater equivalent of 20 mg/L of sulfate, 15 mg/L of iron, and 2 mg/L of manganese will be reoxidized due to amendment addition.

As the treatment process proceeds, we will use groundwater data to assess the accuracy of these generalized assumptions. The amount of OxEA-aq being injected during the first three events will be sufficient to degrade between 200 and 1,000 pounds of petroleum. The actual mass degraded depends on how the hydrocarbons are incorporated into biomass (the species primarily responsible for oxidation) and the extent to which volatile organic compounds are generated as a degradation byproduct (instead of being completely oxidized). Subsequent injections may be modified for optimum remedy performance based on monitoring results.

2.3 Injection Protocol

The injection schedule is presented in Table 3 and is based on quarterly amendment additions. In general, all wells with recent groundwater quality measurements with ORP values lower than the surrounding aquifer ORP values will receive some level of oxidative treatment. These include wells MW-2, MW-4R, MW-5, MW-6, and MW-14 (Figure 7). MW-3 and the air sparging pipe will also be used to introduce amendments and enhance distribution based on their location along the plume axis. The estimated extent of oxidative treatment is presented on Figure 8.

All injections will be under pressure using municipal water pressure or a transfer pump. Pressures will be monitored in-line near the wellhead and will be limited to 15 pounds per square inch. This pressure has been found to preserve well seal integrity while pushing amendment into less-accessible pore spaces. Material will be conveyed to each injection location using a flexible garden hose and a secured high-pressure Furnco compression fitting. In-line valves located up-flow of the pressure gauge will be used to control flow rates and injection pressures. A flow meter will be used to monitor overall injection volumes at each location.

During the injection events, groundwater levels may also be measured in selected wells to help evaluate amendment distribution, overall rise in groundwater levels, and to indicate potential short circuiting of the injected amendments. A contingency plan is provided in Appendix A.

2.3.1 Event 1 - Initial Injection

During the initial injection event, groundwater tracers will be introduced into MW-3 and the air sparging pipe. Also during the initial injection event, PetroBac will be introduced into MW-4R, MW-6, MW-14, and the air sparging pipe since these locations are along the plume axis in areas of known contamination. The initial injection activities are described below for each injection location.

- MW-2. Dissolve 25 pounds of OxEA-aq in 25 gallons of municipal water to create a concentrated amendment solution and inject into the well. After the injection, chase the concentrated solution with municipal water at a rate of 9 gallons of water per 1 gallon OxEA-aq concentrate to maximize amendment distribution away from the injection location and into the surrounding formation.
- MW-3. The order of amendment addition to well MW-3 is tracer, chase water, OxEA-aq, and chase water. Dissolve 25 pounds of sodium chloride in 25 gallons of municipal water and inject into the well. After the tracer injection, add 25 gallons of municipal water to chase and distribute the tracer into the formation. Dissolve 250 pounds of OxEA-aq in 250 gallons of municipal water and inject into the well. After the injection, add chase water at a rate of 9 gallons of water per 1 gallon OxEA-aq concentrate.
- MW-4R. The order of amendment addition to well MW-4R is PetroBac, OxEA-aq, and chase water. Dilute PetroBac in municipal water to 5 percent strength or 5 gallons of PetroBac per 100 gallons of municipal water, and inject into the well. Dissolve 250 pounds of OxEA-aq in 250 gallons of municipal water and inject into the well. After the injection, add chase water at a rate of 9 gallons of water per 1 gallon OxEA-aq concentrate.
- MW-5. Dissolve 25 pounds of OxEA-aq in 25 gallons of municipal water and inject into the well. After the injection, add chase water at a rate of 9 gallons of water per 1 gallon OxEA-aq concentrate.
- MW-6. The order of amendment addition to well MW-6 is PetroBac, OxEA-aq, and chase water. Dilute PetroBac in municipal water to 5 percent strength, or 5 gallons of PetroBac per 100 gallons of water, and inject into the well. Dissolve 50 pounds of OxEA-aq in 50 gallons of municipal water and inject into the well. After the injection, add chase water at a rate of 9 gallons of water per 1 gallon OxEA-aq concentrate.
- MW-14. The order of amendment addition to well MW-14 is PetroBac, OxEA-aq, and chase water. Dilute PetroBac in municipal water to 5 percent strength, or 5 gallons of PetroBac per 100 gallons of water, and inject into the well. Dissolve 200 pounds of OxEA-aq in 200 gallons of municipal water and inject into the well. After the injection, add chase water at a rate of 9 gallons of water per 1 gallon OxEA-aq concentrate.
- Air Sparging Pipe. The order of amendment addition to the air sparging pipe is tracer, chase water, PetroBac, OxEA-aq, and chase water. Dissolve 25 pounds of sodium bromide in 25 gallons of municipal water and inject

into the well. After the tracer injection, add 25 gallons of chase water to distribute the tracer into the formation. Dilute PetroBac in municipal water to 5 percent strength, or 10 gallons of PetroBac per 200 gallons of water, and inject into the well. Dissolve 250 pounds of OxEA-aq in 250 gallons of municipal water and inject into the well. After the injection, add chase water at a rate of 9 gallons of water per 1 gallon OxEA-aq concentrate.

2.3.2 Events 2 Though 4 - Quarterly Injections

During the quarterly injections, the amount of OxEA-aq and Ivey-sol 103 vary each quarter. Please refer to the injection schedule (Table 3) for the rate and frequency of the amendments. The quarterly injection activities are summarized below for each injection location.

- MW-2. No injections are scheduled for Events 2 through 4.
- MW-3. Dissolve the required amount of OxEA-aq in municipal water (at a concentration of 1 pound of amendment to 1 gallon of water) and inject into the well. After the injection, add chase water at a rate of 9 gallons of water per 1 gallon OxEA-aq concentrate.
- MW-4R. Dissolve the required amount of OxEA-aq in municipal water (at a concentration of 1 pound of amendment to 1 gallon of water). Then add lvey-sol 103 to the OxEA-aq concentrated amendment solution. To achieve the manufacturer recommended application strength, add lvey-sol 103 at a rate of 0.4 gallons per 50 pounds of OxEA-aq in 50 gallons of municipal water. Inject the solution into the well and then chase with municipal water at a rate of 1 part concentrate per 9 gallons of chase water.
- MW-5. No injections are scheduled for Events 2 through 4.
- MW-6. Dissolve the required amount of OxEA-aq in municipal water (at a concentration of 1 pound of amendment to 1 gallon of water). Then add the required amount of Ivey-sol 103 to the OxEA-aq concentrated amendment solution. Inject the solution into the well and then chase with municipal water at a rate of 1 part concentrate per 9 gallons of chase water. Note that no injections are scheduled for Event 3.
- MW-14. Dissolve the required amount of OxEA-aq in municipal water (at a concentration of 1 pound of amendment to 1 gallon of water). Then add the required amount of Ivey-sol 103 to the OxEA-aq concentrated amendment solution. Inject the solution into the well and then chase with municipal water at a rate of 1 part concentrate per 9 gallons of chase water.

Air Sparging Pipe. Dissolve the required amount of OxEA-aq in municipal water (at a concentration of 1 pound of amendment to 1 gallon of water). Then add the required amount of Ivey-sol 103 to the OxEA-aq concentrated amendment solution. Inject the solution into the well and then chase with municipal water at a rate of 1 part concentrate per 9 gallons of chase water.

3.0 GROUNDWATER SAMPLING AND ANALYSIS

The monitoring schedule is presented in Table 4. Routine annual sampling was done in November 2010 and will be repeated in October 2011 for all monitoring wells. The October 2011 samples will be collected following the third injection event and will be used to assess oxidative treatment success.

Additional quarterly monitoring of wells MW-3, MW-4R, MW-6, MW-13, and MW-14 is needed to further evaluate treatment progress as the remedial program is implemented. Quarterly groundwater sampling and analysis before and during the quarterly injections is required to: (1) establish groundwater concentration of constituents over the course of seasonal groundwater changes; (2) determine the progress of the treatments; and (3) aid in any necessary adjustments as the injection program proceeds. An additional post-injection quarterly groundwater sampling and analysis event (Event 4) is scheduled to determine constituent concentrations for compliance monitoring. Additional groundwater sampling and analysis for compliance monitoring will be required to determine if petroleum concentrations in Site groundwater are below MTCA Method A groundwater cleanup levels. The scope and schedule for these post-injection compliance monitoring events are summarized in Table 5.

3.1 Groundwater Sampling Procedures

This section describes the procedure to be used for groundwater sampling at the site. Water levels will be measured in all wells before each sampling and/or injection event. Wells will be purged and sampled using low-flow groundwater sampling techniques. Purging and sampling will be conducted at a depth representing the middle of the screened interval of each well.

3.1.1 Sampling Equipment

Equipment used to collect groundwater samples includes:

 Horiba U-22 or equivalent water parameter instrument to collect pH, ORP, turbidity, dissolved oxygen, specific conductivity, and temperature measurements;

- Solinst or equivalent water level indicator;
- Peristaltic pump with disposable polyethylene tubing;
- Laboratory-supplied, pre-cleaned sample containers;
- Coolers with blue ice; and
- Hart Crowser Sample Custody Record and Groundwater Sampling Data forms.

3.1.2 Low-Flow Sampling

Upon arrival at the wellhead, field personnel record well conditions, depth to water, and depth to sediment in the well. Consistent with prior monitoring techniques and to minimize turbidity, low-flow purging methods are employed. Typical target flow rates range from 0.2 to 0.5 L/min. The wells will be purged and sampled with a peristaltic pump. Clean sample tubing is used for each well and disposed of after use.

The field parameters of pH, temperature, specific conductivity, dissolved oxygen, turbidity, and ORP will be measured and recorded periodically during well purging. Once the field parameters remain stable between measurements, groundwater samples will be collected by directly filling the laboratory-provided sample containers.

Blind field duplicate samples will be collected for the annual groundwater sampling events only and at a frequency of one duplicate for every 20 samples.

3.1.3 Field and Laboratory Analysis

During the annual groundwater sampling events, field kit sampling for ferrous iron will be performed on all monitoring wells. During the quarterly groundwater sampling events, field kit sampling for ferrous iron and nitrate will be performed on the five monitoring wells MW-3, MW-4R, MW-6, MW-13, and MW-14.

The groundwater sample containers will be placed in insulated coolers with blue ice. Samples will be transferred under chain of custody procedures to Analytical Resources Inc. in Tukwila, Washington, for laboratory analysis. Groundwater samples will be analyzed for the following analytes:

- TPH-G by Ecology Method NWTPH-Gx;
- Benzene, toluene, ethylbenzene, and xylene (BTEX) by EPA Method 8021B; and
- Nitrate, sulfate, chloride, and bromide by EPA Method 300.0.

3.1.4 Decontamination

All dedicated tubing will be disposed of after sampling. The only nondedicated equipment that will go down the well or come into contact with groundwater and require decontamination is the water level indicator. Decontamination will consist of washing with a detergent solution of Liquinox (or similar) in tap water, followed by a tap water rinse and a distilled water rinse.

3.1.5 Investigation-Derived Waste

Investigation-derived waste (IDW) includes wastewater from well purging and decontamination and used personal protective equipment (PPE), tubing and other solid materials. Wastewater will be temporarily stored on site in sealed, labeled drums. Analytical results from laboratory testing will be used for disposal characterization. Hart Crowser will coordinate off-site transport and disposal of wastewater by a qualified waste handling firm. PPE and other solid wastes will be removed from the site by sampling personnel and placed in a normal refuse container for disposal at a municipal solid waste landfill in accordance with applicable laws.

4.0 CHEMICAL DATA QUALITY REVIEW AND DATA QUALITY ASSURANCE

Data quality will be assessed by the contract laboratory and Hart Crowser to confirm that results meet quality control criteria. Hart Crowser's data review will include the following criteria to evaluate acceptability:

- Sample handling, holding times, and chain of custody;
- Accuracy as evaluated by:
 - Laboratory method blanks;
 - Matrix spike recovery;
 - Laboratory control sample recovery;
- Precision as evaluated by laboratory and field duplicate relative percent differences (RPD); and
- Data completeness.

Data qualifiers may be defined and reported by the laboratory to more completely explain quality control concerns regarding individual sample results. Data qualifiers will be defined in the laboratory's narrative reports associated with each case. Data

qualifier flags, beyond any applied by the laboratory, will also be added by Hart Crowser if sample results fall outside acceptance criteria. Using results of the chemical data quality review, Hart Crowser will determine the acceptability of the data for project use.

4.1 Corrective Actions

If data review by the laboratory or Hart Crowser indicates unacceptable conditions or data (e.g., exceeding holding times, insufficient or inaccurate methods, improper equipment calibration, or other potential field or laboratory deviations from this Work Plan), actions specified in the laboratory SOPs will be taken. Corrective actions may include but are not limited to the following:

- Identifying the source of the violation;
- Reanalyzing samples if holding-time criteria permit;
- Resampling and analyzing;
- Evaluating and amending sampling and analytical procedures; and/or
- Accepting data and flagging it to indicate the level of uncertainty.

If unacceptable conditions occur, the laboratory will contact Hart Crowser to discuss the issues and determine the appropriate corrective action. Any corrective actions taken by the laboratory during sample analysis for this project will be documented by the laboratory in the case narrative associated with the impacted samples.

4.2 Sample Quality Assurance/Quality Control Samples

Field duplicates will be collected for QA/QC during the annual groundwater sampling event. The laboratory will supply trip blanks for those coolers transporting samples destined for volatile fraction analysis during the annual and quarterly groundwater sampling events.

4.2.1 Field Duplicate Samples

Field duplicate samples are used to assess combined field, sampling, and laboratory technique variability. For the annual groundwater sampling events only, one field duplicate will be collected for every 20 groundwater samples collected. The well location selected for a duplicate sample is MW-6. On the chain of custody, the duplicate sample will be designated as MW-KA. The collection time recorded for the duplicate will be one half hour past the actual time the sample was collected. The field duplicate will be analyzed for the same constituents as MW-6.

4.2.2 Trip Blanks

Trip blanks are used to monitor sample cross-contamination during shipment. One trip blank set will be placed in each cooler containing groundwater samples to be analyzed for NWTPH-Gx/BTEX. Sample identifiers for trip blanks will be assigned by the laboratory. Trip blanks will be submitted to the laboratory for NWTPH-Gx/BTEX analysis only.

5.0 EVALUATION AND REPORTING

Data collected during the injection events may be used to modify the program, depending on site-specific conditions. The data will assist with determining if the fourth amendment injection is necessary to achieve remedial goals.

We will prepare a report summarizing results of the remedial program after the Event 4 post-injection groundwater sampling event. The report will include the sample analytical results, groundwater quality data tables, and figures. The report will compare the constituents of concern, migration pathways, and receptors with the appropriate MTCA Method A cleanup levels. We will also provide recommendations for future remedial actions, as necessary. A draft report will be submitted to Ken Peterson, Steve Lathrop, and Sergio Honl at AIG for review and comments. Following incorporation of review comments and document edits, we will submit a revised report to Ecology.

Additionally, annual reports for the groundwater compliance monitoring will be required per the upcoming Agreed Order with Ecology. Therefore, we will prepare and submit these annual reports to Ken Peterson, Steve Lathrop, and Sergio Honl at AIG for review and comments prior to submitting to Ecology.

6.0 SCHEDULE

We expect to begin the field work after receiving approval from Ecology's UIC program. We submitted the UIC registration letter to Ecology on November 12, 2010. We will also submit this Work Plan to Ecology for their review. UIC registration could take 60 days or more after submitting the application to Ecology, depending on Ecology's level of review. For planning purposes, we propose to begin the initial injection event in January 2011. We expect the technical report to be completed three to four weeks after we have received the post-injection laboratory report.

Groundwater compliance monitoring will begin after the post-injection report has been submitted to Ecology. We expect the annual compliance monitoring reports will be completed three to four weeks after we have received laboratory reports for the last scheduled monitoring events for each year (annual fall events).

7.0 LIMITATIONS

Work for this project was performed, and this work plan prepared, in accordance with generally accepted professional practices for the nature and conditions of the work completed in the same or similar localities, at the time the work was performed. It is intended for the exclusive use of Ken's Auto Wash for specific application to the referenced property. This report is not meant to represent a legal opinion. No other warranty, express or implied, is made. If you have any questions regarding this letter, please feel free to contact us.

J:\Jobs\716809\Work Plan\Final Bioremediation WP.doc

		Millivolts	Concentr	ration in mg/L	g/L Concentration in µg/L						
	Date				TPH-			Ethvl-	Total	Total	
Well ID	Sampled	ORP	Nitrate	Sulfate	Gasoline	Benzene	Toluene	benzene	Xylenes	Lead	Diss. Lead
MW-1	4/8/1996	NR			160,000	2,500	19,000	3,000	21,000	65	
	1/5/1998	NR									
	4/6/1998	NR			100,000	180	260	940	9,800	180	
	7/6/1998	NR			93,000	110	200	760	8,800	220	
	10/5/1998	NR									
	12/29/1999	NR			21,600	87.4	47.7	657	3,900		21.3
	3/21/2000	NR			19,800	94.1	59.6	479	2,710		16.5
	6/14/2000	NR			18,800	94.9	26.4	471	2,870		8
	9/12/2000	NR			21,400	111	35.1	496	2,930		6.54
IVIVV-14	1/30/2001				7,450	19.3	14	424	073		
	7/20/2001				20,100	37.2	29.7	318	2,000		
10100-1)	10/27/2001	NR			9 970	46.4	4 55	187	707		
	11/15/2002	NR			8 380	11	25 11	122	357		
	5/9/2003	NR			4.520	2 62	0.5 U	0 775	172	5 33	
	9/30/2003	NR	0.349	0.40 U	6.230 J	11.7 J	1.61 J	151 J	369 J	4.56	
	12/11/2003	NR	0.200	U 1.14	5,890	12.6	5.0 U	5.0 U	271	12.4	
	3/31/2004	NR	0.200	U 1.08	6,270	12.6	5 U	80.4	168.4	4.85	
	6/2/2004	NR	0.200	U 4.24	3,790 J	2.36 J	0.5 U	26.9 J	88.1 J	4.12	
	9/30/2004	NR	0.200	U 0.64	5,700 J	5.52	2.5 U	82.1	256	4.29	
	12/14/2004	NR	0.200	U 0.40 U	5,500 J	4.36	0.643	66.1	178		
	4/4/2005	NR	0.200	U 0.46	8,100 J	6.89	0.746	75.8	221		
	10/6/2005	NR	0.200	U 0.40 U	4,070 J	7.85	0.5 U	43.1	62.8	3.7	
	6/28/2006	NR	0.556	13.4	533	0.545	0.5 U	0.593	5.34	3.41	
	11/13/2006	24	0.200	U 1.40	496	0.933	0.5 U	6.89	5.99	3.03	
	5/25/2007	148	3.12	12.2	54 2 050	0.5 U	0.5 0	0.5 U	1 0		
	6/4/2008	-120	1.010	0 0.90	3,050	7.0	2.00	20.1	20	2.31	
	10/21/2008	-113	0 200	U 0.68	2 040	4 76	0.5 U	16.6	15.1	1 85	,
	10/14/2009	-106	0.900 l	JJ 1.20 U	2,030	12.2 U	0.844 U	18.9	33.8	1.00 2 L	J
MW-2	4/8/1996	NR			50 U	1 U	1 U	1 U	1 U	5 L	J
	1/5/1998	NR			50 U	1 U	1 U	1 U	1 U	15	5 L
	4/6/1998	NR			50 U	1 U	1 U	1 U	1 U	5 L	J
	7/6/1998	NR			50 U	1 U	1 U	1 U	1 U	21	
	10/5/1998	NR			50 U	1 U	1 U	1 U	1 U	34	
	12/29/1999	NR			50 U	0.5 U	0.5 U	0.5 U	1 U		1 L
	3/21/2000	NR			50 U	0.5 U	0.5 U	0.5 U	1 U		1 L
	6/14/2000				50 U	0.5 U	0.5 U	0.55	3.41		1 L
	9/12/2000				50 U	0.5 0	0.5 0	0.5 U	1 U		
	4/26/2001				50 U	0.5 0	0.5 0	0.5 0	1 U		
	7/29/2001	NR			50 U	0.5 U	0.5 U	0.5 U	1 11		
	10/27/2001	NR			50 U	0.5 U	0.5 U	0.5 U	1 U		
	11/15/2002	NR			50 U	0.5 U	0.5 U	0.5 U	1 U		
	5/9/2003	NR			50 U	0.5 U	0.5 U	0.5 U	1 U	1 L	J
	9/30/2003	NR	0.489	3.38	50 U	0.5 U	0.5 U	0.5 U	1 U	2.61	
	12/11/2003	NR	1.080	3.79	50 U	0.5 U	0.5 U	0.5 U	1 U	1 L	J
	3/31/2004	NR	0.912	4.60	13,000	10 U	119	180	2,541 J	1 L	J
	6/2/2004	NR	0.467	3.23	1,480	2.10	0.5 U	0.5 U	11.0	1 L	J
	9/30/2004	NR	0.443	2.93	1,290 J	2.40	0.5 U	0.859	5.11	1 L	J
	12/14/2004		0.922	3.05	50 U	0.5 0	0.5 U	0.5 U	1 U ₄		
	4/4/2005 10/6/2005		0.719	3.52	101	0.5 U	0.5 U	0.5 U	1 U 1 II		
	6/28/2005		0.219	3.13	100	0.741	0.5 0	0.5 U	I U	1 (,
	11/13/2006	230	0 4 1 0	 5 26	50 11	05.0	05 11	05 11			 J
	5/25/2007	162	2.740	8.57	50 U	0.5 U	0.5 U	0.5 U	1 []		
	11/7/2007	15	0.275	4.32	50 U	0.5 U	0.5 U	0.5 U	1 U	1 L	J
	6/4/2008	27	1.440	6.14	50 U	0.5 U	0.5 U	0.5 U	1 U	1 U	J
	10/21/2008	-9	0.200	U 3.21	50 U	0.5 U	0.5 U	0.5 U	1 U	20.8	
	10/14/2009	163	0.900	U 6.50	80 U	0.5 U	0.5 U	0.5 U	1 U	2 L	J

		Millivolts	Concen	tration in mg/L	L Concentration in μg/L						
	Date				TPH-			Ethyl-	Total	Total	
Well ID	Sampled	ORP	Nitrate	Sulfate	Gasoline	Benzene	Toluene	benzene	Xvlenes	Lead	Diss. Lead
MW-3	4/8/1996	NR			50 U	1 U	1 U	1 U	1 U	5 U	
	1/5/1998	NR			50 U	1 U	1 U	1 U	1 U	5 U	
	4/6/1998	NR			50 U	1 U	1 U	1 U	1 U	5 U	
	7/6/1998	NR			50 U	1 U	1 U	1 U	1 U	5 U	
	10/5/1998	NR			50 U	1 U	1 U	1 U	1 U	3.8	
	12/29/1999	NR			50 U	0.5 U	0.5 U	0.5 U	1 U		1 U
	3/21/2000	NR			50 U	0.5 U	0.5 U	0.5 U	1 U		1 U
	6/14/2000	NR			50 U	0.5 U	0.85	0.5 U	1 U		1 U
	9/12/2000	NR			50 U	0.5 U	0.5 U	0.5 U	1 U		1 U
	1/30/2001	NR			50 U	0.5 U	0.5 U	0.5 U	1 U		
	4/26/2001	NR			50 U	0.5 U	0.5 U	0.5 U	1 U		
	7/29/2001	NR			50 U	0.5 U	0.5 U	0.5 U	1 U		
	10/27/2001	NR			50 U	0.5 U	0.5 U	0.5 U	1 U		
	11/15/2002	NR			50 U	0.5 U	0.5 U	0.5 U	1 U		
	5/9/2003	NR			50 U	0.5 U	0.5 U	0.5 U	1 U	1 U	
	9/30/2003	NR	0.228	4.39	50 U	0.5 U	0.5 U	0.5 U	1 U	1 U	
	12/11/2003	NR	0.200	U 4.79	50 U	0.5 U	0.5 U	0.5 U	1 U	1 U	
	3/31/2004	NR	0.812	5.53	50 U	0.2 U	0.2 U	0.2 U	0.5 U	1 U	
	6/2/2004	NR	0.816	5.61	50 U	0.5 U	0.5 U	0.5 U	1 U	1 U	
	9/30/2004	NR	0.253	4.43	50 UJ	0.5 U	0.5 U	0.5 U	1 U	1 U	
	12/14/2004		0.206	4.69	50 U	0.5 U	0.5 U	0.5 U	1 U		
	4/4/2005		0.358	4.23	50 U	0.5 0	0.5 0	0.5 0	10		
	10/6/2005		0.200	U 3.67	50 U	0.5 0	0.5 0	0.5 0	10	1 0	
	0/20/2000	204			 50						
	F/25/2007	140	1 520	6.43	50 0	0.5 0	0.5 0	0.5 U	1 U	10	
	5/25/2007	251	0.168	0.43	50 0	0.5 0	0.5 0	0.5 0	1 U		
	6/4/2008	37	0.100	4.15	50 U	0.5 0	0.5 0	0.5 0	1 1	1 11	
	10/21/2008	28	0.320	3.84	50 U	0.5 U	0.5 0	0.5 0	1 U	1 1	
	10/14/2009	68	0.200	U.I 3 20	80 U	0.0 U	0.5 U	0.5 U	1 U	2 1	
MW/_4	1/5/1998	NR	0.00		200	1 11	27	1	3	10	5 11
10100-4	4/6/1998	NR			400	3	14	1	6	5 11	
	7/6/1998	NR			50 U	1 U	.14	1 U	1 U	5 U	
	10/5/1998	NR			150	1 U	7	1 U	1 U	2	
	12/29/1999	NR			301	51.4	32.5	0.5 U	6.08		1 U
	3/21/2000	NR			414	44.8	28.2	1.92	3.2 U		1 U
	6/14/2000	NR			439	69.7	4.91	2.01	6.8		1 U
	9/12/2000	NR			101	4.49	0.5 U	0.5 U	0.5 U		1 U
	1/31/2001	NR			182	2.22	1.17 U	0.5 U	1.33 U		
	4/26/2001	NR			673	8.79	4.73	4.28	28.6		
	7/29/2001	NR			402	24.3	16.3	2.84	14.8		
	10/27/2001	NR			200	24.9	2.62	1.15	6.57		
	11/15/2002	NR			75.6	0.858	0.5 U	0.5 U	1 U		
	5/9/2003	NR			61.8	0.5 U	0.5 U	0.5 U	1 U	1 U	
	9/30/2003	NR	0.200	U 4.57	161	0.730	0.5 U	2.59	2.59	1 U	
	12/11/2003	NR	1.05	15.3	50 U	0.5 U	0.5 U	0.5 U	1 U	3.22	
	3/31/2004	NR	0.200	U 7.41	267	29.0	1.43	1 U	2.94	1 U	
	6/2/2004	NR	0.200	0 8.32	140	46.4	4.2	0.5 0	1 U	10	
	9/30/2004		0.200	U 4.91	88.7 J	0.5 U	0.5 U	1.83	ïU ∡	ΊU	
	12/14/2004		0.200	U 5.13	50 U	0.5 U	0.5 U		1 U 4 U		
IVIVV-4K	4/4/2000 10/6/2005		0.200		112	1.93	0.5 0	0.3 0	3.57		
	6/28/2000		0.200	U 0.07	744 50 U	0.929		9.31 05 11	ວ.ວ/ 1 II	19	
10100-44)	11/13/2000	50	0.200	11 16 2	107			0.5 0	1 1	5.82	
	5/25/2007	40	2 290	17.6	50 11	0.5 U	0.5 0	0.5 U	1 11		
	11/7/2007	-46	0.031	10.3	75.2	0.5 U	0.5 U	05 U	1 1	0.325	
	6/4/2008	-59	2.030	14.1	50 U	0.5 U	0.5 U	0.5 U	1 U	1 U	
	10/21/2008	-54	0.200	U 6.52	50 U	0.5 U	0.5 U	0.5 U	1 U	6.98	
	10/14/2009	100	0.90	UJ 5.90	80 U	0.5 U	0.5 U	0.5 U	1 U	2 U	
										_ 3	

		Millivolts	Concer	tration in mg/L	/L Concentration in µg/L						
	Date				TPH-			Ethyl-	Total	Total	
Well ID	Sampled	ORP	Nitrate	Sulfate	Gasoline	Benzene	Toluene	benzene	Xylenes	Lead I	Diss. Lead
MW-5	1/5/1998	NR			6200	1	57	3	160	5 U	
	4/6/1998	NR			2800	2	30	2	27	5 U	
	7/6/1998	NR			50 U	1 U	1 U	1 U	1 U	10	
	10/5/1998	NR			4700	2	39	16	94	7.4	
	12/29/1999	NR			779	2.96	0.69	9.03	27.4		10
	3/21/2000				519	0.5 U	13.9	4.95	3.0		1 U
	0/14/2000				708 50 H	3.45 U	0.5 11	0.5 11	1 U		1 1
	4/26/2001	NR			831	7.35	0.5 0	15.3	1 U		
	7/29/2001	NR			53.8	0.5 U	0.5 U	0.5 U	1 U		
	10/27/2001	NR			552	3.29	0.5 U	1.28	1.58		
	11/15/2002	NR			108	0.5 U	0.5 U	0.5 U	0.5 U		
	5/9/2003	NR			78.7	0.5 U	0.5 U	0.5 U	1 U	1 U	
	9/30/2003	NR	0.200	U 8.61	229	0.5 U	0.5 U	0.5 U	1.61	1 U	
	12/11/2003	NR	0.200	U 6.85	50 U	0.5 U	0.5 U	0.5 U	1 U	1 U	
	3/31/2004	NR	1.32	16.1	53	0.2 U	0.2 U	0.2 U	0.5 U	1 U	
	6/2/2004	NR	1.36	11.7	92.8	0.5 U	0.5 U	0.5 U	1 U	1 U	
	12/14/2004		0.200	U 7.57	308	0.5 U	0.5 0	0.5 U	1 0		
	4/4/2005		0.200	U 9.92	020 114	0.5 11	0.5 U	0.5 0	1.07		
	6/28/2006	NR	2.59	16.0	50 U	0.5 U	0.5 U	0.5 U	1 U	1 U	
	11/13/2006	279	2.99	11.7	50 U	0.5 U	0.5 U	0.5 U	1 U	1 U	
	5/25/2007	169	3.400	19.9	50 U	0.5 U	0.5 U	0.5 U	1 U		
	11/7/2007	199	0.110	7.75	50 U	0.5 U	0.5 U	0.5 U	1 U	1 U	
	6/4/2008	52	1.730	11.8	50 U	0.5 U	0.5 U	0.5 U	1 U	1 U	
	10/22/2008	91	0.220	6.35	50 U	0.5 U	0.5 U	0.5 U	1 U	1 U	
	10/15/2009	239	0.90	U 5.20	80 U	0.5 U	0.5 U	0.5 U	1 U	2 U	
MW-6	1/5/1998	NR			2,200	53	17	9	93	5 U	
	4/6/1998	NR			4,200	51	16	25	110	5 U	
	7/6/1998	NR			6,900	11	19	1	510	11	
	10/5/1998				5,800	43	22	48	240	12	
	3/21/2000				2,090	0.75 11	2 1/1 3	28.7	61		1 1
	6/14/2000	NR			2,170	9.78	1 03 U	33.1	101		1 U
	9/12/2000	NR			1.630	12.8	1.2 U	27.9	75.7		1 U
	4/26/2001	NR			1,320	11.3	0.906	1.41	3.37		
	7/29/2001	NR			5,050	8.71	4.99	189	536		
	10/27/2001	NR			1,910	15.3	0.786	1.67	5.49		
	11/15/2002	NR			1,270	9.01	0.5 U	0.594	1.85		
	5/9/2003	NR			1,710	1.79	0.5 U	1.29	21.2	1.29	
	9/30/2003	NR	0.200	U 0.400 U	1,610	16.7	2.50 U	2.91	7.96	1 U	
	12/11/2003		0.200	U 0.685	624	5.67	0.50 U	0.737 J	2.19 J	10	
	6/2/2004		0.200	0 5.02	2 300	0.520 4.78 I	0.2 0	0.350 54.0 I	755	1 20	
	9/30/2004	NR	0.200		1 150	834	0.5 0	0.553	2 92 .1	1.23	
	12/14/2004	NR	0.200	U 0.400 U	672	3.57	0.5 U	0.5 U	1.42		
	4/4/2005 ^b	NR	0.200	U 3.19	1.010	5.91	0.5 U	0.5 U	1.86 °		
	10/6/2005	NR	0.200	U 3.17	1,380 J	8.10	0.5 U	0.632	1.94	1 U	
	6/28/2006	NR	2.6	18.6							
	11/13/2006	-4	0.200	U 1.11	826	3.3	0.5 U	0.5 U	1.89	1 U	
	5/25/2007	-92	0.200	U 2.67	1460	0.5 U	0.5 U	25.6	1.22		
	11/7/2007	-77	0.010	U 2.24	729	3.53	0.5 U	0.5 U	1.69	1 U	
	6/4/2008	-118	0.200	U 3.68	1550	1.93	0.5 U	30.8	2.78	1 U	
	10/22/2008	-118	0.200	U 0.40 U	855	3.1	0.5 U	0.933	3.37	1 U	
	10/14/2009	-102	0.90	UJ 1.2 U	501	7.59 U	0.5 U	1.18 U	1 U	2 U	
MW-12	12/29/1999	NR			50 U	0.5 U	0.5 U	0.5 U	1 U		1 U
	3/21/2000				50 U	0.5 U	0.5 U	0.5 U	1 U ₄		1 U
	0/14/2000				50 U	0.5 U	U.5 U	0.5 U	1 U 1 U		1 U 4 U
	3/12/2000 4/26/2001				50 0	0.5 U 0.5 II	0.5 U		1 U		
	7/29/2001	NR			50 0	0.5 0	0.5 0	1 74	4 83		
			1			0.0 0	5.0 0				

		Millivolts	Concent	ration in mg/L	L Concentration in µg/L						
	Date				TPH-			Ethyl-	Total	Total	
Well ID	Sampled	ORP	Nitrate	Sulfate	Gasoline	Benzene	Toluene	benzene	Xylenes	Lead	Diss. Lead
MW-12	10/27/2001	NR			50 U	0.5 U	0.5 U	0.5 U	1 U		
(cont.)	11/15/2002	NR			50 U	0.5 U	0.5 U	0.5 U	1 U		
	5/9/2003	NR			50 U	0.5 U	0.5 U	0.5 U	1 U	1 U	
	9/30/2003	NR	0.452	5.32	50 U	0.5 U	0.5 U	0.5 U	1 U	1 U	
	12/11/2003	NR	0.200	U 2.77	50 U	0.5 U	0.5 U	0.5 U	1 U	1.47	
	3/31/2004	NR	3.88	8.45	50 U	0.2 U	0.2 U	0.2 U	0.5 U	1 U	
	6/2/2004	NR	3.64	11.7	50 U	0.5 U	0.5 U	0.5 U	1 U	1 U	
	9/30/2004		0.573	5.66	50 UJ	0.5 U	0.5 U	0.5 U	1 U	1 U	
	12/14/2004		0.200	U 2.95	50 U	0.5 0	0.5 0	0.5 0	1 U		
	4/4/2005		0.200	U 3.32	50 U	0.5 0	0.5 U	0.5 0	1 U		
	6/28/2005		2.57	11 5	50 U	0.5 0	0.5 0	0.5 0	1 0	2 98	
	11/13/2006	299	0.590	6.89	50 U	0.5 0	0.5 0	0.5 0	1 1	2.30	
	5/25/2007	183	7 140	18.4	50 U	0.0 U	0.5 U	0.0 U	1 U		
	11/8/2007	168	0.121	11.5	50 U	0.5 U	0.5 U	0.5 U	1 U	1 U	
	6/4/2008	108	6.020	16.4	50 U	0.5 U	0.5 U	0.5 U	1 U	1 U	
	10/22/2008	69	0.330	10.1	50 U	0.5 U	0.5 U	0.5 U	1 U	1 U	
	10/14/2009	75	0.90	UJ 5.20	80 U	0.5 U	0.5 U	0.5 U	1 U	2 U	
MW-13	12/29/99	NR			50 U	0.5 U	0.5 U	0.5 U	1 U		1 U
	3/21/2000	NR			50 U	0.5 U	0.5 U	0.5 U	1 U		1 U
	6/14/2000	NR			50 U	0.5 U	0.5 U	0.5 U	1 U		1 U
	9/12/2000	NR			50 U	0.5 U	0.5 U	0.5 U	1 U		1 U
	4/26/2001	NR			50 U	0.5 U	0.5 U	0.5 U	1 U		
	7/29/2001	NR			50 U	0.5 U	0.5 U	0.5 U	1 U		
	10/27/2001	NR			50 U	0.5 U	0.5 U	0.5 U	1 U		
	9/30/2003	NR	0.455	4.91	50 U	0.5 U	0.5 U	0.5 U	1 U	10	
	12/11/2003		0.477	5.56	50 U	0.5 U	0.5 U	0.5 U	1 U	1.56	
	3/31/2004		1.60	8.04	50 U	0.2 0	0.2 0	0.2 0	0.5 U	1 0	
	0/2/2004		0.406	0.52	50 U	0.5 0	0.5 0	0.5 U	1 U	1 0	
	9/30/2004 12/14/2004		0.490	4.49	50 00	0.5 0	0.5 0	0.5 0	1 0		
	4/4/2005	NR	0.412	4 99	50 U	0.5 0	0.5 0	0.5 0	1 1		
	10/6/2005	NR	0.348	3 68	50 U	0.5 U	0.5 U	0.5 U	1 U	1 U	
	6/28/2006	NR									
	11/13/2006	270	0.940	6.18	50 U	0.5 U	0.5 U	0.5 U	1 U	1 U	
	5/25/2007	175	1.670	7.57	50 U	0.5 U	0.5 U	0.5 U	1 U		
	11/8/2007	254	0.490	4.09	50 U	0.5 U	0.5 U	0.5 U	1 U	1 U	
	6/4/2008	110	1.280	5.51	50 U	0.5 U	0.5 U	0.5 U	1 U	1 U	
	10/22/2008	219	0.440	3.56	50 U	0.5 U	0.5 U	0.5 U	1 U	1 U	
	10/15/2009	231	0.90	U 3.3	80 U	0.5 U	0.5 U	0.5 U	1 U	2 U	
MW-15	1/30/2001	NR			161	1.53	0.5 U	0.5 U	1.18 U		
	4/26/2001	NR			50 U	0.5 U	0.5 U	0.5 U	1 U		
	7/29/2001	NR			50 U	0.5 U	0.5 U	0.5 U	1 U		
	10/27/2001	NR			50 U	0.5 U	0.5 U	0.5 U	1 U		
	11/15/2002	NR			50 U	0.5 U	0.5 U	0.5 U	1 U		
	5/9/2003	NR			50 U	0.5 U	0.5 U	0.5 U	1 U	1 U	
	9/30/2003	NR	0.282	5.02	50 U	0.5 U	0.5 U	0.5 U	1 U	1 U	
	3/31/2003		0.415	8.52	50 U	U.5 U	0.5 U	0.5 U		1 U	
	5/31/2004 6/2/2004		0.200	0.42	50 U	0.2 U 0.5 II	0.2 U		U.5 U 1 II	I U ₄ ⊔	
	0/20/2004		1.07	0.32	50 U	0.5 U	0.5 U	0.5 U 0.5 U	1 U 1 II	1 U	
	12/14/2004	NR	0.429	4.50 U 6.68	50 03	0.5 0	0.5 0	0.5 0	1		
	4/4/2005	NR	0.200	U 745	50 0	0.5 0	0.5 0	0.5 0	1 1		
	10/6/2005	NR	0.340	4.14	50 U	0.5 U	0.5 U	0.5 U	1 U	1 []	
	6/28/2006	NR									
	11/13/2006	229	0.450	6.48	50 U	0.5 U	0.5 U	0.5 U	1 U	1 U	
	5/25/2007	113	3.070	10.4	50 U	0.5 U	0.5 U	0.5 U	1 U		
	11/7/2007	435	0.220	5.21	50 U	0.5 U	0.5 U	0.5 U	1 U	1 U	
	6/5/2008	109	2.010	8.02	50 U	0.5 U	0.5 U	0.5 U	1 U	1 U	
	I				-					-	

		Millivolts	Concent	ration in mg/L		Concentration in µg/L						
	Date				TPH-			Ethyl-	Total	Total		
Well ID	Sampled	ORP	Nitrate	Sulfate	Gasoline	Benzene	Toluene	benzene	Xylenes	Lead	Diss. Lead	
MW-15	10/22/2008	86	0.280	3.81	50 U	0.5 U	0.5 U	0.5 U	1 U	1 L	J	
(cont.)	10/14/2009	160	0.90	UJ 3.1	80 U	0.5 U	0.5 U	0.5 U	1 U	2 L	J	
MTCA Metho	d A Groundwa	ater Cleanu	p Level		800/1,000 ^a	5	1000	700	1000	15	15	

Notes:

TPH-G analyzed by EPA Method 8015 prior to 1999. After that, TPH-G was analyzed by NWTPH-G; BTEX Analyzed by EPA Method 8021B, except in March 2004 where BTEX was analyzed by EPA Method 8260B.

Total and Dissolved Lead analyzed by EPA Method 6010 or 6020.

ORP = Oxidative-reductive potential.

mg/L = Milligrams per liter.

 μ g/L = Micrograms per liter.

-- Not analyzed.

U = Not detected at specified reporting limit.

J = Estimated concentration.

R = Rejected concentration.

ND = Not detected.

NR = Not reported.

Bolded concentrations exceed MTCA Method A cleanup levels.

Access to well MW-13 obstructed in November 2002 and May 2003.

Access to well MW-5 obstructed in September 2004.

Data from 1996 and 1998 collected by Sage Environmental.

Well MW-1 was removed during the October 2000 excavation. Wells MW-14 and MW-15 were installed in January 2001 after the excavation. Well MW-4 was replaced as well MW-4R by Hart Crowser in October 2005, following removal of the well during UST removal activities in April 2005.

Dashed line indicates date of excavation.

a) Cleanup level for TPH-G with/without detectable benzene.

b) Values shown are the average of the results for the sample and its field duplicate.

c) The value is the result for the field duplicate. The result for the sample was ND (not detected at the detection limit of 1.0 µg/L).

Table 2 - Hydrogen Balance CalculationsKen's Auto WashEllensburg, Washington

Treatment Target Area Specificati	ons			
Vertical Treatment (ft)	10			
Treatment Width (ft)	80			
Treatment Length (ft)	400			
Effective Porosity	0.25			
Foc	0.010			
Treatment Area Pore Volume	2,264,000	L	597,361	Gallons
Hydrogen/Electron Donor Availab	ility			
Constituent	Groundwater Concentration (mg/L)	Molecular Weight (g/mol)	Moles of H ₂ to Oxidize / Mole Analyte	Moles of H ₂ Donor In Treatment Area
Native Electron Donors				
Groundwater TPH-G	1.5	100	22	747
Approximate % Aromatic Estimated Total Soil and GW TPH-G	1%			3,896
Groundwater TPH-Dx	0.005	226	49	2
Estimated Total Soil and GW TPH-Dx				13,011
Estimated M	loles of Petrol	eum Hydrogen I	Donor Available	e 16,907
Hydrogen/Electron Acceptors				
Constituent	Groundwater Concentration (mg/L)	Molecular Weight (g/mol)	Moles of H ₂ to Reduce Mole Analyte	Moles of H ₂ Acceptor In Treatment Area
Native Electron Acceptors				-
Sulfate	-20.0	96.1	4	-1,885
Fe ⁺² Formation from Fe ⁺³	-15.0	55.8	0.5	-304
Mn ⁺² Formation from Mn ⁺⁴	-2.0	54.9	1	-82
Methane Formation	-5.0	16	4	-2,830
	Na	ative Hydrogen A	cceptor Subtotal	-5,101
Added Oxidative Amendment	Added Amenn	nent Mass (lbs.)	Moles H ₂ /lb.	
OxEA-aq™	2,0		10.6	21,160
	Ac	ded Hydrogen A	cceptor Subtotal	21,160
	Estimated	a woles of Hydr	ogen Acceptor:	16,059
Estimated Oxidative Treatn	nent Progress I	Based on Desig	n Assumptions	: 95%

Notes:

L = liters; gal = gallons; Fe = iron; Mn= manganese; $1ft^3 = 7.48$ gals = 28.3L; 3.79L = 1 gal.

Negative values in Native Electron Acceptors indicates formation/reoxidation of reduced atomic forms within the treated plume.

Physical constants per Oregon DEQ Risk-Based Decision Making Guidance.

Electron and hydrogen equivalents per Principles and Practices of Enhanced Anaerobic Bioremediation of Chlorinated Solvents, Air Force Center for Environmental Excellence (August 2004) and Bioremediation Specialists product data for OxEA-aq.

OxEA-aq is a blend of oxidants with macro- and micronutrients to improve amendment use.

ft = feet; L = liters; mg/L = milligrams per liter; gal = gallons; $\frac{1}{2}$ = molecular hydrogen equivalents.

Table 3 - Enhanced Bioremediation Injection ScheduleKen's Auto WashEllensburg, Washington

Injection	J	Event 1 lanuary 201	1	Ever April 2	nt 2 2011	Ever July 2	3-Event OxEA-aq	Oc	Ever tobe	nt 4 r 2011	4-Event OxEA-aq	
Location	OxEA-aq	PetroBac	Tracer	OxEA-aq	lvey-sol	OxEA-aq	lvey-sol	Subtotal	OxEA	-aq	lvey-sol	Total
MW-2	25 lbs							25 lbs				25 lbs
MW-3	250 lbs		CI 25 lbs	200 lbs		150 lbs		600 lbs	250	lbs		850 lbs
MW-4R	250 lbs	5 gal		125 lbs	1.0 gal	25 lbs	0.2 gal	400 lbs	200	lbs	1.60 gal	600 lbs
MW-5	25 lbs							25 lbs				25 lbs
MW-6	50 lbs	5 gal		25 lbs	0.2 gal			75 lbs	25	lbs	0.20 gal	100 lbs
MW-14	200 lbs	5 gal		100 lbs	0.8 gal	25 lbs	0.2 gal	325 lbs	75	lbs	0.60 gal	400 lbs
Sparge	250 lbs	10 gal	Br 25 lbs	200 lbs	1.6 gal	100 lbs	0.8 gal	550 lbs	250	lbs	2.00 gal	800 lbs
Event Total	1,050 lbs	25 gal	50 lbs	650 lbs	3.6 gal	300 lbs	1.2 gal	2,000 lbs	800	lbs	4.40 gal	2,800 lbs

Notes:

PetroBac contains biodegradable surface-active agents and petroleum-degrading microbes to enhance amendment consumption and petroleum destruction.

OxEA-aq is a soluble blend of oxidants with macro- and micronutrients to enhance petroleum degradation.

Ivey-sol is a biodegradable, nonionic surfactant formulated to improve bioremediation of petroleum hydrocarbons.

Table presents maximum injection masses. Actual mass may be reduced based on performance and monitoring results.

Abreviations:

Br = Food-grade sodium bromide salt

CI = Food-grade sodium chloride salt

lbs = pounds

gal = gallons

Table 4 - Enhanced Bioremediation Monitoring ScheduleKen's Auto WashEllensburg, Washington

Monitoring	No	Bas ovem	eline ber 20	10		Eve April	ent 2 2011			Eve July	ent 3 2011			E Oct	Event ober 2	4 2011		Post J	-Injeo anua	tion E ry 201	vent 2
Well	G	V	lons	F	G	V	lons	F/N	G	V	lons	F/N	G	V	lons	F	Ν	G	V	lons	F/N
Injection We	lls																				
MW-2	Х	Х	Х	Х									Х	Х	Х	Х					
MW-3	Х	Х	Х	Х	Х	Х	Х	Х	Х	Х	Х	Х	Х	Х	Х	Х	Х	Х	Х	Х	Х
MW-4R	Х	Х	Х	Х	Х	Х	Х	Х	Х	Х	Х	Х	Х	Х	Х	Х	Х	Х	Х	Х	Х
MW-5	Х	Х	Х	Х									Х	Х	Х	Х					
MW-6	Х	Х	Х	Х	Х	Х	Х	Х	Х	Х	Х	Х	Х	Х	Х	Х	Х	Х	Х	Х	Х
MW-14	Х	Х	Х	Х	Х	Х	Х	Х	Х	Х	Х	Х	Х	Х	Х	Х	Х	Х	Х	Х	Х
Downgradie	nt We	ells																			
MW-12	Х	Х	Х	Х									Х	Х	Х	Х					
MW-13	Х	Х	Х	Х			Х	Х			Х	Х	Х	Х	Х	Х	Х			Х	Х
MW-15	Х	Х	Х	Х									Х	Х	Х	Х					

Notes:

Monitoring will be performed prior to any injection activities.

Water level elevations will be measured prior to well purging and sampling.

Samples will be collected using low-flow techniques and a flow-through cell, consistant with recent monitoring events.

Abreviations:

G = Gasoline-range petroleum hydrocarbons by Ecology Method NWTPH-Gx.

V = Volatile organic compounds benzene, toluene, ethylbenzene, and xylene by EPA Method 8021B.

lons = Nitrate as nitrogen, sulfate, bromide, and chloride by EPA Method 300.0.

F = Field kit testing of ferrous iron.

N = Field kit testing of nitrate.

Table 5 - Groundwater Compliance Monitoring ScheduleKen's Auto WashEllensburg, Washington

Well	Purpose	2013-2017 ^a	2018-2021	2022 ^b
MW-2	Bound Plume - East	Annual ^c		Quarterly
MW-3	Background	Annual ^c	Biannual ^e	Quarterly
MW-4R	Source Area (Upgradient Edge)	Quarterly ^d	Biannual ^e	Quarterly
MW-5	Bound Plume - West	Annual ^c		Quarterly
MW-6	Plume Extent	Quarterly ^d	Biannual ^e	Quarterly
MW-12 ^f	Bound Plume - Southwest	Annual ^c		Quarterly
MW-13	Downgradient Point of Compliance	Quarterly ^d	Biannual ^e	Quarterly
MW-14	Source Area	Quarterly ^d	Biannual ^e	Quarterly
MW-15	Bound Plume - Southeast	Annual ^c		Quarterly

Notes:

Monitoring will include groundwater level measurements, field parameter measurements, and groundwater sample collection for chemical analyses specified below. If injections are to be performed into the well, the well will be field-tested for nitrate, nitrite, ammonium, and ferrous iron.

^a Timeline assumes injections are performed and amendment concentrations have not reduced to background levels. If the site enters monitored attenuation, sampling defaults to 2018 scope.

^b Final compliance monitoring would include analysis for NWTPH-Gx, BTEX, and total lead.

^c Annual monitoring includes analysis of NWTPH-Gx, BTEX, nitrate, sulfate, and total lead.

- ^d Quarterly monitoring includes analysis of NWTPH-Gx, BTEX, nitrate, and sulfate.
- ^e Biannual refers to twice per year and would be based on typical high and low groundwater elevations at the site. Includes analysis of NWTPH-Gx, BTEX, nitrate, sulfate, and total lead.

^f Well not located in May, August, and November 2012 and possibly destroyed.

Schedule assumes 5-year review by Ecology following 2013 sampling round.

Schedule after 2018 is tentative pending Ecology 5-year review.

Monitoring schedule after 2022, if necessary, will be based on review of previous data.

Blank entries indicate no monitoring planned in specific wells.













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APPENDIX A NONENDANGERMENT CONTINGENCY PLAN

APPENDIX A NONENDANGERMENT CONTINGENCY PLAN

This contingency plan complies with the requirement of the Underground Injection (UIC) Program, which requires that discharge(s) from UIC injection wells must meet the nonendangerment standard of WAC 173-218-080. Contingency Plan elements include actions to be taken in case of spills, system failures, equipment breakdowns, and/or unforeseen environmental degradation caused by the cleanup activities. The Washington State Department of Ecology (Ecology) required preparation of a written Contingency Plan for the UIC Program at the Ken's Auto Wash site in its January 24, 2011, letter to Ken Peterson. Ecology's letter rule authorized the site for the planned underground injections of *in situ* remediation amendment products.

The site is registered as UIC site number 31210.

SITE LOCATION	Ken's Auto Wash
	Ellensburg, Washington
	Phone: (509) 925-7920
EMERGENCY	Police Department911
RESPONDERS	Fire Department
	Ambulance911
EMERGENCY	Hart Crowser, Seattle Office
CONTACTS	• Mike Ehlebracht (206) 369-5164 (cell)
	• Angie Goodwin (206) 954-2549 (cell)
	• Troy Fowler (971) 678-0843 (cell)
	Client Contact – Ken Peterson (509) 929-0027 (cell)
	National Response Center
	Washington Emergency Management (800) 258-5990
	Ecology Central Regional Office (509) 575-2490
	EPA Emergency Response Team (908) 321-6660
	Poison Control Center
	Chemtrec(800) 424-9300

Emergency Contingency Information

Spill Containment

Chemical amendments that will be used on site include:

- OxEA-aq manufactured by Bioremediation Specialists, LLC;
- PetroBac manufactured by ETEC, LLC;
- Sodium bromide and potassium chloride salt tracers purchased from Cascade Columbia Distribution Company; and
- Ivey-sol 103 manufactured by Ivey International Inc.

OxyEA-aq and the tracers are dry powder products and will be picked up from the manufacturer in 50-pound bags. PetroBac is a liquid and will be picked up from the manufacturer in 5-gallon pails. Ivey-sol 103 is a liquid and will be shipped from the manufacturer to Hart Crowser in a 55-gallon drum. Ivey-sol 103 will be stored off site and only the volume needed during the quarterly injections will be transported to the site. The products will be mixed with water in small batches directly prior to injecting. Any vessel that contains products and water will be vented due to potential pressure buildup from fermentation gases. Material Safety Data Sheets (MSDS) for these products are included in Appendix B of this document. The use and control of the chemical amendments to be injected is described in Hart Crowser's Work Plan.

Container labels will contain the following information:

- Identity of hazardous chemicals;
- Appropriate hazard warnings; and
- Name and address of the chemical manufacturer and/or distributor.

Labels will not be removed or defaced. If a label is damaged or is not legible, a replacement label will be requested from the manufacturer.

When on site, Hart Crowser personnel will be equipped with sorbent pads and other materials capable of containing a relatively small spill. Dry material will be handled with care and any spill will be swept immediately. In the event of a tank rupture or other material spill of the liquid mixture, we will attempt to stop and contain the flow of material using absorbents, booms, dirt, or other appropriate material. The spill material will be drummed for proper disposal. As necessary, we will contact a spill removal firm through Chemtrec for material cleanup and disposal, as required. Sufficient empty drums will be kept on site as a contingency for spill containment.

Injectate Migration

Although the amendments are mobile products and travel with groundwater, the injections are designed to balance the amendment amounts to avoid over application and migration beyond the historically impacted area downgradient of the former USTs, near wells MW-14 and MW-6.

Preferential flow path formation will be minimized by using controlled flows and injection pressures as described in the Work Plan to the extent practical. During the injection events, groundwater levels will also be measured in selected wells to help evaluate amendment distribution, fluctuations in groundwater levels, and to determine whether potential short circuiting of the injected amendments could be occurring. Any apparent preferential flow path formation or short-circuiting will be documented and minimized by decreasing flow rates and injection pressures.

Groundwater Monitoring

Routine annual sampling was completed in November 2010 and will be repeated in October 2011 for all monitoring wells. Quarterly performance monitoring events are scheduled for April and July 2011 to monitor treatment progress in selected wells. The October 2011 samples will be collected following the third bioremediation amendment injection event and will be used to assess overall oxidative treatment success. A post-injection quarterly monitoring event is scheduled for January 2012 to determine groundwater concentrations for compliance monitoring.

Based on estimated groundwater travel times from the center of the treatment area to monitoring wells MW-6 and MW-13, we anticipate that quarterly groundwater monitoring frequency will provide sufficient data to evaluate treatment performance. Additional quarterly monitoring of wells MW-3, MW-4R, MW-6, MW-13, and MW-14 will also be conducted.

To confirm that over application or migration of amendment products beyond targeted area is not occurring, field sampling for nitrate will be performed in selected wells prior to each injection. Field nitrate results will determine whether conditions are appropriate for additional amendment application.

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APPENDIX B MATERIAL SAFETY DATA SHEETS (MSDS)

Material Safety Data Sheet

Material Name: OxEA-aq

*** Section 1 - Chemical Product and Company Identification ***

Manufacturer Information Bioremediation Specialists

8910 SW Gemini Drive Beaverton, OR 97008-7123 Phone: (971) 327-9113 Fax: (503) 620-6918 EMERGENCY: (971) 678-0843 or (971) 781-2232

** Section 2 - Hazards Identification ***

Emergency Overview

This product is not expected to produce any unusual hazards during normal use. Exposure to high dust levels may irritate the skin, eyes, nose, throat, or upper respiratory tract.

Potential Health Effects: Eyes

Airborne dust or direct contact can cause mechanical irritation of eyes. If burning, redness, itching, pain or other symptoms persist or develop, consult physician.

Potential Health Effects: Skin

Direct, prolonged or repeated contact with the skin may cause irritation.

Potential Health Effects: Ingestion

Not considered a likely route of exposure under normal product use conditions. May be harmful or fatal if swallowed.

Potential Health Effects: Inhalation

Dust exposures generated during the handling of the product may irritate eyes, skin, nose, throat, and upper respiratory tract. Persons subjected to large amounts of this dust will be forced to leave area because of nuisance conditions such as coughing, sneezing and nasal irritation. Labored breathing may occur after excessive inhalation. If respiratory symptoms persist, consult physician.

HMIS Ratings: Health: 1 Fire: 0 HMIS Reactivity 1

Hazard Scale: 0 = Minimal 1 = Slight 2 = Moderate 3 = Serious 4 = Severe * = Chronic hazard

*** Section 3 - Composition / Information on Ingredients ***

CAS #	Component	Percent
Proprietary	Trade Secret #1	10-50
Proprietary	Trade Secret #2	0-50
Proprietary	Trade Secret #3	0-30
Proprietary	Trade Secret #4	0-30
Proprietary	Trade Secret #5	0-20

* * * Section 4 - First Aid Measures * * *

First Aid: Eyes

In case of contact, do not rub or scratch your eyes. Flush thoroughly with water for 15 minutes to remove particles. If irritation persists, consult physician.

First Aid: Skin

Wash with mild soap and water. A commercially available hand lotion may be used to treat dry skin areas. If skin has become cracked, take appropriate action to prevent infection and promote healing. If irritation persists, consult physician.

First Aid: Ingestion

This product is not intended to be ingested or eaten. May be harmful or fatal if swallowed. If ingested, call physician.

First Aid: Inhalation

Remove to fresh air. Leave the area of dust exposure and remain away until coughing and other symptoms subside. Other measures are usually not necessary, however if conditions warrant, contact physician.

Material Safety Data Sheet

Material Name: OxEA-aq

*** Section 5 - Fire Fighting Measures ***

General Fire Hazards

See Section 9 for Flammability Properties.

Not expected to burn.

Hazardous Combustion Products

Not Determined

Extinguishing Media

Use extinguishing media appropriate for surrounding fire.

Fire Fighting Equipment/Instructions

Firefighters should wear full protective gear.

NFPA Ratings: Health: 1 Fire: 0 Reactivity: 1 Other: Oxidizer

Hazard Scale: 0 = Minimal 1 = Slight 2 = Moderate 3 = Serious 4 = Severe

*** Section 6 - Accidental Release Measures ***

Containment Procedures

Wear appropriate personal protection. Stop the flow of material if this is without risk.

Clean-Up Procedures

Use normal clean up procedures. Wear appropriate protective equipment. Ventilate area. If dry, shovel or sweep up material from spillage and place collected material into a container for recovery or waste disposal. Avoid dust generation. Avoid inhalation of dust and contact with eyes and skin. Maintain proper ventilation. If vacuum is used to collect dust, use an industrial vacuum cleaner with a high efficiency air filter. If sweeping is necessary, use dust suppressant. Do not use compressed air for clean up. These procedures will help minimize potential exposures. Dispose in accordance with all local, state, provincial and federal regulations.

Evacuation Procedures

Isolate area. Keep unnecessary personnel away.

Special Procedures

None

*** Section 7 - Handling and Storage ***

Handling Procedures

Minimize dust generation and accumulation. Avoid breathing dust. Wear the appropriate respiratory protection against dust in poorly ventilated areas and if exposure limits are exceeded (see Sections 8). Avoid dust contact with eyes. Wear the appropriate eye protection against dust (See Section 8). Use good safety and industrial hygiene practices.

Storage Procedures

Store at room temperature in a dry location. Keep containers closed when not in use

*** Section 8 - Exposure Controls / Personal Protection ***

Engineering Controls

If user operations generate airborne dust, use ventilation to keep dust concentrations below permissible exposure limits. Where general ventilation is inadequate, use process enclosures, local exhaust ventilation, or other engineering controls to control dust levels below permissible exposure limits.

PERSONAL PROTECTIVE EQUIPMENT

Personal Protective Equipment: Eyes/Face

Wear eye protection (safety glasses or goggles) to avoid particulate irritation of the eye.

Personal Protective Equipment: Skin

Gloves or protective clothing are usually not necessary but may be desirable in specific work situations. For brief contact, no precautions other than clean body-covering clothing should be needed. Wear gloves and protective clothing to prevent repeated or prolonged skin contact. Barrier creams or skin lotion may be applied to face, neck, wrist and hands when skin is exposed to help prevent drying of skin.

Personal Protective Equipment: Respiratory

Use NIOSH approved respiratory protection if airborne dust is generated.

Personal Protective Equipment: General

None

*** Section 9 - Physical & Chemical Properties ***

Appearance:	White to off white	Odor:	I ow to no odo
Physical State:	Solid (powder)	pH:	ca. 6.5-7.0
Vapor Pressure:	NA	Vapor Density:	NA
Boiling Point:	NA	Melting Point:	NA
Solubility (H2O):	32 g/100 g solution	Specific Gravity:	2.3-2.5
Evaporation Rate:	NA	VOC:	0
Octanol/H2O Coeff.:	ND	Flash Point:	None Known
Flash Point Method:	NA	Upper Flammability Limit	NA
		(UFL):	
Lower Flammability Limit	NA	Burning Rate:	NA
(LFL):			
Auto Ignition:	NA		

*** Section 10 - Chemical Stability & Reactivity Information ***

Chemical Stability

This is a stable material under ordinary conditions of use and storage.

Chemical Stability: Conditions to Avoid

Heat, flames, ignition sources, and contact with incompatibles.

Incompatibility

Oxidizing and reducing agents, metals, metal hydrides, alkalis, moisture, strong acids, strong bases, and magnesium. Keep away from flammable and combustible substances.

Hazardous Decomposition

Not Determined

Possibility of Hazardous Reactions

Will not occur.

*** Section 11 - Toxicological Information ***

Acute Dose Effects

A: General Product Information

No information available for the product.

B: Component Analysis - LD50/LC50

Trade Secret #1 (Proprietary)

Oral LD50 Rat 3015 mg/kg

Trade Secret #2 (Proprietary)

Oral LD50 Rat 1267 mg/kg

Trade Secret #3 (Proprietary)

Oral LD50 Rat 6500 mg/kg; Dermal LD50 Rabbit >7950 mg/kg

Trade Secret #4 (Proprietary)

Oral LD50 Rat 2000 mg/kg

Carcinogenicity

A: General Product Information

No information available for the product.

B: Component Carcinogenicity

None of this product's components are listed by ACGIH, IARC, OSHA, NIOSH, or NTP.

*** Section 12 - Ecological Information ***

Ecotoxicity

A: General Product Information

No information available for the product.

Material Safety Data Sheet

Material Name: OxEA-aq

B: Component Analysis - Ecotoxicity - Trade Secret #2 (Proprietary) Test & Species	Aquatic Toxicity	Conditions
96 Hr LC50 Lepomis macrochirus 96 Hr LC50 Oncorhynchus mykiss	2000 mg/L [static] 994.4-1107 mg/L [static]	
Trade Secret #3 (Proprietary) Test & Species 96 Hr LC50 Oncorhynchus mykiss 96 Hr LC50 Oncorhynchus mykiss 96 Hr LC50 Pimephales promelas 96 Hr LC50 Pimephales promelas	26.5 mg/L 24.8-29.4 mg/L [flow-through] 3.3 mg/L 33 mg/L [static]	Conditions
Test & Species96 Hr LC50 Leuciscus idus96 Hr LC50 Brachydanio rerio96 Hr LC50 Brachydanio rerio96 Hr LC50 Brachydanio rerio96 Hr LC50 Cyprinus carpio96 Hr LC50 Pimephales promelas96 Hr LC50 Oncorhynchus mykiss96 Hr LC50 Poecilia reticulata	460 - 1000 mg/L [static] 250 mg/L 480 mg/L [flow- through] 420 mg/L [semi- static] 18 mg/L >100 mg/L 32.2-41.9 mg/L [flow-through] 5.2-8.2 mg/L [static] 123-128 mg/L [semi- static]	Conditions
96 Hr LC50 Poecilia reticulata 24 Hr EC50 Daphnia magna 48 Hr LC50 Daphnia magna	126 mg/L 423 mg/L 14 mg/L	

*** Section 13 - Disposal Considerations ***

US EPA Waste Number & Descriptions

Component Waste Numbers

No EPA Waste Numbers are applicable for this product's components.

Disposal Instructions

All wastes must be handled in accordance with local, state and federal regulations.

See Section 7 for Handling Procedures. See Section 8 for Personal Protective Equipment recommendations.

*** Section 14 - Transportation Information ***

US DOT Information

Shipping Name: Not Regulated

*** Section 15 - Regulatory Information ***

US Federal Regulations

Component Analysis

None of this products components are listed under SARA Section 302 (40 CFR 355 Appendix A), SARA Section 313 (40 CFR 372.65), or CERCLA (40 CFR 302.4).

Material Safety Data Sheet

Material Name: OxEA-aq

State Regulations

Component Analysis - State

The following components appear on one or more of the following state hazardous substances lists:

Component	CAS	CA	MA	MN	NJ	PA	RI
Trade Secret #1	Proprietary	No	Yes	No	Yes	Yes	Yes
Trade Secret #2	Proprietary	No	Yes	No	No	Yes	Yes
Trade Secret #4	Proprietary	No	Yes	No	No	Yes	Yes

Component Analysis - WHMIS IDL

The following components are identified under the Canadian Hazardous Products Act Ingredient Disclosure List:

Component	CAS #	Minimum Concentration
Trade Secret #2	Proprietary	1 %
Trade Secret #4	Proprietary	1 %

Additional Regulatory Information

Component Analysis - Inventory

Component	CAS #	TSCA	CAN	EEC
Trade Secret #1	Proprietary	Yes	DSL	EINECS
Trade Secret #2	Proprietary	Yes	DSL	EINECS
Trade Secret #3	Proprietary	Yes	DSL	EINECS
Trade Secret #4	Proprietary	Yes	DSL	EINECS

*** Section 16 - Other Information ***

Other Information

The information herein is presented in good faith and believed to be accurate as of the effective date given. However, no warranty, expressed or implied, is given. It is the buyer's responsibility to ensure that its activities comply with Federal, State or provincial, and local laws.

Key/Legend

EPA = Environmental Protection Agency; TSCA = Toxic Substance Control Act; ACGIH = American Conference of Governmental Industrial Hygienists; IARC = International Agency for Research on Cancer; NIOSH = National Institute for Occupational Safety and Health; NTP = National Toxicology Program; OSHA = Occupational Safety and Health Administration., NJTSR = New Jersey Trade Secret Registry, DSL= Domestic Substance List, EINECS= European Inventory of Existing Commercial Chemical Substances

MATERIAL SAFETY DATA SHEET

This MSDS complies with OSHA'S Hazard Communication Standard 29 CFR 1910.1200 and OSHA Form 174				74			
	NTITY AND MAN	UFA	TURER'S I	NFORM	ATION		
NFPA Rating: Health-1; Flammability-0	; Reactivity-0; Special		HMIS Rating: Heal	th-1; Flammab	ility-0; Reactiv	ity-0; Personal	Protection-D
Manufacturer's Name:	ETEC, LLC		DOT Hazard (Classificati	on: None		
Address	6635 NE 59 th Place		TRADE NAM	F• EZ′	Τ-Δ2		
	Portland, OR 97218	8					
Date Prepared: 2/1/07	Prepared By: BT		MSDS Numbe	er: 010		Revision	- 2
Information Calls:	(503) 260-3799		NOTICE: JUDG	MENT BAS	SED ON IN	DIRECT TE	ST DATA
SECTIO	N 1 - MATERIAL II	DENTI	FICATION A	AND INFO	ORMATI	ON	
COMPONENTS-CHEMICAL NAM	IES AND COMMON NAME	ES	ACS Number	SARA	OSHA PEL	ACGIH	Carcinogen
(Hazardous Components 1% or greater	; Carcinogens 0.1% or greater	r)	NI/A	III LIST No	(ppm)	TLV (ppm)	Ref. Source **
Bacterial Consortium			IN/A	NO	N/E	IN/E	IN/E
SEC	TION 2 - PHYSICAI	CHE	MICAL CHA	RACTEI	RISTICS		
Boiling Point:	212. F		Specific Gravity ($H_2O=1$):			1.00
Vanor Pressure · PSIC @ 70 F (Aer.	nsols): N/A		Vanor Pressure (N	International States	(mm Hg and	Temnerature	1.00
Vapor Density (Air = 1):	62		Evanoration Rate	(BIJAC = 1)	·	remperature	1.00
Solubility in Water:	Complete		Water Reactive:	(Belle = 1)	•		No
Annearance and Odor	Thin brown / tan liquic	1 with slig	shtly sour odor				110
	CTION 3 - FIRE AN		PI OSION H	AZARDI	лтл		
FLAMMABILITY as per USA flame	projection test (aerosols). N	I/A Auto	Dignition Temperat	ure: N/A Fla	mmability Lir	nits in Air by	% in Volume:
FLASH POINT AND METHOD US	ED N/A		- 8		LEL: N/A	∧ %UF	EL: N/A
EXTINGUISHER MEDIA: Non-Co	mbustible	SPE	CIAL FIRE FIGH	ITING PROC	CEDURES: N	None // Cl	
Unusual Fire & Explosion Hazards:	None						
	SECTION 4 - RE	ACTI	νίτν μαζα	RDDATA			
STABILITY: I XI STABLE I	1 UNSTABLE	HAZAR	POUS POLYME	RIZATION:		XIWIIIN	OT OCCUR
Incompatibility (Mat to avoid): Not	ne Identified	Conditi	ans to Avoid Non	e Identified		A J WILL I	OTOCCOR
Hazardous Decomposition Products	• None	Contaiti					
riazardous Decomposition r roducts	SECTION 5 - I	JEAT'	гн на 7арг	рата			
PRIMARY ROLITES OF ENTRY-	[] INHALATION [X]]	ILAL	ON [X]SKINA	BSORPTION	J [X] FYF		AZARDOUS
ACUTE EFFECTS: None							
Inhalation: None							
Eve Contact: May be an irritant Shin Contact: May be an irritant							
Ingestion: Gastrointe	stinal irritant		phill contact				
CHRONIC EFFECTS: None kno	wn.						
Medical Conditions Generally Ager	avated by Exposure:	None Ide	entified				
SE	CTION 6 . EMERG	ENCY	FIRST AID	PROCED	URES		
Eve Contact: Flush with water for 14	minutes If irritation persists	s get medi	ical attention	INCOLD	UKLD		
Skin Contact: Wash with soap and w	ater.	, get mea					
Inhalation: Move to fresh air No.	adverse effects noted.						
Ingestion: DO NOT INDUCE V	OMITING. Drink large quar	ntity of wa	ater. Get immediate	e medical atter	ntion.		
SEC	TION 7 - CONTRO) PROTECT	IVE MEA	SURES		
Respiratory Protection (specify type	: None normally needed				beneb		
Protective Gloves:	Rubber if desired		Eve Prote	tion: Safet	v glasses.		
Ventilation Requirements: Normal room ventilation							
Other Protective Clothing & Equipment: Apron and boots if desired							
Hygienic Work Practices: Wash with soap and water after contact.							
SECTION 8 - PRECAUTIONS FOR SAFE HANDLING AND USE							
Stens To Be Taken If Material Is Spilled Or Released: Dike and contain Collect and re-use							
Waste Disposal Methods: Rinse container with water and dispose of accordingly							
Precautions To Be Taken In Handling & Storage: Recommended storage temperature is 50 F. Seal container after use.							
Other Precautions &/or Special Hazards: KEEP OUT OF REACH OF CHILDREN.							

We believe the statements, technical information and recommendations contained herein are reliable, but they are given without warranty or guarantee of any kind. ** Chemical Listed as Carcinogen or Potential Carcinogen. [a] NTP [b] IARC Monograph [c] OSHA [d] Not Listed [e] Animal Data Only

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NFPA Rating: Health-1: Elammability-0	· Reactivity-0: Special		MIS Rating: Healt	h-1. Flammah	ility-0. Reactiv	vity-0: Personal	Protection-G
Manufacturer's Name:	FTEC IIC	1	DOT Hazard ('lassificati	on• None	ity o, Tersona	
	6635 NE 59 th Place			Jassincau			
Address:	Portland, OR 97218	r	FRADE NAM	E: En	zyme Acc	elerator	
Date Prepared: 2/1/07	Prepared By: BT	l	MSDS Numbe	er: 011	Revisio	n - 0	
Information Calls:	(503) 260-3799	1	NOTICE: JUDG	MENT BA	SED ON IN	DIRECT TE	EST DATA
SECTIO	N 1 - MATERIAL IDE	NTI	FICATION A	ND INFO	ORMATI	ON	
COMPONENTS-CHEMICAL NAM	IES AND COMMON NAMES		ACS Number	SARA	OSHA PEL	ACGIH	Carcinogen
(Hazardous Components 1% or greater	; Carcinogens 0.1% or greater)		5000 05 5	III LIST	(ppm)	TLV (ppm)	Ref. Source **
MONOCYCLIC TERPENE			5989-27-5	No	N/E	N/E	d
NATURAL ENZYMES			N/A		N/E	N/E	N/E
SEC.	TION 2 - PHYSICAL/C		MICAL CHA	RACTE	RISTICS		1.00
Boiling Point:	220 F		Specific Gravity (H	20=1):	II 17		1.00
Vapor Pressure: PSIG @ 70 F (Aerose	DIS): N/A		Vapor Pressure (No	n-Aerosols)(r	nm Hg and Te	emperature):	18 @ /5° F
Vapor Density (Air = 1):	.62	1	Evaporation Rate (BUAC = 1):			1.20
Solubility in water:		<u> </u>	water Reactive:				NO
Appearance and Odor:	Tan colored liquid with cit	trus od	or.				
	CTION 3 - FIRE AND) EX	PLOSION H		JAIA mmahility Lis	nite in Ainhr	0/ in Volumou
FLAMMABILITY as per USA flame	e projection test (aerosols): N/A	Auto	Ignition Temperat	ure: N/E Fla		nits in Air dy	% in volume:
EXTINCUISHED MEDIA: Non Co	mbustible	SDF	CIAL FIDE FICH	TINC PRO	LEL: IN/E	Lone	L: IN/E
Laugual Eiro & Europian Haganda	Drovido shielding to protect pore		PECIAL FIRE FIGHTING PROCEDURES: None				
Unusual Fire & Explosion Hazarus:	SECTION 4 DEA		717777 11 4 77 4 1				
	SECTION 4 - KEAU					X 1 X / I X / I X X	OTOCOUD
STABILITY: [X] STABLE [J UNSTABLE HA	AZAR	DOUS POLYMEI	RIZATION:	[] WILL	X J WILL N	OFOCCUR
Incompatibility (Mat. to avoid): Not	ne Identified	onaitio	ons to Avoid: None	e Identified			
Hazardous Decomposition Products				DATA			
DIMARY DOUTES OF ENTRY.	SECTION 5 - HE		H HAZARD		N I VIEV	P	
ACUTE EFEECTS: None		GESTI		ADSOKPTIO	IN [A]EI	E	
Inhalation: Can cause	haadacha dizzinass						
Innatation: Can cause headache, dizziness. Fue Contact: Invitation							
Ingestion: Chemical	nneumonitis if aspirated into lung		Skii Contact	•			
CHRONIC EFFECTS: None know	wn	50.					
Medical Conditions Generally Agar	avated by Exposure: As	sthma					
SF	CTION 6 - EMERGEN		FIRST AID	PROCED	URES		
Eve Contact: Flush with water for 14	minutes. Get medical attention		FIRST AID	KOCED			
Skin Contact: Wash with soap and w	ater.						
Inhalation: Move to fresh air.							
Ingestion: DO NOT INDUCE V	OMITING. Drink large quantity	v of wa	ter. Get immediate	e medical atter	ntion.		
SECTION 7 - CONTROL AND DEATEOTIVE MEASURES							
Respiratory Protection (specify type	: None normally needed						
Protective Gloves:	Solvent resistant		Eve Protec	tion: Safet	v glasses.		
Ventilation Requirements:	Normal room ventilation			Suret	,		
Other Protective Clothing & Equip	nent: None						
Hygienic Work Practices:	Wash with soap and water	after c	ontact.				
SECTION 8 - PRECAUTIONS FOR SAFE HANDLING AND USE							
Steps To Be Taken If Material Is Spilled Or Released: Cover with absorbent material and scoop up. Flush area with water							
Waste Disposal Methods: Dispose of in accordance with local, state and federal regulations.							
Precautions To Be Taken In Handling & Storage: Keep away from temperatures above 130 F.							
Other Precautions &/or Special Hazards: KEEP OUT OF REACH OF CHILDREN.							

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MSDS NUMBER: 100609-01

MATERIAL SAFETY DATA SHEET DECON-IT All-Purpose Cleaner

SECTION 1: CHEMICAL PRODUCT AND COMPANY IDENTIFICATION

Chemical Name: D Chemical Name: N Chemical Family: N Formula: N Synonym(s): D

DECON-IT All-Purpose Cleaner Not Applicable (mixture) Non-ionic Surfactants Not Applicable (mixture) DECONIT

COMPANY IDENTIFICATION United States:

Enviro Supply & Services Inc.	1791 Kaiser Ave., Irvine, California, United States 92614
Prepared By:	Technical Products Department (Ivey International Inc.)
Telephone Number:	+ 1 250 923 6326 or Toll Free + 1 800 246 2744
Prepared (Last Updated):	June 11, 2010

Ivey International Inc. (IVEY) urges each customer or recipient of this MSDS to study it carefully to become aware of and understand the proper use and handling of the subject product. The reader should consider consulting reference materials, and/or IVEY technical support personnel, and/or other recognized experts, as necessary or appropriate to the use and understanding of the data contained in this MSDS. To promote the safe handling, storage and use of this product, each customer or recipient should (1) notify his employees, agents, contractors, and others whom he knows or believes will use this product, of the information in this MSDS and any other information regarding product use, storage and handling, (2) furnish this same information to each of his customers for the product, and (3) request his customers to notify their employees, customers, and other users of the product, and of this information.

SECTION 2: COMPOSITION INFORMATION

Components:	DECONT
Generic Description: DECON-IT [™] Technology-Stor listed.	Water based biodegradable wetting agents and surfactants. ck Mixtures. Patented and/or proprietary blends. Information in this MSDS is applicable for all products
SECTION 3: HAZ	ZARDS IDENTIFICATION
Effects of a Single Expo	DSURE
Swallowing:	Non to slightly toxic. May cause abdominal discomfort and nausea for some individuals.
Skin Absorption:	No evidence of harmful effects
Inhalation:	No evidence of harmful effects
Skin Contact:	Brief contact should not result in any significant effects. Prolong exposure may cause mild irritation with local itching and redness for individuals with sensitive skin
Eye Contact:	May cause mild to moderate initiation
Effects of Repeated Exp	Dosure: Repeated skin contact may course mild the sub-
Medical Conditions:	Existing dermatitis may be aggrevated through some light dermatitis (dryness of skin).
Section 4: FIRST	AID MEASURES
Swallowing:	If patient if fully conscious, give two classes of the
Skin Absorption:	Wash exposed skin with soap and water Other in the line
Inhalation: Eve Contact:	dermatitis persists. Wash exposed clothing before reuse. Not applicable.
-/	introductely flush eyes with water and continue to flush as required. By

any contact lenses, if worn. Obtain medical attention if deemed necessary.

Note to Physician: There is no required antidote. Treatment should be directed at the control of symptoms and the clinical condition of the patient.

Section 5: FIRE FIGHTER MEASURES

Flammability:	Not Flammable
Auto Ignition Temp.	Not Available
Upper Flammable Limit	Not Established
Lower Flammable Limit	Not Established
Explosive Date:	Explosive Power - Not Available
	Rate of Burning - Not Available
Hazardous Combustion Products:	Not applicable
Special Protective Equipment:	Not Applicable
Extinguishing Media:	Not Applicable
Extinguishing Media To Be Avoided:	Not Applicable
Special Fire Fighting Procedures:	Not Applicable
Section & ACCIDENTAL DE	LEACE MEACUIDEC

Section 6: ACCIDENTAL RELEASE MEASURES

Step to be Taken if Material is Released or Spilled:

d: Eliminate and/or contain source with inert material (sand, earth, absorbent pads, etc.). Wear basic eye and skin protection. Floor may be slightly slippery; so use care to avoid falling. Avoid discharge to natural waters, and/or dilute with water. Transfer liquids to suitable containers for recovery, re-use or disposal. Contact IVEY for technical assistance if required.

Section 7: HANDLING AND STORAGE

Handling Procedures:Avoid contact with eyes, skin, and clothing. Do not swallow. Keep containers
closed or sealed when not in use. Wash thoroughly after handling.Storage:Keep closed or sealed when not in use. Do not allow to freeze.

Ventilation: General mechanical room ventilation is expected to be satisfactory.

Section 8: EXPOSURE CONTROLS/PERSONAL PROTECTION

Latex, or similar would be sufficient.
None expected to be needed. However, if an engineered /
industrial application where vapors and/or misting may occur,
wear MSHA/NIOSH approved half mask air purifying respirator.
Mono Goggles or similar.
No special requirements.
Wear an apron and /or coveralls.
Eye bath.
General mechanical room ventilation is expected to be satisfactory.

Section 9: PHYSICAL AND CHEMICAL PROPERTIES

Physical state:	Water Based Liguid
Appearance:	Clear to slightly Cloudy White Color
Odor:	Mild
Molecular Weight:	Mixture (Not Applicable)
Boiling Point:	Not Applicable
Freezing Point:	Around 0°C (32 °F)
Pour Point:	Not Applicable
Melting Point:	Not Applicable
Specific Gravity:	0.99-1.04 (Water = 1.0)
Vapor Pressure:	<0.01 mm Hg
Vapor Density:	> 1 (Air = 1.00)
pH:	Not Available (Typically 6.5-7.5 Range)
Solubility In Water:	100%
Evaporation Rate:	<0.01
Coefficient of Oil/Water Distribution	Not Determined

Section 10: STABILITY AND REACTIVITY

CL . 1. 191		
Stability	; h_ h	Stable
Conditions to Avoid:		Prolonged excessive heat may cause product decomposition.
		Freezing should also be avoided as it may cause product
Income	tible Materials,	decomposition. In some cases it may cause irreversible changes.
mcompa	audie Materials;	Normally un-reactive; however avoid strong bases at high
		temperatures, strong acids, strong oxidizing agents, and
		materials with reactive hydroxyl compounds. These compounds
		would damage the mixture and reduce its effectiveness during
Hazardo	us Decomposition Brody	application.
Hazardo	us Polymerization	A will not applicable.
Sactio		
Secuo		GICAL INFORMATION
Exposure	e limit of Material:	Not Established
LD/50:		Not Available
LC/50:		Not Available
		Not Established
Carcinog	jenicity of Material:	None Known
Reproductive Effects:		Not Available
Irritancy of Material:		See Section 3
Sensitizing Capability:		Not Available
Synergis	tic Materials:	Not Available
LD: Letha	al Dose LC: Lethal Concen	tration EL: Exposure Limit
<u>Sectio</u>	n 12: ECOLOGIC	AL CONSIDERATIONS
Environn	nental Toxicity:	Low Potential to affect aquatic organisms*
Biodegra	idability:	>90% in 28 days**
LC/50:		48 Hour: 0.11 %, Species: Daphnia magna
LC/50:		96 Hour: 0.07695%, Species: Rainbow Trout
* \	When used in accordance	e with Ivey International Inc. In-situ and Ex-situ Remediation Application
(Guidelines.	
**	Based on actual testing	or on data for similar material(s). Degradation Biodegradation reached in
,	Medified OFCD Company	The Article The Article State and Article Stat

Modified OECD Screening Test (OECD Test No.301 E) after 28 days: 90 %. Biodegradation reached in CO2 Evolution Test (Modified Sturm Test, OECD Test No. 301 B) after 28 days: 70 %.

All available ecological data have been taken into account for the development of the hazard and precautionary information contained in this Material Safety Data Sheet.

Section 13: DISPOSAL CONSIDERATIONS

Waste Disposal Method: For aqueous DECON-IT mixture solutions; aerobic biological wastewater treatment systems are effective in treating said mixtures. DECON-IT does not have any known negative affect on coagulant or flocculent water treatment processes.

Section 14: TRANSPORTATION INFORMATION

UN Number:Not ApplicableTDG Classification:Not RequiredShipping Name:DECON-ITPacking Group:Not ApplicableSpecial Shipping Instructions:Do not allow to freeze

Section 15: REGULATORY INFORMATION

WHMIS Classification:Not Controlled as per WHMIS Regulation.CPR Compliance:This product has been classified in accordance with the hazard criteria of
the CPR, and the MSDS contains all the information required by the CPR.CEPA Compliance:All ingredients of this product are listed on the DSL.

Ref:DECON-IT/MSDS/100609-01(Revised June 11, 2010)