CALIFORNIA REGIONAL WATER QUALITY CONTROL BOARD LOS ANGELES REGION

MONITORING AND REPORTING PROGRAM NO. 7104 for ELIXIR INDUSTRIES (CA0062537)

I. Reporting Requirements

A. Elixir Industries, (hereinafter Discharger) shall implement this monitoring program on the effective date of this Order. All monitoring reports shall be submitted quarterly and must be received by the Regional Board by the dates in the following schedule. All monitoring reports should be addressed to the Regional Board, Attention: <u>Information Technology Unit</u>. The first monitoring report under this Program is due by April 15, 2004.

Reporting Period	Report Due
January – March	April 15
April – June	July 15
July-September	October 15
October-December	January 15
Annual Summary Report	March 1

If there is no discharge during any reporting period, the report shall so state.

- B. The Discharger shall submit an annual summary report (for both dry and wet weather discharges), containing a discussion of the previous year's effluent and receiving water monitoring data, as well as graphical and tabular summaries of the data. The data shall be submitted to the Regional Board on hard copy and on a 3 ½ " computer diskette. Submitted data must be IBM compatible, preferably using EXCEL software. This annual report is to be received by the Regional Board by March 1 of each year following the calendar year of data collection.
- C. Each monitoring report shall contain a separate section titled "Summary of Non-Compliance" which discusses the compliance record and corrective actions taken or planned that may be needed to bring the discharge into full compliance with waste discharge requirements. This section shall clearly list all non-compliance with waste discharge requirements, as well as all excursions of effluent limitations.
- D. The Discharger shall inform the Regional Board well in advance of any proposed construction activity that could potentially affect compliance with applicable requirements.

II. Effluent Monitoring Requirements

- A. A sampling station shall be established for each point of discharge and shall be located where representative samples of that effluent can be obtained. The Discharger shall collect the effluent sample at the exit of the PACT reactor, prior to the effluent entering the receiving water (i.e., storm drain).
- B. This Regional Board shall be notified in writing of any change in the sampling stations once established or in the methods for determining the quantities of pollutants in the individual waste streams.
- C. Pollutants shall be analyzed using the analytical methods described in 40 CFR sections 136.3, 136.4, and 136.5 (revised May 14, 1999); or, where no methods are specified for a given pollutant, by methods approved by this Regional Board or the State Board. Laboratories analyzing effluent samples and receiving water samples shall be certified by the California Department of Health Services Environmental Laboratory Accreditation Program (ELAP) or approved by the Executive Officer and must include quality assurance/quality control (QA/QC) data in their reports. A copy of the laboratory certification shall be provided each time a new certification and/or renewal of the certification is obtained from ELAP.

The monitoring reports shall specify the analytical method used, the Method Detection Limit (MDL), and the Minimum Level (ML) for each pollutant. For the purpose of reporting compliance with numerical limitations, performance goals, and receiving water limitations, analytical data shall be reported by one of the following methods, as appropriate:

- 1. An actual numerical value for sample results greater than or equal to the ML; or,
- 2. "Detected, but Not Quantified (DNQ)" if results are greater than or equal to the laboratory's MDL but less than the ML; or,
- 3. "Not-Detected (ND)" for sample results less than the laboratory's MDL with the MDL indicated for the analytical method used.

Current MLs (Attachment A) are those published by the State Water Resources Control Board in the *Policy for the Implementation of Toxics Standards for Inland Surface Waters, Enclosed Bays, and Estuaries of California, March 2, 2000.*

D. Where possible, the MLs employed for effluent analyses shall be lower than the permit limitations established for a given parameter. If the ML value is not below the effluent limitation, then the lowest ML value and its associated analytical method shall be selected for compliance purposes. At least once a year, the Discharger

shall submit a list of the analytical methods employed for each test and associated laboratory QA/QC procedures.

The Regional Board, in consultation with the State Board Quality Assurance Program, shall establish a ML that is not contained in Attachment A to be included in the Discharger's permit in any of the following situations:

- 1. When the pollutant under consideration is not included in Attachment A;
- 2. When the Discharger and Regional Board agree to include in the permit a test method that is more sensitive than that specified in 40 CFR Part 136 (revised May 14, 1999);
- 3. When the Discharger agrees to use an ML that is lower than that listed in Attachment A;
- 4. When the Discharger demonstrates that the calibration standard matrix is sufficiently different from that used to establish the ML in Attachment A, and proposes an appropriate ML for their matrix; or,
- 5. When the Discharger uses a method whose quantification practices are not consistent with the definition of an ML. Examples of such methods are the USEPA-approved method 1613 for dioxins and furans, method 1624 for volatile organic substances, and method 1625 for semi-volatile organic substances. In such cases, the Discharger, the Regional Board, and the State Board shall agree on a lowest quantifiable limit and that limit will substitute for the ML for reporting and compliance determination purposes.
- E. Laboratory analyses all chemical, bacteriological, and toxicity analyses shall be conducted at a laboratory certified for such analyses by the California Department of Health Services Environmental Laboratory Accreditation Program (ELAP). A copy of the laboratory certification shall be submitted with the Annual Report.
- F. Water/wastewater samples must be analyzed within allowable holding time limits as specified in 40 CFR section 136.3. All QA/QC items must be run on the same dates the samples were actually analyzed, and the results shall be reported in the Regional Board format, when it becomes available, and submitted with the laboratory reports. Proper chain of custody procedures must be followed, and a copy of the chain of custody shall be submitted with the report.
- G. All analyses shall be accompanied by the chain of custody, including but not limited to data and time of sampling, sample identification, and name of person who performed sampling, date of analysis, name of person who performed analysis, QA/QC data, method detection limits, analytical methods, copy of laboratory

certification, and a perjury statement executed by the person responsible for the laboratory.

H. For parameters that both monthly average and daily maximum limits are specified and the monitoring frequency is less than four times a month, the following shall apply. If an analytical result is greater than the monthly average limit, the sampling frequency shall be increased (within one week of receiving the test results) to a minimum of once weekly, if possible, at equal intervals, until at least four consecutive weekly samples have been obtained, and compliance with the monthly average limit has been demonstrated. The Discharger shall provide for the approval of the Executive Officer a program to ensure future compliance with the monthly average limit.

III. Effluent Monitoring Program

A. The effluent monitoring program for the discharge of treated groundwater through Discharge Serial No. 001 (Latitude 33° 51' 00" and Longitude 118° 16' 75") is:

Constituent	Units	Sampling Frequency
Weekly Flow ^{1/}	gallons/week	wekly
Daily Average Flow ^{1/}	gallons/day	weekly
рН	S.U.	weekly
Temperature	°F	quarterly
Oil and grease	mg/L	quarterly
Total suspended solids	mg/L	quarterly
BOD	mg/L	quarterly
Settleable solids	ml/L	quarterly
Turbidity	NTU	quarterly
Lead ^{2/}	ì g/L	quarterly
Mercury ^{2/}	ì g/L	quarterly ^{3/}
Benzene	ì g/L	quarterly
Toluene	ì g/L	quarterly
Xylene	ì g/L	quarterly
Ethylbenzene	ì g/L	quarterly
Ethylene dibromide	ì g/L	quarterly
Vinyl chloride	ì g/L	quarterly
1,1-Dichloroethane	ì g/L	quarterly
1,2-Dichloroethane	ì g/L	quarterly

Constituent	Units	Sampling Frequency
Dichloroethylene	ì g/L	quarterly
1,1,1-Trichloroethane	ì g/L	quarterly
Trichloroethylene	ì g/L	quarterly
Tetrachloroethylene	ì g/L	quarterly
Ethanol	mg/L	quarterly
Methanol	mg/L	quarterly
Isopropanol	mg/L	quarterly
Butanol	mg/L	quarterly
Remaining Priority Pollutants (see page T-17)	ì g/L	semi-annually
Methyl tertiary butylether (MTBE)	ì g/L	semi-annually
Total petroleum hydrocarbons	ì g/L	semi-annually
Perchlorate	ì g/L	semi-annually
Toxicity-acute ^{4/}	% survival	annually
Toxicity-chronic ^{4/}	% survival	annually

- 1/ The Discharger shall measure flow using the flow meter on-site. The Discharger shall record a total flow at the conclusion of the week (i.e., Friday), which shall represent the weekly flow (gallons/week). The Discharger shall also calculate the daily average flow for each week by dividing the total weekly flow by the number of days in that week that the system was operational; this shall represent the daily average flow (gallons/day).
- 2/ Measured as total recoverable.
- 3/ If the pollutant is within the acceptable WQBEL (or non-detect) for 4 consecutive months, this frequency shall be reduced to quarterly. If the pollutant is detected after this reduction, the frequency shall revert back to monthly.
- 4/ Refer to Item IV.

The mass emission (in lb/day) for the discharge shall be calculated and reported using the limitation concentration and the actual flow rate measured at the time of discharge, using the formula:

 $m = 8.34 C_i Q$

where: m = mass discharge for a pollutant, lb/day $C_i = limitation$ concentration for a pollutant, mg/L Q = actual discharge flow rate, mgd

IV. Toxicity Monitoring Requirements

- A. Acute Toxicity Effluent Monitoring Program
 - The Discharger shall conduct acute toxicity tests on effluent grab samples by methods specified in 40 CFR Part 136 which cites USEPA's *Methods for Measuring the Acute Toxicity of Effluents and Receiving Waters to Freshwater and Marine Organisms*, Fifth Edition, October 2002, USEPA, Office of Water, Washington D.C. (EPA/821-R-02-012) or a more recent edition to ensure compliance in 100 % effluent.
 - 2. The fathead minnow, *Pimephales promelas*, shall be used as the test species for fresh water discharges and the topsmelt, *Atherinops affinis*, shall be used as the test species for brackish effluent. The method for topsmelt is found in USEPA's *Short-term Method for Estimating the Chronic Toxicity of Effluents and Receiving Waters to West Coast Marine and Estuarine Organisms*, Third Edition, October 2002 (EPA/821-R-02-014).
 - 3. In lieu of conducting the standard acute toxicity testing with the fathead minnow, the Discharger may elect to report the results or endpoint from the first 48 hours of the chronic toxicity test as the results of the acute toxicity test.
 - 4. Effluent samples shall be collected after all treatment processes and before discharge to the receiving water.
- B. Chronic Toxicity Effluent Monitoring Program
 - The Discharger shall conduct critical life stage chronic toxicity tests on 24-hour composite 100 percent effluent samples in accordance with EPA's Short Term Methods for Estimating the Chronic Toxicity of Effluents and Receiving Waters to Freshwater Organisms, Fourth Edition, October 2002 (EPA/21-R-02-013) or EPA's Short Term Methods for Estimating the Chronic Toxicity of Effluents and Receiving Waters to Marine and Estuarine Organisms, Third Edition, October

2002 (EPA/821-R-02-014).

2. Effluent samples shall be collected after all treatment processes and before discharge to the receiving water.

Test Species and Methods:

- a. The Discharger shall conduct tests as follows: with a vertebrate, an invertebrate, and a plant for the first three suites of tests. After the screening period, monitoring shall be conducted using the most sensitive species.
- b. Re-screening is required every 15 months. The Discharger shall re-screen with the three species listed above and continue to monitor with the most sensitive species. If the first suite of re-screening tests demonstrates that the same species is the most sensitive than re-screening does not need to include more than one suite of tests. If a different species is the most sensitive or if there is ambiguity then the Discharger shall proceed with suites of screening tests for a minimum of three, but not to exceed five suites.
- c. The presence of chronic toxicity shall be estimated as specified using West Coast marine organisms according to EPA's *Short-Term Methods for Estimating Chronic Toxicity of Effluent and Receiving Water to Fresh Water Organisms*, Fourth Edition, October 2002 (EPA-821-R-02-013).
- C. Quality Assurance
 - 1. Concurrent testing with a reference toxicant shall be conducted. Reference toxicant tests shall be conducted using the same test conditions as the effluent toxicity tests (e.g., same test duration, etc).
 - 2. If either the reference toxicant test or effluent test does not meet all test acceptability criteria (TAC) as specified in the test methods manuals (EPA/600/4-91/002 and EPA/821-R-02-014), then the Discharger must re-sample and re-test at the earliest time possible.
 - 3. Control and dilution water should be receiving water or laboratory water, as appropriate, as described in the manual. If the dilution water used is different from the culture water, a second control using culture water shall be used.

D. Accelerated Monitoring

- 1. If toxicity exceeds the limitations (as defined in Order No. R4-2003-0149, Sections I.B.3.a.i. and 1.B.3.b.i), then the Discharger shall conduct six additional tests, over a six-week period. The Discharger shall ensure that they receive results of a failing acute toxicity test within 24 hours of the close of the test and the additional tests shall begin within 3 business days of the receipt of the result. If the accelerated testing shows consistent toxicity, the Discharger shall immediately implement the Initial Investigation of the Toxicity Reduction Evaluation (TRE) Workplan.
- 2. If implementation of the initial investigation TRE Workplan indicates the source of toxicity (e.g., a temporary plant upset, etc.), then the Discharger may discontinue the Toxicity Identification Evaluation (TIE).
- 3. The first step in the initial Investigation TRE Workplan for downstream receiving water toxicity can be a toxicity test protocol designed to determine if the effluent from Discharge Serial No. 001 causes or contributes to the measured downstream chronic toxicity. If this first step TRE testing shows that the Discharge Serial No. 001 effluent does not cause or contribute to downstream chronic toxicity, using EPA' sShort Term Methods for Estimating the Chronic Toxicity of Effluents and Receiving Waters to Fresh Water Organisms, October 2002, (EPA-821-R-02-013). Then a report on this testing shall be submitted to the Board and the TRE will be considered to be completed. Routine testing in accordance with MRP No. 7104 shall be continued thereafter.
- E. Steps in TRE and TIE procedures:
 - Following a TRE trigger, the Discharger shall initiate a TRE in accordance with the facility's initial investigation TRE workplan. At a minimum, the Discharger shall use EPA manuals EPA/600/2-88/070 (industrial) or EPA/833B-99/002 (municipal) as guidance. The Discharger shall expeditiously develop a more detailed TRE workplan for submittal to the Executive Officer within 30 days of the trigger, which will include, but not be limited to:
 - a. Further actions to investigate and identify the cause of toxicity;
 - b. Actions the Discharger will take to mitigate the impact of the discharge and prevent the recurrence of toxicity;
 - c. Standards the Discharger will apply to consider the TRE complete and to return to normal sampling frequency; and,

- d. A schedule for these actions.
- 2. The following is a stepwise approach in conducting the TRE:
 - a. Step 1 Basic data collection. Data collected for the accelerated monitoring requirements may be used to conduct the TRE:
 - b. Step 2 Evaluates optimization of the treatment system operation, facility housekeeping, and the selection and use of in-plant process chemicals;
 - c. If Steps 1 and 2 are unsuccessful, Step 3 implements a TIE and employment of all reasonable efforts and using currently available TIE methodologies. The objective of the TIE is to identify the substance or combination of substances causing the observed toxicity;
 - d. Assuming successful identification or characterization of the toxicant(s), Step 4 evaluates final effluent treatment options;
 - e. Step 5 evaluates in-plant treatment options; and,
 - f. Step 6 consists of confirmation once a toxicity control method has been implemented.

Many recommended TRE elements parallel source control, pollution prevention, and storm water control program best management practices (BMPs). To prevent duplication of efforts, evidence of implementation of these control measures may be sufficient to comply with TRE requirements. By requiring the first steps of a TRE to be accelerated testing and review of the facility's TRE workplan, a TRE may be ended in its early stages. All reasonable steps shall be taken to reduce toxicity to the required level. The TRE may be ended at any stage if monitoring indicates there is no longer toxicity (or six consecutive chronic toxicity results are less than or equal to 1.0 TU_c).

- The Discharger may initiate a TIE as part of the TRE process to identify the cause(s) of toxicity. The Discharger shall use the EPA acute and chronic manuals, EPA/600/6-91/005F (Phase I)/EPA/600/R-96-054 (for marine), EPA/600/R-92/080 (Phase II), and EPA-600/R-92/081 (Phase III) as guidance.
- 4. If a TRE/TIE is initiated prior to completion of the accelerated testing schedule required by Part I.B.3.a.ii and Part I.B.3.b.ii of this permit, then the accelerated testing schedule may be terminated, or used as necessary in performing the TRE/TIE, as determined by the Executive Officer.

- 5. Toxicity tests conducted as part of a TRE/TIE may also be used for compliance, if appropriate.
- 6. The Board recognizes that toxicity may be episodic and identification of causes of and reduction of sources of toxicity may not be successful in all cases. Consideration of enforcement action by the Board will be based in part on the Discharger's actions and efforts to identify and control or reduce sources of consistent toxicity.
- D. Reporting
 - 1. The Discharger shall submit a full report of the toxicity test results, including any accelerated testing conducted during the month as required by this permit. Test results shall be reported as % survival with the discharge monitoring reports (DMR) for the month in which the test is conducted.
 - 2. If an initial investigation indicates the source of toxicity and accelerated testing is unnecessary, then those results also shall be submitted with the DMR for the period in which the investigation occurred.
 - a. The full report shall be submitted on or before the end of the month in which the DMR is submitted.
 - b. The full report shall consist of (1) the results; (2) the dates of sample collection and initiation of each toxicity test; (3) the acute toxicity average limit or chronic toxicity limit or trigger.
 - 3. Test results for toxicity tests also shall be reported according to the appropriate manual chapter on Report Preparation and shall be attached to the DMR. Routine reporting shall include, at a minimum, as applicable, for each test:
 - a. Sample date(s);
 - b. Test initiation date;
 - c. Test species;
 - d. End point values for each dilution (e.g., number of young, growth rate, percent survival);
 - e. NOEC value(s) in percent effluent;
 - f. IC₁₅, IC₂₅, IC₄₀ and IC₅₀ values in percent effluent;
 - g. TU_c values $\left(TU_c = \frac{100}{NOEC}\right)$;
 - h. Mean percent mortality (<u>+</u>standard deviation) after 96 hours in 100% effluent (if applicable);

- i. NOEC and LOEC values for reference toxicant test(s);
- j. C₂₅ value for reference toxicant test(s);
- k. Any applicable charts; and
- I. Available water quality measurements for each test (e.g., pH, D.O., temperature, conductivity, hardness, salinity, ammonia).
- 4. The Discharger shall provide a compliance summary, which includes a summary table of toxicity data from all samples collected during that year.

The Discharger shall notify by telephone or electronically, this Regional Board of any toxicity exceedance of the limit or trigger within 24 hours of receipt of the results followed by a written report within 14 calendar days of receipt of the results. The verbal or electronic notification shall include the exceedance and the plan the Discharger has taken or will take to investigate and correct the cause(s) of toxicity. It may also include a status report on any actions required by the permit, with a schedule for actions not yet completed. If no actions have been taken, the reasons shall be given.

V. Receiving Water Monitoring Requirements

A. To conduct reasonable potential analysis (RPA) receiving water monitoring data is required. RPA will determine: (1) if water quality-based effluent limitations for priority pollutants are required, and (2) to calculate effluent limitations, if required.

Receiving monitoring station shall be within 50 feet upstream from or near the discharge point (of storm drain) into Receiving Water (Dominguez Channel). The receiving water monitoring shall be conducted for the first two years on an annual basis. The two time annual monitoring of the receiving water shall be conducted at the same time as annual effluent monitoring of priority pollutants.

The receiving water monitoring for toxic pollutants are listed below:

Constituent	Units	Type of Sample	Monitoring Frequency
PH	Standard units	grab	annually
Hardness (as CaCO ₃)	mg/L	grab	annually
PAHs	μg/L	grab	annually
Antimony	μg/L	grab	annually
Arsenic	μg/L	grab	annually
Beryllium	μg/L	grab	annually
Cadmium	μg/L	grab	annually
Chromium (III)	μg/L	grab	annually
Chromium (VI)	μg/L	grab	annually

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Constituent	Units	Type of Sample	Monitoring Frequency
Lead	μg/L	grab	annually
Mercury	μg/L	grab	annually
Nickel	μg/L	grab	annually
Selenium	μg/L	grab	annually
Silver	μg/L	grab	annually
Thallium	μg/L	grab	annually
Zinc	μg/L	grab	annually
Cyanide	μg/L	grab	annually
Asbestos	μg/L	grab	annually
Acrolein	μg/L	grab	annually
Acrylonitrile	μg/L	grab	annually
Benzene	μg/L	grab	annually
Bromoform	μg/L	grab	annually
Carbon tetrachloride	μg/L	grab	annually
Chlorobenzene	μg/L	grab	annually
Chlorodibromomethane	μg/L	grab	annually
Chloroethane	μg/L	grab	annually
2-Chloroethylvinyl ether	μg/L	grab	annually
Chloroform	μg/L	grab	annually
Dichlorobromomethane	μg/L	grab	annually
1,1-Dichloroethane	μg/L	grab	annually
1,2-Dichloroethane	μg/L	grab	annually
1,1-Dichloroethylene	μg/L	grab	annually
1,2-Dichloropropane	μg/L	grab	annually
1,3-Dichloropropylene	μg/L	grab	annually
Ethylbenzene	μg/L	grab	annually
Methyl bromide	μg/L	grab	annually
Methyl chloride	μg/L	grab	annually
Methylene chloride	μg/L	grab	annually
1,1,2,2-Tetrachloroethane	μg/L	grab	annually
Tetrachloroethylene	μg/L	grab	annually
Toluene	μg/L	grab	annually
1,2-Trans-dichloroethylene	μg/L	grab	annually
1,1,1-Trichloroethane	μg/L	grab	annually
1,1,2-Trichloroethane	μg/L	grab	annually
Trichloroethylene	μg/L	grab	annually
Vinyl chloride	μg/L	grab	annually
2-Chlorophenol	μg/L	grab	annually
2,4-Dichlorophenol	μg/L	grab	annually
2,4-Dimethylphenol	μg/L	grab	annually

Constituent	Units	Type of Sample	Monitoring Frequency
2-Methyl-4,6-Dinitrophenol	μg/L	grab	annually
2,4-Dinitrophenol	μg/L	grab	annually
2-Nitrophenol	μg/L	grab	annually
4-Nitrophenol	μg/L	grab	annually
3-Methyl-4-Chlorophenol	μg/L	grab	annually
Pentachlorophenol	μg/L	grab	annually
Phenol	μg/L	grab	annually
2,4,6-Trichlorophenol	μg/L	grab	annually
Acenaphthene	μg/L	grab	annually
Acenaphthylene	μg/L	grab	annually
Anthracene	μg/L	grab	annually
Benzidine	μg/L	grab	annually
Benzo (a) Anthracene	μg/L	grab	annually
Benzo (a) Pyrene	μg/L	grab	annually
Benzo (b) Fluoranthene	μg/L	grab	annually
Benzo (g,h,l) Perylene	μg/L	grab	annually
Benzo (k) Fluoranthene	μg/L	grab	annually
Bis (2-Chloroethoxy) Methane	μg/L	grab	annually
Bis (2-Chloroethyl) Ether	μg/L	grab	annually
Bis (2-Chloroisopropyl) Ether	μg/L	grab	annually
Bis (2-Ethylhexyl) Phthalate	μg/L	grab	annually
4-Bromophenyl Phenyl Ether	μg/L	grab	annually
Butylbenzyl Phthalate	μg/L	grab	annually
2-Chloronapthalene	μg/L	grab	annually
4-Chlorophenyl Phenyl Ether	μg/L	grab	annually
Chrysene	μg/L	grab	annually
Dibenzo (a,h) Anthracene	μg/L	grab	annually
1,2-Dichlorobenzene	μg/L	grab	annually
1,3-Dichlorobenzene	μg/L	grab	annually
1,4-Dichlorobenzene	μg/L	grab	annually
3,3'-Dichlorobenzidine	μg/L	grab	annually
Diethyl Phthalate	μg/L	grab	annually
Dimethyl Phthalate	μg/L	grab	annually
Di-n-Butyl Phthalate	μg/L	grab	annually
2,4-Dinitrotoluene	μg/L	grab	annually
2,6-Dinitrotoluene	μg/L	grab	annually
Di-n-Octyl Phthalate	μg/L	grab	annually
1,2-Diphenylhydrazine	μg/L	grab	annually
Fluoranthene	μg/L	grab	annually
Fluorene	μg/L	grab	annually

Constituent	Units	Type of Sample	Monitoring Frequency
Hexachlorobenzene	μg/L	grab	annually
Hexachlorobutadiene	μg/L	grab	annually
Hexachlorocyclopentadiene	μg/L	grab	annually
Hexachloroethane	μg/L	grab	annually
Indeno (1,2,3-cd) Pyrene	μg/L	grab	annually
Isophorone	μg/L	grab	annually
Napthalene	μg/L	grab	annually
Nitrobenzene	μg/L	grab	annually
N-Nitrosodimethylamine	μg/L	grab	annually
N-Nitrosodi-n-Propylamine	μg/L	grab	annually
N-Nitrosodiphenylamine	μg/L	grab	annually
Phenanthrene	μg/L	grab	annually
Pyrene	μg/L	grab	annually
1,2,4-Trichlorobenzene	μg/L	grab	annually
Aldrin	μg/L	grab	annually
alpha-BHC	μg/L	grab	annually
beta-BHC	μg/L	grab	annually
gamma-BHC	μg/L	grab	annually
delta-BHC	μg/L	grab	annually
Chlordane	μg/L	grab	annually
4,4'-DDT	μg/L	grab	annually
4,4'-DDE	μg/L	grab	annually
4,4'-DDD	μg/L	grab	annually
Dieldrin	μg/L	grab	annually
Alpha-Endosulfan	μg/L	grab	annually
Beta-Endosulfan	μg/L	grab	annually
Endosulfan Sulfate	μg/L	grab	annually
Endrin	μg/L	grab	annually
Endrin Aldehyde	μg/L	grab	annually
Heptachlor	μg/L	grab	annually
Heptachlor Epoxide	μg/L	grab	annually
Polychlorinated Biphenyls ¹	μg/L	grab	annually
Toxaphene	μg/L	Grab	annually

The sum of Aroclors 1242, 1254, 1221, 1232, 1248, 1260, and 1016.

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- B. Please note that the report for this required monitoring must be submitted with the self-monitoring reports in accordance with the schedule provided in Section I.A of this *M&RP* No. CI-7104.
- C. SWRCB-approved laboratory methods and the corresponding MLs for the examination of each priority pollutant are listed in <u>Attachments A and B</u>. Reporting requirements for the data to be submitted are listed in <u>Attachment B</u>. We recommend that you select the analytical method from Attachment A capable of achieving the lowest ML for each pollutant. ML is necessary for determining compliance for a priority pollutant when an effluent limit is below the MDL.
- D. The laboratory analytical data shall include applicable MLs, MDL, quality assurance/quality control data, and shall comply with the reporting requirements contained in the Attachments B & C.

VI. Interim Monitoring and Reporting

A. Monitoring for TCDD Equivalents –The Discharger shall conduct effluent/receiving water monitoring for the presence of the 2,3,7,8-tetrachlorodibenzo-p-dioxin (TCDD or Dioxin) congeners. The monitoring shall be a grab sample with a minimum frequency of once during dry weather and once during wet weather for 1 year. The Discharger shall calculate Toxic Equivalence (TEQ) for each congener by multiplying its analytical concentration by the appropriate Toxicity Equivalence Factors (TEF). Compliance with the dioxin limitation shall be determined by the summation of the 17 individual TEQs.

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<u>Congeners</u>	TEF
2,3,7,8-tetra CDD	1.0
1,2,3,7,8-penta CDD	1.0
1,2,3,4,7,8-hexa CDD	0.1
1,2,3,6,7,8-hexa CDD	0.1
1,2,3,7,8,9-hexa CDD	0.1
1,2,3,4,6,7,8-hepta CDD	0.01
Octa CDD	0.0001
2,3,7,8-tetra CDF	0.1
1,2,3,7,8-penta CDF	0.05
2,3,4,7,8-penta CDF	0.5
1,2,3,4,7,8-hexa CDF	0.1
1,2,3,6,7,8-hexa CDF	0.1
1,2,3,7,8,9-hexa CDF	0.1
2,3,4,6,7,8-hexa CDF	0.1
1,2,3,4,6,7,8-hepta CDF	0.01
1,2,3,4,7,8,9-hepta CDF	0.01
Octa CDF	0.0001

Ordered by: ____

Dennis A. Dickerson Executive Officer Date: December 4, 2003