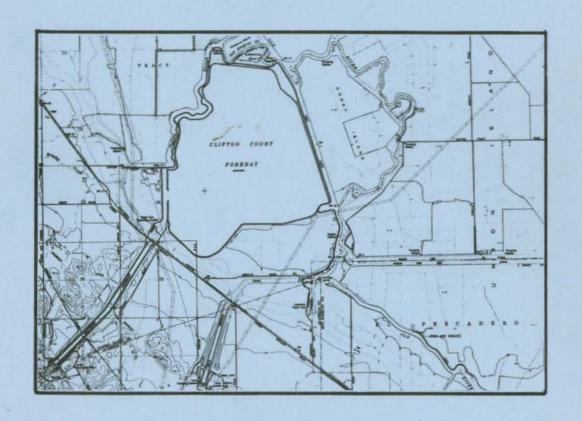
State of California The Resources Agency DEPARTMENT OF WATER RESOURCES Central District

## INTERAGENCY DELTA HEALTH ASPECTS MONITORING PROGRAM PROJECT REPORT



December 1986

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Central District

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December 1986



Portions of this report were prepared with the assistance of

Marvin Jung and Associates

Contract B-55923

Technical assistance provided in the amount of \$5,100

#### FOREWORD

In 1982, the Department of Water Resources appointed a panel of scientists to evaluate the human health aspects of using Delta water supplies. The panel concluded that there was insufficient data on many important factors and contaminant sources that could affect water quality. Some of these factors include tidal action and riverflows, agricultural drainages, pesticide use, waste water discharges, and water movement within the Delta. The panel recommended a program to develop a comprehensive analytical model that would incorporate and analyze these factors.

In April 1983, the Interagency Delta Health Aspects Monitoring Program was initiated in response to the panel's recommendation. The program is now in its fourth year of monitoring and examining the effects of natural and man-related events on the quality of Delta water supplies.

The 1986 Project Report describes program activities and presents findings for data collected between January 1985 and June 1986. Study results indicate that Delta water supplies are generally of acceptable quality with respect to the levels of chemical contaminants and minerals that may affect human health.

The program should continue to provide needed information on sources of degradation of Delta water supplies. The program's combined activities of monitoring and investigating water quality changes are invaluable to water resource planning and protection.

James U. McDaniel

Chief, Central District

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#### ACKNOWLEDGMENTS

The Department of Water Resources has been the lead agency conducting the Interagency Delta Health Aspects Monitoring Program and has been provided with a combination of funding and technical assistance from the Department of Health Services, City of Stockton, East Bay Municipal Utility District, U. S. Bureau of Reclamation, and water contractors of the State Water Project. A Technical Advisory Group has guided the program staff in setting project priorities and activities.

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#### SUMMARY

In 1982 the Department of Water Resources appointed a scientific advisory panel to examine human health aspects related to the use of water supplies from the Sacramento-San Joaquin Delta. The panel submitted a report with several long-term recommendations for monitoring and studying the water quality of the Delta. The panel had specific concerns about the effects from waste discharges, riverflow conditions, pesticides, ocean water intrusion, agricultural drainage, and water project operations. The Interagency Delta Health Aspects Monitoring Program was initiated in July 1983 in response to the panel report.

Several tasks were undertaken to meet specific concerns of the 1982 scientific advisory panel. Separate tasks were developed to:

- Monitor pesticide contaminants in water,
- Study effects of tides and riverflow on export water quality,
- Characterize and track water sources and movement,
- Identify and study agricultural drainages,
- Examine contributing sources of total trihalomethane formation potential,
- ° Examine waste discharges, and
- Test computer models that might help predict impacts from both point (e.g. sewage outfalls) and non-point (e.g. land runoff) sources of pollutants.

This project report of the monitoring program describes activities from January 1985 through June 1986 and presents current findings.

Monitoring data showed selenium, pesticides, and sodium levels in Delta water supplies are below drinking water standards or Department of Health Services action levels.

By examining the electrical conductivity and chloride to sodium ion ratios, water sources were identified. The data showed that the interaction of tidal excursions and Sacramento River flow during the summer had a significant effect on the quality of water exported by the Delta-Mendota Canal and Banks Pumping Plant facilities. During the last half of 1985, export water was predominantly a mixture of Sacramento River water blended with San Francisco Bay tide water. The effect of San Joaquin River water on export water quality was not apparent.

Agricultural drain water and irrigation return water could be significant sources of trihalomethane precursor material to Delta waters. They may also affect mineral content of receiving waters by increasing the concentration of salts. Efforts are in progress to quantify these loads to the Delta.

Discharge of treated effluent from the Sacramento Regional Wastewater Treatment Plant did not significantly affect the quality of water downstream at Hood and Greene's Landing. This may be attributed to the high level of treatment and dilution at the discharge site at Freeport.

A pesticide monitoring selection scheme was developed to target monitoring efforts. The scheme eliminated the approach of conducting expensive and numerous laboratory tests for chemicals with a low probability of detection in water. Emphasis was placed on siteand time-specific monitoring for chemicals in high use or with a relatively high potential of being carried by water.

Computer models developed by the U. S. Environmental Protection Agency are

being tested for predicting the effects of pollutants on Delta water quality. The models could improve monitoring efforts and identify additional types of measurements or studies needed to help predict the effects of natural and man-induced events on water quality in the Delta.

The Department will continue the program to meet the long-term objectives of the scientific advisory panel and the needs of its Technical Advisory Group.

#### Chapter 1. INTRODUCTION

This is the second project report of the Interagency Delta Health Aspects Monitoring Program. Five semiannual progress reports and a project report were published earlier.

The program began in July 1983; initial focus was on monitoring raw water supplies in the Delta for contaminants that could affect human health. The scope of work has expanded to collection of data on specific factors that can affect the water quality and quantity of exported water supplies. These factors include riverflows, agricultural related practices, and tidal movements.

These new activities were initiated to meet the recommended long-term objectives of a scientific advisory panel that investigated human health aspects of Sacramento-San Joaquin Delta water supplies. This panel was appointed by the Department of Water Resources because of concerns about the quality of raw water supplies diverted from the Delta for domestic use. Findings of the panel were submitted to the Department on December 31, 1982, in a report titled, Public Health Aspects of Sacramento-San Joaquin Delta Water Supplies /1/.

The panel had recommended the following long-term objectives to the Department:

1. Establish a monitoring program that identifies sources of contaminants to the Delta, how contaminants from each source are transported through the system, and how they affect concentration at points of withdrawal.

- 2. Obtain information on factors that affect the movement and fate of contaminants in the Delta.
- 3. With such information, develop a comprehensive analytical model to incorporate and analyze the following elements:
- Location and magnitude of sodium, asbestos, and organic material, including inflows to the Delta, agricultural drainage, waste water discharges, and ocean water intrusion.
- Factors affecting contributions from each important source such as riverflow, season, level of waste water treatment, and reservoir release patterns.
- Ovariability of constituent concentrations at critical points of the Delta as affected by sources and flow patterns.
- effects of Delta water quality, storage, transport, blending, and treatment on the quality of treated drinking water.
- 4. This model would provide information for making decisions on how to manage the water resources of the State.

Separate tasks are being performed to address some of the concerns expressed in the long-term objectives. These tasks are described in subsequent sections of the report and are summarized in Table 1.

#### Table 1

#### PROGRAM TASKS TO ADDRESS SPECIFIC CONCERNS

#### Effects from Tidal Excursions and Riverflows

- <u>Task WO1. Health Aspects Water Quality Monitoring</u>. Sampling is conducted monthly at key Delta locations for sodium, trihalomethane formation potential, minerals, and other parameters to determine if raw water supplies can be treated to meet drinking water standards and to identify potential treatment and human health problems.
- <u>Task WO2, Characterization of Water Sources</u>. Water sources are being characterized by comparing constituents at key Delta stations. The data will be used to help track general water movement and water quality trends in the Delta.
- Task WQ3. Tidal Effects Study on Exported Water Quality. During different summer tidal stages, the direction and mixing of water along Old and Middle rivers and other channels are being studied. The data will be used to help quantify the effects on water quality from different sources of water to the Clifton Court Forebay and Delta-Mendota Canal intakes during low Delta outflow and various tidal conditions.

#### Effects from Agriculture Related Activities

- Task AG1, Drainage Water Quality Monitoring. Irrigation return flows from drainages at Empire Tract,
  Grand Island, and Tyler Island are being monitored for salts, pesticides, trihalomethane (THM)
  formation potential, and other constituents. The data will be used to assess the loading effects
  of drainage on receiving water quality during the year.
- <u>Task AG2, Locating Irrigation Return Water Discharges</u>. Discharge points of irrigation return water on leveed Delta islands are being identified and mapped. This information will be used to identify sources of contaminants and plan upcoming work to assess the impact of agricultural drainages on Delta water quality.
- Task AG3, San Joaquin River Monitoring. Comprehensive water quality monitoring near Vernalis for total and dissolved trace inorganics, pesticides, and other constituents has been initiated to study the effects of San Joaquin River water quality on exported water. There is concern about selenium and other trace elements that are discharged into the San Joaquin River from agricultural drainage.
- Task AG4. Selected Pesticide Monitoring. Through a selection protocol based on pesticide usage patterns and environmental behavior, water samples are collected for specific pesticide analyses. The data are used to identify potential contamination problems for raw water supplies and treatment plants. Sampling is conducted more frequently during chemical application periods.
- Task AG5, Modeling Pesticide Fate and Transport. Existing computer models developed by the U.S.

  Environmental Protection Agency (EPA) to predict the fate and movement of organic pesticides in an aquatic system are being tested to help assess the threat of contamination to drinking water supplies. The models are used to evaluate the pesticide monitoring selection protocol for

Task AG4 and to study the effects of riverflow and other environmental conditions on the distribution of pesticide contaminants.

Task AG6, Health Effects Database on Selected Chemicals. Drinking water standards now exist for only a few pesticides. A computer literature search for human health effects data is underway for chemicals appearing on the selected pesticide monitoring task (AG4) for which there are no drinking water standards. The data will be used to assess the degree of risk to users of Delta water supplies found with traces of these contaminants.

#### Effects from Waste Water Discharges

<u>Task WD1, Survey Major Waste Water Dischargers</u>. The Central Valley Regional Water Quality Control Board provided effluent monitoring data on major municipal and industrial waste dischargers within the program study area. The data will be examined to estimate total waste loads and to determine if special studies on receiving water impacts are needed.

#### Effects of Raw Water Quality and Treatment

Task TR1, Assess THM (Trihalomethane) Formation Potential. As part of the monthly water quality monitoring activities (Task WQ1), parameters such as total organic carbon and color are measured. In addition, water samples are chlorinated and analyzed for total THM formation potential and THM species to identify potential THM treatment problems. Water quality parameters related to the extent of THM formation during disinfection are also being studied.

#### Modeling Water Quality in the Delta

Task MOD1. Use Existing Water Quality Models. Computer models developed by EPA to study the distribution, fate, and transport of waste waters and spilled materials are being tested for use in studying Delta water quality as affected by waste water discharges and pesticide usage. EPA recently made several models available for personal computer use. Models under study include EXAMS (exposure analysis modeling system), QUAL2E (a stream quality routing model), and WASP3P (a chemical transport and fate model).

### Chapter 2. FINDINGS

Results of the Interagency Delta Health Aspects Monitoring Program for 1985 through June 1986 showed:

- 1. Selenium concentrations in the Delta are meeting the (10 ug/L) drinking water standard. The highest concentrations have been observed in the lower San Joaquin River in Mud and Salt sloughs. Subsequent dilution and natural removal processes result in concentrations of 2 ug/L or less at the San Joaquin River near Vernalis. The data indicate that selenium does not constitute a health threat to consumers of Delta water supplies.
- 2. Pesticides concentrations have been far below Department of Health Services action levels or drinking water criteria. When found, the levels were barely above the analytical limit of detection (generally 1 ug/L or less). The data indicate a wide margin of safety in the drinking water quality with respect to harmful pesticide concentrations.
- 3. Irrigation return flow drainage can have major effects on water quality. Preliminary data indicate that drainage from Delta islands is a major contributing source of trihalomethane precursor materials and may have the most significant effect on the total trihalomethane formation potential of Delta water supplies exported by the State and Federal water projects.
- 4. Asbestos analyses of surface waters need to be improved to obtain reproducible results. Until the methodology is refined, asbestos data cannot be interpreted.

- 5. Sodium levels in Delta channels met the National Academy of Sciences recommended limit of 270 mg/L for persons on moderately restricted sodium diets. However, the levels exceeded the 20 mg/L limit for persons on severely restricted sodium diets. Persons on severely restricted sodium diets generally drink sodium-free water.
- 6. The quality of export water was significantly affected by Sacramento River flows and tidal influences during the last half of 1985. Comparisons of chloride and sodium ratios showed the direction and predominant source of water to the export pump intakes. Electrical conductivity measurements alone were insufficient "tracers" of water movement.
- 7. The quality of export water was reflective of Sacramento River water mixed with saline bay water. The effects of San Joaquin River quality and flows on export water were not detectable.
- 8. The drinking water quality of the Sacramento River downstream of the Sacramento Regional Wastewater Treatment Plant outfall does not appear to be greatly affected by the waste discharge.
- 9. The use of water quality models to study the fate and transport of constituents in surface waters and discharges may help predict water quality changes and improve monitoring effectiveness.

#### Chapter 3. RECOMMENDATIONS

The following recommendations are offered as a result of monitoring to date.

- 1. Efforts should be continued to meet the long-term objectives of a 1982 Department appointed scientific advisory panel that examined human health factors of Delta water supplies.
- 2. Monitoring possible effects of San Joaquin River flows and quality on export waters should continue in view of public concern over selenium, pesticides, and agricultural drainage constituents.
- 3. The potential effect of Delta island irrigation return waters on Delta water quality should be examined, as preliminary data suggest these drainages are major sources of trihalomethane precursors and may have the most important effect on the total

trihalomethane formation potential of Delta water exported by the State and Federal water projects.

- 4. The monitoring program and special tasks should be performed to meet the information requirements of computer water quality models developed to predict the effects on water quality from spills, waste discharges, project operations, and riverflow.
- 5. Standard mineral analyses should be included in the monitoring program to improve the characterization of water sources. Ionic ratios proved to be more useful than electrical conductivity measurements alone.
- 6. Asbestos monitoring should be discontinued until the analytical method for quantifying asbestos can provide confidence in the interpretation of results.

### Chapter 4. PROGRAM TASKS AND RESULTS

Department of Water Resources staff has been responsible for conducting the Interagency Delta Health Aspects Monitoring Program. Program activities are developed to meet the recommendations of a Technical Advisory Group and the long-term objectives recommended by the 1982 Department appointed scientific advisory panel.

Laboratory support is provided by the Department's Bryte Laboratory and through contractual agreement with McKesson Environmental Services in Pleasanton. The Bryte Lab conducted standard water quality measurements (e.g. conductance, mineral content), trihalomethane testing, and on occasion, pesticide analyses. McKesson laboratory primarily performed pesticide and priority pollutant analyses. Bromide and dissolved copper testing were also performed on request. Bromide and copper analyses were later stopped because detection limits were above sample concentrations. Performance of both laboratories was evaluated by duplicate sample splitting, internal quality control measurements, and spiked samples. Details are discussed in Appendix D, Laboratory Performance 1985-86.

This second project report presents findings and progress of various tasks associated with the program from January 1985 through June 1986. Activities prior to 1985 were reported in an earlier project report /2/.

### Water Quality and Tidal Effects Studies

Three tasks are underway to study the effects of tidal excursions and riverflows on Delta water quality.

## Health Aspects Water Quality Monitoring (Task WQ1)

Monthly sampling is conducted at key Delta locations for sodium, trihalomethane formation potential, minerals, and other parameters. The data are used to determine if raw water supplies are meeting drinking water standards and to identify potential treatment and human health problems. The study area and locations of key monitoring stations are shown in Figure 1.

Field measurements of conductivity, pH, dissolved oxygen, and temperature are made on site. Water samples are collected with a stainless steel Kemmerer-type sampler. Samples are appropriately treated and stored in clean containers provided by the laboratories for the type of analyses to be conducted. Glass bottles and vials are used for water samples undergoing pesticide, trihalomethane, and priority pollutant analyses. Plastic containers are used for standard mineral analyses. Samples undergoing pesticide analysis are delivered to the laboratories on the day of collection.

Results of the field and laboratory measurements are presented in Appendix A, Monitoring Program Data. Field sampling methodology is described in Appendix B, Field Sampling Procedures.

The National Academy of Sciences (NAS) has recommended a 270 mg/L limit in drinking water for persons on moderately restricted sodium diets and a 20 mg/L limit for those on severely restricted sodium diets. Sodium levels in all water samples were below the 270 mg/L limit except for agricultural

drain water samples taken from Empire Tract. Since agricultural drainage is not used as a drinking water supply, the NAS limit would not apply. Sodium levels were occasionally above the 20 mg/L limit at some stations; however, most persons on a severely restricted sodium diet use sodium-free bottled water.

The current drinking water standard for selenium is 10 ug/L. EPA is currently proposing a 45 ug/L drinking water criterion. Water samples collected from all stations had selenium concentrations at or below 2 ug/L. Most results were below the analytical detection limit of 1 ug/L.

Starting in 1985, measurement of water samples for asbestos was reduced to a sampling frequency of twice a year because the interpretative value is in question due to high variability in the data. Asbestos analyses done in triplicate on the same water samples differed significantly. Until improvements are made in the determination of asbestos in water, high confidence in asbestos data cannot be obtained.

Pesticide monitoring for a select group of chemicals meeting specific behavioral characteristics was also conducted. A complete description of the pesticide monitoring task is presented later in this report. In general, most pesticides monitored were below the analytical limit of detection (1 ug/L or less). Of those chemicals detected, trace amounts were found near the limit of detection.

Tests for trihalomethane formation potential and trihalomethane species that are formed when raw water samples are chlorinated were also conducted. These tests do not reflect the actual trihalomethane concentrations in finished (treated) drinking water available to the public. The tests were conducted to identify when and where modified water treatment operations may be necessary when water is

withdrawn from a specific area in the Delta.

Complete descriptions of the pesticide monitoring results and the trihalomethane studies are presented separately in this report.

## Characterization of Water Sources (Task WQ2)

Constituents are being compared to characterize water sources and mixing at key stations. Comparisons of electrical conductivity, major ion concentrations, and specific ion ratios by molarity and weight are some of the methods being used to follow general water movement and water quality trends in the Delta.

The quality of water exported by the Delta-Mendota Canal and the State Water Project is affected by a complex variety of sources and conditions. Primary water sources include fresh water of the Sacramento and San Joaquin rivers. These waters, in turn, are affected by agricultural drainage, sea water intrusion, waste discharges, and land runoff. The proportion of Sacramento and San Joaquin river waters entering the State and Federal water projects has been estimated by salinity measurements (electrical conductivity or total dissolved solids). However, salinity measurements may not accurately reflect water movement and mixing, as waters of similar salinity may differ significantly in ionic composition.

As a first step in studying water movement and quality changes in the Delta that affect the State and Federal water projects, the characteristics of water at the intakes and major channels leading to the intakes were examined. For January 1985 through June 1986, the data showed:

1. Exported waters, measured at the Banks Pumping Plant headworks and

Station Location Number Station Name Station Number Station Location Number Station Name Station Number

American River at Water Treatment Plant A0714010 (IO)

Clifton Court Intake KA000000

Sacramento River at Greene's Landing B9D82071327

Delta-Mendota Intake at Lindeman Road B9C74901336

Cache Slough at Vallejo
Pumping Plant
B9D81781448

(12)

H. O. Banks Delta Pumping Plant at Headworks KA000331

Lindsey Slough at
Hastings Cut
B9D81581462

(13)

Middle River at Borden Highway (Middle River at Highway 4 Bridge) B9D75351293

Agricultural Drain on Grand Island
B9V81171369

(14)

San Joaquin River near Vernalis B0702000

Agricultural Drain on
Tyler Island
B9V80801348

(15)

Lake Del Valle Stream Release DV004000

Little Connection Slough at Empire Tract (end of 8-Mile Road) B9D80361299

(16)

Mallard Slough at CCWDPP B8X80221556

Agricultural Drain on Empire Tract (west end of 8-Mile Road) B9V80361274

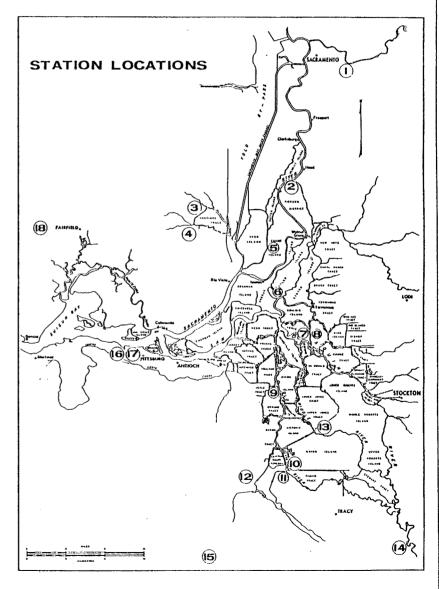
**(17)** 

Sacramento River at Mallard Island E0B80261551

Rock Slough at Old River B9D75841348

(18)

North Bay Interim Pumping Plant Intake KE000000



Delta-Mendota Canal intake, were higher in EC and chloride to sodium (Cl:Na) molar ratios during the last half of 1985 than in 1984 (Figure 2).

- 2. The rise and fall of EC and Cl:Na molar ratios at the Banks headworks and Delta-Mendota Canal intake corresponded with similar observations at the Rock Slough at Old River station and Middle River station (Figure 3).
- 3. The rise and fall of EC and C1:Na molar ratios at the Rock Slough and Middle River stations corresponded to that of the Sacramento River at Mallard Island station, which is subject to tidal excursion and bay salinity intrusion during low riverflows. Rock Slough station water was more affected by the Mallard Island water quality than was Middle River (Figure 4).
- 4. Observed water quality at the aforementioned stations corresponded to reduced Sacramento River flows, which were lower than flows during the same period in 1984. San Joaquin River flows were essentially unchanged from the previous year. (Figure 5). Molar chloride to sodium ratios indicated fairly constant composition of river water quality at Greene's Landing and Vernalis (Figure 5).
- 5. San Joaquin River EC measured near Vernalis resembled export water conductivity, but falsely suggested that the waters were similar in composition. The molar Cl:Na ratios differentiated between the water types during July 1985 through January 1986 (Figure 6). Quality of exported water was more similar to water flowing into the southern Delta through Old and Middle rivers. The value of using ionic ratios over salinity values was demonstrated in

this study of water movement and characterization.

Standard mineral analyses have been added to the list of laboratory determinations to be performed on water samples. Ionic ratios will also be analyzed to improve the ability to track water sources and changes with time. Results of recent mineral analyses are shown in Table 2.

### Study of Tidal Effects on Export Water Quality (Task WQ3)

The direction and mixing of water along Old and Middle rivers and other channels will be studied during different tidal stages. The data will be used to help quantify water quality effects of different sources of water to the Clifton Court and Delta-Mendota Canal intakes.

Three sampling runs were conducted during high slack tide (Figure 7). On August 7, Old River was sampled; on August 21, both Old River and Middle River were sampled; and on August 22, the east and west ends of Potato Slough were sampled. Water samples were collected at the 6-foot depth for standard mineral analyses. Results of laboratory analyses are not yet available, but depth profiles of field conductivity and temperature measurements are presented in Appendix E, Tidal Effects Study Field Measurements. Sampling during other hydrologic conditions and at other reaches is being planned. The studies will provide information on the proportion of Sacramento and San Joaquin river waters and bay water diverted to the Federal and State water project intakes.



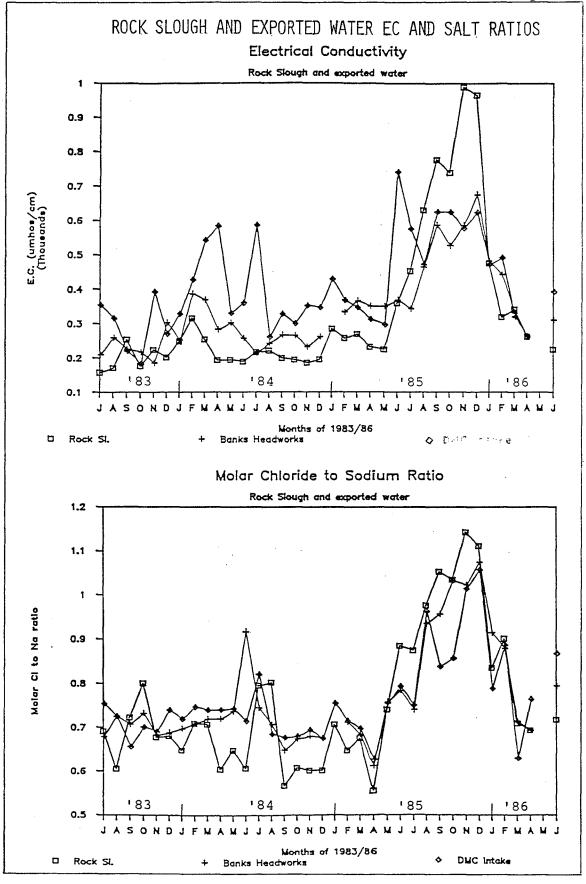
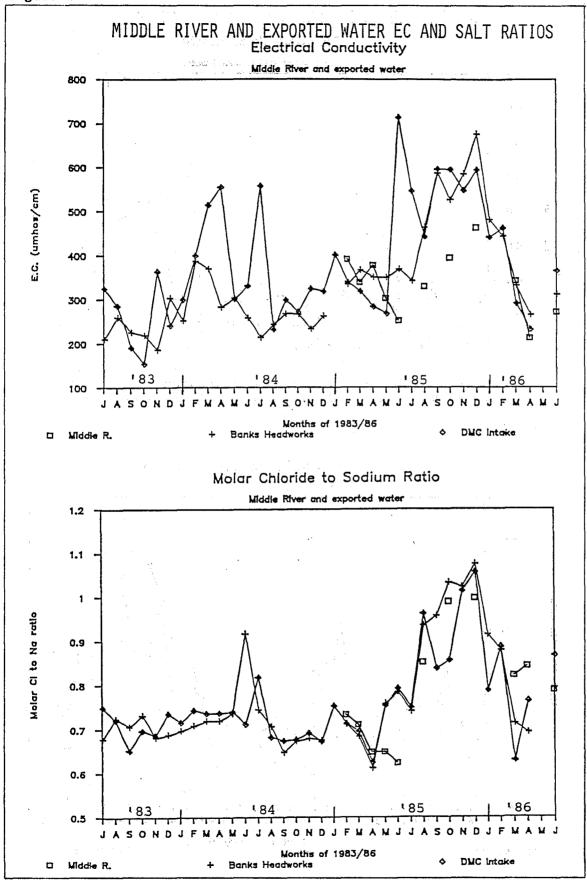
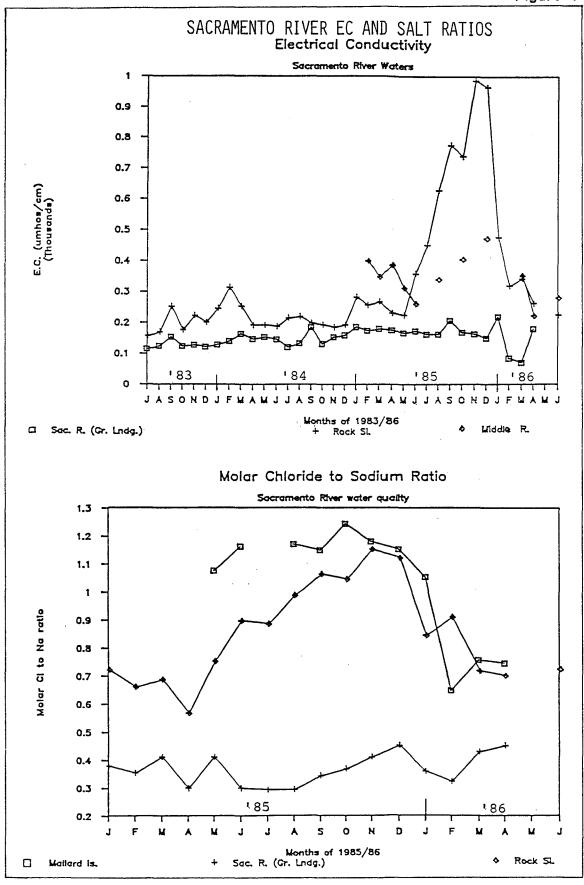
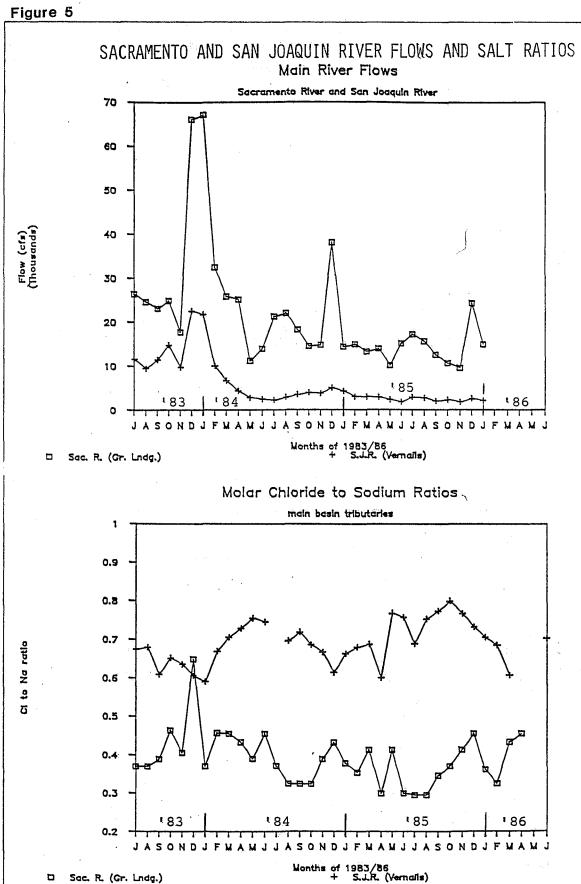


Figure 3









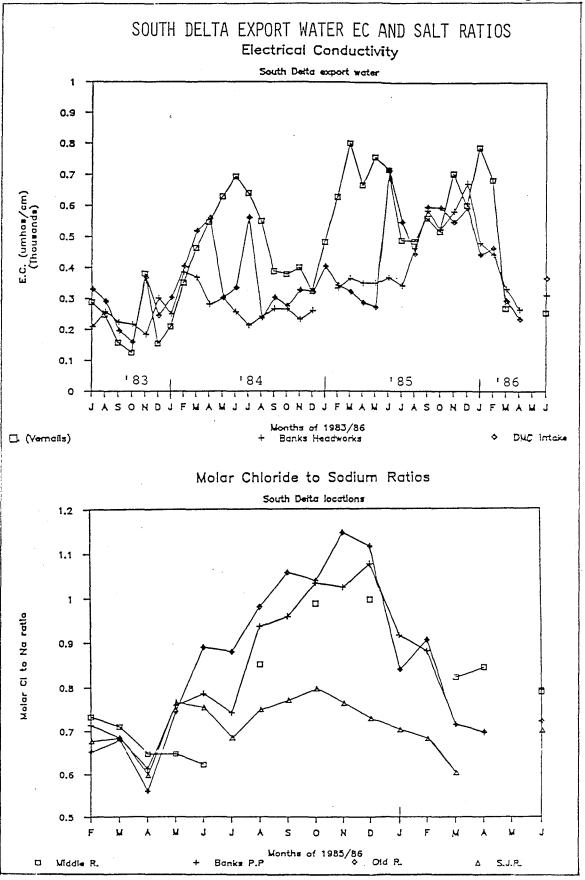


Table 2 MINERAL DATA pН EC STATION DATE TEMP A1k% S04 DO CA MG TDS THICK TURB NA ĸ C1 NO3 R FIELD (uS) <----- Milligrams per Liter----- (FTU) NAME (C) 03/04/86 19.5 7.3 AGDEMPIRE 2840 8.0 205 100 2.7 7 233 127 345 595 138.0 0.4 1860 924 04/17/86 15.0 7.4 AGDEMPIRE 1610 8.8 90 47 148 3.3 202 62 357 5.3 0.3 996 418 10 05/13/86 21.5 7.5 AGDEMPIRE 2000 6.6 108 56 204 2.7 217 50 506 0.8 0.3 1190 500 15 AGDEMPIRE 06/11/86 22.0 8.1 2760 150 296 830 0.0 0.4 5.7 84 2.5 215 18 1630 720 14 AGDGRAND 02/27/86 17.5 7.0 602 4.4 46 29 35 4.0 118 132 27 27.0 0.4 419 235 24 15 CLIFTON 03/04/86 16.5 7.3 306 7.8 **7** 29 2.1 50 29 3.1 0.2 177 21 41 66 04/09/86 16.5 7.2 CLIFTON 197 8.8 11 5 20 1.5 39 24 20 1.2 0.2 121 48 14 CLIFTON 05/07/86 15.5 7.3 280 8.8 16 7 27 1.8 55 36 28 3.2 0.2 171 69 13 CLIFTON 06/04/86 20.5 7.3 303 8.2 16 R 29 52 3.8 0.2 177 73 26 1:7 39 33 DMC 07/02/86 24.5 7.3 530 7.0 28 14 2.6 78 5.2 0.3 338 128 13 54 65 62 BANKS 07/02/86 24.0 7.3 6.4 9 2.3 0.2 25 305 16 31 59 33 1.6 231 77 34 ROCKSL 07/02/86 25.5 7.3 225 6.3 13 .∵8 19 1.9 56 21 19 1.0 0.1 144 66 15 GREENES 03/13/86 11.5 7.3 3 70 11.0 6 3 0.8 30 4 2 0.9 0.0 49 28 58 GREENES 04/23/86 18.5 179 8.5 7.3 8 0.0 13 10 1.2 64 12 7 3.1 114 66 14 **GREENES** 05/28/86 23.5 7.3 188 7.5 13 8 1.4 65 14 2.1 0.0 109 66 14 GREENES 06/25/86 24.5 7.3 161 7.8 11 7 11 1.2 52 11 8 1.5 0.1 106 56 13 02/27/86 MALLARDIS 14.5 7.0 169 8.8 12 6 12 2.0 43 18 12 5.8 0.1 102 54 58 MALLARDIS 03/13/86 13.0 7.3 161 9.4 10 6 12 1.8 42 18 14 2.6 0.1 108 50 51 MALLARDIS 04/23/86 16.5 7.3 226 8.9 12 7 20 1.6 48 22 23 2.6 0.1 136 59 22 MALLARDIS 05/28/86 17.0 7.6 4160 8.6 41 90 680 29.0 65 193 1240 1.4 0.4 2340 473 26 MALLARDIS 06/25/86 21.0 7.7 4250 8.1 40 689 28.0 0.9 0.4 2430 487 94 65 197 1280 36 **VERNALIS** 03/04/86 15.0 7.3 268 8.3 28 1.9 26 0.2 14 6 50 38 2.6 166 60 26

04/09/86

05/07/86

06/04/86

7.3

7.3

169

257

254

595

9.2

8.8

8.0

7.9

10

15

15

31

5

7

7

16

18

27

2.6

65

1.5

1.8

1.6

3.0

39

54

49

90

24

37

37

82

18

27

28

75

1.5

4.9

3.3

5.6

0.1

0.2

0.2

0.3

114

168

160

390 144

45

66

66

20

17

22

9

15.0

14.5

07/02/86 23.0 7.5

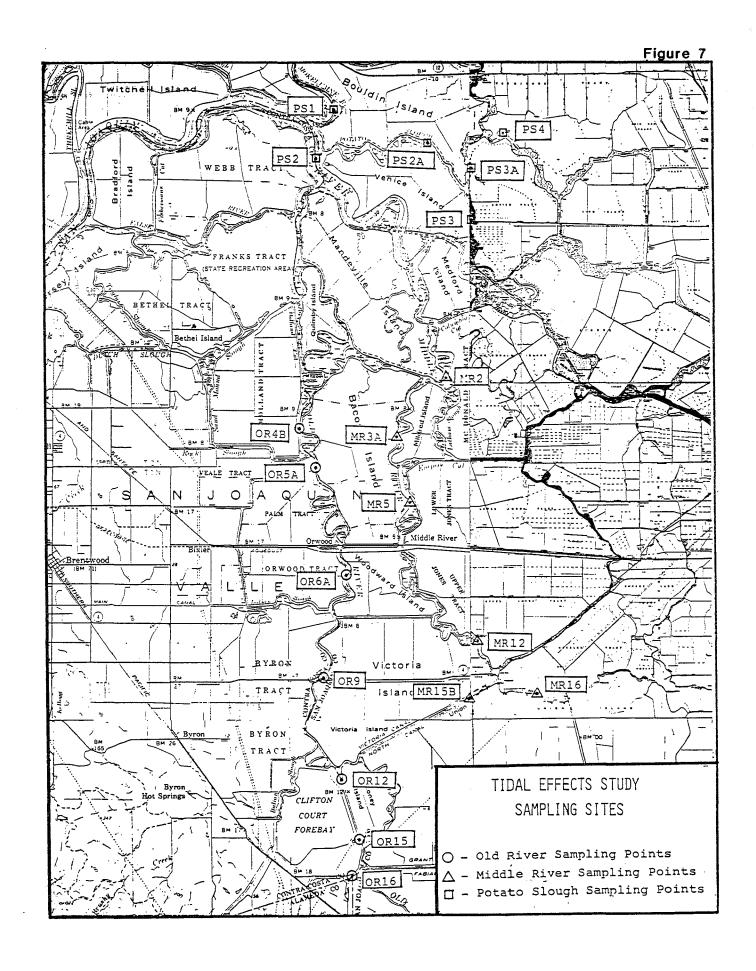
20.5 7.3

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# Reffects of Agriculture on Water Quality Studies

There are six program tasks to study effects of agriculture and related activities on Delta water quality.

## Drainage Water Quality Monitoring (Task AG1)

Irrigation return flows from drainages at Empire Tract, Grand Island, and Tyler Island are being monitored for salts, pesticides, THM formation potential, and other constituents. The data will be used to assess the loading effects of drainage on receiving water quality during the year.

The sampling locations and estimated size of each island are shown in Figures 8 and 9. Data to compute monthly loadings of drainage constituents have been requested from the island managers. Monthly loadings will be estimated by multiplying constituent concentrations by pumping rates. Electrical energy usage and data on pump efficiencies will be used to estimate pumping rates.

Monthly conductivity and molar chloride to sodium ratios are shown in Figure 10. During the historic flood of February 1986, the levee at Tyler Island failed, resulting in the inundation of the island. No samples were collected from Tyler Island until after the levee was repaired and water was pumped back into the channels. Sampling resumed in June 1986, but the data may not reflect typical drainage. Debris and many dead fish were observed in the drain, attributed to the receding water and to clean-up operations on The drainage was highly the island. turbid, deeply colored, and odorous (hydrogen sulfide gas).

Monthly sampling at drainages on Empire Tract and Grand Island was not interrupted.

Drainage quality at Empire Tract is distinctly different than drainage from Tyler and Grand islands. Empire Tract drainage exhibits chloride to sodium ratios similar to sea water. The electrical conductivity value of the drainage is about 1,000 uS/cm higher than that of the other two islands. Also, the laboratory analyses show a greater fraction of brominated trihalomethanes in Empire Tract drainage compared to the other islands. These differences may be due to a connate water source.

At all three drainages, similar patterns in conductivity were observed. Peak levels generally occurred in October through March, followed by progressively decreasing values in April and May, with annual lows in June through August. The shifts in the values result from application of river water during the irrigation months and leaching of soils during winter.

Pesticide concentrations were below detection or in trace amounts when detected at these drains. Sodium levels and conductance would exceed health standards if used for domestic purposes. Trihalomethane formation potentials were exceptionally high and indicate a significant contribution of THM precursor material to Delta waters (Table 3).

# Locating Irrigation Return Water Discharges (Task AG2)

Discharge points of irrigation return water are being identified and mapped. This information will be used to identify sources of contaminants and to plan upcoming work to assess their impact on water quality.

Figure 11 is a map showing drainage discharge points near the Clifton Court and Delta-Mendota Canal intakes. A request for data to estimate loadings

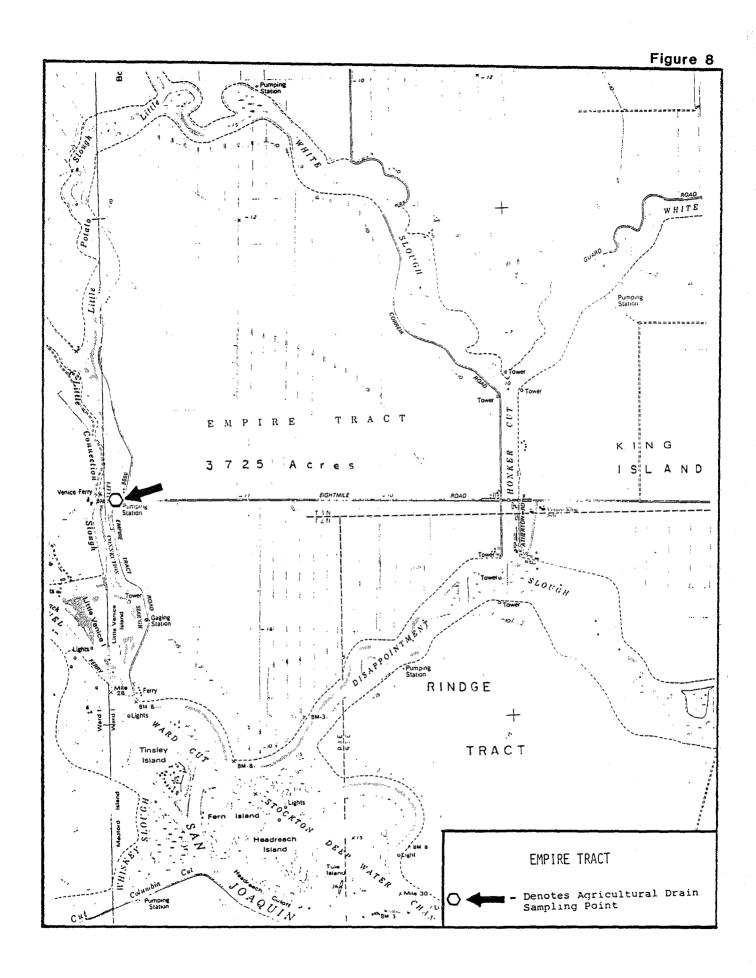


Figure 10

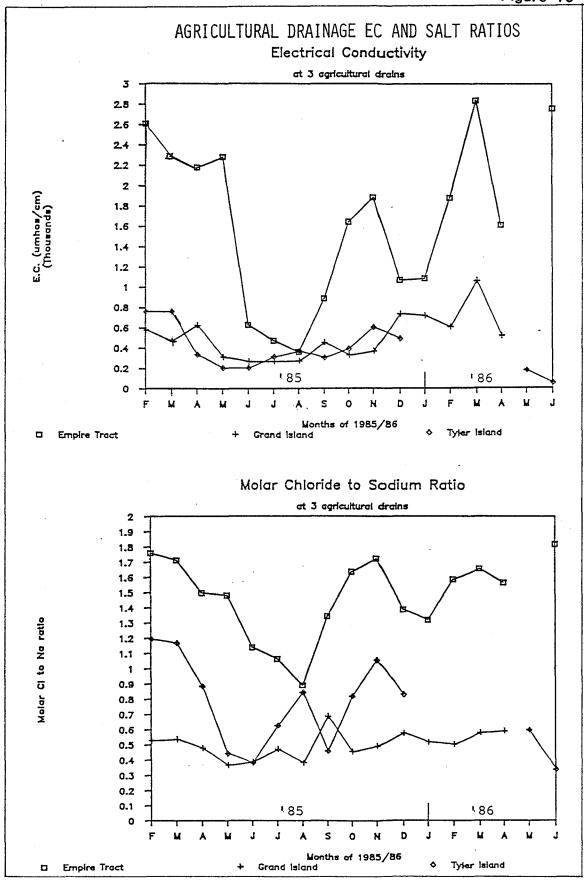
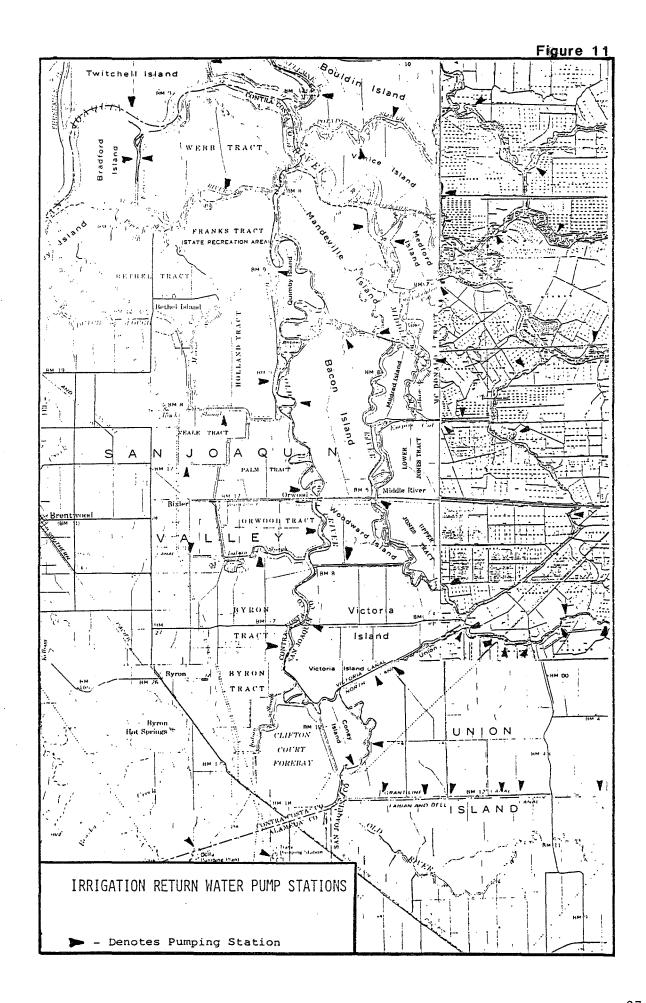


Table 3

AGRICULTURAL DRAINAGE QUALITY

				-	70		a 1		CH	CH	CH	CH	MITTER CO.
STATION	DATE (	Na	C1	Se )	EC (uS)	(FTU)	Color	Asbest (MF/L)	C13	C12Br	C1Br2 ug/L		TTHMFP
STATION				,					`	=======			
AGDEMPIRE	02/06/85	252		0.000	2610	26	25		1500	920	930	81	3400
AGDEMPIRE	03/06/85	226		0.000	2330	14		92					
AGDEMPIRE	04/05/85	224	517		2180	10	75		1800	920	370	31	3100
AGDEMPIRE	05/01/85	248	566	0.000	2280	14	160		1800	900	440	29	3200
AGDEMPIRE	06/05/85	54	95		629	15	75		1800	280	25	0	2100
AGDEMPIRE	07/24/85	42	69		472	10	40		2100	140	19	0	2300
AGDEMPIRE	08/01/85	32	44	0.000	360	8	100		2100	150	10	0	2300
AGDEMPIRE	09/11/85	83	172		886	4	150		3000	460	48	2	3500
AGDEMPIRE	10/02/85	149	376	0.000	1640	10	50		2200	790	330	26	3300
AGDEMPIRE	11/13/85	170	452	0.000	1880	4	80		2100	920	390	40	3400
AGDEMPIRE	12/03/85	87	186		1070	8	200	76	2900	360	44	1	3300
AGDEMPIRE	01/16/86	112	228		1087	3	160		6900	490	67	1	7500
AGDEMPIRE	02/13/86	162	396		1880	11	150		2600	650	170	8	3400
AGDEMPIRE	03/04/86	233	595		2840	7	200		1500	660	210	14	2400
AGDEMPIRE	04/17/86	148	357	0.000	1610	10	160	•	1900	830	320	13	3100
AGDEMPIRE	05/13/86			0.001			150		570	330	160	15	1100
AGDEMPIRE	06/11/86	296		0.000	2760	14	80				_	_	
AGDGRAND	02/06/85	43		0.000	576	34	25		2100	32	4	0	2100
AGDGRAND	03/06/85	35		0.000	468	21		630				_	
AGDGRAND	04/05/85	53	39		625	30	80		2000	100	4	0	2100
AGDGRAND	05/01/85	23		0.000	310	26	50		1000	41	0	0	1000
AGDGRAND	06/05/85	20	12 16		265 267	22 70	35 80		840 1800	37 60	0 2	0	880 1900
AGDGRAND AGDGRAND	07/24/85 08/01/85	22 22		0.000	273	30	50	_	1300	60 49	1	0	1400
AGDGRAND	08/01/85	31	33	0.000	451	28	30		1100	94	8	0	1200
AGDGRAND	10/02/85	27		0.000	327	25	30	ef	820	56	3	0	880
AGDGRAND	11/13/85	29		0.000	368	16	35		890	69	3	0	960
AGDGRAND	12/03/85	55		0.000	735	31	100	2100	2800	160	5	Ō	3000
AGDGRAND	01/16/86	64	51		716	26	80		3500	130	6	0	3600
AGDGRAND	02/27/86	35	27		602	24	100		1700	83	2	0	1800
AGDGRAND	03/13/86	64		0.001	1060	22	160		3200	180	5	0	3400
AGDGRAND	04/23/86	32		0.000	513	54	50		1700	82	2	0	1800
AGDGRAND	05/28/86	21	16		323	36	50		640	29	3	1	670
AGDGRAND	06/25/86	20	15		290	35	40						
AGDTYLER	03/27/85	46	84	0.000	743	29		530					
AGDTYLER	04/24/85	56	100		743	28	100		2100	260	27	0	2400
AGDTYLER	05/22/85	23	31	0.000	320	17	70		1800	91	4	0	1900
AGDTYLER	06/26/85	15	10		188	18	50		1400	45	3	0	1400
AGDTYLER	07/10/85	14	8		189	17	100		1600	51	1	0	1600
AGDTYLER	08/28/85	21		0.000	299	9	100		2100	78	3	0	2200
AGDTYLER	09/11/85	24	31		354	10	50		2200		6	0	
AGDTYLER	10/02/85	26		0.000	289	14	100		1200	70	2	0	1300
AGDTYLER	11/13/85	28		0.000	376	11	160		2000	120	2	0	2100
AGDTYLER	12/03/85	36		0.000	587	12	100	190	2100	85	2	0	2200
AGDTYLER	01/16/86	38	48		476	9	120		3500	83	8	0	3600
AGTYLER	06/11/86	10	9	0.000	158	768							



discharged into the rivers has also been made. Monthly loadings will be estimated by multiplying constituent concentrations by pumping rates, which will be estimated using information on electrical energy usage and pump efficiencies.

Actual pumping efficiencies may be lower than estimated, as the program's field crew found one return flow drain (east side of Orwood Tract) to be poorly maintained (e.g. intense algal mats and hyacinth growth at the pump station). This particular drain appeared to be discharging mostly air, rather than drainage, into Old River because of suction problems at the pump station inlet. Other drains are probably in a similar state.

# San Joaquin River Monitoring (Task AG3)

Comprehensive water quality monitoring near Vernalis has been initiated to study the effects of San Joaquin River on exported water quality. There are concerns about pesticides, selenium, and other trace elements that are being discharged from agricultural drainage into the San Joaquin River.

The drinking water standard for selenium is 10 ug/L. Maximum selenium concentrations so far have been 2 ug/L or less at the Vernalis station. Concentrations at some sites in the San Joaquin River upstream of Vernalis have been higher than 2 ug/L. Substantial amounts of farm drainage are discharged

into the lower reach of the river. The highest selenium concentrations are in Salt and Mud sloughs, where selenium-laden agricultural drainage enters from the Grasslands area (Figure 12 and Table 4).

In February 1985, an experiment was conducted to reduce selenium concentrations in the South Grasslands area. Participants included the U. S. Bureau of Reclamation, U. S. Fish and Wildlife Service, Department of Fish and Game, Central Valley Regional Water Quality Control Board, and Department of Water Resources. The study consisted of two steps: (1) rerouting subsurface drainage flows around the Southern Grasslands into Mud Slough, a tributary of the San Joaquin River, and (2) diverting Delta-Mendota Canal water into the South Grasslands area for dilution and flushing.

Department of Water Resources staff participated in monitoring selenium levels in the San Joaquin River during the experiment. The short-term experiment showed a reduction in selenium concentrations in surface waters with distance from the discharge point (Table 5 and Figure 12). Overall, data indicate the selenium drinking water standard is being met at Vernalis.

Pesticide levels have generally been below laboratory detection limits, except for methyl parathion (2.5 ug/L). Overall, none of the monitored pesticides is affecting drinking water quality of the San Joaquin River near Vernalis (Table 6).

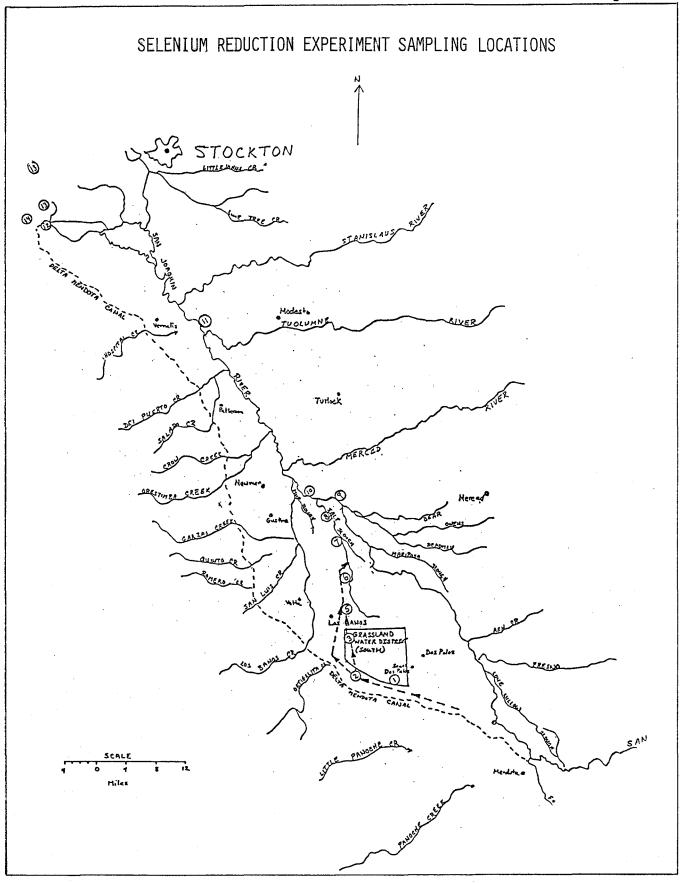


Table 4 MINOR ELEMENTS IN LOWER SAN JOAQUIN RIVER AND TRIBUTARIES STANAME Ν÷ DATE TIME TEMP pН DO FLOW EC TDS Cr Cin Mn Hg Mo (PST (oC) (mg/L) (cfs) (uS) - Milligrams per Liter MERCED 03/11/86 1215 17.0 7.0 10.8 3400 69 0.0 56 0 0.00 0.00 0.00 0.02 0.000 0.000 0.01 0.00 0.00 MERCED 04/21/86 1200 16.0 8.4 9.8 1000 0.0 45 0.00 0.01 0.00 0.03 0.000 0.000 0.01 0.00 0.00 53 0 MERCED 05/06/86 925 13.0 7.1 1000 0.0 42 0 0.00 0.00 0.01 0.03 0.000 0.000 0.01 0.00 0.00 9.5 52 MUDSL 0.04 0.000 0.002 0.00 0.00 03/12/86 16.0 8.5 1600 736 0.7 447 0.00 0.00 0.00 0.01 0.12 0.000 0.006 0.01 MUDSI. 04/21/86 1310 24.0 7.8 8.4 200 1030 1.1 656 O 0.000.00 0.00 0.00 0.00 MUDSL 05/06/86 1030 17.0 7.8 7.0 50 2150 1.9 1420 0 0.00 0.00 0.01 0.50 0.000 0.006 0.01 0.01 0.00 ORESTIMBA 03/12/86 1140 14.0 8.0 10.0 509 445 0.2 283 0 0.00 0.00 0.00 0.02 0.000 0.001 0.00 0.00 0.00 ORESTIMBA 04/21/86 24.0 337 0.2 213 0.00 0.01 0.00 0.04 0.000 0.000 0.01 0.00 0.00 1345 7.6 8.1 60.9 ORESTIMBA 05/06/86 1100 15.0 8.3 9.8 615 0.2 401 O 0.00 0.00 0.00 0.00 0.000 0.001 0.00 0.00 0.00 SALTSL 0.00 0.01 0.17 0.000 0.010 0.01 0.01 0.01 03/11/86 1030 13.5 7.3 889 1770 1160 0 0.00 8.7 2.2 SALTSL 04/21/86 1105 23.0 5.7 740 1380 1.4 898 0 0.00 0.02 0.00 0.22 0.000 0.008 0.01 0.01 0.01 SALTSL 05/06/86 0.00 0.18 0.000 0.010 0.01 0.00 830 15.0 7.7 1.3 809 0.00 0.01 0.01 7.4 525 1230 0 FREMONTFD 03/12/86 1300 16.0 9.5 5000 89 0.0 68 0.00 0.00 0.00 0.02 0.000 0.000 0.00 0.00 0.00 FREMONIFD 04/21/86 0.000 0.00 0.00 0.05 0.000 0.00 0.01 1245 22.0 7.4 8.8 3772 128 0.1 87 O 0.00 0.01 0.12 FREMONTFD 05/06/86 1015 16.0 7.4 7.9 1900 538 0.4 316 0.00 0.00 0.00 0.000 0.003 0.00 0.00 0.00 GRAYSON 03/12/86 1030 14.0 7.1 8.2 22000 294 0.2 182 0 0.00 0.00 0.00 0.05 0.000 0.001 0.00 0.00 0.00 GRAYSON 04/21/86 21.0 0.2 154 0.00 0.00 0.00 0.06 0.000 0.001 0.02 0.01 0.00 1515 7.4 8.7 11000 249 GRAYSON 05/06/86 0.00 0.00 0.08 0.000 0.002 0.00 0.00 0.00 1245 15.0 8.5 10000 403 0.3 0.00 MAZE 03/12/86 12.0 145 0.00 0.01 0.07 0.000 0.001 0.02 0.00 0.00 930 7.1 8.5 17830 234 0.2 0 0.00 MAZE 0.04 04/22/86 845 16.0 7.2 8.4 10790 213 0.1 134 0.00 0.01 0.00 0.000 0.000 0.01 0.01 0.00 0.00 MAZE 05/06/86 1400 15.0 7.3 8.5 8600 317 0.2 190 n 0.00 0.00 0.03 0.000 0.001 0.01 0.00 0.00 PATTERSON 03/11/86 1330 14.5 7.3 0.2 179 0.00 0.00 0.00 0.04 0.000 0.001 0.01 0.00 0.00 8.8 9090 290 PATTERSON 04/21/86 1430 22.0 7.4 8775 0.2 155 0.00 0.01 0.00 0.05 0.000 0.001 0.01 0.00 0.00 8.5 249 PATTERSON 0.00 0.01 0.05 0.002 05/06/86 1200 0.3 244 0.00 0.000 0.01 0.00 0.00 15.0 7.4 8.2 6300 392 STEVINSON 0.0 0.00 0.00 0.03 0.000 0.000 0.01 0.00 03/11/86 1130 14.0 9.1 12010 91 69 0 0.00 0.00 STEVINSON 04/21/86 22.0 0.00 0.00 0.03 0.000 0.000 0.00 1130 7.3 8.9 6431 69 0.0 56 O 0.00 0.00 0.00 STEVINSON 05/06/86 945 16.0 8.8 1600 170 0.0 110 0.00 0.00 0.01 0.04 0.000 0.000 0.00 0.00 0.00 STANISLAUS 03/12/86 0.02 840 11.0 7.1 10.5 4849 80 0.0 58 O 0.00 0.00 0.00 0.000 0.000 0.00 0.00 0.00 STANISLAUS 04/22/86 750 16.0 89 0.0 65 o 0.00 0.01 0.00 0.03 0.000 0.000 0.01 0.00 0.00 7.4 9.4 1670 STANISLAUS 05/06/86 1545 13.0 7.1 -0--0-0.00 0.00 0.00 0.00 0.000 0.000 0.00 0.00 0.00 11.2 1560 -0-0 TUOLUMNE 0.00 0.00 0.03 0.000 0.00 03/11/86 0.0 0.000 0.01 0.00 1445 13.5 7.1 53 0.00 9.5 6100 68 0 TUOLUMNE 0.01 0.00 0.03 0.000 0.000 0.01 0.00 0.00 04/21/86 1540 17.0 11.0 4550 54 0.0 0.00 TUOLUMNE 48 0.00 0.00 0.00 0.000 0.000 0.01 0.00 0.00 05/06/86 1315 13.0 2900 0.0 0.00 7.3 9.9 68 0 Sampling Stations Location Station Number Station Name Merced River at Milliken Bridge B0513100 MERCED B0040000 MIDSI. Mud Slough near Stevinson Orestimba Creek below Highway 33 B0873500 ORESTIMBA Salt Slough near Stevinson 80047000 SALTSL FREMONTFD San Joaquin River at Fremont Ford Bridge B0737500 B0708000 GRAYSON San Joaquin River near Grayson MAZE San Joaquin River at Maze Road Bridge B0704000 PATTERSON B0720000 San Joaquin River at Patterson Bridge

San Joaquin River near Stevinson

Stanislaus River at Koetitz Ranch

Tuolumne River at Tuolumne City

B0740000

B0311500

B0410500

STEVINSON

STANISLAUS

TUOLUMNE

## Table 5

## SELENIUM REDUCTION EXPERIMENT (February 27, 1986)

C+ o	tion Number and Description	Selenium	
عده	ictor number and bescription	Board	DWR
L	Agatha Canal @ Helm Canal (No Drainwater)	<5	
2	Camp 13 Ditch @ CCID Main Canal (Blended Drainwater)	42	
3	Camp 13 Ditch (Mud Slough 100 feet North of Mallard Road)	21	
4	Mud Slough @ Santa Fe Grade	18	
5	Santa Fe Canal @ Highway 152	18	
6	Santa Fe Canal Discharge to Mud Slough (100 feet West of Mud Slough)	29	31
7	Salt Slough (South Boundary of San Luis Island NWR)	12	
8	Salt Slough @ Lander Ave	10	
9	San Joaquin River @ Lander Ave	<5	
10	San Joaquin River @ Highway 140	8	
11	San Joaquin River Vernalis station		2
12	Delta-Mendota Canal Intake (Lindeman Road)		<1
13	Clifton Court Intake		<1
14	Harvey O. Banks Delta Pumping Plant		<1
15	Rock Slough @ Old River		<1

Board = Central Valley Regional Water Quality Control Board
DWR = Department of Water Resources Bryte Laboratory

## TABLE 6

## VERNALIS STATION PESTICIDE MONITORING DATA

	7/16	8/20	12/4	5/21
Chemical	<u> 1985</u>	<u> 1985</u>	<u> 1985</u>	1985
2,4-D salt	ND	ND		ND
bentazon	ND	ND		ND
carbofuran	ND	ND		ND
chloropicrin	NĐ	ND		ND
copper			8	
dacthal	ND	ND		ND
D-D mixture	ND	ND	ND	ND
MCPA	ND	ND		ND
metalaxyl	ND	ND		ND
methamidophos	ND	ND		ND
methyl bromide	ND	ND	ND	ND
methyl parathion	2.5	ND		ND
molinate	ND	ND		ND
paraquat dichloride	ND	ND		ND
thiobencarb	ND	ND		ND
xylene	ND	ND	ND	ND

ND = Not detected above analytical detection limit of 1 ug/L or less, or not reported when less than twice the background quality control blanks.

Analyses performed by McKesson Environmental Services.

# Selected Pesticide Monitoring (Task AG4)

Through a selection protocol based on pesticide usage patterns and environmental behavior, water samples are collected for specific pesticide analyses. The data are used to identify potential contamination to raw water supplies and at treatment plants. Attention is focused on chemicals that might present treatment difficulties, such as the highly water soluble compounds. Less soluble compounds tend to be removed more readily by flocculation, settling, and filtration processes because they are generally associated with suspended particulate matter such as silt and clays.

The selection protocol produces a siteand time-specific target list of pesticides for monitoring to improve chances of detecting any chemicals in the water and to eliminate the need for broad scans for hundreds of chemicals. Instead, the target list includes

specifically named chemicals and those detectable under the same analytical method. The target lists are developed from the California Department of Food and Agriculture annual pesticide use database, which was sorted by counties and chemicals. Chemicals that are water soluble or in high use are identified for each watershed where sampling stations are located. The period of application or use of each chemical is also included in the database. Identified chemicals then appear on the monthly target lists for each sampling station. A more complete description of the pesticide monitoring selection scheme is provided in Appendix F, Pesticide Monitoring Selection Scheme.

Results of the pesticide monitoring are shown in Table 7. Sampling primarily focused on the application period (summer), with a sampling run in winter (runoff months) and a run in early spring (pre-emergent herbicide applications). Most of the targeted chemicals were below the analytical limit of detection. Reported chemicals were generally below State Action Levels for drinking water or were near the low level detection limits of the laboratories. These results indicate Delta water supplies are acceptable for domestic uses.

## Modeling Pesticide Fate and Transport (Task AG5)

Computer models developed by EPA to predict the fate and movement of organic pesticides in an aquatic system are being tested to help assess the potential of contamination to drinking water supplies. The models are used to test the pesticide monitoring selection protocol for Task AG4 and to study the effects of changing riverflow and other environmental conditions on the distribution of pesticide contaminants.

One model under review is EXAMS (Exposure Analysis Modeling System).

EXAMS is a steady state and dynamic model designed for rapid evaluation of the behavior of synthetic organic chemicals in aquatic ecosystems. The program computes:

- Exposure (the ultimate expected environmental concentrations resulting from a long-term steady pattern of pollutant loadings),
- Fate (the distribution of the chemical in the environment and the fraction of the loadings consumed by each transport and transformation process), and
- Persistence (the time required for effective purification of the system once the loadings cease).

1

\* 4773 1 12 71

A model such as EXAMS could be used to assess the likelihood of contamination to water supplies in a given reach in the Delta. Other possibilities are to help target monitoring to those environmental compartments (sediment, water, biota) where the chemicals will most likely be distributed and, thereby, allow more effective monitoring of their presence.

Brief descriptions of the EPA computer models are presented in Appendix G.

## Health Effects Database on Selected Chemicals (Task AG6)

Drinking water standards currently exist for only a few pesticides. A computer literature search for human health effects data is underway for those chemicals appearing on the selected pesticide monitoring target lists that do not have drinking water standards. The data will be used to help assess the degree of risk presented to Delta water supplies found with traces of these contaminants.

A summary of current information on the toxicity of some of these chemicals appears in Table 8.

Table 7 PESTICIDE MONITORING DATA, 1985 AND 1986 (All Units in ug/L)

Target pesticide	Sampling date			-		_	Grand Is.S ag. dr.nr.						main dr	CliftonC .intake	Leve
,4-D salt	07/16/85	0.1			ND	ND	ND	ND	0.1	ND	ND ND	ND			
	08/20/85	0.01			ND	ND	ND	ND	ND	NID	ND	ND	300	,	
	12/04/85	0.01				ND						. ND	ND		
	05/21/86	0.5				1	NTO.	ND	0.2	MD	M	M		ND	
entazon	07/16/85	0.1			1.6	ND	ND	ND	0.3	ИD	ND	ND			
	08/20/85	0.2			ND	ND	ND	ND	0.5	ND	ND	ND			
	12/04/85	0.5				ND						ND	ND		
	05/21/86	1				ND		ND						ND	
arbofuran	07/16/85	0.5			ND	ND	ND	ND	ND	NTO	ND	ND			
	08/20/85	0.5			ND	ND	ND	ND	ND	ND	ND	ND			
	12/04/85	0.1				ND						ND	ND		
	05/21/86	0.2				ND		ND						ND	
nloropicrin	07/16/85	0.1	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND			30
	08/20/85	0.1	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND			25.0
	12/04/85	0.1			ND	ND		ND	ND		,	ND	ND		26.5
	05/21/86	0.1				ND		ND						ND	*5
opper dacthal	12/04/85	5			5	13		8	ND			8	10		
	07/16/85	0.01			ND	ND	ND	ND	ND	ND	ND	ND			
	08/20/85	0.05			ND	ND	ND	ND	ND	ND	ND	ND			
	12/04/85	0.3				ND						ND	ND		
	05/21/86	0.01				ND		ND						ND	
-D mixture	07/16/85	0.1	ND	ND	. ND	ND	ND	ND	ND	ND	ND	ND			
	08/20/85	0.1	ND	ND	ND	ND	ND	. ND	ND	ND	ND	ND			
	12/04/85	0.5			ND	ND		ND	ND			ND	ND		
	05/21/86	0.2				ND		ND						ND	
CPA	07/16/85	1	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND			
	08/20/85	10	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND			
	12/04/85	2				ND						ND	ND		
	05/21/86	20				ND		ND		٠.				ND	
etalaxyl	07/16/85	1			ND	ND	ND	ND	ND	ND	ND	ND			
	08/20/85	10			ND	ND	ND	ND	ND	ND	ND	ND			
	12/04/86	0.1				ND						ND	ND		
	05/21/86	0.05				ND		ND					<b></b>	ND	
ethamidophos	07/16/85	2			ND	ND	ND	ND	ND	ND	ND	ND			
ourmand broom	08/20/85	0.5			ND	ND	ND	ND	ND	ND	ND	ND			
	12/04/85	5			ND	ND	ND	112	1.0	112		ND	ND		
	05/21/86	5				ND		ND					ND	ND	
ethyl bromide	07/16/85	0.5	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND		ND	
saly1 blomide	08/20/85	0.5	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND			
	12/04/85		ND.	שא			ND	ND	ND	110	IID.	ND	ND		
		0.7			ND	ND			ND			ND	KD		
	05/21/86	0.5			***	ND	170	. ND	300	3773	MD	100		ND	
ethyl parathion	07/16/85	2.5			ND	ND	ND	2.5	ND	ND	ND	ND			3
	08/20/85	1			ND	ND	ND	ND	ND	ND	ND	ND	1770		3
	12/04/85	0.01				ND						ND	ND		3
	05/21/86	0.005				ND		ND						0.03	3
olinate	07/16/85	1	ND	1	ND	ND	ND	ND	ND	ND	ND	ND			2
	08/20/85	0.5	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND			2
	12/04/85	0.05				ND						ND	ND		2
	05/21/86	0.05				ND		ND						ND	2
raquat dichloride		10			ND	ND	ND	ND	ND	ND	ND	ND			
	08/20/85	10			ND	ND	ND	ND	ND	ND	ND	ND			
	12/04/85	20				ND						ND	ND		
	05/21/86	10				ND		ND						ND	
niobencarb	07/16/85	8	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND			×
	08/20/85	1	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND			*:
•	12/04/85	0.05				ND						ND	ND		*
	05/21/86	0.05				ND		ND						ND	*
ylene	07/16/85	0.2	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND			. 62
	08/20/85	0.5	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND			62
	12/04/85	0.4			ND	ND		ND	ND			ND	ND		62
	05/21/86	0.2				ND		ND						ND	62
	-,,	***				עונג		110						M	02

Tentative recommended action level. The recommended action level for taste and odor threshold is 1.0 ug/L for thiobencarb and 37 ug/L for chloropicrin.

Note: Blanks indicate no analysis performed for that chemical. ND = Not detected when less than twice the blank value.

Analyses performed by McKesson Environmental Services.

## Table 8 PESTICIDE TOXICITY INFORMATION

			Acute		Subacut	e Toxicit	У		Chr	onic Tox:	lcity
		Symptoms	Toxicity		Dose	Length		Animal	Dose	Length	j.
_Chemical_	<u>Characteristics</u>	of Poisoning	(LD50)*	<u>Tested</u>	(mgq)	<u>of Test</u>	Results	Tested	(ppm)	of Test	<u>Results</u>
2,4-D Salt	White crystals	No characteristic symptoms for humans	375			·		Rat Dog (Hi	500	2 yrs 2 yrs vels not	No effect No effect tested)
Bentazon	White crystalline solid	Apathy, ataxia, prostration, tremors anorexia, vomiting	1100	Rat Dog	1600 3000	90 days 90 days	Minimum effect level	Rat	350	2 yrs	Minimum effect level
Carbofuran	White crystalline solid**	,	8								
Chloro- picrin	Slightly oily liquid** Reduced amenities, Faint odor-0.0073mg/L**	n's c	250								
Dacthal	White crystalline solid		3000	Adult mallard	5000	100 days	No effect	Rat Dog	10000 10000	2 yrs 2 yrs	No effect No effect
D-D Mixture			140		•						,
Methyl bromide	Colorless liquid or gas Threshold limit 20 ppm Cumulative poison*	ytestes	lmg/L								- / - - 
Paraquat dichloride	White crystalline solid; faint ammoni-	Vomiting, diarrhea, general malaise	150				-	Rat	170	2 yrs	No significant abnormalities
	acal odor							Dog	85, 170	27 mos	Effects apparent
Thio- bencarb	Light yellow or brownish yellow liquid		920	Rat Dog	660 660	90 days 90 days	Minimum effect level	Rat Dog	30 30	2 yrs 2 yrs	Minimum effect level

Unless otherwise noted, information is from "Herbicide Handbook of the Weed Science Society of America" (4th Edition), 1979.

<sup>\*</sup> W. T. Thomas, "Agricultural Chemicals"; LD50 values resulted from tests performed on white rats, in milligrams per kilogram of body weight.
\*\* "The Merck Index" (9th Edition)

<sup>\*\*\*</sup>Karel Verschueren, "Handbook of Environmental Data on Organic Chemicals" (2nd Edition)

# Waste Waters, THMs, and Modeling Studies

Three tasks are underway to examine waste water discharges, trihalomethane formation potential, and water quality modeling.

# Survey Major Waste Water Dischargers (Task WD1)

The Central Valley Regional Water Quality Control Board provided effluent monitoring data on major municipal and industrial waste dischargers within the program study area. The data will be examined to estimate total waste loads and to determine if special studies on receiving water impacts are needed.

Data on the Sacramento Regional Wastewater Treatment Plant are shown in Table 9 for some months of 1985 and This is the largest publicly owned waste water treatment facility in the Delta. The effluent outflow is significantly small in comparison to the Sacramento River outflow. suggests that there is sufficient mixing and dilution of this highly treated effluent. For comparison, water quality downstream in the Sacramento River at Hood and Greene's Landing appears to be well within drinking water standards (Table 10). Effluent data for other waste dischargers are also being tabulated for review.

MONTHI V AVIT			
SACRAMENTO REGIONAL V	WASTE WATER	TREATMENT	PLANT
3	Table 9	•	

<u>Date</u>	Effluent Discharge (MGD)	Suspended Matter (mg/L)	Suspended Matter (1bs/day)	TDS (ppm)	EC (uS/cm)	River Flow (MGD)
1/85	120	9	9,007	330	590	17,100
2/85	123	38	38,981	370	650	18,700
3/85	124	20	25,837	339	597	14,500
4/85	109	9	8,182	364	620	12,600
9/85	124	10	10,342		640	12,500
10/85	127	13	13,769		600	9,900
11/85	143	38	45,320		610	10,900
12/85	143	17	20,275		600	16,600
1/86	143	15	17,889		590	19,300
2/86	198	8	14,356		570	67,250
3/86	175	8	11,676		610	74,403
4/86	142	8	9,474		630	26,100

1 million gallons per day (MGD) equals 1.55 cubic foot per second (cfs).

SOURCE: Central Valley Regional Water Quality Control Board

Table 10
SACRAMENTO RIVER WATER QUALITY AT GREENE'S LANDING AND AT HOOD

	v .											СН	CH	Сн	СН	TTHMF	,	
STATION		TEMP .		DO	Na	C1	Se	EC	Turb	Color	Asbest		C12Br		Br3		TOC	FLOW
NAME	DATE	(C)	ън	(						00101				-ug/L				(CFS)
	>=====================================											-		-				
	07/21/83			8.7	7	4		115	9	2		190	8	1	0	200	1.6	26400
	08/18/83			8.2	7	4	4	124	8	8.		200	14	1	0	220	1.6	24600
	09/13/83			8.3	10	6		154	12	8		600	18	2	0	620	1.8	23100
	10/04/83			9.0	7	5		124	10	5	380	200	9	0	0	210	1.6	24800
	11/01/83			9.1	8	5		128	6	5	340	210	. 8	0	. 0	220	1.7	17700
	12/06/83			10.6	4	4		122	30	30.		300	9	Ö	0	310	4.1	66100
	01/10/84			10.7	7	4		129	19	20	3200	220	10	1	0	230	1.7	67200
	02/01/84			10.8	7	5		140	14	12	740	190	11	ī	0	200	1.5	32400
10.00	03/07/84	in the	1	10.8	10	. 7		164	8	8	540	230	28	1		260	1.6	25800
	04/04/84			10.4	9	. 6		148	8	5	680	250	14	1	0	260	1.6	25100
	05/02/84			9.4	10	6	5.4	154	8	8	110	180	13	1	0	190	2.0	11200
	06/06/84			8.7	10	7	• •	146	9	8	200	250	15	1	. 0	270	2.0	13900
	07/10/84			8.2	7	4		121	11	5	150	260	10	. 0	. 0	270	1.6	21200
	08/01/84			7.9	8	4		133	11	5	730	300	10	1	0	310	1.6	22000
	09/05/84			7.7	12	6	0.000		11	.8		390	20	1	0	410	2.4	18240
	10/04/84			9.0	8	4	0.000	132	7	.ņ 5		170	13	.1	0	180	1.6	14500
	11/08/84			9.7	10	6	0.000	154	11	8		210	11	0	. 0	220	2.1	14800
	12/05/84			10.9	9	6	0.000	160	24	15	1100	240	14	1	0	260	2.6	38100
	01/30/85		7.4	11.9	12	. 7	0.000	186	3	13	1100	240	14	_	v	200	2.0	14300
	02/06/85			12.1	11	6	0.000	174	8	10		360	14	1	0	380		14900
	03/06/85			10.5	11	. o. 7	0.000	180	5	10	180	300	14	_	Ū	360		13200
	04/05/85			9.3	13	6	0.000	176	7	2	100	160	13	0	0	170		13900
	05/01/85			8.8	11	7	0.000		11	10		210	12	1	0	220		10200
	06/05/8			8.5	13	6	0.000	173	9	10		290	19	1	- 0	310		15100
	07/24/85			8.0	11	5	0.000	163	8	10		230	17	-		310		17200
	08/01/85			7.9	. 11	5	0.000	163	10	10		480	14	2	0	500	3.9	15600
	09/04/85			7.8	15	8	0.001	W	8	5		220	22	2	0	240	3.5	12500
	10/02/85			8.2	14	8	0.000	168	7	5		200	14	1	0	220	1.6	10600
	11/13/85			9.7	11	. 7	0.000		. 6	5		290	20	1	0	310	2.8	9500
	12/03/85			9.3	10	7	0.000	149	28	35	380	690	21	1.	0	710	16	24200
4	01/16/86			10.6	18	10	0.000		9	15	260	660	22	1	0	680	2.3	14900
	02/27/86			10.5	4	2	0.000	84	64	20		340	7	0	0	350	4.2	14900
	03/13/8			11.0	3	2	0.000		58	10		430	8	0	0	440	2.4	
	04/23/86			8.5	3 10	7	0.000		38 14	10		310	22	1 .	0	330	1.9	
	05/28/8			7.5	10	,		1/9	14	10		170	12	2	1			
	06/25/86			7.8			0.000			10		1/0	12	2	T	190	2.9	
HOOD	03/30/83			10.7		1.	0.000		20	5		310	9	0	0	320	7.00	40000
HOOD	06/29/8			8.5		4		131 128	20 6	3		230	12	7.44	0	240	S (1)	40000
HOOD	08/26/8			8.1		5 5		149	10			280	13	0	0	290		20000 23200
HOOD	10/21/8			8.7		5 4		122	. 10			260	10	0	0	270		16300
HOOD	12/29/82			10.9		-			33			480	16	1	. 0	500		
HOOD	02/24/83			10.9		4		130 113	33 30			120	4	0	0	120		71700
HOOD	04/27/8			10.0										4	4			74000
HOOD			7.3	9.1		3		112 101	26 17			166 200	6 8		0	180 210	-	54600
11000	06/22/8:	74.2	7.3	7.1				101	1/			200		J	· · ·	210		43540

# Assess THM (Trihalomethane) Formation Potential (Task TR1)

Trihalomethanes (THMs) are a group of compounds that can be formed in drinking water during the disinfection process of chlorination. Organic substances such as fulvic and humic acids occurring naturally in the water react with chlorine to form THMs. There are four species of THMs normally found in drinking water: chloroform (CHC13), bromodichloromethane (CHC12Br), dibromochloromethane (CHBr2C1), and bromoform (CHBr3).

As part of the monthly water quality monitoring (Task WQ1), parameters such as total organic carbon, bromide, and color are measured. In addition, water samples are chlorinated and analyzed for THM species and total THM formation

potential to determine if THM control might be a problem.

Total trihalomethane formation potential (TTHMFP) measurements are summarized in Table 11. The TTHMFP test is designed to estimate the maximum levels of THMs that could be produced from a water supply and, accordingly, does not predict actual concentrations of THMs in finished drinking water. Many factors, including temperature, pH, and chlorine contact time and dosage, affect actual THM formation in water treatment facilities. Treated drinking water contains lower THM concentrations than the maximum potential estimated by this assay procedure. There are also methods such as ammonia addition after chlorination to reduce THM levels in finished drinking water supplies.

T	able 11	
TRIHALOMETHANE	FORMATION	POTENTIALS

Station (Raw Water Supply)	Number Samples	Maximum	Average	Std. Dev.	Minimum	Maximum Bromoform
Ag Drain at Empire Tract	15	7500	3160	1332	1100	81
Ag Drain at Grand Island	15	3600	1779	904	670	1
Ag Drain at Tyler Island	9	3600	2078	648	1300	0
American River at WTP	28	380	238	57	150	0
Banks Pumping Plant	41	1900	550	267	220	13
Cache Slough	15	920	707	171	380	2
Clifton Court Intake	29	710	467	120	170	13
Cosumnes River	18	840	252	153	140	0
Delta-Mendota Canal Intake	29	860	465	138	230	10
Sacramento River at Greene's Landing	32	710	309	140	170	1
Honker Cut	12	570	402	104	250	5
Sacramento River at Hood	8	500	266	107	120	4
Little Connection Slough	11	710	404	165	170	0
Lindsey Slough	19	2300	1049	408	420	2
Mallard Slough	7	810	446	182	210	280
Sacramento River at Mallard Island	11	1400	904	255	510	990
Middle River	10	880	603	138	390	5
Mokelumne River	18	420	250	66	120	0
North Bay Interim Pumping Plant	27	780	370	100	280	Ö
Rock Slough at Old River	29	770	460	132	220	36
San Joaquin River at Vernalis	42	1500	519	237	220	12
OVERALL	425	7500	645	712	120	990

test is used to compare the relative level of THM forming materials in raw water supplies. The EPA Maximum Contaminant Level for total THM is 100 ug/L in finished drinking water supplies. There is no standard for TTHMFP in raw waters.

The diverse sources of natural organic matter include biological productivity in the water column and watershed, waste discharges, and farm drainage. Total organic carbon (TOC) analyses were performed to measure the potential amount of available organic matter for THM formation. However, due to laboratory quality control problems with TOC analyses, data analysis to correlate TOC measurements with TTHMFP values are temporarily suspended. Laboratory staff are examining lab workbooks and correcting erroneously computed results (see Appendix D).

TTHMFP results from the three agricultural drains indicate that Delta soils may be a major source of organic trihalomethane precursors. Bromoform, a THM that is more difficult to treat, appeared at some stations. Bromide, a common sea water constituent, combines with trihalomethane precursors during chlorination to form brominated THM species such as bromoform, dichlorobromomethane, and dibromochloromethane. Brominated THMs were highest in Mallard Slough and in the Sacramento River near Mallard Island. Water quality here is significantly controlled by riverflow and tidal influences, as seen by conductance, sodium, chloride, and brominated THM data. The trace amounts of bromoform at the Banks Pumping Plant headworks, Clifton Court intake, Delta-Mendota Canal intake, Rock Slough at Old River, and Middle River reflect diluted tidal waters that are exported by pumping operations at the State and Federal facilities. Bromoforms at Vernalis and other areas probably reflect the application of Project waters diverted from the Delta that contained bromides or leaching of

bromide deposits in soils from ancient marine deposits.

Total THM formation potential was greater in the southern Delta than in the Sacramento River at Greene's Landing and at the American River Water Treatment Plant stations. The potential is most likely higher because of lower channel flows, agricultural drainages, and higher biological productivity within the southern region as compared to stations on the Sacramento and American rivers.

The TTHMFP at Lindsey Slough was also high and may be attributed to local agricultural drainage and extensive riparian vegetation at the sampling station and a long water retention time that might increase the concentration of precursors in the water from decaying matter.

# Use of Existing Water Quality Models (Task MOD1)

Computer models developed by the Environmental Protection Agency to study the distribution, fate, and transport of waste waters and spilled materials are being tested for use in studying Delta water quality as affected by waste water discharges and pesticide usage. EPA recently made several models available for use on microcomputers.

One model under study is QUAL2E (Enhanced Stream Water Quality Model), a stream quality routing model. QUAL2E is a steady state model for conventional pollutants in branching streams and well mixed lakes. It includes conservative substances, temperature, coliform bacteria, biochemical oxygen demand, dissolved oxygen, nitrogen, phosphorus, and algae. The model is widely used for waste load allocation and permitting in the United States and other countries.

Another model is WASP (Water Quality Analysis Program). WASP is a generalized modeling framework for contaminant fate and transport in surface waters. Based on a flexible compartment modeling approach, WASP can be applied in one, two, or three dimensions if desired. Problems studied using WASP

include biochemical oxygen demanddissolved oxygen dynamics, nutrients and eutrophication, bacterial contamination, and toxic chemical movement.

These EPA models are described in Appendix G.

## REFERENCES

- Public Health Aspects of Sacramento-San Joaquin Delta Water
  Supplies -- A Panel Report for the California Department of Water
  Resources. California Department of Water Resources.

  December 31, 1982. 59 pp.
- Interagency Delta Health Aspects Monitoring Program, Project Report. California Department of Water Resources, Central District. May 1985.

## Appendix A

## MONITORING PROGRAM DATA

#### DELTA HEALTH ASPECTS PROGRAM MONITORING DATA

TO THE MEDICAL COLOR OF THE COLOR COLOR

THM FORMATION POTENTIAL CH CH CH CH Station Date Time Temp. pН D.O. Na C1EC Se Turb. Color TOC Asbest. C13 BrCl2 Br2Cl Br3 TTHMFP FLOW Name (PST) (oC) - mg/L. (--------(uS/cm) (TU) (CU) (mg/L) (mF/L) (-----) (cfs.) AGDEMPIRE 02/06/85 905 6.0 7.3 9.8 252 685 0.000 2610 26 25 1500 920 930 81 3400 AGDEMPIRE 03/06/85 945 10.5 7.3 7.6 226 597 0.000 2330 14 ------92 AGDEMP1RE 04/05/85 7.3 517 850 21.5 3.9 224 2180 10 75 1800 920 370 31 3100 \_\_\_ AGDEMPIRE 05/01/85 830 20.0 7.6 6.5 248 566 0.000 2280 14 160 1800 900 29 3200 440 AGDEMP I RE Ø6/Ø5/85 807 20.0 7.3 95 4.0 54 629 15 75 -------1800 280 25 0 2100 \_\_\_ AGDEMP I RE 07/24/85 907 23.0 €.8 4.1 42 69 472 10 40 2100 140 19 0 2300 AGDEMPIRE 08/01/85 825 22.0 6.8 5.5 32 44 Ø. 000 360 8 100 22.0 150 ---2100 10 0 2300 6.9 AGDEMP I RE 09/11/85 1020 19.5 4.5 83 172 886 150 4 19.0 \_--3000 460 48 2 3500 AGDEMPIRE 10/02/85 700 376 0.000 18.0 7.6 7.6 149 1640 10 50 18.0 --2200 790 330 26 3300 AGDEMPIRE 11/13/85 8ଡଡ 7.0 7.3 9.0 170 452 0.000 1880 4 80 34.0 ---2100 920 390 40 3400 AGDEMPIRE 12/03/85 1710 14.0 7.0 5.4 87 186 44.0 1070 8 200 76 2900 360 44 1 3300 AGDEMPIRE 01/16/86 1145 12.0 €.8 5.8 112 228 \_\_\_ 1087 31.0 6900 3 160 ---490 67 1 7500 AGDEMPIRE Ø2/13/86 1200 14.0 6.8 6.7 162 396 ---1880 11 150 40.0 --2600 650 170 8 3400 \_\_\_ AGDEMPIRE 03/04/86 19.5 233 1330 7.3 8.0 595 ---2840 7 200 65.0 \_\_ 1500 660 210 14 2400 --AGDEMP I RE Ø4/17/86 15.0 357 915 7.4 8.8 148 0.000 1610 10 160 47.0 1900 830 320 13 3100 AGDEMPIRE 05/13/86 1000 21.5 7.5 6.6 204 506 0.001 2000 15 150 61.0 570 330 160 15 1100 \_\_\_ AGDEMPIRE 06/11/86 800 22.0 8.1 5.7 296 830 0.000 2760 14 80 65.0 ------------AGDGRAND 02/06/85 1030 11.5 7.1 7.5 43 35 Ø. 000 57€ 34 25 \_\_ 2100 32 4 0 2100 ------AGDGRAND 03/06/85 1100 12.5 6.9 5.3 35 29 0.000 468 21 .... ---630 ------------AGDGRAND 04/05/85 1000 18.5 39 7.3 5.0 53 625 30 8Ø ------\_\_\_ 2000 100 0 2100 AGDGRAND 05/01/85 945 18.5 6.9 5.7 0.000 23 13 310 26 50 ------1000 41 0 1000 AGDGRAND 06/05/85 915 21.0 7.3 6.6 20 12 \*\*\* 265 22 35 ------840 37 Ø 880 AGDGRAND 07/24/85 715 22.5 7.2 5.5 22 16 ---267 70 80 1800 60 2 0 1900 AGDGRAND 08/01/85 945 21.5 7.1 0.000 6.5 20 13 273 30 50 17.0 1300 49 0 1400 1 AGDGRAND 09/11/85 1150 19.5 7.2 €.1 31 33 451 28 30 1100 14.0 94 8 0 1200 AGDGRAND 10/02/85 300 19.0 7.2 6.0 27 19 0.000 327 25 30 4.5 ---820 56 3 0 880 AGDGRAND 11/13/85 945 12.5 7.3 4.5 29 22 0.000 368 35 9.0 \_\_\_ 890 16 69 3 Ø 960 13.0 AGDGRAND 12/03/85 1845 7.0 3.8 55 49 0.000 735 31 100 39.0 2100 2800 160 5 Ø 3000 @1/16/86 51 AGDGRAND 1315 13.5 7.3 7.3 64 . -- ---716 26 80 20.0 ---3500 130 3600 6 Ø AGDGRAND 02/27/86 1130 17.5 7.0 4.4 35 27 ---602 24 100 28.0 \_--1700 83 2 (2) 1800 AGDGRAND @3/13/86 1300 14.5 6.6 5.8 64 57 0.001 1060 22 160 56.0 3200 180 5 0 3400 \_\_\_ AGDGRAND 04/23/86 1200 18.5 7.3 7.6 29 513 32 0.000 54 50 23.0 --1700 82 2 0 1800 ---AGDGRAND 05/28/86 1115 22.5 7.3 7.4 21 16 323 36 50 38.0 29 \_\_\_ 3 640 670 \_\_ AGDGRAND 06/25/86 1200 24.5 6.8 15 290 35 7.2 20 40 43.0 ---------AGDTYLER 03/27/85 1245 11.5 6.8 7.8 46 84 0.000 743 29 \_\_\_ 530 ~~ ---AGDTYLER 04/24/85 1230 19.5 7.3 100 5.8 56 743 28 100 ---2100 260 27 0 2400 AGDTYLER 21.5 05/22/85 1130 7.2 4.7 23 31 0.000 320 70 17 1800 91 0 1900 AGDTYLER 06/26/85 1115 24.0 6.8 5.5 15 10 188 18 50 1400 45 \_\_\_ 3 0 1400 AGDTYLER 07/10/85 1200 25.5 7.0 В 4.5 14 189 17 100 51 1600 0 1600 --**AGDTYLER** 08/28/85 1200 23.5 7.3 6.7 21 20 0.000 299 9 100 38.0 ---78 2100 3 0 2200 ---**AGDTYLER** 09/11/85 1115 19.5 7.2 31 354 6.1 24 ---10 50 27.0 \_\_\_ 2200 \_\_\_ 6 Ø \_\_ **AGDTYLER** 10/02/85 800 17.5 6.9 3. ≥ 26 18 Ø. ØØØ 289 14 100 15.0 \_--1200 70 2 0 1300 \_\_\_ **AGDTYLER** 11/13/85 900 6.0 6.8 8.1 28 35 0.000 376 11 160 19.0 ---2000 120 2 0 2100 AGDTYLER 12/03/85 1800 12.5 7.0 3.7 36 58 0.000 587 100 12 64.0 190 2100 85 2 0 2200 AGDTYLER 01/16/86 1245 11.0 €.9 4.6 38 48 9 ---47E 120 35.0 3500 83 8 0 3600 **AGDTYLER** 06/11/86 915 19.5 7.3 7.9 Э 10 0.000 158 768 AMERICAN 07/21/83 945 17.0 7.3 10.0 2 1 35 2 1.2 \_\_\_ 230 3 21 Ø 230 5000.0 1 **AMERICAN** 08/18/83 1400 19.0 7.3 10.1 2 1 36 2 --Ø 1.2 210 2 230 4500.0 1 16 **AMERICAN** 09/13/83 1000 19.5 7.2 9.2 2 1 39 Ø 2 \_\_\_ 21 4000.0 1.0 220 220 AMERICAN 10/04/83 1215 20.0 7.1 9.1 2 1 ---42 5 1.8 110 160 Ø (2) 170 3500.0 1 11 AMERICAN 11/01/83 1205 17.0 2 7.1 9.0 1 ---4 i n 2 5 1.2 110 150 Ø 0 150 2500.0 AMERICAN 12/06/83 1025 11.0 9

46

1

12

2.3

270

1100

Ø

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8570.0

7.2

11.8

					DEL	-18 06	4L I I	HOPELI	5 PRUBRI	HIM MILIN	TICKTUE	DHIH							
	,														1 FORMA			TIAL	
Station	Date	Time	Temp.	Ηά	D. O.	N1	ci	C	EC	T	0-1	TO0	00000	CH	CH	CH	CH		
Name	Date	(PST)	(oC)	hu		Na mg/L		Se \	(uS/cm)	(TU)	Color (CU):	TOC	Asbēst.					TTHMEP	
								•	(us/em/ =======				(mF/L)						(cfs)
AMERICAN	01/10/84	1130	9.0	7.0	11.9	2	1		50	10	10	1.1	2200	200	 4	a	0	200	8380.0
AMERICAN		1220	9.5	7.1	11.9	5	ź		53	4	5	1.0	490	- 500	. 4	- 0	Ø	200	3080.0
AMERICAN	03/07/84	1030	9.5	7.3	11.6	5	1		57	. 3	2	1.3	260	260	17	.0	2	280	3980.0
AMERICAN	04/04/84	1035	11.0	7.1	11.4	. 5	1		55	5	2	1.2	190	200	ີ່ຮ	و .	Ø	200	4370.0
AMERICAN	05/02/84	810	12.5	7.1	11.7	2	1		54	1	2	1.3	18	160	4	ø	Ø	160	2440.0
AMERICAN	06/06/84	1045	15.0	7.3	10.3	2	2	-	52	. 3	2	1.0	12	270	10	1	Ø	280	4070.0
AMERICAN	07/10/84	950	18.0	7.3	9.4	ē	ī	- <u></u> -	.48	1	0	1.2	18	290	4	ø	0	290	4920.0
AMERICAN	08/01/84	1050	19.5	7.2	9. 1	2	1		46	: 1	2	1.2		310	4	~ Ø	Ø	310	4890.0
AMERICAN	09/05/84	915	22.0	7.2	8.6	2	î		51	1	2	1.3		320	5	ø	121	320	1470.0
AMERICAN	10/04/84	1130	19.5	7.1	9.1	. 5	1		42	. 2	5	1.2		160	5	ø	Ø	160	2220.0
AMERICAN	11/08/84	1120	16.0	7.0	9.3	2	2		51	11	15	3.2		- 280	. 5		0	2 6	
AMERICAN	12/05/84	1120	11.0	7.3	11.2	2	2		59	. 6	5	1.5	110 -		- 3	Ø	Ø	280	1730.0 5020.0
AMERICAN	02/13/85	1320	10.0	7.3	11.9	2	2		63	. 2	15	1.0	110	180 230	•	. 0		180	
AMERICAN	03/13/85	1215	12.0	7.3	11.2	5	2		63	5	<u> </u>		82	- 230	6		0	240	1740.0 1280.0
AMERICAN	04/10/85	1130	14.5	7.3	10.5	3	2		67	2	0		. <del>-</del> -	41.5	6		0	190	
AMERICAN	05/08/85	1120	14.0	7.3	10.7	1:3	2	0.000	-62	1		,	* = <del>-</del>	180 240	3	2	20	240	1270.0 3730.0
AMERICAN	06/12/85	1200	18.5	7.3	9.9	2	: 2		60	. 2	. 0			290	5	: 1	- 0	300	2800.0
AMERICAN	08/14/85	1115	20.0	7.2	9. 1	2	. 2		56	1	2	1.5		210	8	. i	20	220	3350.0
AMERICAN	10/09/85	1130	16.5	7.2	9.2	5	2	0.000		1	0	1.4		180	. 5	Ø	0	180	1460.0
AMERICAN	12/03/85	2030	12.5	7.2	10.5	- 3	. 5		64	Ė	5	2.0	70	260	6	ő	0	270	1440.0
AMERICAN	03/11/86	1315	12.0	7.1	12.0	- 2	1		56	76	-25	3.3		370	. 5	Ø	20	380	
AMERICAN	04/17/86	1130		7.3	11.2	2	1	. a. aaa		- 6	15	1.4		300	5	0	Ø.	300	
AMERICAN	05/13/86	1145		7.3	10.0	. 2	2	0.000		: 3	25	1.4		190	6	1	Ø	200	
AMERICAN	06/11/86		16.5	7.3	10.0	2	5	0.000		3	15	2.7		190				حيونوا	
CACHE	01/31/84	1045	11.5	8.3	12.4	85	88		976	13	. 8	5.5	980	300	85	31	- 2	420	
CACHE	02/22/84	1055	12.5	8.1	10.4	82	82		896	76	15	6.4	2500	360	87	26	1	470	
CACHE	03/14/84	1030	16.5	8.1	8.4	79	80		897	14	15	7.6	650	- 270	82	27	Ø:	380	
CACHE	04/11/84		15.5	8.6	10.1	59	57		720	20	10	8.0	1700	500	81	18	(2) (2)	- 500	
CACHE	05/23/84		21.0	8.3	9.0	36	: 34		-						-		_		
CACHE	06/13/84	815	19.0	8.2	8.5	. 42	42		488 595	34 52	30 30	6.7	1100	57Ø	63 83	8	Ø	640	
CACHE	07/11/84	900	24.5	8.3	8.5	36	34	, <del></del>	541			7.0	4000	750		. 8	-	850	
CACHE	08/22/84	1040	21.5	8.1	7.5	32	29			46	25	8.4	1400	800	<b>64</b>	4	0	870	
CACHE	09/12/84	1100	23.0	8.1	8.9	39	- 3B		495 577	92	50	7.1		600	- 51	- 4	0	660	
CACHE		930					42	0.001	577	20	.30	8.4		630	64	5	Ø	700	
CACHE	10/11/84		19.5	8.2	7.8	44			594	29	25	-6.0		850	. 69	6	Ø	920	
CACHE	11/15/84 12/06/84	950	12.5	7.4	7.7	38	.38	Ø. ØØØ		95 50	30	9.0	7000	730	47	4	0	780	
CACHE	04/10/85	935	10.5 16.0	7.9 8.3	8.8 9.5	64	64	0.001	744	50	50	8.5	3200	720	87	10	0	820	
CACHE	05/08/85	935			9.4	63	62	0.001	713	24	10			640	88	16	0	7.40	
CACHE	05/29/85		16.5 17.5	8.4 8.4	9.5	44 36	38 33	0.001	560 - 512	28 22	25			760	77 	. E	- 0	840	
CACHE	06/12/85	1000	24.0	8.1	7.1	35	33	0.001	499	50	20		<del></del>	870		5	 Ø	920	
CLIFTON	07/26/83	1135	21.0	7.3	7.9	20	22		208	22					43	- 7	Q Q	*	
CLIFTON	08/23/83	1000	21.5	7.3	7.7	27	31				- 8	3,2		310	-42 70		_	360	1481.0
CLIFTON									283	20	8	3.1		360	72	12	0	440	2242.0
	09/14/83	1035	22.5	7.3	. 7.8	17	17		180	11	.10	3.3		330	-23	4	. 0	360	0.0
CLIFTON	10/12/83 11/08/83	910 945	20.0 16.0	7.1 7.3	8.3 8.5	12	13		137	: 12	-12	2.8	530	310	-27	2	(0	340	0.0
CLIFTON	12/13/83		12.0	7.1	9.6	33 16	36 16		324 171	10	20 25	3.3	910	270	63	17	- Ø	350	652.0
CLIFTON	01/24/84	940	10.0	7.3	10.8	22 -	55			13		2.9	510	380	30	3	. 0	410	0.0
CLIFTON	02/28/84		13.0	7.5	10.8	39.	42		226 389	12	.25	3.1	510	300	39	6	0	340	0.0
CLIFTON	03/27/84	945	16.5	7.4	9.4	35	42 40		362	7 10	18	3.1	410	280	67 .70	18	Ø Ø	360	2367.0
CLIFTON	04/25/84	1040		7.4	9.3		30				25	3.8	480 ,	380	79 50	17	_	480	2452.7
CLIFTON	05/30/84	1040 820	16.5 24.0	7.1	7.4	27 29	33		288 307	12 19	15 20	3.8	890	320	56. 67	13	Q1 Ø	390	4199.1
CLIFTON	Ø6/27/84		25.5	7.2		50:	- 56 ·					4.9	650	420	-	15		500	2779.4
GEAT TON	<b>6</b> 6767764	249 ·	- E.J. 1)	7.6	6.3	าด (	JD		472	28	30	5.4	500	350	110	31	1	490	2994.7

Note: -- = no data

#### DELTA HEALTH ASPECTS PROGRAM MONITORING DATA

					DEL	- 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1			5 / KOOK		2   0   1   1	, prilly		THM	1 FORMA	ATION A	OTEN:	TIAL	
														CH	CH	CH	CH		
Station	Date		Temp.	, pH	D. O.	Na	C1	Se	EC		Color		Asbest.			Br2C1			
Name		(PST)			<b>\</b>				(uS/cm)		(CU)	(mg/L)				- ug/L		)	(cfs)
CLIFTON	07/25/84	940	24.0	7.5	8.6	18	21	0.000		18	 25	4.4	960	420	52	 8	: 0	480	4753.7
CLIFTON	08/29/84	815	24.5	7.3	7.6	20	23		222	11	15	3.2		390	54	10	ø	450	3827.1
CLIFTON	09/27/84	1040	22.0	7.5	8.3	24	24	0.000	261	Ê	15	3.2		390	49	12	ø	450	1704.6
CLIFTON	10/25/84	1045	17.0	7.5	10.0	27	29		284	7	18	3.4		300	54	14	ø	370	0.0
CLIFTON	11/29/84	1245	12.0	7.3	10.2	20	21		233	11	30	3.7		460	48	6	ø	510	2400.0
CLIFTON	12/12/84	1055	11.5	7.3	10.0	21	22	0.000	252	16	35	4.7	420	390	52	5	é	450	5150.0
CLIFTON	01/30/85	925	7.0	7.1	10.5	32	37	0.000	348	8									2500.0
CLIFTON	02/27/85	1100	13.0	7.3	9.8	26	28	0.000	303	14	40			410	64	8	Ø	480	4200.0
CLIFTON	03/27/85	1030	12.5	7.4	9.6	33	34	0.000	334	8			670						3620.0
CLIFTON	04/24/85	1030	18.0	7.6	9.6	24	24	0.000	277	8	8			470	56	7	<b>Ø</b> 1	530	4200.0
CLIFTON	05/22/85	930	21.5	8.1	9.2	25	29	0.000	264	21	15			610	65	11	Ø	690	2490.0
CLIFTON	06/26/85	915	24.5	7.5	7.7	37	40	Ø. ØØØ	314	17	15			550	88	24	1	660	5290.0
CLIFTON	07/10/85	900	25.5	7.5	6.5	43	50	0.001	386	15									4500.0
CLIFTON	08/28/85	1000	23.5	7.4	7.7	51	69	0.000	458	10	10	4. Ø		460	110	47	3	620	5770.0
CLIFTON	09/25/85	940	22.5	7.4	6.6	64	80	0.000	602	12									3000.0
CLIFTON	10/23/85	915	17.5	7.5	8.9	52	77	0.000	484	9	10	2.3		330	130	59	4	520	3490.0
CLIFTON	11/15/85	1045	12.0	7.4	10.2	92	143	0.000	679	12									1800.0
CLIFTON	12/03/85	1305	12.0	7.4	10.1	98	162	0.000	744	10	8	3.7	230	310	220	170	13	710	5960.0
CLIFTON	01/23/86	1045	11.5	7.3	9.0	48	60	0.000	410	8									5510.0
CLIFTON	02/13/86	950	11.5	7.3	10.4	41	55	0.000	423	17									
CLIFTON	03/04/86	1045	16.5	7.3	7.8	29	29	0.001	306	21	20	8.0		520	64	7	Ø	590	
CLIFTON	04/09/86	1100	16.5	7.2	8.8	20	20	0.000	197	14	20	3.9		570	62	5	0	640	
CLIFTON	05/07/86	850	15.5 20.5	7.3	8.8 8.2	27 29	28 33	0.001 0.001	28Ø 3Ø3	13 26	20	6.3		350 140	51 28	7 6	Ø Ø	410 170	
CLIFTON CLIFTON	06/04/86 07/02/86	945 920	24.5	7.3 7.3	6.5	55	66	e. wei	534	11		10.0		140			<b>₩</b> ,	170	
COSUMNES	07/21/83	830	22.5	7.3	8.5	3	2		67	1	2	1.0		200	6	0	Ø	210	257.0
COSUMNES	08/18/83	1255	28.0	7.7	8.3	4	5		85	i	5	1.2		190	9	ø	ø	200	102.0
COSUMNES	09/13/83	900	25.0	7.3	7.8	4	2		90	1	2	1.2		210	ē	ø	ø	220	76.0
COSUMNES	10/04/83	1105	21.5	7.3	8.9	4	2		80	2	5	1.2	140	150	6	ē	ø	160	102.0
COSUMNES	11/01/83	1110	18.0	7.3	9.3	4	2		82	9	8	1.6	180	170	5	ē	ø	180	378.0
COSUMNES	12/06/83	935	8.5	7.2	12.0	7	2		81	7	18	2.4	230	830	7	Ø	Ø	840	1420.0
COSUMNES	01/10/84	1030	8.0	7.2	11.8	3	2		78	4	8	1.0	300	160	4	0	Ø	160	1230.0
COSUMNES	02/01/84	1115	9, 5	7.0	11.5	4	2		93	2	5	0.9	18	140	. 5	Ø	0	140	561.0
COSUMNES	03/07/84	935	11.5	7.3	11.4	4	2		86	1	5	1.3	91	190	11	Ø	Ø	200	766.0
COSUMNES	Ø4/Ø4/84	940	14.0	7.1	10.7	3	2		80	1	5	1.6	95	200	9	Ø	Ø	210	794.Ø
COSUMNES	05/02/84	720	14.0	7.3	10.6	4	1		76	1	2	1.0	25	130	5	Ø	Ø	140	597.0
COSUMNES	Ø6/Ø6/84	950	19.0	7.3	9. 1	3	2		74	2	5	1.2	33	230	11	1	Ø	240	294.0
COSUMNES	07/10/84	900	27.5	7.7	7.6	4	2		86	2	5	1.6	10	240	. 9	0	Ø	250	74.0
COSUMNES	08/01/84	1003	27.0	7. E	8.1	4	2		93	1	10	2.1		320	9	Ø	Ø	330	48.0
COSUMNES	09/05/84	820	25.5	7.3	7.1	4	. 2		96	1	5	2.0		300	11	Ø	Ø	310	
COSUMNES	10/04/84	1025	21.0	7.4	9.0	4	2		90	. 2	2	1.5		160	7	0	0	170	
COSUMNES	11/08/84	1015	13.5	7.2	10.2	4	2		82	12	25	2.5		280	6	Ø	0	290	
COSUMNES	12/05/84	1040	10.5	7.3	11.3	5	4		129	2	8	2.2	9	280	9	0	0	290	
DMC	07/26/83	1045	23.0	7.3	7.5	33	38		322	31	5	3.6		290	54	10	0	350	4723.0
DMC	08/23/83	905	21.5	7.3	7.7	28	31		283	22	5	3.2		400	59 26	9 4	Ø	470 340	3573.0 3245.0
DMC DMC	09/14/83 10/12/83	940 835	21.0 18.5	7.3 7.3	7.8 8.5	18 14	18 15		188 151	19 18	12 12	2.4 3.2	~- 76ø	310 200	26 26	2	Ø	340 230	2439.0
DMC	11/08/83	915	16.5	7.2	8.2	37	39		361	11	50	3.4	1100	270	48	14	Ø	330	153.0
DMC	12/13/83	1035	12.0	7.2	9.5	37 23	26		238	18	35	3.5	570	320	37	6	Ø	360	3725.0
DMC	01/24/84	915	10.5	7.3	10.7	30	33		297	16	35	3.2	1600	340	52	11	ø	400	1198.0
DMC	02/28/84	1025	12.5	7.5	10.0	42	48		397	11	18	3.1	370	280	76	25	1	380	4309.0
DMC	03/27/84		16.0	7.3	9.5	53	60		511	24	15	3.8	700	270	90	35	è		4402.0

THM FORMATION POTENTIAL

			e := r .											TH		HIIUN		IHL	
Station	Date	Time	Temp.	рH	D. O.	Na	C1.	Se	EC	Turb.	Color	TOC	Asbest.	CH.	CH.	CH: Br2Cl	CH English	TUMED	FLOW
Name	2002	(PST)	(oC)	р.,	ζ				(uS/cm)	(TU)	(CU)	(mg/L)		₹ <u></u> -					(cfs)
=========		=====								2							=====	.=====	======:
DMC	04/25/84	955	15.5	7.5	9.3	60	68		552	18	10	4.7	1800	300	120	45	2	470	4071.0
DMC	05/30/84	750	23.5	7.4	7.6	29	33		298	24	20	4.7	380	380	66	14	2	460	2390.0
DMC	06/27/84	905	25.5	7.3	6.0	32	35		328	30	35	5.0	730	380	70	15	2	460	3313.0
DMC	07/25/84	910	24.0	7. 7	7.4	58	73		554	28	15	4.4	1100	450	150	57	4	660	4688.0
DMC	Ø8/29/84	740	24.5	7.3	7.3	21	55		229	16	18	3.7		330	48	. 9	Ø	390	3027.0
DMC	09/27/84	1005	22.0	7.4	8.2	28	29	0.000		13	15	3.8		330	55	12	0	400	3150.0
DMC	10/25/84	1000	16.0	7.8	9.8	25	26	Ø. 000		8	20	3.3		360	66	12	Ø	440	3959.0
DMC	11/29/84	1215	11.0	7.4	10.2	32	34	0.000		9	25	4. 1		400	64	12	Ø	480	3901.0
DMC	12/12/84	1015	11.5	7.2	9.3	31	32	0.000		18	25	4.9	590	370	60	8	Ø	440	4004.0
DMC	01/30/85	850	7.5	7.3	10.6	38	44	0.001		7									4003.0
DMC	02/27/85	1015	13.0	7.5	9.9	31	34	0.000		11	35	77.		410	75	12	0	500	4221.0
DMC	03/27/85	945	12.0	7.4	9.8	29	31	0.000		8		***	980						3144.0
DMC	04/24/85	1000	17.5	7.5	9.5	25	24	0.000		9	5	<del></del>		340	57	5	Ø	400	3997.0
DMC	05/22/85	900	20.5	8.3	9.1	25	29	0.000		22	20		<b></b> .	550	71	10	0	630	3136.0
DMC DMC	06/26/85	830	24.5	7.6	7.1	78	95	0.001		23	10			580	180	9	10	860	2877.0
DMC	07/10/85 08/28/85	920 920	24.5 23.0	7.4 7.4	6.7 7.7	59 50	68 74	0.001 0.000	54 <u>4</u> 441	24 17	20	9.7		410	120	70		600	4578.0 4160.0
DMC	09/25/85	915	22.5	7.5	6.8	66	85	0.001		15	20	3. / 		410	120	70	3		3980.0
DMC	10/23/85	840	16.5	7.4	7.2	60 60	79	0.000		13	5	3.6		270	110	58	5	440	3890.0
DMC	11/15/85	1015	12.0	7.4	10.5	68	106	0.000		11		J. 0	==,.	E 7 6	110	<u>تار</u> ت		~	4040.0
DMC	12/03/85	1305	12.0	7.4	10.1	72	117	0.000		10	15	6.3	370	360	190	120	6	680	3940.0
DMC	01/23/86	1000	11.5	7.3	8.8	52	63	0.000		8									3310.0
DMC	02/13/86	915	11.5	7.5	10.2	44	60	0.000		16									
DMC	03/04/86	1015	16.5	7.3	7.9	29	28	0.001		25	25	7.8		580	€1	6	Ø	650	
DMC	04/09/86	945	16.0	7.3	9.0	23	27	0.000		55	25	4, 2		600	58	7	ō	660	·
DMC	05/07/86	815	16.0	7.2	8.3	27	28		278	15	10	6.2		260	40	5	Ø	300	
DMC	06/04/86	900	21.5	7.3	7.7	36	48	0.000	2	31		9.5		250	54	8	Ø	310	
DMC	07/02/86	845	24.5	7.3	7.0	54	62	0.001		13									
BANKS	03/30/82	900	12.5	7.3	9.7	38	35		315	9				930	66	7	Ø	1000	6300.0
BANKS	06/29/82	720	20.0	8.0	8.3		41	, <del></del>	322	1.1		-		490	83	14	Ø	590	240.0
BANKS	08/26/82	905	21.0	7.9	8.3		19		213	19				430	34	4	Ø	470	4240.0
BANKS	10/21/82	845	18,5	7.2	8.0		23		212	€				370	45	7	Ø	420	2779.0
BANKS	12/29/82	1200	10.0	7.1	9. 7		23		225	9		į——		630	49	4	0	680	645.0
BANKS	02/24/83	1210	14.0	7.4	9.3		30	··	288	10				190	26	4	0	220	6119.0
BANKS	04/27/83	910		7.3	8.4		42		367	6				360	. 69	10	E	440	125.0
BANKS	06/22/83	830	20.5	7.2	8.4		1.4	,	143	11				350	28	4	Ø	380	2262.0
BANKS	07/26/83	1000	23.0	7.3	8.3	21	22		211	17	В	2.8		300	38	6	0	340	1306.0
BANKS	08/23/83	830	22.5	7.3	8.0	25	28		261	17	В	3.5		420	58	9	Ø	490	2179.0
BANKS	09/14/83	850	22.0	7.3	7.0	55	24	, <del></del>	226	8	20	2.9		330	38	8	0	380	61.0
BANKS	10/12/83	755	20.5	7.3	7.6	53	56	, <del></del> -	219	6	20	3.1	860	260	47	8	•	320	306.0
BANKS	11/08/83	850	16.5	7.2	8.6	19	20	- <del></del> -	186	7	25	2.8		310	40	7	Ø	360	1154.0
BANKS	12/13/83	940	12.0	7.3	10.2	32	34		305	13	40	3.3		360	42	7	Ø	410	326.0
BANKS	01/24/84	850	9.5	7,3	11.2	26	28		252 200	5 5	20 20	2.9 3.2	21	320	44 75	8 20	Ø Ø	370 400	267.0 2563.0
BANKS	02/28/84 03/27/84	940 840	12.0 16.5	7.5 7.3	10.0 9.8	42 36	4 <u>6</u> 40		388 370	. 5 20	20 30	3.2 4.2		310 460	80	16	, e	560	104.0
BANKS	04/25/84	915	15.0	7.3	9.3	27	30		283	37	25	3.9		570	62	12	20	640	3925.0
BANKS	05/30/84	725	23.0	7.5	7.1	29	33		304	16	12	4.7		400	72	18	ø	490	1865.0
BANKS	06/27/84	820	24.5	7.3	6.6	24	34		364 258	29	40	4.7		410	7 <i>5</i> 59	. 8	Ø	480	2884.0
BANKS	07/25/84	830	23.0	7.4	8.1	20	23	- n <u></u>	214	16	20	4.7		420	57	9	é	490	4359.0
BANKS	08/29/84	715	23.0	7.3	7.4	55	24		244	7	18	3.1		360	55	10	0	420	3438.0
BANKS	09/27/84	925	22.5	7.3	8.6	25	25	0.000		7	15	3.3		370	55	10	ø	440	1723.0
BANKS	10/25/84	920	16.5	7.7	9.3	25	56	0.000		á	20	2.9	and the second second	300	59	9	ø	370	903.0
201110	10/10/04		10.0	, , ,	٦. ٦		LO	0.000		٠	0	L. 7		250		-		J. J.	200. C

Note: -- = no data

## DELTA HEALTH ASPECTS PROGRAM MONITORING DATA

																ATION F		TIAL	
Station	Date	Timo	Tono	-u	D. O.	Nin	C1	Se	cc	T	C-1	TOC	0	CH	CH	CH	CH	TTUMES	F1. O
Name	Date	Time (PST)	Temp.	,pH	(	Na. mg/l	C1		EC (uS/cm)		Color (CU)	TOC (mg/L)	Asbest. (mF/L)			- ug/L		TTHMFP	rwow (cfs)
========								•											
BANKS	11/29/84	1130	11.5	7.5	10.5	20	21	0.000	233	11	30	3.3		430	44	6	Ø	480	2797.0
BANKS	12/12/84	945	11.5	7.3	10.0	23	24		263	10	25	4.3		380	50	6	Ø	440	4258.0
BANKS	02/27/85	945	13.5	7.5	9.5	30	33	Ø. ØØØ	335	8	35			310	71	10	Ø	390	4151.0
BANKS	03/27/85	900	12.5	7.4	10.1	36	38	0.000	367	11			520						3486.0
BANKS	04/24/85	915	17.5	7. E	8.7	36	34		351	11	5			410	81	17	0		4520.0
BANKS	05/22/85	815	19.5	8.1	8.6	35	41	Ø. ØØØ	351	26	5	***		580	90	17	20	690	1917.0
BANKS	06/07/85	850	23.5	7.5	7.4	32	37		355	30									2619.0
BANKS BANKS	06/26/85	800	23.5	7.7	7.5	38	46		370	32	20			550	110	24	_ 1		5222.0
BANKS	07/10/85	800	24.5	7.5	7.5	42	48	0.000	343	16	15			590	160	35	~ 2		4572.0
BANKS	08/28/85 09/25/85	830 820	22.5 22.5	7.4 7.5	7.8 7.9	54 69	78 102	Ø. ØØØ Ø. ØØØ	466 588	10 6	10 10	6.4		390	140	69	5		5260.0
BANKS	10/23/85	800	17.0	7.6	8.9	59	94	0.000	527	7	5	2.7 4.0		340 290	89 150	40 90	10 13		3020.0
BANKS	11/15/85	930	12.0	7.4	9.5	71	112	0.000	586	6	10	2.9		260	160	100		540	3200.0 2150.0
BANKS	12/03/85	1415	11.5	7.4	10.1	85	141	0.000	676	10	10	3.6	230	240	210	150	10		6320.0
BANKS	01/23/86	920	12.0	7.3	9.2	56	79	0.000	482	12	25	7.2		1700	170	47	2		5170.0
BANKS	02/13/86	845	11.5	7.7	10.5	45	61	0.000	444	17	25	8.6		780	140	es.	1		
BANKS	03/04/86	930	16.5	7.3	8.2	30	33	0.000	332	14	30	5.8		600	70	6	ø		
BANKS	04/09/86	915	17.5	7.5	9.4	29	31	Ø. ØØØ	265	13	20	5.0		630	76	10	Ø	720	
BANKS	05/07/86	745	15.5	7.3	8.9	28	31		284	11	15	5.0		460	74	10	Ø	540	
BANKS	<b>06/04/86</b>	815	19.5	7.5	8.6	31	38	0.001	312	32				340	45	9	Ø	390	
BANKS	07/02/86	805	24.0	7.3	6.4	31	33	Ø. 000	305	25									
HONKER	<b>0</b> 2/23/83	1045	13.0	7.3	8.9		27		233	13				210	33	6	0	250	
HONKER	Ø4/27/83	1030		7.3	8.8		33		303	9				300	72	10	5	390	
HONKER	06/22/83	1000	23.5	7.3	7.6		20		184	11				370	43	7	0	420	
HONKER	08/17/83	1000	24.5	7.3	7.1	8	8		126	6	8	2.5		310	25	5	Ø		
HONKER	10/04/83	700	20.5	7.3	8.0	. 7	7		114	6	12	2.1	190	290	14	1	0		
HONKER	12/06/83	820	10.0	7.2	10.0	17	26		232	18	60	6.4	620	520	47	7	0		
HONKER	02/01/84	755	10.0	7.1	9.7	27	32		302	11	25	5.8	380	450	68	10	Ø		
HONKER	04/04/84 06/06/84	815 740	15.0 19.0	7.3 7.5	9.6 7.6	12	14		171	9	12	3.0	500	310	32	4	0		
HONKER	Ø8/Ø1/84	702	23.0	7.3	7.8	13	12		178	10	10	3.8	260	340	40	7	0		
HONKER	10/04/84	750	18.5	7.3	8.8	11 7	12 5		166	8 5	15 5	2.8		460	34	4	0		
HONKER	12/05/84	85Ø	10.5	7.2	9.8	12	15		120 184	13	35	1.8		240	14	1	0		-
DVGH	08/10/83	1200	23.5	8.5	8.4	19	16		466	13	33 5	5.0 3.2	770	480 310	37 32	4 4	0		
DVGH	08/10/83	1145	12.5	7.8	3.9	14	11		395	3	2	2.9	140 744	360	26	2	21		
DVSR	09/20/83	720	14.5	7.3	5.3	15	12		414	5	8	2.9		450	16	2	0		
DVSR	10/18/83	1150	18.0	8.0	7.0	17	13		430	1	8	2.9	54		10			470	
DVSR	11/21/83	1150	15.5	7.9	8.4	18	15		469	4	15	3.6	310	230	29	4	0		
DVSR	03/11/86	845	13.0	8.1	11.3	14	12	Ø. ØØØ	322	90	30	6.6		660	33	i	0		
DVSR	05/13/86	700	16.0	8.2	6.4	15	11	0.000	356	4	20	4.8		510	24	2	. 0		
LINDSEY	Ø7/11/84	940	24.5	8.4	6.7	37	29		426	36	35	6.3	2700	770	57	6	2		
LINDSEY	08/22/84	1105	21.5	8.0	7.6	35	26		411	65	50	7.1		950	65	4	2		
LINDSEY	09/12/84	1155	22.5	7.6	7.0	34	25	0.000	424	27	50	7.5		930	59	3	121		
LINDSEY	10/11/84	950	19.5	7. B	8.0	32	21		383	28	50	5.6		840	59	4	0		
LINDSEY	11/15/84	1045	12.5	7.5	8.6	31	23	0.000	353	28	25	4.7		570	45	Š	ō		
LINDSEY	12/06/84	1050	11.0	7.3	8.3	44	34	0.000	441	37	50	9.7	3500	1000	59	ē	ē		
LINDSEY	Ø1/25/85	1045	6.0	7.4	9.2	56	46	0.000	558	12									
LINDSEY	02/13/85	1150	10.5	7.3	6.7	43	35	0.000	381	110	50			1200	65	3	Ø	1300	
LINDSEY	<b>0</b> 2/22/85	1030	11.0	7.4	8.6	57	39	0.000	445	65									
LINDSEY	03/13/85	1145	12.5	7.6	9.1	51	41	0.000	482	60			7500						
LINDSEY	04/10/85	1015	18.0	7.7	8.6	61	44	0.000	531	20	15		-,-	580	86	9	Ø	680	
LINDSEY	05/08/85	1000	17.0	8.1	8.8	60	47	0.000	574	18	20			660	88	4	(2)		

THM FORMATION POTENTIAL

CH CH CH CH Station Date Time Temp. pH D.O. Na Ci Se EC Turb, Color TOC Asbest. C13 BrC12 Br2C1 Br3 TTHMFP FLOW (PST) (oC) Name (-----) (uS/cm) (TU) (CU) (mg/L) (mF/L) <----- ug/L -----> (cfs) LINDSEY 05/29/85 1030 20.0 7.9 8.6 55 47 \_\_\_ 571 27 LINDSEY 06/12/85 1045 25.0 7.9 51 45 0.000 30 900 97 7.1 541 28 --6 0 1000 07/24/85 610 36 LINDSEY 22.0 7.6 7.0 40 33 Ø. ØØØ 421 ---------\_\_\_ ---\_\_\_ \_\_\_ ~--LINDSEY 08/14/85 955 750 21.0 7.8 8.6 38 32 0.000 405 48 30 8.2 \_\_\_ 69 5 ø 820 ---LINDSEY 09/11/85 900 19.5 7.7 7.5 40 37 0.000 443 30 25 9.8 --820 54 4 Ø 880 LINDSEY 10/09/85 1005 16.5 7.6 42 41 מממ. 496 31 17.0 1500 0 1600 8.1 38 \_\_\_ 66 3 ---7.5 LINDSEY 11/19/85 820 8:5 10.0 40 37 0.000 442 18 15 7.7 \_--\_\_ \_\_\_ \_--------LINDSEY 12/03/85 720 11.5 7.4 8.7 56 6.3 0.000 569 25 60 15.0 1160 1300 70 2 0 1400 LINDSEY 01/16/86 745 15.0 2200 0 2300 10.5 7.3 6.7 65 58 0.000 458 38 80 \_---56 2 790 LINDSEY 02/27/86 750 16.5 €.8 3.0 21 16 0.000 208 46 60 10.0 \_\_ 35 Ø 0 820 LINDSEY 03/13/86 730 13.5 7.1 6.2 23 20 0.000 221 68 1300 47 0 1300 100 15.0 - 1 LINDSEY 04/23/86 730 18.5 7.6 5.3 44 39 0.000 387 48 70 12.0 \_\_\_ 1100 84 6 Ø 1200 LINDSEY 05/28/86 600 20.0 8.0 6.0 52 47 0.000 528 26 25 \_\_\_ 380 38 5 2 420 8.0 ---LINDSEY 06/25/86 635 2105 8.0 7. 2 43 37 0.000 4F.1 38 ലമ 22.0 \_\_ \_\_\_ \_-\_\_\_ LCONNECTSL 02/06/85 845 7.0 7.4 11.2 20 22 252 5 15 Ø ---660 46 6 710 \_--\_\_ ----LCONNECTSL 03/06/85 915 18 \_--11.0 7.4 10.0 14 218 7 ------140 \_\_ \_\_\_ \_\_ \_\_ LCONNECTSL 04/05/85 815 17.5 7.3 9.5 13 11 \_\_\_ 188 5 \_\_\_ \_--230 26 2 Ø 260 LCONNECTSL 05/01/85 800 19.0 7.4 9. 1 13 11 0.000 175 5 5 \_--\_\_\_ 280 27 0 310 \_\_\_ LCONNECTSL 06/05/85 745 20.5 7.5 8.7 10 7 Ø 330 13 \_\_\_ 180 5 300 26 2 LCONNECTSL 06/07/85 700 23.0 7.7 8.7 9 --178 7 -13 --\_\_ \_\_ ---LCONNECTSL 08/01/85 800 22.5 7.4 8.0 13 10 5 -3.B 360 32 2 Ø 390 --186 10 LCONNECTSL 10/02/85 640 20.0 7.5 7.8 \_\_\_ 209 .5 240 26 Ø 270 18 11 4 3.1 3 LCONNECTSL 11/13/85 730 11.5 7.3 9.0 12 11 \_\_\_ 183 3 25 3.4 340 34 Ø 380 --LCONNECTSL 12/03/85 1645 11.5 733 10.2 15 15 --204 5 15 6. A 68 380 36 3 Ø 420 ---LCONNECTSL 03/11/86 1145 14.5 7.3 9.0 12 19 ----192 22 25 17.0 ---650 -51 3 Ø 700 ---7 ø 500 --LCONNECTSL 04/17/86 945 15.5 7.2 8.5 17 20 0.001 195 11 20 4.2 440 51 LCONNECTSL 05/13/86 945 19.5 7.3 8.4 12 15 ---162 14 25 4.2 \_\_ 150 16 2 Ø 170 ---LCONNECTSL 06/11/86 745 9 8 25 5.8 \_\_\_ 21.5 7.3 7.9 ---136 12 \_\_\_ \_-----MALLARD 07/28/83 1045 24.2 7.3 8.6 11 11 ,... 137 18 5 3.3 . ----260 26 3 Ø: 290 \_\_ MALLARD 950 21.0 Ø 08725783 7.6 8.0 21 27 \_\_ 216 19 15 3.4 \_\_\_ 300 65 13 380 MALLARD 09/20/83 900 21.0 7.3 7.7 15 16 181 13 15 3.4 410 21 3, Ø 430 \_\_ MALLARD 10/18/83 910 17.5 7.3 8.5 13 13 152 q 3Ø 3.2 690 MALLARD 11/21/83 1005 12.5 7.2 9.5 15 16 \_\_\_ 1:80 16 40 4.5 1400 170 36 4 Ø 210 MALLARD 12/28/83 930 10.0 7.3 - 13 13 168 38 30 3.7 26000 390 30 5 0 430 ---10.3 MALLARD 750 7.7 02/13/85 11.5 11.9 96 155 0.000 749 12 25 \_------250 190 130 28. 570 MALLARD 03/13/85 815 320 558 0.000 \_\_\_\_ 1300 <u>--</u> 14.0 8.4 13.5 2160 10 \_\_\_ MALLARD 04/10/85 730 16.0 7.5 8.0 348 569 2210 25 5 90 180 260 280 810 -------MIDDLER 02/06/85 830 6.5 7.3 11.2 38 43 0.000 391 13 25 ---780 84 20 Ø 880 MIDDLER 03/06/85 900 10.0 34 0.000 339 12 7.4 10.0 31 \_\_\_ 210 \_\_ \_\_ \_--\_\_\_ MIDDLER 04/05/85 730 17.0 7.5 40 378 6 5 300 76 Ø 390 8.9 40 \_\_ \_\_ \_\_ 16 MÍDDLER 05/01/85 650 19.0 7.6 9.3 29 29 0.001 303 9 10 410 68 10 Ø 490 \_\_ MIDDLER 06/05/85 640 20.0 7.8 9.0 25 252 17 550 67 Ø 620 ---26 \_\_ 5 8 MIDDLER 06/07/85 805 23.5 7.7 8.9 23 25 ----256 16 \_\_\_ .... \_\_\_ MIDDLER 08/01/85 700 22.0 7.4 7.8 35 46 0.000 331 12 20 3.9 ---660 110 26 1 800 \_-MIDDLER 10/23/85 1115 7.5 7 550 18.0 9.4 40 61 0.000 396 10 2.2 \_\_\_ 380 120 45 2 MIDDLER 12/03/85 1215 11.5 7.4 10.3 54 83 0.000 464 8 12 4.6 100 340 160 68 5 570 MIDDLER 03/11/86 1030 14.5 7.3 8.2 3Ø 38 0.001 343 24 25 6.2 530 110 12 ø 650 \_\_ ---MIDDLER 04/17/86 730 14.0 7.3 8.8 20 86 0.001 213 12 25 3.5 \_\_ 440 60 9 Ø 510 \_\_ MIDDLER 05/13/86 830 19.5 7.3 8.1 26 30 0.000 270 13 30 4.0 480 76 Ø 570 ---\_\_ 11 06/11/86 615 28 34 0.000 272 20 7.5 MIDDLER 22.5 7.3 7.8 14 ------------230 Ø 230 1550.0 2 .3 \_\_ 3 Ø MOKELUMNE 07/21/83 715 18.0 7.2 9.6 1 \_\_\_ 34 2 1.4 MOKELUMNE Ø8/18/83 Ø 0 250 928.0 800 19.0 6.6 9.2 2 34 2 5 1.2 240 8

Note: -- = no data

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#### DELTA HEALTH ASPECTS PROGRAM MONITORING DATA

					DEL	"IH HE	#LIH	HRAFC L	S PRUGR	AM MUN	TIDKING	3 DATA							
														1HT	4 FORMA	NOITE	POTEN	TIAL	
														CH	CH	CH	CH		
Station	Date	Time	Temp.	ρН	D. O.	Na	Cl	Se	EC	Turb.	Color	TOC	Asbest.	C13	BrC12	Br2C1	Br3	TTHMEP	FLOW
Name		(PST)		•	<b>(</b> -				(uS/cm)		(CU)		(mF/L)	<b>(</b>		- ug/L		>	(cfs)
			=====:	=====							======					=====			.====:
MOKELUMNE	09/13/83	750	19.0	7.1	8.8	2	1		33	2	2	1.3		250	6	2	Ø	260	1040.0
MOKELUMNE	10/04/83	815	17.5	6.8	9.5	2	1		32	2	5	1.4	17	240	4	Ø	0	240	1210.0
MOKELUMNE	11/01/83	750	16.5	6.6	8.3	1	1		31	6	8	1.6	31	190	3	ē	ō	190	1420.0
MOKELUMNE	12/06/83	740	12.0	6.8	10.4	ā	1		38	6	8	4.6	200	190	3	ø	ě	190	2990.0
MOKELUMNE	01/10/84	925	10.5	6.9	11.0	ē	1		42	9	12	1.8	170	220	3	ø	ø	220	3790.0
MOKELUMNE	02/01/84	850	9.5	6.7	11.2	2	î		44	é	10	1.4	32	110	5	2	0	120	
MOKELUMNE	03/07/84	830	11.0	7.2	11.5	2	1		45	3	8					_			1210.0
MOKELUMNE	04/04/84	735	13.0	7.3	10.9	5	i		47		2	1.5	26	260	5	0	Ø	260	907.0
MOKELUMNE	05/02/84	625	14.0	7.2	10.7	5	1		46	2	5	1.5	44	230	5	0	0	240	439.0
MOKELUMNE			15.5							2		1.7	10	200	4	0	0	200	270.0
	06/06/84	825		7.3	10.2	2	1		47	2	2	1.5	53	230	7	Ø	Ø	240	265.0
MOKELUMNE	07/10/84	755	17.5	7.3	9.5	2	1		48	1	2	1.6	12	360	5	<b>Ø</b>	0	360	333.0
MOKELUMNE	08/01/84	820	23.5	7.2	9.5	2	1		47	1	0	1.7		310	5	Ø	Ø	320	303.0
MOKELUMNE	09/05/84	720	18.5	7.3	9.3	2	1		48	1	5	1.5		420	5	Ø	Ø	420	
MOKELUMNE	10/04/84	915	17.5	7.2	9.4	2	1		44	2	2	1.6		290	5	Ø	Ø	300	
MOKELUMNE	11/08/84	920	16. Ø	7.0	9.6	2	1		45	7	8	2.3		260	4	Ø	Ø	260	
MOKELUMNE	12/05/84	945	12.0	7.2	10.9	2	2		46	4	5	1.8	19	200	4	Ø	Ø	200	
NOBAY	07/28/83	830	21.0	7.9	9.0	10	5		301	4	5	2.7		290	15	1	Ø	310	5.0
NOBAY	08/25/83	725	19.0	8.5	8.9	10	5		301	4	5	2.7		340	26	2	Ø	370	5.0
NOBAY	09/20/83	1120	20.0	7.6	9.7	9	5		301	2	5	3. 1		350	9	Ø	21	360	5.0
NOBAY	10/18/83	720	17.0	8.9	9.5	10	5		298	2	12	3.2	200						11.1
NOBAY	11/21/83	845	11.0	7.8	10.4	11	7		312	11	25	3.0	1600	280	18	1	2	300	1.0
NOBAY	12/28/83	815	11.5	7.6	10.2	11	6		279	22	20	2.6	6000	270	17	5	Ø	290	1.0
NOBAY	01/31/84	850	11.5	8.2	11.3	12	7		322	4	8	2.6	2600	300	18	1	Ø	320	1.0
NOBAY	02/22/84	925 .	12.0	8.2	10.7	12	6		314	6	8	3.1	2900	290	18	1	0	310	0.5
NOBAY	03/14/84	850	16.0	8.3	8.2	13	€		333	4	5	3.0	1500	340	21	1	Ø	360	0.0
NOBAY	04/11/84	840	15.0	8.4	10.4	10	6		310	4	2	2.8	2000	290	18	1	Ž.	310	1.0
NOBAY	05/23/84	925	20.0	8.4	9.3	10	5		312	4	5	3.2	370	400	18	1	ē	420	1.5
NOBAY	06/13/84	640	17.5	8.5	9.5	9	5		306	1	5	2.8	1100	400	18	1	ō	420	4.0
NOBAY	07/11/84	735	19.5	7.5	9.1	9	5		308	4	- 5	2.9	1200	340	17	1	ō	360	4.5
NOBAY	08/22/84	917	19.0	8.4	9.2	10	S		314	ė	8	2.8		340	17	î	ő	360	5.0
NOBAY	09/12/84	930	19.5	8.4	9.0	. 9	5		321	2	ē	3.0		380	20	î	ø	400	4.5
NOBAY	10/11/84	815	18.0	8.2	9.1	9	5		312	3	5	2.5		470	20	î	ø	490	7.0
NOBAY	11/15/84	845	13.0	8.0	9.4	10	6		296	4	10	2.6		310	. 15	ī	ě	330	11.0
NOBAY	12/06/84	825	10.5	8.1	10.1	15	10		339	12	18	3.6	1600	400	23	í	ø	420	11.0
NOBAY	02/13/85	920	10.5	8.0	8.7	18	10	0.000	321	60	50			750	31	_	Ø	780	
NOBAY	03/13/85	930	13.0	8.3	10.0	13	8	0.000	350	4			1100	7.30		1		780	13.0
NOBAY	04/10/85	830	17.5	8.4	9.5	14	8		371	3	•								1.0
NOBAY	05/08/85	830	16.0	8.1	9.8		5		334		Ø 1Ø			260	22	2	Ø	280	4.5
NOBAY	Ø6/12/85	845	20.0		9.2	11	5 5	0.000	334 325	4				300	22	1	Ø	320	4.5
NOBAY	08/14/85	900	18.0	8.2 8.3	10.1	10 10	5			4	10			320	26	1	Ø	350	6.5
									336	2	5	3.4		250	27	1	0	280	5.5
NOBAY	10/09/85	900	16.0	8.3	9.7	9	5	0.001	330	1	5	3.2		310	20	2	Ø	330	6.0
NOBAY	12/03/85	840	11.5	8.0	10.3	10	6	0.000	320	7	5	3.9	430	300	24	1	0	320	13.0
NOBAY	Ø3/13/86	915	14.0	8.0	9.5	11	6	0.000	278	30	20	3.7		520	22	1	0	540	
NOBAY	04/23/86	1045	18.0	8.2	9. 1	13	7	0.000	336	7	10	2.7		320	24	2	Ø	350	
NOBAY	05/28/86	945	19.5	8.3	9.6	10	5	0.000	306	7	5	3. 1		300	15	1	0	320	
NOBAY	Ø6/25/86	845	19.0	8.3	9.2	9	5	0.000	293	5	10	7.2							
ROCKSL	07/26/83	1240	23.Ø	7.0	7.0	15	16		158	16	8	3.4		310	34	5	Ø	350	
ROCKSL	08/23/83	1100	24.5	7.2	6.9	15	14		171	17	8	2.6		440	35	4	Ø	480	
ROCKSL	09/14/83	1145	25.0	7.1	6.1	26	29		254	15	35	4.6		440	43	9	20	490	
ROCKSL	10/12/83	1005	21.0	7.1	7.7	17	21		177	11	20	2.8	950	270	39	6	6	320	
ROCKSL.	11/08/83	1030	17.0	7.2	8.4	22	23		224	10	25	3.5	570	260	37	7	Ø	300	
ROCKSL	12/13/83	1220	12.0	6.9	9.8	20	21		202	11	30	3.0	560	270	36	4	ē	310	
																•	-		

THM FORMATION POTENTIAL

14300.

CH CH CH CH Time Temp. pΗ Station Date D. O. Na Cl Se EC Turb. Color TOC Asbest. Cl3 BrCl2 Br2Cl Br3 TTHMFP FLOW <---- mg/L Name (PST) (oC) ----> (uS/cm) (TU) (CU) (mg/L) (mF/L) (----) (cfs) ROCKSL 01/24/84 1025 10.0 7.3 10.8 25 25 248 35 500 ---16 3.3 320 42 8 0 370 ROCKSL 1205 32 35 02/28/84 13.5 7.5 10.0 316 500 65 ----11 30 3.6 340 12 ıΛ 420 ---ROCKSL 9.8 03/27/84 1030 16.5 7.5 22 24 \_\_\_ 254 17 30 3.2 480 370 54 8 Ø 430 ---ROCKSL Ø4/25/84 1135 16.5 15 193 7.3 9.6 14 14 15 3.4 1100 310 31 Ø 340 \_\_ ROCKSL 05/30/84 905 24.0 7.5 8.1 15 15 194 16 12 3.8 140 360 39 5 400 ROCKSL 06/27/84 1050 26.0 7.2 €.8 16 15 189 12 30 3.5 430 380 39 4 Ø 420 ROCKSL 07/25/84 1045 .24.0 7.7 8.1 22 27 217 10 15 2.5 600 320 63 17 400 171 ROCKSL 08/29/84 900 24.0 7.4 8.2 21 26 221 5 12 2.6 \_\_\_ 310 60 16 0 390 ROCKSL 09/27/84 1130 23.0 7.8 8.3 16 14 199 9 10 2.8 \_--310 31 3 Ø 340 ROCKSL 10/25/84 1130 17.0 8.0 10.9 16 15 194 8 12 3.2 \_\_\_ 330 32 4 0 370 ROCKSL 11/29/84 1330 12.0 7.4 10.5 14 13 186 10 30 3.7 580 32 2 Ø 610 ROCKSL 12/12/84 1145 11.0 7.3 9.7 14 13 ---195 1 1 30 4.4 540 410 31 2 0 440 \_\_\_ ROCKSL 01/30/85 1015 8.0 7.2 10.8 22 24 0.001 284 - 3 \_\_ \_\_ ---ROCKSL 02/27/85 0.000 258 25 \_\_ 350 45 Ø 400 ---1145 14.0 7.5 . 10.3 21 21 5 6 ROCKSL 03/27/85 1115 12.0 7.4 10.1 24 25 0.000 269 6 590 ---\_\_-ROCKSL 2 Ø 480 04/24/85 1123 18.0 7.8 10.1 21 18 0.000 232 7 430 42 5 \_\_ \_\_\_ ROCKSL 05/22/85 1020 21.5 8.2 9.2 21 24 0.000 225 17 15 \_\_\_ 520 56 11 Ø 590 ROCKSL 06/07/85 930 23.0 7.9 9. 1 25 30 252 \_\_\_ 16 ROCKSL 06/26/85 1000 23.0 770 7.6 8.0 41 56 0.000 360 19 10 ------600 110 60 3 ROCKSL 25.0 07/10/85 955 7.3 7.6 60 81 0.000 453 8 \_\_ \_\_\_ ---------\_\_\_ ROCKSL 08/28/85 1045 23.5 7.6 8. 1 81 122 0.000 630 8 10 2.8 340 160 100 19 620 \_\_\_ ROCKSL 09/25/85 1032 22.5 7.6 101 164 0.000 776 \_\_ \_\_\_ \_\_ 8. 1 8 \_\_\_ ROCKSL 10/23/85 17.5 36 600 1015 7.8 10.0 .99 158 0.000 738 5 2.1 \_--210 210 140 ROCKSL - 11/15/85 1140 12.5 7.5 - 10.4 135 238 0.000 988 \_\_ ----4 \_\_\_ ROCKSL 12/03/85 1125 11.5 -10.5 133 228 0.000 965 . 6 10 260 140 200 210 24 570 --7.4 3. 1 ROCKSL 01/23/86 1145 11.0 7.3 9.6 66 85 0.000 476 E \_-\_\_\_ --\_\_ -------\_\_ ROCKSL 02/13/86 1045 11.5 10.2 36 50 0.000 319 \_\_ \_\_\_ \_\_\_ -----------\_\_\_ ~-7.4 13 ROCKSL 0.000 35 670 67 0.740 03/04/86 32 35 8.4 \_\_\_ 6 1140 17.5 7.3 6.2 342 16 ROCKSL 04/09/86 1215 17.0 7.3 8.5 29 31 0.000 262 11 20 3.5 \_\_ 520 11 610 --ROCKSL 05/07/86 945 17.0 7.2 21 23 227 13 20 7.8 \_\_ 510 48 5 O1 560 \_\_ 7.4 225 Ø 220 ROCKSL 06/04/86 1040 22.5 7.3 7.6 19 21 0.000 21 \_\_\_ 12.0 ---200 23 2 ROCKSL 07/02/86 1000 25.5 7.3 19 225 15 6.3 19 ------. ---\_\_ \_\_ \_\_\_ 1.6 GREENES 07/21/83 600 19.5 7.3 8.7 7 4 ---115 Э . 2 --190 8 1 0 200 26400. 220 24600. GREENES 08/18/83 645 21.0 7.5 8.2 7 124 8 - 8 \_\_ 200 14 1 Ø 4 1.6 GREENES 09/13/83 2 620 23100. 640 20.5 7.3 8.3 10 F. 154 12 8 1.8 600 18 2 GREENES 10/04/83 925 18.0 7.3 9.0 7 5 \_\_\_ 124 10 5 1.6 380 200 9 0 Ø 210 24800. GREENES 11/01/83 650 17.0 7.3 9.1 8 5 --128 6 5 1.7 340 210 8 ø Ø 220 17700. 9 Ø Ø 310 66100. GREENES 12/06/83 635 10.5 7.4 10.6 122 30 30 2200 300 4 4.1 GREENES 01/10/84 7 3200 220 10 1 **Ø**1 230 67200. 815 9.0 10.7 4 \_\_ 129 19 20 1.7 7.3 0 200 32400. GREENES 02/01/84 950. 10.0 7.1 10.8 7 5 \_\_ 140 14 12 1.5 740 190 11 1 GREENES 03/07/84 735 12.0 7.5 10.8 10 .7 164 8 8 1.6 540 230 .28 1 Ø 260 25800. 260 25100. GREENES 04/04/84 635 13.5 7.5 10.4 9 148 8 5 1.6 680 250 14 1 Ø. 6 GREENES \_\_ Ø 190 11200. 05/02/84 530 -16.0 7.3 9.4 154 2.0 110 180 13 1 10 € 8 8 270 13900. GREENES 06/06/84 625 18.0 7.5 8.7 10 7 ---146 9 8 2.0 200 250 15 1 Ø GREENES 07/10/84 650 22.5 7.4 8.2 7 4 \_\_ 121 11 5 1.6 150 260 10 Ø Ø 270 21200. GREENES 600 \_\_\_ 5 0 310 22000. 08/01/84 21.5 7.4 -7.9 8 4 133 11 .1.6 --300 10 1 \_\_ 270 17800. GREENES 08/21/84 1040 23.0 7.3 8.2 11 6 164 12 10 1.8 250 1€ 1 Ø GREENES 09/05/84 6.05 22.0 7.7 12 0.000 8 2.4 \_\_ 390 20 1 Ø 410 18240. 7.4 6 185 11 180 14500. GREENES 10/04/84 620 17.5 7.4 9.0 8 4 0.000 132 7 5 1.6 170 13 1 0 GREENES 11/08/84 820 14.0 7.3 9.7 10 6 0.000 154 8 2.1 210 11 Ø Ø 220 14800. 11. 14 GREENES 745 10.5 10.9 9 Ø. ØØØ 160 24 15 2.6 1100 240 1 Ø 260 38100. 12/05/84 7.4 6

Note: -- = no data

01/30/85

1145

9.0

7.4 11.9

12

7 0.000

186

.3

GREENES

## DELTA HEALTH ASPECTS PROGRAM MONITORING DATA

																HOITE		IAL	Z \ (1)
Station	Date	Time	Temp.	pН	D. O.	Na	Cl	Se	EC	Tueb	Color	TOC	Asbest.	CH	CH ByCl2	CH Br2Cl	CH Boz T	TUMED	- (
Name	Date	(PST)	(oC)	· Pri		mg/			(uS/cm)		(CU)		(mF/L)			- uo/L		>	(cfs)
				*====												-		=====	
GREENES	02/06/85	1130	8.0	7.5	12.1	11	6	Ø. ØØØ	174	8	10			360	14	1	Ø	380	14900.
GREENES	03/06/85	1200	11.0	7.4	10.5	11	7	0.000	180	5			180						13200.
GREENES	04/05/85	1035	19.0	7.4	9.3	13	6	0.000	176	7	2			160	13	0	Ø	170	13900.
GREENES	05/01/85	1030	19.0	7.3	8.8	11	7	0.001	167	11	10			210	12	1	Ø	220	10200.
GREENES	05/29/85	510	18.0	7.4	9.15	13	7		178	10							~~~	740	15200.
GREENES GREENES	06/05/85	955	21.0 22.5	7.4 7.3	8.5 8.0	13	6 5	Ø. ØØØ Ø. ØØØ	173 163	9 8	10			290	19	1	0	310	15100. 17200.
GREENES	07/24/85 08/01/85	800 1035	22.5	7.5	7.9	11	5	Ø. ØØØ	163	10	10	3.9		480	14	2		500	15600.
GREENES	Ø9/Ø4/85	930	22.0	7.3	7.8	15	8	0.001	207	8	5	3.5		220	22	2	ø	240	12500.
GREENES	10/02/85	1015	21.5	7.5	8.2	14	8	0.000	168	7	5	1.6		200	14	1	ø	220	10600.
GREENES	11/13/85	1040	12.0	7.3	9.7	11	7	0.000	163	6	5	2.8		290	20	1	2	310	9500.0 ✓
GREENES	12/03/85	1930	11.5	7.3	9.3	10	7	0.000	149	28	35	16.0	. 380	690	21	1	Ø	710	24200.
GREENES	01/16/86	1400	10.0	7.3	10.6	18	10	0.000	218	9	15	2.3		660	22	1	0	680	14900.
GREENES	02/27/86	1240	12.5	7.1	10.5	4	2	0.000	84	64	20	4.2		340	7	Ø	Ø	350	74.600
GREENES	Ø3/13/86	1345	11.5	7.3	11.0	3	2	0.000	70	58	10	2.4		430	8	Ø	Ø	440	90,900
GREENES	04/23/86	1245	18.5	7.3	8.5	10	7	Ø. 000	179	14	10	1.9		310	22	1	Ø	330	17,500
GREENES	Ø5/28/86	1200	23.5	7.3	7.5	12	Э	Ø. ØØØ	188	14	10	2.9		170	12	2	1	180	14,000 11,200
GREENES	06/25/86	1250	24.5	7.3	7.8	11	8	Ø. ØØØ	161	13									
аоон	03/30/82	1050	11.0	7.3	10.7		4		131	20	5			310	9	0	Ø	320	40000:
HOOD	06/29/82	905	20.0	7.9	8.5		5		128	- 6				230	12	0	0	240	20000.
HOOD	08/26/82	1100	22.0	7.5	8.1 8.7		5 4		149 122	10 4				280 260	13 10	Ø	Ø	290 270	23200: 16300:
H00D 400D	10/21/82 12/29/82	1150 1400	18.0 9.5	7.5 7.2	10.9		4		130	33				480	16	1	Ø	500	71700.
HOOD	02/24/83	1410	12.0	7.5	10.6		2		113	30				120	4	ō	ø	120	74000.
HOOD	04/27/83	540		7.3	10.0		3		112	26				166	6	4	4	180	54600.
HOOD	06/22/83	1100	19.5	7.3	9. 1		3		101	17				200	8	ø	ø	210	43540.
MALLARDIS	05/08/85	700	16.0	7.8	8.7	1740	2890	0.000	9290	14	10			12	84	330	650	1100	7170.0
MALLARDIS	05/29/85	835	17.0	7.7	8.7	454	736		2720	26									8520.0
MALLARDIS	06/12/85	700	21.5	7.8	8.0	469	840		2980	19	5			65	170	340	300	880	4480.0
MALLARDIS	Ø8/14/85	730	19.0	8.0	8.5	1390	2510	0.000	8480	19	5	3.7	<del></del>	61	54	250	680	1000	1910.0
MALLARDIS	09/11/85	735	18.5	7.9	8.2	1230	2180	Ø. ØØØ	7320	12	5	3.0		21	94	370	500	980	3580.0
MALLARDIS	10/09/85	735	17.0	8.0	8.4	980	1880	0.000	6330	10	5	4.5		21	140	340	520	1000	1860.0
MALLARDIS	11/19/85	1015	11.5	8. 1	9.6		4260	ଡ. ଉପପ	13100	9	5	3.1							4610.0
MALLARDIS	12/03/85	1010	12.0	7.5	9.9			0.000	9970	8	8	3.4		11	72	340	640	1100	17200.
MALLARDIS	01/16/86	940	10.0	7.7	10.2	2180	3540	0.000	10700	16	20	4.6		5	44	320	990	1400	8270.0
MALLARDIS	02/27/86	955	14.5	7.0	8.8	12	12	0.000	169	58	25	5.3		490	29	1	Ø	520	
MALLARDIS	03/13/86	1130	13.0	7.3	9.4	12	14	0.000	161	51	30	5. 4		670	38	2	0	710	
MALLARDIS	04/23/86	915	16.5	7.3	8.9	20	23	0.000	226	22	20	3.5		440	64 88	8	Ø 35Ø	510 740	
MALLARDIS MALLARDIS	Ø5/28/86 Ø6/25/86	815 1035	17.0 21.0	7.6 7.7	8.6 8.1	680 689	1240 1280	0.000 0.000	4160 4250	26 36	15 10	7.1 10.0		39 		260	330	740	
VERNALIS	03/30/82	715	10.5	7.7	9.9		36	<b></b>	341	14	13	16.6		1400	67	9	Ø	1500	9720.0
VERNALIS	Ø6/29/82	530	18.0	7.7	8.4		30		267	15				470	93	12	ø	580	7400.0
VERNALIS	08/25/82	710	21.0	7.7	7.3		50		392	22				390	71	19	Ø	480	3750.0
VERNALIS	10/21/82	715	16.0	7.3	9.0		17		166	8				330	37	ž.	Ø	370	7420.0
VERNALIS	12/29/82	800	9.0	7.0	9.3		12		152	28				770	37	ø	ø	810	21500.
VERNALIS	02/24/83	1040	13.0	7.5	9.6		26		264	18				190	24	4	0	220	29100.
VERNALIS	04/27/83	740		7.1	9.7		11		150	12				310	20	6	5	340	36600.
VERNALIS	06/22/83	630	21.0	7.0	8.5		10		117	23				380	23	2	Ø	400	24100.
VERNALIS	Ø7/26/83	815	20.0	7.3	7.7	29	30		288	29	5	3.5		290	54	12	Ø	360	11300.
VERNALIS	08/23/83	700	20.0	7.2	8.0	23	24		247	19	5	3.0		420	39	7	Ø	470	9170.0
VERNALIS	Ø9/14/83	715	20.0	7.4	8.2	15	14		158	16	10	2.8		350	21	3	Ø	370	11200.
VERNAL IS	10/12/83	625	17.5	7.1	8.5	11	11		126	12	10	2.8	780	270	24	3	Ø	300	14500.

THM FORMATION POTENTIAL

														CH	CH	CH	CH	THE	
Station	Date	Time	Temp.	pН	D. D.	Na	C1;	Se	EC.		Color	TOC	Asbest.	C13	BrC1≥	Br2Cl	Br3 T		
Name	.========	(PST)			<b></b>				(uS/cm)	(TU)	(CU)		(mF/L)			- ug/L			(cfs)
VERNALIS	11/08/83	730	15.0	7.3	8.2	-39	38		381	18	25	4.2	1300	300	62	12	.Ø	370	9370.0
VERNALIS	12/13/83	825	11.0	7.1	10.0	14	13		155	14	30	3.2	740	330	22	2	Ø	350	22200.
VERNALIS	01/24/84	735	10.0	7.0	10.0	21	19		210	14	25	3.1	870	340	32	4	Ø	380	21400.
VERNALIS	02/28/84	815	12.0	7.5	9.7	38	3,9		352	10	15	3.2	270	250	60	15	2	320	9640.0
VERNALIS	03/27/84	720	14.5	7.3	9.4	48	52		464	34	15	3.9	1800	280	86	23	2	390	6300.0
VERNALIS	04/25/84	755	14.0	7.3	8.8	59	66		547	24	8	4.8	1700	290	110	42	2	440	3980.0
VERNALIS	05/30/84	620	24.5	7.9	7.3	69	80		629	75	10	6.1	1300	380	120	56	3	560	2440.0
VERNALIS	06/27/84	650	25.5	7.3	6.3	77	88		694	50	25	5.8	1300	360	130	58	3	550	2050.0
VERNALIS	07/25/84	705	23.0	7.5	6.5		92	0.001	640		15	5. 4	3300	450	150	72	7.	680	1840.0
VERNALIS	08/29/84	620	24.0	7.6	7.1	58	62		549	24	20	4.8		350	110	48	2	510	2520.0
VERNALIS	09/27/84	725	20.0	7.4	8.3	39	43	Ø. ØØØ	388	17	10	4.2		280	79	21	Ø	380	3140.0
VERNALIS	10/25/84	810	15.5	7.4	7.9	39	41	0.000	378	15	12	3.9		260	64	23	1	350	3580.0
VERNALIS	11/29/84	940	11.5	7.1	9.2	43	44	0.000	400	10	25	4.4		380	68	15	Ø	460	3440.0
VERNALIS	12/12/84	830	11.0	7.3	9.2	34	32	0.000	324	€	12	3.6	510	240	50	12	Ø	300	4700.0
VERNALIS	01/30/85	750	8.0	7.4	10.5	54	55	0.001	483	3									3850.0
VERNALIS	02/22/85	1310	12.0	7.4	6.4	75	69	0.001	598	10	20							,,	3170.0
VERNALIS	02/27/85	815	12.5	7.4	9.6	70	73	0.002	629	8	25			220	97	48	E	370	2640.0
VERNALIS	03/27/85	845	12.0	7.4	9.0	92	97	0.002	801	17			810					- <del></del>	2580.0
VERNALIS	04/24/85	745	17.0	7.4	7.9	87	80	0.002	667	19	5	77		360	140	61	3	560	2520.0
VERNALIS	05/22/85	700	20.5	7.4	7.2	84	99	0.002	756	31	10			400	160	68	12	640	1920.0
VERNALIS	05/29/85	645	18.0	7.7	7.9	89	98		774	28									1900.0
VERNALIS	06/26/85	645	-23.0	7.5	7.3	81	94	0.001	717	52	10			540	160	66	7	770	1420.0
VERNALIS	07/10/85	645	22.5	7.4	7.1	55	58	0.001	490	28	5			520	130	41	3	690	2500.0
VERNALIS	08/28/85	715	19.5	7.7	7.4	52	60	0.001	487	18	5	3.9		410	100	34	2	550	2400.0
VERNALIS	09/25/85	707	21.5	7.4	6.8	59	70	0.000	563	21	5	3.1		380	98	30	4	510	1600.0
VERNALIS	10/23/85	700	15.5	7.4	7.4	53	65	0.000	519	12	5	2.4		320	110	29	2	460	1950.0
VERNALIS	11/15/85	820	8.5	7.5	9.7	80	94	0.001	706	7	15	2.9		220	130	71	7	430	1400.0
VERNALIS	12/03/85	1530	13.5	7.4	8.9	66	74	0.001	604	18	18	6.5	560	590	140	32	120	760	2250.0
VERNALIS	01/23/86	745	12.0	7.5	8.8	99	107	0.000	790	18	15	3.2		930	160	7E		1200	1750.0
VERNALIS	02/13/86	730	11.5	7.3	9.0	82	86	0.002	686	15	_ 5	4.3		450	140	56	3	650	
VERNALIS	03/04/86	800	15.0	7.3	8.3	28	26	0.001	268	26	35	7.8		540	56	6	Ø	600 700	
VERNALIS	04/09/86	800	- 15.0	7.3	9.2	18	18	. 0. 000	169	20	25	5.3		650	47 51	4 6	Ø	390	
VERNALIS	05/07/86	630	14.5	7.3	8.8	27 26	27 28	0.001 0.001	257 254	17 22	15	6.0		330 220	41	E	121	270	
VERNALIS VERNALIS	06/04/86 07/02/86	745 650	20.5 23.0	7.3 7.5	8.0 7.9	65	75	e. ee:	595	9		- ==							
					9.0		1760			1	 5	9.2		19	140	500		1200	11.6
SEDCKS	07/20/83 08/17/83	1125 650	25.0 28.0	8.6 7.9	8.0		1640		12600 11600	1	8	9.3		26	110	420	280	840	9.6
SLDCKS	09/06/83	915	26.5	7.8	8.0		1660		11900	1	10	9.5		67	340	720		1500	8.8
SLDCK5	10/06/83	815	21.5	8.4	8.3		1610		11900	2	25	28.0		36	260	710		1600	7.4
SLDCKE	11/15/83	1425	15.5	8.8	13.0	2140			11300	E	45	30.0		39	280	710		1700	8.5
SLDCK2	12/20/83	1110	13.5	8.2	10.3		1380		10500	1	18	7.5		42	190	410	330	970	15.5
SLDCK17	07/20/83	915	23.5	8.5	9.0		1590		11500	1	5	9.5		34	160	520		1300	11.6
SLDCK17	08/16/83	1240	- 30.5	7.9	9.4	2120			11500	ė	8	10.0		30	140	750		1300	10.3
SLDCK17	09/06/83	800	25.5	7.9	8.0	2180			11700	5	12	18.0		70	310	600		1400	9.5
SLDCK17	10/05/83	1340	23.0	8.6	12.5	2160	,		11800	2	30	29.0		31	210	750		1700	
SLDCK17	11/15/83	1330	16.5	8.6	11.5	2300			11700	3	25	19.0		35	230	580		1600	6.9
SEDCK17	07/20/83	805	21.5	8.3	9.5	1970			11000	1	5	7.3		37	150	480		1200	11.6
SLDCK41	Ø8/16/83	1130	25.0	7.6	7.5	2020			11100	4	8	10.0		18	130	420	250	820	9.5
SLDCK41	09/06/83	700	23.5	7.9	11.6	2070			11400	3	15	11.0		100	330	350	180	960	9. 1
SLDCK41	10/05/83	1245	22.0	8.3	7.7	2040			11400	1	15	13.0		30	160	370	280	840	7.5
SLDCK41	11/15/83	1240	16.5	8.6	15.5	2700			13400	4	25	21.0		25	200	480	230	930	6.9
SLDCK41	12/20/83	955	15.0	8.1	10.8	1760			9320	ż	8	9.8		32	140	310	230	710	13.6
							· <del>-</del>			_	_								

Note: -- = no data

### DELTA HEALTH ASPECTS PROGRAM MONITORING DATA

														THM FORMATION POTENTIAL					
														CH	CH	CH	CH		
Station	Date	Time	Temp.	рΗ	D.O.	Na	C1	Se	EC	Turb.	Color	TOC	Asbest.	C13	BrC12	Br201	Br3 T	THMFP	FLOW
Name		(PST)	(oC)		<	mg/	/L	}	(uS/cm)	(TU)	(CU)	(mg/L)	(mF/L)	( <del></del>		- ug/L		·)	(cfs)
	=======================================	ar 1 im 1/2 az ar	1.0 254 524 515 515 515 517		=======================================	23 22 22 22 22 24 <b>2</b>	======	<b>2</b> 122 52 52 52 52 52			======	. = = = = :: = :	======================================	=====	=====	****		=====	
SLDPD5	Ø7/2Ø/83	1210	22.0	8.6	4.0	2940	2160	*** ****	14700	Ø	12	11.0		21	180	780	950	1900	
SLDPD5	Ø8/17/83	715	25.0	7.5	1.4	2980	2250		15200	1	12	11.0		20	190	720	520	1400	
SLDPD5	09/06/83	950	24.0	7.5	1.5	2540	1960		13600	0	8	8.7		76	340	750	490	1700	
SLDPD5	10/06/83	855	20.0	7.7	3.3	2300	1780		12500	Ø	25	11.0		58	270	660	1300	2300	
SLDPD5	11/15/83	1455	13.0	8.€	10.8	2120	1520		11200	2	35	26.0		59	320	750	960	2100	
SLDPD5	12/20/83	1135	13.0	8.0	8.7	2020	1390	<del></del>	10200	1	20	11.0		63	220	470	380	1100	
SLDPC	Ø7/28/83	945	23.0	7.5	8.4	944	865		5890	3	2	4.2		36	120	190	140	490	
SL.DPC	Ø8/25/83	845	20.0	8.1	8.8	940	860		5900	3	5	4.0		42	170	260	140	610	
SLDPC	09/20/83	1000	22.5	7.6	8.5	1120	1010		6910	96	5	4.3		38	110	290	160	600	

Note: -- = no data

## Appendix B

## FIELD SAMPLING PROCEDURES

### APPENDIX B

#### APPARATUS AND METHODS EMPLOYED

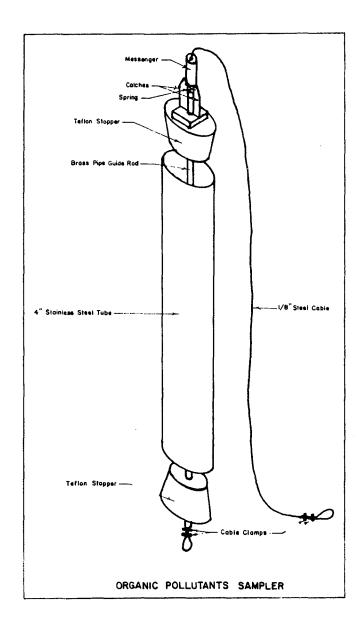
This appendix describes sampling apparatus, sampling methods, and analytical methods employed in the Interagency Delta Health Aspects Monitoring Program.

## Sampling Apparatus

Prior to January 1984, samples were collected in a 1.5-liter steel bucket with a 1-meter chain attached; the bucket and chain were prepared for sampling by detergent washing and drying. The equipment was transported in detergent-washed aluminum foil. Sampling involved attaching a small diameter nylon rope to the end of the chain and dipping the bucket into the water to collect the sample. To avoid contamination, the rope was not allowed to enter the water.

Beginning in January 1984, and continuing since then, samples have been collected using a specially constructed device developed by the Department of Water Resources (see Figure 1). A stainless steel tube with Teflon closures and a triggering mechanism are the main components of the device, which was produced using parts from old Kemmerer samplers. The important feastures of the device are: (1) it enables subsurface sampling, and (2) the water being sampled is not in contact with potentially contaminating materials.

Before being used for the first time, the device was soaked for about a week in water containing detergent. This procedure was intended to cleanse the equipment of any surface contaminants that may have been present.



Prior to sampling, the device was washed in detergent, rinsed, dried, and wrapped in detergent-washed foil. A nylon rope attached to a short length of steel cable was used to suspend and operate the sampler. As was the case with the sampling bucket, the rope was not allowed to contact the water.

## Sampling Methods

Samples for Total Trihalomethane Formation Potential analyses were filtered through 0.45uM Millipore membranes, using a stainless steel filtration apparatus that was washed in detergent, rinsed, dried, and wrapped in detergent-washed foil prior to sampling. The purpose of the filtration was to simulate the clarification and filtration processes employed in water treatment.

Filtration apparently has only a minor effect on trihalomethane formation potential of most fresh water samples. Twenty-five fresh water samples were analyzed in duplicate, one sample being filtered and the other unfiltered. The average difference between the filtered and unfiltered samples was 14 percent; this difference is in the order of magnitude of the analytical variation of the test method. Filtered water was poured into 40 mL screw-top vials with Teflon septa, leaving no airspace, as specified by the U. S. Environmental Protection Agency /1/.

Water samples for total organic carbon analyses were poured into acid-fixed 30 mL glass bottles with tapered glass stoppers, then sealed with washed foil.

Samples for the above analyses were transported iced to the DWR Bryte Laboratory within 24 hours of sampling.

Field analyses were performed at the time of sampling. Temperatures were taken by means of a radial thermometer graduated in intervals of 0.5 degrees. Celsius. Measurements of pH were performed by use of a Hellige colorimetric pH comparator. Dissolved oxygen concentrations were determined in the field by the modified Winkler titration method, and electrical conductivity was determined by use of a Beckman SoluBridge for conductivities less than 8,000 umhos/cm and a Beckman Model RC-19 electrical conductivity bridge for higher conductivities.

Asbestos samples were collected in pint-sized polyethylene bottles and shipped on the day of collection via express mail to the EMS Laboratory in Hawthorne, California. Priority pollutant samples were collected in gallon containers, three per sample (for extractables). Also, 40 mL samples were collected in glass containers (five per sample) for volatile organic analyses. The sample containers were completely filled, eliminating headspace. Volatilization losses during filling were minimized by tilting sample vials and allowing the sample to run down the inside of the vial without causing turbulence. caps of the sample containers were Teflon-lined. These samples were delivered to McKesson Environmental Services laboratory in Dublin, California, within 24 hours of collection.

## Analytical Methods

Upon delivery to the DWR Bryte Laboratory, raw water samples for trihalomethane formation potential analyses were chlorinated at about 50 milligrams per liter (mg/L) chlorine dosage. This high dosage was used to assure a chlorine residual after the 7day incubation period at 25 degrees Celsius. This procedure should be acceptable, as studies have determined that ultimate trihalomethane formation is independent of dosage, where the dosage exceeds the chlorine demand of the sample /2/. At the end of seven days, samples were dechlorinated using sodium thiosulfate and analyzed by the purge and trap method of gas chromatographic analysis established by EPA /1,3/. Asbestos samples and priority pollutant samples were likewise analyzed by methodology established by EPa /4,5/. Selenium was analyzed by a method developed by the U S. Geological Survey for its low detection level work /6/. All other analyses were performed according to Standard Methods /7/.

### APPENDIX B REFERENCES

- 1 Federal Register. 44:231 (November 29, 1979) Appendix C, Part III, 68690-68691.
- R. A. Minear and C. M. Morrow. Raw Water Bromide Levels and Relationship to Distribution of Trihalomethanes in Finished Drinking Water. Tennessee Water Resources Research Center, Knoxville. Prepared for Office of Water Research and Technology, Washington, D.C. Contract DI-14-34-001-1145. PB83-256735. March 1983. 271 pp.
- 3 Federal Register. 44:231 (November 29, 1979) Appendix C, Part I, 68672-68682.
- 4 <u>Interim Method for Determining Asbestos in Water</u>. EPA-600/4-80-005. January 1980.
- J. E. Longbottom and J. J. Lichtenberg. <u>Test Methods -- Methods</u> for Organic Analysis of Municipal and Industrial Wastewater. EPA-600/4-82-057. July 1982.
- J. G. Crock and F. E. Lichte. An Improved Method for the Determination of Trace Levels of Arsenic and Antimony in Geological Materials by Automated Hydride Generation Atomic Absorption Spectroscopy. Analytica Chimica Acta. 144:223-233. 1982.

## Appendix C

## LABORATORY QUALITY CONTROL PROCEDURES

M-Kesson

## LIMITS OF DETECTION

Pollution of Delta waters by industrial or agricultural chemicals can occur from both point sources and non-point sources. In either case, the high degree of dilution afforded by the high volume of run-off water entering the Delta is expected to result in very low concentrations of synthetic organic chemicals in Delta waters. These expected low concentrations challenge the analytical methodologies available for the detection and measurement of compounds of interest.

For the present program, primary emphasis has been placed on analysis for the EPA "Priority Pollutants." For this purpose we have used the following EPA Test Methods:

Method 624 - Purgeables
Method 601 - Purgeable Halocarbons
Method 625 - Base/Neutrals and Acids
Method 608 - Organochlorine Pesticides
and PCBs.

When certain non-priority pollutant compounds have been determined, other EPA methods were employed; for example, Method 614 - Organophosphorous Pesticides.

Each of the EPA methods includes values for method detection limits for many of the compounds covered by the specific method. The GC/MS methods (624 and 625) are the methods of choice for an initial survey, since the mass spectrometer is a universal detector which also provides positive identification of the analyte. However, the sensitivity of this detector is such that the method detection limit is generally higher than the expected level of those organic pollutants in Delta waters. Consequently, some procedural modifications have been used and some additional analyses using more sensitive detectors have been completed.

For purgeable priority pollutants, the initial analysis was by Method 624, for which method detection limits of 1 - 10  $\mu$ g/L are reported. This was supplemented by use of Method 601, for which detection limits of 0.1 - 1  $\mu$ g/L can be achieved. This latter method uses a halogen specific detector of high sensitivity.

For extractable organics, Method 625 offers detection limits in the range of 2 to 20 µg/L. In order to improve on this, the procedure has been modified slightly to increase the concentration factor by x10 and consequently to lower the detection limit by a factor of 10. This was achieved by increasing the sample volume to 2 liters (from 1 liter) and concentrating the extract to 0.2 ml (instead of 1 ml).

A similar treatment of sample extraction has been used with Method 608 and Method 614. These methods employ highly sensitive detectors, with very low reported method detection limits (0.002  $\mu g/L$  for dieldrin and 0.012  $\mu g/L$  for diazinon, for example).

The method detection limits (MDL) as quoted above may be considerably lower than the actual limit of detection (LOD) for any real sample since the MDL is determined without consideration of matrix interferences, sample blanks, etc. For the present project, matrix interferences are the limiting factor, restricting the amount by which the LOD can be lowered by increasing the concentration factor. Values quoted for LOD in this report (for non-GC/MS methods) are analysts' estimates of analyte concentrations needed for determination of that analyte above the matrix interference level.

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## QUALITY CONTROL / QUALITY ASSURANCE

McKesson Environmental Services laboratories operate under a thorough program of quality assurance/quality control.

## Sample Receipt, Handling, Storage and Control

When a sample arives from the field, the sample custodian performs the following functions:

- Receipt of sample is recorded.
- Package is inspected and any damage recorded.
- Package contents are verified.
- Chain-of-Custody document is completed and discrepancies reported.
- Sample is logged in, number assigned and sample tagged.
- Laboratory sample sheet is initiated.
- Sample is assigned to storage.

## Security, Chain-of-Custody and Document Control

In order to maintain a clear record for sample traceability and document accountability, the following procedures are enforced:

- Environmental Services laboratories and sample storage areas are maintained as secure facilities at all times.
- Chain-of-Custody procedures are rigorously followed.
- A document control officer is appointed.
- Documents are numbered and a document inventory maintained to include log books, sample sheets, and quality assurance documents.

## Laboratory Operations

The laboratory performs adequate quality control on samples to assure the precision and accurancy of the data. The following are the minimum quality control requirements:

- One sample analyzed in duplicate for every ten samples or batch of samples.
- One spiked sample for every ten samples or batch of samples. Spikes shall be made at two to three times the detection limit, or at the analyte level.
- Surrogate compounds for volatile organic, base/neutral, and acid extractables.
- Method and field blanks, as apropriate, especially for aqueous samples.

For the present program, Methods 601, 624 and 625 employ surrogate spike compounds with the analysis of each sample. An internal standard is used with each sample for Method 608 and individual compound recoveries have been determined for typical compounds covered by other methods used.

# QUALITY CONTROL PROCEDURES DEPARTMENT OF WATER RESOURCES BRYTE LABORATORY

- 1. Laboratory blanks are run on each analytical day.
- 2. Travel blanks are run along with each group of samples.
- 3. Standards are run at the beginning and end of each group of analyses.
- 4. Sample aliquot volumes are adjusted so standards bracket concentration of analyte, or are within 10 percent of sample peak height for each compound being analyzed.
- 5. Duplicate spiked samples are analyzed for precision and accuracy determinations on approximately 10 percent of samples.

# LIMITS OF DETECTION DEPARTMENT OF WATER RESOURCES BRYTE LABORATORY

DEPARTMENT OF WATER RESOURCES DATTE LABOR	AIORI
	Detection Limit
Compound	<u>(ug/L)</u>
chloroform	0.1
bromodichloromethane	0.1
dibromochloromethane	0.2
bromoform	0.5
Alachlor	0.01
Atrazine	0.01
Azinphosmethyl (Guthion)	0.01
Bentazon	<b>-,-</b> , ∞
Chlorothalonil	0.01
2,4-D, Alkanolamine Salts	0.01
D-D Mixture	0.1
DEF	0.01
Diazinon	0.01
2,6-Dichloro-4-Nitroaniline	0.01
Dicofol	0.01
Dimethoate	0.01
Dimethyl Tetrachloroterephthalate	
(Dacthal)	0.01
DNBP (Dinoseb)	0.01
Disulfoton	0.01
Diuron	0.01
Ethylene Dibromide	0.2
Malathion	0.01
Methyl Bromide	0.1
Methyl Parathion	0.01
Parathion	0.01
Simazine	0.01
Toxaphene	0.5
Trifluralin	- <b></b>
Xylene	4

# GAS CHROMATOGRAPHIC CONDITIONS EMPLOYED FOR VOLATILE HALOCARBON ANALYSES

Gas Chromatograph:

Tracor 565

Detectors:

Hall 700A Electrolytic Conductivity Detector

Tracor 703 Photoionization Detector

Column:

6-foot glass tube, 2 mm I.D.

Column Packing:

1% SP-1000 on Carbopack B 60/80 mesh (Supelco, Inc.)

Confirmation: n-octane on Porisil-C 100/200 mesh (Supelco, Inc.)

Temperatures:

Injector: 200°C

Column:

1% SP-1000; 100°C - 4 min.; 8°C/min to 200°C; hold 8 min.

n-octane; 60°C - 4 min; 6°C/min to 170°C; hold 4 min.

Detector Base: 250°C

Reactor: 825°C

Carrier Gas: He; Flow 30 mL/min

Reaction Gas: H2; Flow 50 mL/min

Recorder Chart Speed: 0.5 inch/min

Sampler: 5mL - Tekmar Liquid Sample Concentrator, Model LSC-2.

Purge 11 min; Desorb 4 min; Bake 10 min.

Trap: As specified in EPA Method 601 /1/

Approximate Retention Time (min) /2	?/: n-octane	SP-1000
Chloroform	7.0	7.4
Bromodichloromethane	9.8	10.4
Dibromochloromethane	12.4	13.6
Bromoform	15.0	16.6

<sup>/1/</sup> Reference: Federal Register. 44:233 - Purgeable Halocarbons Method 601

/2/ Standards: Trihalomethane Mixture 4-8746. Supelco, Inc., Bellefonte, PA 16823

# ANALYSIS OF TRIHALOMETHANE REFERENCE SAMPLE, MARCH 1982

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			28 17								
<u>Organization</u>	CHC13	Trihal	omethane	Concentra	ation (ug/L <u>CHBr3</u>						
DWR Bryte Laboratory	3.1	3	3.3	8.6	36: 36: 3 10: 10: 3	51					
DOUG Canitation and		F	4.4		the first the second	7					
Radiation Laboratory	2.8	2	2.8	6.4	31.7	43.7					
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## Appendix D

#### LABORATORY PERFORMANCE 1985-86

#### Appendix D

#### LABORATORY PERFORMANCE 1985-86

The performance of the Department's Bryte Laboratory and of McKesson Environmental Services was measured by the analytical results of internal quality control and inter-laboratory quality assurance samples. Several methods were used for appraisal; they included the analysis of spiked samples, field replicates, and laboratory replicates.

#### **Bryte Laboratory**

Measurements of standard water quality parameters and volatile organic trihalomethane compounds were performed by the Department's Bryte Laboratory. Results of duplicate sample analyses are shown in Table 1. The laboratory experienced repeated difficulties in controlling the quality of TOC (total organic carbon) analyses. There were unacceptable differences in the results of duplicate samples.

Bryte Laboratory staff identified one cause for the TOC discrepancies. Two analysts had used incorrect multiplication factors to report values based on the sample aliquot sizes they had used. Four different volumes (1, 2, 3, or 5 mL) were used in the TOC analyses. Laboratory worksheets are being examined and checked for computation errors.

							Ta	able 1									
			AN	ALYSIS	OF :	SPLIT	SAMP	LES BY	DWR B	RYTE	LABOR	ATORY					
Station_		Time PST	Temp °C	рН	EC uS	DO <	Na 	Cl -mg/L-	Se	TOC							TTHMFP >
Clifton	04/09/86	1100 1115		7.2 7.2	197 195		20 20		0.000 0.001		14 14	20 30	570 610	62 53	5 5	0 0	640 670
Banks	09/25/85	0820 0820	22.5 22.5		588 584	7.9 7.9	69 70		0.000		6 6	10 5	340 290	89 170	40 63	10 13	400 540
Lindsey	06/25/86	0635 0600	21.5 20.0	8.0 7.9	461 480		43 44	37 38	0.000		38 38	20 10	350 270	36 34	4 8	1 3	390 320
No.Bay	05/28/86	0945 1045	19.5 19.5		306 300	9.6 9.5	10 9	· 5	0.000		7 6	5 10	300 120	15 8	1 3	0 2	320 130
Mallard Island	12/03/85	1010 1010			9970 9950			3130 3130	0.000		8 8	8 5	11 9	72 78	340 280	640 540	1100 910
Greene's	02/27/86	1240 1240	12.5 12.5			10.5 10.5	4 4	_	0.000		64 63	20 10	340 320	7 8	0 0	0 0	350 330
Vernalis	11/15/85	0820 0820	8.5 8.5	7.5 7.5	706 709		80 80	94 94	0.001 0.001		7 7	15 5	220 240	130 130	71 71	7 8	430 450

The cause for disparities in TOC results for duplicate samples when aliquot volumes were the same was traced to instrumentation problems. A new TOC analyzer will be purchased. Until the accuracy of the TOC data can be established, program staff will not interpret or correlate TOC data with other parameters.

The differences in chloroform values between duplicate samples were considered small and acceptable to the monitoring program. However, duplicate sample analyses for TTHMFP and some THM species had significant discrepancies. Color measurements were also rarely repeatable in the laboratory, for no known reason. The results of duplicate samples for other water quality parameters were found to be acceptable.

The installation of several new analytical instruments has disrupted work at the Bryte Laboratory, and a large backlog of samples has built up. As a result of the backlog and because of the problems with TOC analyses, the TTHMFP and TOC samples will be analyzed by the McKesson Environmental Services laboratory starting in November 1986.

#### McKesson Environmental Services

McKesson Environmental Services (MES) is a commercial laboratory facility located in Pleasanton, California. MES is under contract to the Department to provide pesticide and priority pollutant analyses of water for the Health Aspects Monitoring Program. On occasion, MES conducted bromide and dissolved copper analyses when requested. However, these two analyses were discontinued because sample concentrations were often much lower than the MES laboratory detection limits. Quality control procedures are presented in Appendix C.

MES conducted spike recovery tests on each chemical requested for analysis by the Department. Table 2 shows the results of these tests for field samples collected in June through August 1985, December 1985, and May 1986. Both distilled water and field samples were spiked to conduct these recovery measurements.

In general, method spike recoveries varied between sampling runs and among analytes, but overall recoveries were better than 70 percent. Exceptions were analyses for methamidophos (24%, 46%, and 60% at 40 ug/L); 2,4-D salt (50% at 20 ug/L); MCPA (52% at 60 ug/L); and methyl parathion (42% at 1 ug/L). The method spikes represent the achievable recovery and variation with the analytical method used by the laboratory. Extraction methods to improve the recovery of methamidophos were initiated by MES as a result of the consistently low recoveries.

Spiked samples prepared by the Department's Bryte Laboratory were also submitted with each batch of field samples. These spiked samples were coded and "blind" to MES. The samples consisted of tap water spiked with pesticides. Results are presented in Table 3. The results pinpointed some errors in identifying compounds and reporting laboratory results. Upon notification, MES conducted an investigation to correct the problems. The problems and corrective actions were /1/:

- 1. MES reported no detection of 2,4-D in the August 20 and 21, 1985 QA spike. Upon re-examination of the data, MES discovered an incorrect transcription of results from laboratory worksheets to the final report form. The chemical 2,4-D was actually found and confirmed on a second gas chromatography (GC) column.
- 2. MES reported 1.7 ug/L of metalaxyl in a reference spike that was not spiked with metalaxyl. MES re-examined the chromatogram and noted that the retention time for metalaxyl was outside the retention time range set

Table 2

McKESSON ENVIRONMENTAL SERVICES RECOVERIES OF SPIKED SAMPLES FOR IN-HOUSE QUALITY CONTROL MEASUREMENTS

		June 198	5 analyses	July 198	5 analyses	August 1	985 analyses	Dec. 1989	5 analyses	May 1986	analyses
Lab		Spiked	Percent	Spiked	Percent	Spiked	Percent	Spiked	Percent	Spiked	Percent
Method	Chemical	Amount	Recovery	Amount	Recovery	Amount	Recovery	Amount	Recovery	Amount	Recovery
-======			=========		**********			======		======	
622	2,4-D salt	10	71	10	50	10.4	72	20	32	20	50 *
HPLC	Bentazon	20	107 *	20	93 *	30	38	30	75	30	140 *
614	Carbofuran	10	110	10	97	5	58	5.3	107 *		
GC-ECD	Chloropicrin	11	100	11	27	1.14	62	1.1	73	1.0	120 *
608	Dacthal	10	137	10	140	1	100	1	110 *	1.5	150 *
601/602	D-D mixture	12	97 ×	8	101 *	20	95	20	88	20	76
622	MCPA	30	74	30	60	31	80	60	42	60	52 *
614	Metalaxyl	30	81 %	30	81 *	5	80	5	54		
614	Methamidophos	315	10	315	10	40	46 *	40	60 <b>*</b>	40	26 *
614	Methyl bromide	12	. 98 ★	8	105 *	20	145	20	93	20	119
614	Methyl parathion	1	42 *	10	40	5	100 *	5	120 *		
614	Molinate	10	119	10	140	5	74 *	5.1	82 *		
wet chem	Paraquat dichloride	200	85 *	20	77	20	98	20	99	20	75 *
614	Thiobencarb	10	110	10	98	5	44	5	94 *		
601/602	Xylene	12	98 *	8	114 *	60	127	40	93	20	74
AAS	Copper	50	106 *	50	96	15	111	10	107		
	Bromide	8	91	8	88	0.8	81				

The "\*" designates these were recoveries of spikes in distilled water samples.

Values without "\*" in percent recovery column were recoveries of spikes to actual field samples.

All units in ug/L except for bromide which is in mg/L.

Table 3

ANALYSIS OF SPLIT SAMPLES BY McKESSON ENVIRONMENTAL SERVICES LABORATORY

Sample <u>Number</u>	Date Sampled	Date Reported	Spike Compound	Spike Concentration ug/L	Recovered* Concentration ug/L	Percent Recovery
RP52	6/17/85	8/20/85	2,4-D Salt Dacthal Molinate	5.0 4.9 5.4	2.7 (6.0) 5.9	54 122 109
RP67	7/16/85	9/29/85	2,4-D Salt Dacthal Molinate	4.9 4.7 5.3	1.0 (4.2) 3.6	20 85 68
RP71	8/20/85	9/20/85	2,4-D Salt Dacthal Molinate	5.0 4.9 5.4	(1.6) (1.7) 3.7	32 35 69
RP86	9/4/85	10/2/85	2,4-D Salt Dacthal Methyl Parathion	5.0 4.9 5.1	2.7 4.2 3.2	54 86 63

All spike samples were prepared by Department of Water Resources Bryte Laboratory.

<sup>\*</sup> Where numbers are in parentheses, the spike was not detected.

Numbers are revised values after discovering the cause of the errors.

for its identification. The chromatogram also contained numerous large peaks, making interpretation difficult, and probably resulted from dirty glassware.

MES did not detect Dacthal in three reference spikes that contained about 5 ug/L Dacthal. It was later determined that MES had been led to an incorrect identification of the retention time for Dacthal peaks on the gas chromatograms because of a contaminated reference standard used by MES for internal spikes. The contaminated Dacthal standard produced two peaks, one for Dacthal and the other for the contaminant chlorobenzene. When further analyses were performed with a new, pure Dacthal standard, the misidentifications were corrected and the chromatograms showed the correct retention time for Dacthal. All chromatograms of DWR samples were then re-examined and corrected with the proper Dacthal results.

The Department of Health Services (DOHS) was particularly concerned about future misidentifications and failure of reporting unidentified peaks. DOHS offered the following suggestions /2/:

- "1. For each analysis requested by DWR from its support laboratories, information pertaining to all unidentified peaks should be reported. When such peaks occur, retention times may offer qualitative information; quantitative data is accessible by one or both of the following two formats:
- (i) Using the retention time of the standard chemical compound nearest the unknown as a reference, calculate, and report the unknown chemical's concentration on the basis of relative peak heights between standard and unknown.

  Reagent- and method blanks should be used and compared as well.

(ii) Using the lowest relative response for any standard of known concentration analyzed by the method in question, calculate the maximum concentration for any unknown peak observed. Make proper accounting for blank contributions and report this also."

MES agreed to follow the DOHS recommendations on tabulating retention time and peak area data for the Department, but cautioned against attempts to quantify the data from reported unidentified peak areas. MES experience had led to the observation that /3/:

Electron capture of flame photometric detectors common to pesticide analysis have sensitivities which may vary by several orders of magnitude from one compound to another. Without some knowledge of the compound producing the peak, no quantitative information can be inferred."

DWR requested MES to follow the DOHS recommendations on a trial basis thereafter.

Field duplicates were also submitted to MES. These are samples taken from the same location and time and split into more than one sample set for analysis. When field duplicates were not obtained, field replicates were taken. Replicate samples are those taken from the same location within a short time period, with each replicate stored in its own sample container. When large volumes of water are needed for analysis, field replicates are more convenient to obtain than proportioning water samples among several containers. There was good agreement in the results (Table 4).

Field duplicate samples were also split between MES and the DWR Bryte Lab for comparison. The results are shown in Table 4.

Table 4

McKESSON ENVIRONMENTAL SERVICES LABORATORY ANALYSES

OF DUPLICATE SAMPLES

	6-17-85	5		7-16-8	5		8-20-8	5		9-4-85		
	San Joa	quin Rive	er	San Joa	aquin Riv	er	San Jo	aquin Riv	er	Colusa	Basin	
	near Ve	ernalis		near V	ernalis		near V	ernalis		Drain		
Chemical	RP 56	RP 57	1.o.d.	RP 66	RP 68	1.o.d.	RP 75	RP 76	1.o.d.	RP 82	RP 85	1.o.d.
	======		.======	******	=======			=======				
2,4-D salt	ND	ND	.1	ND	ND	.1	ND	ND	.01	ND	ND	.5
Bentazon	ND	ND	. 2	ND	ND	.1	ND	ND	.2	0.9	0.8	.2
Carbofuran	ND	ND	.5	סמ	ND	.5	ND	ND	.5	ND	0.08	.02
Chloropicrin	ND	ND	.1	ND	ND	.1	ND	ND	.1	ND	ND	.1
Dacthal	ND	ND	.01	ND	ND	.01	ND	ND	.05	ND	ND	.01
D-D mixture	ND	ND	.1	ND	ND	.1	ND	ND	.1	NTD	ND	.2
MCPA	ND	ND	1.0	ND	ND	1.0	ND	ND	10	NTD	ND	20
Metalaxyl	ND	ND	1.0	ND	ND	1.0	ND	ND	10	ND	ND	.05
Methamidophos	ND	ND	2.0	ND	ND	2.0	ND	ND	.5	ND	ND	13
Methyl bromide	ND	ND	.5	ND	ND	.5	ND	ND	.5	ND	ND	.5
Methyl parathion	ND	ND	2.5	2.5	ND	2.5	ND	ND	1	ND	ND	.01
Molinate	ND	ND	1.0	ND	ND	1.0	ND	ND	.5	0.09	0.08	.01
Paraquat dichloride	ND	ND	20.0	ND	ND	20.0	ND	ND	10	ND	ND	10
Thiobencarb	ND	ND	8.0	ND	ND	8.0	ND	ND	1	0.08	0.07	.01
Xylene	ND	ND	1.0	ND	ND	. 2	ND	ND	.5	ND	ND	.2
Copper	ND	ND	5.0	6.0	16	5.0	5	ND	5			
Bromide	ND	ND	0.6	2.6	ND	0.6	ND	ND	.1			
Chloride	87	85	.2	64	64	.1	130	120	.02			

1.o.d. = limit of detection

ND = not detected

All units in ug/L except for bromide and chloride values which are in mg/L.

RP numbers (e.g. RP 56) are sample identification codes.

# Department of Health Services Evaluation

In May 1986, the Department of Health Services was asked to evaluate the performance of pesticide analyses by McKesson Environmental Services and the DWR Bryte Laboratory. River water was collected from the Sacramento River at Greene's Landing and spiked with a variety of pesticides. The staff of the DOHS Sanitation and Radiation Laboratory in Berkeley performed the spiking. The amounts and materials placed into the water samples were unknown to the DWR staff and to the laboratories. Duplicate sets of the spiked samples were delivered to MES and the Bryte Lab by the monitoring program staff. The Bryte Lab does not routinely perform pesticide analyses for the Health Aspects Monitoring Program because of limited

capabilities. However, samples were sent to Bryte to assess its current limitations and assist the laboratory in upgrading its capabilities.

Duplicate samples from three Delta locations were also submitted to each laboratory. These samples were not spiked.

Both laboratories were requested to analyze for specific compounds and report unidentified peaks in the chromatograms.

The reports of MES and the Bryte Lab were submitted to DOHS for review. The initial cursory review suggested major reporting discrepancies in the analysis for some compounds in the spiked reference samples and raised many points that needed clarification (Attachment 1) /4, 5/.

A meeting among DOHS, DWR, and MES representatives was held on October 3, 1986 to discuss and clarify the results. The meeting revealed a misunderstanding between MES and DWR on the reporting requirements that were expected and MES' reporting policy on trace contaminants and limits of detection by the laboratory. The full text of these discussions is presented in Attachment 2 /6/.

In summary, the qualitative assessment of the QA study indicated MES is capable of detecting the compounds spiked in the samples. Compounds spiked by DOHS but not reported by MES resulted when analyses or the

appropriate analytical methodology were not requested by DWR.

Another QA study will be conducted with the inclusion of the laboratories of The Metropolitan Water District of Southern California and East Bay Municipal Utility District.

The QA program has been effective in identifying laboratory problems and miscommunication between the Monitoring Program staff and the laboratories. In all cases, investigations have been fruitful and corrective actions have been taken. The limited QA activities thus far clearly demonstrate the importance of continuing QA as an integral part of the program.

#### APPENDIX D REFERENCES

- Michael Larson and Dr. Warren Steele, McKesson Environmental Services. Letter to Richard Woodard, Department of Water Resources. October 9, 1985.
- Dr. Michael Volz, Department of Health Services. Memorandum to B. J. Archer, Department of Water Resources. March 27, 1986.
- Dr. Warren Steel, McKesson Environmental Services. Letter to B. J. Archer, Department of Water Resources. April 28, 1986.
- Dr. Michael Volz, Department of Health Services. Memorandum to B. J. Archer, Department of Water Resources. September 15, 1986.
- Dr. Ben Tamplin and Dr. Michael Volz, Department of Health Services. Memorandum to B. J. Archer, Department of Water Resources. October 1, 1986.
- Dr. Michael Volz, Department of Health Services. Memorandum to B. J. Archer, Department of Water Resources. October 8, 1986.

Via:

## Memorandum

To : Mr. B. J. Archer, Chief

Water Quality and Reuse Section, Central District

Department of Water Resources (DWR)

Department of Water Resources (DWR)

B. R. Tamplin, Ph.D., Chief & for Sanitation and Radiation Laboratory

Michael G. Volz, Ph.D. Michael G. Volz, Ph.D.

From : Environmental Biochemist

Quality Assurance Officer

Sanitation and Radiation Laboratory

Date: September 15, 1986

Subject: QA Evaluation of

SRL Spike Sample Study with MES and

DWR/Bryte

Attached find a qualitative summary of analytical results (Table 1) and pertinent information (Table 2) generated by the Sanitation and Radiation Laboratory of the Department of Health Services (SRL), McKesson Environmental Services (MES), and DWR's Bryte Laboratory (DWR/Bryte) in support of the recent QA activity involving spikes of selected organic chemicals by SRL into river water supplied by DWR.

SRL attempted to meet as many as possible of DWR's requests for spiked samples pertaining to specific analytical groups in this study. However, as indicated in Table 2, we were limited by the breadth of our supply of stock reference samples and chronic problems with instrumentation requisite to substantiate spiked sample composition. Despite these inhibitions, the precision over 4 replications of the combined spiking and analytical protocols for many analytes was exceptionally good (Table 2). This suggests that each laboratory received representative spikes.

After an examination of the results, SRL recommends the following:

- (1) MES and DWR/Bryte should reevaluate their analytical data in support of the QA activity taking into account the information presented in Tables 1 and 2.
  - (a) Some spiked compounds originally not reported actually may have been seen on chromatograms but were not correctly identified.
  - (b) Other compounds not spiked into river water by SRL but reported by one or both of the other laboratories may simply be misidentifications in conjunction with (a) above or, in the case of analytes associated with those analyses not performed by SRL, may be reflective of actual contamination of the river water.
- (2) MES and DWR/Bryte should clarify their reporting procedure for laboratory data. We do not know if some spiked compounds were not reported simply because method and/or matrix "blank" concentrations were accounted for internally prior to the data reporting phase. We also do not know if Limits of Detection were nominal such as the MDLs in the EPA 600 series or whether the reported Limits of Detection were actually attained by the laboratories.

- (3) MES and DWR/Bryte should consider the impact(s) of knowing what chemicals specifically mentioned by DWR as requiring quantitation in this activity or as part of DWR's regular IDHAMP monitoring program may have influenced data interpretation following generic laboratory methodology. For example, if it was assumed for one or more reasons that certain substances were expected to be present, was it the convention to assume that the peaks found were "close" enough to warrant a "positive" finding in the absence of more substantive confirmatory information?
- (4) MES and DWR/Bryte should address how <u>previous</u> information and chromatographic characteristics in their respective data bases characterizing river water quality may have influenced qualitative interpretations of the data generated in this study.
- (5) <u>Quantitative</u> assessments regarding relative laboratory performance on specific analytical methodologies should be addressed in future communications.

Please contact us should you need further assistance at 8-571-2201 or (415) 540-2201.

- cc: G. W. Fuhs, Dr. sci. nat., DL/DHS
  - M. Jung, DWR
    - R. Woodard, DWR
    - A. del Rosario, SRL/DHS
    - S. Khalifa, Ph.D., SRL/DHS

Table 1
Qualitative Summary

Analytical Method	(ug/L)	Pres	Presence Reported*				
Chemical Compound	Spiked** by SRL	_SRL	MES	DWR/Brvte			
EPA 601/602	(0.5-3)						
Methylene chloride	(+)	+					
1,1-Dichloroethylene	(+)	+					
1,1-Dichloroethane	(+)	+	+				
Chloroform	(+)	+	+	+			
Carbon Tetrachloride	(+)	+	+	+			
1,2-Dichloropropane	(+)	+	+				
Trichloroethylene	(+)	+	+	+			
1,1,2-Trichloroethane	(+)	+	•				
Dibromochloromethane	(+)	+	+				
Tetrachloroethylene	(+)	+	+	+			
Chlorobenzene	(+)	+	+	+			
2-Chloroethyl vinyl ether	(+)#						
trans-1,2-Dichloroethylene	(+)	+	+	+			
1,2-Dichloroethane	( <del>^</del> )	· +	+				
1,1,1-Trichloroethane	(+)`	+ .	+	+			
Bromodichloromethane	(+)	+		+			
trans-1,3-Dichloropropene	(+)	+					
cis-1,3-Dichloropropene	(+)	+					
Benzene	(+)	(N/A)	+				
Bromoform .	(+)	+	+				
1,1,2,2-Tetrachloroethane	(+)	÷					
Toluene	(+)	(N/A)	+				
Ethylbenzene	(+)	(N/A)	+				
Dichlorobenzene	(-)	(N/A)		+ .			
EPA 608	(0.2-6)						
Dacthal	(+)	+	+	+			
Heptachlör	(+)	+					
Heptachlor Epoxide	(+)	+					
Lindane	(+)	+	•				
DDE	(+)	. +					
Endrin	(+)	+		+			
DDD	(+)	+		+			
DDT .	(+)	+					
Methoxychlor	(+)	+					

Analytical Method	ne vide	(ug/L)	Pr	esence Reported*
Chemical Compound	 	Spiked** by SRL	SRL	MES DWR/Bryte
EPA 614 Diazinon Methyl Parathion Ethyl Parathion Molinate Carbofuran Malathion		(0.6-0.9) (+) (+) (+) (-) (-) (-)	+ + +	+ + + + + +
EPA 622 2,4-D		(12-18) (+)	+	
EPA 632-HPLC Carbaryl Methomyl	* 3 * *	(5-8) (+) (+)	(N/A) (N/A)	
GC-ECD Chloropicrin		(N.S.) (-)	(N/A)	e de la composition de la composition La composition de la composition de la La composition de la
<u>Wet Chemistry</u> Paraquat		(N.S.) ('-)	(N/A)	
Others Atrazine/Simazine EDB		(N.S.) (-) (-)	(N/A) (N/A)	

- \* (+) denotes presence of chemical compound was reported; no entry denotes presence of chemical compound was not reported; data is from Summary Tables in memo of 8/14/86 from B. J. Archer (DWR) to Dr. B. R. Tamplin (SRL/DHS).
- \*\* (+) denotes chemical compound spiked into river water;
  - (-) denotes chemical compound was not spiked into river water.
- (N/A) Chemical compound was not analyzed for. See Table 2 for additional information.
- # Manufacturer cannot guarantee stability of this compound in standard mixture.
- N.S. Not spiked by SRL.

Table 2

SRL Analytical Support Information<sup>a</sup>

Analytical Method	Limits of Detection	Comment(s)
EPA 601	0.5 ug/L (Nominal)*	See *
EPA 602	(N/A)-0.5 ug/L (Nominal)*	Spiked with Benzene, Toluene, Ethylbenzene. See **
EPA 608	0.01-0.20 <sub>u.g</sub> /L	Method Spike Recoveries: 80 - 90 %: Range in precision for each analyte over all analytes: 1.3 - 11.1 %
EPA 614	0.02 - 0.05 ug/L	Method Spike Recoveries: Range in precision for each analyte over all analytes: 1.3 - 3.2 %
EPA 622	0.08 $_{u}g/L (2,4-D)^{\#}$	Method Spike Recovery:81% precision: ± 9.9%
EPA 632-HPLC	N/A	Spiked with Carbaryl and Methomyl. See **.
GC-ECD	N/A	Did not spike with Chloropicrin.
Wet Chemistry	N/A	Did not spike with Paraquat.

a - Analytical results derived from mean of 4 separate analyses (4 spiked bottles of river water.

<sup>\* -</sup> For purposes of reporting as per AB 1803 policy; for EPA 601 instrumental limits of detection (areal integration) range: 0.003 - 0.19 ug/L.

<sup>\*\* -</sup> Instrument non-operational.

 $<sup>{\</sup>rm N/A}$  - Analysis not performed by SRL/DHS.

<sup># -</sup> Analytical method (SRL/DHS) was from Application Scientist Vol. 1 (J. T. Baker) as per S. Khalifa, Ph.D.

### Memorandum

Mr. B.J. Archer, Chief
Water Quality and Reuse Section
Central District
Dept. of Water Resources (DWR)
P.O. Box 160088
3251 "S" St.
Sacramento, CA 95816

Date: October 8, 1986

Subject: QA Evaluation of MES' Performance on Spiked River Water Samples

Via B.R. Tamplin, Ph.D., Chief **Bax** Sanitation and Radiation Lab

From:

M.G. Volz, Ph.D.
Environmental Biochemist
Sanitation and Radiation Lab

On October 3, 1986, in Pleasanton, CA, Rick Woodard and Marvin Jung of your staff and I met with Dr. Warren Steele of DWR's contract laboratory, McKesson Environmental Services (MES). The purpose of the meeting was to discuss results of the recent Quality Assurance (QA) Study designed to evaluate the analytical proficiency of MES when DWR provided them with river water samples which had been previously spiked with selected organic compounds by the Sanitation and Radiation Laboratory (SRL) of the Department of Health Services (DHS). See attached memo of M. Volz to B. Archer, 9/15/86, for details.

Our discussion has revealed that, rather than analytical methodologies being highly suspect as might be concluded from a superficial evaluation of the attached results, the following statements better describe the data.

- 1. Some compounds like methylene chloride (a widely used organic solvent in extraction protocols) were not reported because of inherent contamination problems with both samples and blanks that are typical of commercial laboratory operations.
- 2. Certain compounds co-elute with others on chromatograms, e.g., several of the EPA m601 analytes, thus preventing definitive compound identification and subsequent reporting.
- 3. Many analytes in the EPA m608 scan were apparently detected on chromatograms by MES staff but were not reported except as "unidentified peaks" pursuant to prior agreement with DWR.
- 4. <u>Additional</u> compounds reported by MES in the EPA m614 methodology may be reflective of the actual presence of these pesticides in unspiked

Mr. B. J. Archer, Chief Page 2 October 8, 1986

river water. A similar argument could be made for Atrazine, Simazine, and EDB.

5. A compound like Bentazon (specifically requested as an analyte by DWR) would not have been seen using EPA m632. Hence, MES utilized an alternate procedure. However, the SRL spikes of Carbaryl and Methomyl then were not quantifiable by MES and not reported.

As a result of the above, SRL/DHS recommends the following:

- A. Each chemical which was spiked into river water by SRL but was not reported by MES should be evaluated as an individual analyte and be commented upon by MES to DWR.
- B. Similarly each chemical reported by MES but not spiked by SRL should be addressed as in (A). Those instances where the actual presence of compounds in unspiked river water may have been expected to occur should be differentiated from those where suspected or confirmed compound misidentification and reporting has taken place. In the future, unspiked river water also should be provided to participating laboratories to help resolve this issue.
- C. Careful evaluation of what truly was expected of MES by DWR and DHS with respect to each and every analyte and/or analytical method under consideration should be made. There appeared to be several instances of miscommunication in the QA Study. Resolution of these discrepancies is essential for future program-and cost effective QA activities in support of the IDHAMP.
- D. The performance of DWR's Bryte laboratory also should be carefully evaluated using criteria (A)-(C) above. Proficient laboratory support from this source is essential for the IDHAMP.

Mr. B. J. Archer, Chief Page 3 October 8, 1986

> Quantitative assessment of the present QA Study should be made only after the qualitative aspects described above have been resolved. Perhaps any quantitative assessment should be held in abeyance until EBMUD and MWD have entered future QA evaluations. They both indicated such an interest in our September 26, meeting.

For further information please contact this office at 8-571-2201 or (415) 540-2201.

G.W. Fuhs, Dr. sci. nat. P.R. Rogers, SEB
J. Crook, Ph.D., SEB cc:

D.P. Spath, Ph.D., SEB

F. Baumann, SCL

A. del Rosario, SRL

S. Khalifa, Ph.D., SRL

## Appendix E

TIDAL EFFECTS STUDY FIELD MEASUREMENTS

#### OLD RIVER TIDE CYCLE STATIONS

Program Station <u>Number</u>	DWR Station <u>Number</u>	Station Name
OR 4b	B9D75891348	Old River North of Rock Slough
OR 5a	B9D75821343	Old River opposite Rancho del Rio
OR 6a	B9D75571335	Old River South of Orwood
OR 9	B9D75351342	Old River near Byron (Highway 4)
OR 12	B9D75111331	West Canal at Old River
OR 15	B9D74971332	West Canal at Clifton Court Forebay Intake
OR 16	B9D74901334	Delta-Mendota Canal near Intake from Old River

#### MIDDLE RIVER TIDE CYCLE STATIONS

Program Station Number	DWR Station <u>Number</u>	Station Name
MR 2	B9D80011306	Middle River at Latham Slough
MR 3a	B9D75881321	Middle River North of Empire Cut
MR 5	B9D75741317	Middle River at Bacon Island Bridge
MR 12	B9D75431293	Middle River North of Highway 4 Bridge
MR 15b	B9D75311300	Victoria Canal near Middle River
MR 16	B9D75311282	Middle River West of Tracy Blvd.

#### POTATO SLOUGH TIDE CYCLE STATIONS

Program Station Number	DWR Station <u>Number</u>	Station Name
PS 1	B9D80611333	Mokelumne River near Mouth
PS 2	B9D80501344	San Joaquin River at Mouth of Potato Slough
PS 2a	B9D80531311	Potato Slough near Little Potato Slough
PS 3	B9D80371300	Little Connection Slough at Venice Ferry
PS 3a	B9D80481300	Little Potato Slough near Potato Slough
PS 4	B9D80561291	White Slough near Little Potato Slough

FIELD MEASUREMENTS DURING HIGH SLACK TIDE
Old River Sampling Run -- August 7, 1986

Time Depth Temp. E.C. Bottom

Station I.D.	Time PST	Depth ft.	Temp. F.	E.C. uS/cm	Bottom depth (ft.)	Comments	
OR-4B	0708	1	72	220	27	midchannel	
		3	72	220			
		6	72	225	100		*1
the state of the		9	72	235	10 miles 10	•	1.7
	0705	12	71	240			
	· .	15	71	240		à	
	0700	18	71 71	245			
the first of	0702	24	71	245	Territoria de la compansión		
OR-4B	0716	3	72	230	21.5	dragged anchor	
		6		230		to south	
		9	<sup>18</sup> - valestrika	235		•	
		12		235	A constant	slack water	
	0719	15		240		ended 0820	
OR-5A	0741	1		235	15	slack water	en en produce. Se se
		·· 3		230		ended 0840	. Interest
		6		230			
4.7		9		230	1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1		
******	0744	12		230			
OR-6A	0800	3		230	17	slack water	$\epsilon = \epsilon_i$
		6		230		ended 0910	
	0810	12	•	230	. 4		
OR9	0819	3.		230	24	sampled at	
		6		225	tan in the in	center support	
		9		225	e de	of Highway 4	
		12		225		bridge on	
		15	74	225		southside	na e e e
OR-12	0841	3	74	240	32	negative flow	y <sup>†</sup>
		6		245		observed	
		6 <b>9</b>		245	A	W. Canal site	100
	11. 1	18	$\gamma = \gamma^{-1/2} - T - n$	245	* * * * * * * * * * * * * * * * * * *		
	0845	24	74.5	250	•	· 6	$\mathcal{F}_{i}^{(\mathcal{F})} = \{\mathcal{F}_{i}^{(\mathcal{F})}\}$
ing the state of t	in district	147			English Control		
OR-15	0905	3	75	240	19	Clifton Ct.	
		9 18	75	240 240		gates all open	
OR-16C	0927	3	75	245	8	DMC intake	
		6		245			
OR-16S	0938	3	75	240	24	Old River	
		9		240		side	
		15		240		strong current	
		21		240		sampled depths are less than noted	

FIELD MEASUREMENTS DURING HIGH SLACK TIDE Old River Sampling Run -- August 21, 1986

Station I.D.	Time PST	Depth ft.	Temp. F.	E.C. uS/cm	Bottom depth (ft.)	Comments
OR-4B	0720	3	70	260	35	
		6		260		
		9		265		
		12		265		
		15		265		
	0726	18		265		
OR-4B	0810	3		245	35	At slack
		6		250		
		9		260		
		12		265		
		15		265		
		18		265		
		21		270		
		24	70	270		
OR-5A	0819	3	70	260	19	
		6		260		
		9		260		
		12		260		
		15		260		
	0821	18	71	260		
OR-6A	0836	3	71	260	21	
		6		260		
		9		260		
		12		260		
		15		260		
	0840	18		260		
OR-9	0853	3	72	265	20	
		6		265		
		9		265		
		12		265		
		15		265		
	0856	18		265		
OR-12	0917	3	72	280	28	strong current
		6		280		sampled depths
		9		280		less than noted
		12		280		
	0920	15		280		
OR-15	0925	3	72	285	20	Clifton Ct.
		6		285		gates closed
		9		280		
		12		280		
		15		280		
OR-16S	0935	3	72	290	14	Old River
		6		290		side of
		9		290		DMC intake
		12		290		
•	0937	15		290		
OR-16C	0943	3	72	300	17	Canal side of
		6		300		DMC intake
		9		300		
	0946	12		300		

FIELD MEASUREMENTS DURING HIGH SLACK TIDE Middle River Sampling Run -- August 21, 1986

Station	Time	Depth ft.	Temp.	E.C.	Bottom	Clammant =
I.D.	PST	IC.	F.	uS/cm	depth (ft.)	Comments
MR-1	0730	i	$C_{2}(X_{i}^{*})$	174	Programme Annual Control	Materials.
	0734	6		171		
	0,54	9	1''.	171		
		12	71, 10°	170		
	0725				/	
	0735	15	$t_{\rm s}$	170		
		18		169		
		21	4.00	169		
		24		168		
		27		168		
		30		168	:	
		_			8.5	
MR-3A	0747	1	: U	182	* # * **	
		6		180	e de la companya de l	
		9		180		
		12	1,111	180		· · · · · · · · · · · · · · · · · · ·
		15	Shark.	179		
		18		178	•	
		21	1.7	179		
		24	2	178		
		27	ì	178		
MR-5	0802	1	V. 4	258		
		6		259		
		9		260		
		12		260	No.	
		15		261	dia	e"
		18	100	260		
		21	1.0	260		
		24	,13.	260	0	
		27		260		
		30		260	*	
		. 30	3	200	2.00	
e visto de la composición del composición de la						
MR-12	0842	1		237		
IM IZ	0042	6		237	1.1	
		9				
				236		
The transfer of Value		12	-15	235	•	
		15	£ 45.	244	s *	
			i gir.			
MD 155	0007	-		05-		
MR-15B	0907	1	£n€-	251		
f 9 30, 1 4.		6	र्वतम् -	246		\$
1 - 1 - 1 - 1 - 1 - 1 - 1 - 1 - 1 - 1 -		9	184 (384	246		
a was directly and		12		246		
			56. C			
MR-16	0933	, 1	, jár	247	1877	
a trade and the second and the second		. 6	10, 1	249	7 X AM	127 44 7

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FIELD MEASUREMENTS DURING HIGH SLACK TIDE Potato Slough Sampling Run -- August 22, 1986

Station I.D.	Time PDT	Depth ft.	Temp. F.	Ē.C. uS/cm	Bottom depth (ft.)	Comments
PS-1	0800	 3	70	200		
10 1	0000	6	, ,	200		
		9		200		
		12		200		
		15		200		
		18		200		
	0802	21		200		
PS-2	0821	3	68	210		Windy and chopp
		6		210		,
		9		210		Sampled depths
		12		210		less than noted
	0822	21		210		
PS-2A	0843	3	70	180		•
		6		178		
		9		178		
		12		178		
	0844	15		178		
PS-3	0857	3	72	175		South of
		6		175		ag drain
		9		175		discharge
		12		175		on Empire Tr.
	0859	18		180		
DC 24	0000	2	70	100		
PS-3A	0908	3	70	180		
		6		180		
		9		183		
		12 15		185 185		
	0910			185		
	0910	18		103		
PS-4	0919	3	· 71	195		
		6		195		
		9		195		
		12		195		
	0920	15		195		

## Appendix F

## PESTICIDE MONITORING SELECTION SCHEME

#### PESTICIDE MONITORING SELECTION SCHEME

As part of the Interagency Delta Health Aspects Monitoring Program, surface waters were monitored for agricultural chemicals that might be difficult to control using conventional water treatment practices. In general, such chemicals are water soluble and have a low affinity for adsorption onto particulate matter. Consequently, flocculation, settling, and filtration processes are ineffective in removing these dissolved substances. On the other hand, chemicals with sparingly low water solubilities tend to be readily attracted to solid media and can be controlled in a typical treatment facility.

Selection of chemicals and timing for monitoring at a site can be difficult. Broad scans for hundreds of chemicals are expensive (thousands of dollars per sample) and do not produce significantly more information than does taking a sensible and rational approach. The continued practice of limiting analyses to traditionally monitored chemicals such as banned chlorinated pesticides may even be less productive in assessing current water quality conditions.

The Department chose to develop and use a selection scheme based on a combination of quantitative information (e.g. reported chemical usage patterns and properties) and judgmental assessments (e.g. major activities upstream of a sampling site). A database of the quantitative information was compiled for the selection process.

The objective of the scheme was to develop a list of those chemicals with the highest probability of posing treatment difficulties to public water supplies in the Delta. Chemicals on this list would be monitored.

The selection scheme produced site- and time-specific target lists of chemicals for monitoring. The scheme and database can also be used in other types of monitoring programs (e.g. ground water, biological contamination surveys) by using different selection criteria values (e.g. ranges of water solubilities and partition coefficients). Target lists could be developed for different environmental compartments (e.g. sediment, water, biota).

#### Method

Pesticide and crop pattern data of the State Department of Food and Agriculture were compiled to determine the amount and period of usage. Data were obtained for 1983, the most recent database containing a full year of record at the time of the compilation. Data for pesticide usage were ranked for each county and then combined for watersheds of interest

to this program (those encompassing our sampling sites). The chemicals were then ranked by usage for each watershed.

Information was compiled for each chemical on water solubility, log P (octanol/water partition coefficients), log Koc (soil activity coefficients), estimated half-life in water, period of use by month, type of use, and whether it was on the AB-1803 list. (The AB-1803 list is the California Assembly Bill 1803 list of chemicals that must be monitored in ground water by the Department of Health Services).

The octanol/water partition coefficient is defined as the ratio of a chemical's concentration in the octanol phase to that in the aqueous phase of a two-phase octanol/water system. The ratios are often reported in logarithmic units (log P). Values of P are meaningful since they represent the tendency of a chemical to partition itself between an organic phase (e.g. soil, fish) and an aqueous phase. Chemicals with low P values are relatively hydrophilic (i.e. water soluble) and have small soil/sediment absorption coefficients, and small bioconcentration factors for aquatic life. Chemicals with high P values (e.g. log P greater than 4) are very hydrophobic. The P values can be measured in the laboratory or estimated from water solubility relationships, knowledge of chemical structure, and other solvent/water partition coefficients.

The soil adsorption coefficient, Koc, is the ratio of the amount of chemical adsorbed per unit weight of organic carbon (oc) in the soil or sediment to that amount in solution at equilibrium. Logarithmic values, log Koc, are reported because of the high range of values. The degree of adsorption affects the chemical's mobility, volatilization, photolysis, hydrolysis, and biodegradation. Koc can be measured in the laboratory and estimated from empirical relationships with other chemical properties (e.g. solubility, log P).

Information on the chemical properties was compiled from numerous recent publications /1-11/ and the ISHOW (Information System for Hazardous Organics in the Water Environment) computer database of EPA. When conflicting values were found, the lower values were entered into the database. An excellent discussion of the degree of error associated with measurements of chemical properties is presented in Lyman et al /12/.

The chemicals were grouped by selected ranges of reported or calculated water solubilities and specified ranges of partition coefficients as measured by their affinities for water or organic-laden soil (e.g. by log P and log Koc values). Eight groups were created from the following criteria:

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Group	Water Solubility	log P and log Koc
1	> 999 mg/L	equal to or <2
2	> 999 mg/L	>2 but < or equal to 3
3	100-999 mg/L	equal to or <2
4	100-999 mg/L	>2 but < or equal to 3
5	10-99  mg/L	equal to or <2
6	10-99 mg/L	>2 but < or equal to 3
7	< 10  mg/L	equal to or <2
8	< 10 mg/L	>2 but < or equal to 3

A ninth group that would comprise those chemicals of log P or Koc values above 3 was not pertinent because it represented the very hydrophobic chemicals generally controllable in a modern water treatment plant.

Chemicals that had certain water solubilities and both log P and log Koc values were sorted and placed into the appropriate groups. However, those chemicals missing solubility data, log P, or Koc data were read as zero values by the computer software program, Lotus Symphony.

The groups represented those chemicals more likely to be dissolved in water (Groups 1 and 2) and those more likely to be in suspended material and organic particles in the water column (increasingly hydrophobic in order of group number).

The selection process for developing a list of candidate chemicals to be monitored consisted of inclusion of the most water soluble chemicals (Group 1 and 2 chemicals) and those with moderate water solubilities and partition coefficients (Groups 3 and 4). Additional pesticides, regardless of solubilities and partition coefficients, were added to the list when applied amounts were significant (among the top in ranked usage for the watershed) and the application method might lead to water contamination. For example, rice herbicides were added to the list because of the large quantities used and because they are applied to rice ponds just a few days before pond water and surface agricultural drainage are discharged into nearby rivers. To eliminate selection bias, each chemical was given a unique code for identification during the sorting and selection of pesticides for inclusion in the candidate lists. This step was taken to avoid inclusion of chemicals that technically might not meet the selection criteria but that were popular or traditional chemicals in other monitoring studies.

A final target list of chemicals to be monitored at specific sampling stations was developed after site location data on riverflow direction and upstream pesticide use and cropping pattern data were considered. This step reduced the list to those chemicals with the higher probability of contaminating waters upstream of the sites. For example, pesticide use data for the watershed where the American River water treatment plant is located represented use data for Sacramento, El Dorado, and Placer counties. The rice chemicals molinate and thiobencarb ranked high in use and were on the list of candidate

chemicals for monitoring. However, rice fields are not located upstream of this site and therefore these two chemicals were not on the final target list of chemicals to be monitored at the American River water treatment plant site.

Site- and time-specific target lists were developed, since information on the months of application (based on cropping patterns) were included in the database. The monthly target lists provided information on which water soluble chemicals would more likely be detected in water (dissolved phase) at the Delta sampling stations.

#### Conclusion

The database will be revised as new information on pesticide use, application, and physical-chemical properties is received. The success in developing target lists depends on the reliability and accuracy of such data. The resulting tabulations and information can also be used to predict which chemicals would be found in different compartments of an aquatic system (e.g. sediment, water, biota).

The described protocol illustrates the need to combine numerical selection criteria (e.g. usage, solubilities, and partition values) and non-numerical information (e.g. station location and upstream activities) to improve the possibility of detecting chemicals in the aquatic system.

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## Appendix G

# U. S. ENVIRONMENTAL PROTECTION AGENCY WATER QUALITY MODELS

#### APPENDIX G

## U. S. ENVIRONMENTAL PROTECTION AGENCY WATER QUALITY MODELS

Selected water quality models are now available through the Center for Water Quality Modeling for the personal computer. These models were taken from mainframe or minicomputer systems and are designed for the DOS environment on the IBM PC XT/AT family of microcomputers and compatible systems. The models are EXAMS, QUALZE, WASP3, DYNHYD3, PRZM, and MINTEQ.

#### **EXAMS**

The Exposure Analysis Modeling System is a steady state and dynamic model designed for rapid evaluation of the behavior of synthetic organic chemicals in aquatic ecosystems. EXAMS computes exposure (the ultimate expected environmental concentrations resulting from a long-term steady state pattern of pollutant loadings), fate (the distribution of the chemical in the environment and the fraction of the loadings consumed by each transport and transformation process), and persistence (the time required for effective purification of the system once the loadings cease). EXAMS is an interactive program and allows the user to specify and store the properties of chemicals and ecosystems, modify the characteristics of either via simple English-like commands, and conduct rapid, efficient evaluations of probable fate of chemicals.

#### QUAL2E

The Enhanced Stream Water Quality Model QUAL2E is a steady state model for conventional pollutants in branching streams and well mixed lakes. It includes conservative substances,

temperature, coliform bacteria, biochemical oxygen demand, dissolved oxygen, nitrogen, phosphorus and algae. QUAL2E is widely used for waste load allocation and permitting in the United States and other countries.

#### WASP3

The Water Quality Analysis Simulation Program is a generalized modeling framework for contaminant fate and transport in surface waters. Based on the flexible compartment modeling approach, WASP can be applied in one, two, or three dimensions. Problems that have been studied using WASP include biochemical oxygen demand-dissolved oxygen dynamics, nutrients and eutrophication, bacterial contamination, and toxic chemical movement.

A variety of water quality problems can be addressed with the selection of appropriate kinetic subroutines that may be either selected from a library or written by the user. Toxics WASP (TOXIWASP) combines a kinetic structure adapted from EXAMS with the WASP transport structure and simple sediment balance algorithms to predict dissolved and sorbed chemical concentrations in the bed and overlying waters. Eutrophication WASP (EUTROWASP) combines a kinetic structure adapted from the Potomac Eutrophication Model with the WASP transport structure. EUTROWASP predicts dissolved oxygen, carbonaceous biochemical oxygen demand, phytoplankton, carbon, and chlorophyll a, ammonia, nitrate, organic nitrogen, and ortho-phosphate in the bed and overlying waters.

#### DYNHYD3

DYNHYD3 is a simple 2 dimensional hydrodynamic model capable of handling variable tidal cycles, wind, and unsteady inflows. DYNHYD3 was updated from the Potomac Dynamic Estuary Model (DEM). This model has the ability to produce an output file that can be linked with WASP3 to supply the flows and volumes to the water quality model.

#### PRZM

The Pesticide Root Zone Model simulates the vertical movement of pesticides in unsaturated soil, within and below the plant root zone, and extending to the water table using generally available input data that are reasonable in spatial and temporal requirements. The model consists of hydrology and chemical transport components that simulate runoff, erosion, plant uptake, leaching, decay, foliar wash off, and volatilization (implicity) of a pesticide. Predictions can be made daily, monthly or annually.

#### **MINTEQ**

MINTEQ is a geochemical model that is capable of calculating equilibrium aqueous speciation, adsorption, gas phase partitioning, solid phase saturation states, and precipitation-dissolution of 11 metals (arsenic, cadmium, chromium, copper, lead, mercury, nickel, selenium, silver, thallium and zinc). MINTEQ can solve a broad range of chemical equilibrium problems for surface and ground waters. MINTEQ contains an extensive thermodynamic data set and contains 6

different algorithms for calculating adsorption.

#### SWMM and DYNTOX

Two other water quality models are under development for the PC environment. The projected release date for the distribution of these models was July 1, 1986. A brief description of each model is given below.

The Storm Water Management Model (SWMM) is a comprehensive model for simulation of urban runoff quantity and quality. All aspects of the urban hydrologic and quality cycles are simulated including surface runoff, transport through the drainage network, and storage and treatment (including cost). A choice of techniques is available for simulation in a sewer system, a kinematic wave procedure for most problem assessments, and a full equation routing method of surcharged systems. SWMM can be used for both single event and continuous simulation. It has been used in a planning context as well as for detailed design studies. SWMM also has a long history of use in the United States and Canada for urban drainage design.

DYNTOX is a waste load allocation procedure based upon dilution of whole effluent toxicity using a probabilistic modeling technique. It is a simple, interactive program using ANNIE as the user interface. DYNTOX can perform three types of simulations -- Continuous, Monte Carlo and Log Normal -- that, based on probabilities, can aid in deriving a waste load allocation limit.

Reference: U. S. Environmental Protection Agency. 1986. "Research and Development NEWSLETTER -- Water Quality Exposure and Risk Modeling". U. S. Environmental Protection Agency. EPA/600/M-86/018. July 1986.

## **CONVERSION FACTORS**

Quantity	To Convert from Metric Unit	To Customary Unit	Multiply Metric	o Convert to Metric Unit Multiply Customary Unit By
Length	millimetres (mm)	inches (in)	0.03937	25.4
	centimetres (cm) for snow depth	inches (in)	0.3937	2.54
	metres (m)	feet (ft)	3.2808	0.3048
	kilometres (km)	miles (mi)	0.62139	1.6093
Area	square millimetres (mm²)	square inches (in²)	0.00155	645.16
	square metres (m²)	square feet (ft²)	10.764	0.092903
	hectares (ha)	acres (ac)	2.4710	0.40469
	square kilometres (km²)	square miles (mi²)	0.3861	2.590
Volume	litres (L)	gallons (gal)	0.26417	3.7854
	megalitres	million gallons (10 <sup>6</sup> gal)	0.26417	3.7854
	cubic metres (m³)	cubic feet (ft³)	35.315	0.028317
	cubic metres (m³)	cubic yards (yd³)	1.308	0.76455
	cubic dekametres (dam³)	acre-feet (ac-ft)	0.8107	1.2335
Flow	cubic metres per second (m³/s)	cubic feet per second (ft³/s)	35.315	0.028317
	litres per minute (L/min)	gallons per minute (gal/min)	0.26417	3.7854
	litres per day (L/day)	gallons per day (gal/day)	0.26417	3.7854
	megalitres per day (ML/day)	million gallons per day (mgd)	0.26417	3.7854
	cubic dekametres per day (dam³/day)	acre-feet per day (ac- ft/day)	0.8107	1.2335
Mass	kilograms (kg)	pounds (lb)	2.2046	0.45359
	megagrams (Mg)	tons (short, 2,000 lb)	1.1023	0.90718
Velocity	metres per second (m/s)	feet per second (ft/s)	3.2808	0.3048
Power	kilowatts (kW)	horsepower (hp)	1.3405	0.746
Pressure	kilopascals (kPa)	pounds per square inch (psi)	0.14505	6.8948
	kilopascals (kPa)	feet head of water	0.33456	2.989
Specific Capacity	litres per minute per metre drawdown	gallons per minute per foot drawdown	0.08052	12.419
Concentration	milligrams per litre (mg/L)	parts per million (ppm)	1.0	1.0
Electrical Conductivity	microsiemens per centimetre (uS/cm)	micromhos per centimetre	1.0	1.0
Temperature	degrees Celsius (°C)	degrees Fahrenheit (°F)	(1.8 × °C)+3	12 (°F-32)/1.8