# 1. Title and Approval Page

# SOUTHERN CALIFORNIA MARINE INSTITUTE VOLUNTEER MONITORING QUALITY ASSURANCE PROJECT PLAN

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# 3. Distribution List

All group leaders, and technical advisors will receive copies of this Quality Assurance Project Plan (QAPP), and any approved revisions of this plan. Once approved, this QAPP will be available to any interested parties by requesting a copy from <u>the project management</u>.

# 4. **Project Organization**

The Southern California Marine Institute Volunteer Monitoring Project is a multi-organization project. Nine volunteer monitoring groups will monitor and assess natural streams, storm drains, harbors, estuaries, and nearshore ocean waters in the Los Angeles Basin. These organizations are:

- 1. Algalita Marine Research Non-Profit Foundation,
- 2. Bolsa Chica Conservancy,
- 3. Cabrillo Marine Aquarium (CMA),
- 4. Divers Involved Voluntarily in Environmental Rehabilitation and Safety (D.I.V.E.R.S.),
- 5. Heal the Bay,
- 6. Santa Monica BayKeeper,
- 7. Southern California Marine Institute (SCMI),
- 8. Surfrider- Huntington/Seal Beach Chapter, and
- 9. Surfrider- Long Beach Chapter

See Appendix A for a "Description of the Participating Organizations".

Several resource agencies have assisted in the development of this project from its conception. Additional partnerships will be developed to ensure adequate technical support to all participating volunteer monitoring groups. The QAPP reflects the diversity of monitoring and organizational support involved in this project. For the elements of this QAPP, we have addressed aspects that are shared with all groups as well as those aspects that are unique to individual groups. While the goals of monitoring may vary, the data quality objectives are consistent allowing us to compare data collected by different organizations.

### 4.1. Technical Advisory Committee for all Groups

Erick Burres, State Water Resources Control Board Rick Pieper, Southern California Marine Institute Rich Gossett, CRG Marine Labs

# 5. Problem Definition/Background

#### 5.1 Problem Statement

Many volunteer monitoring organizations exist in the Los Angeles Region. Prior to SCMI's involvement in citizen monitoring efforts, there had been little or no coordination between these organizations regarding sampling methods, parameters, sampling locations, data sharing, and other aspects. Education and coordination between all citizen monitors and regulatory agencies is necessary to provide valuable data and to maximize the results of monitoring efforts. SCMI has and will continue to coordinate citizen monitoring group efforts by holding training sessions, Quality Assurance/Quality Control (QA/QC) sessions, helping with data collection and upload to the state water board, and any other aspect that becomes necessary to maximize efficiency between groups.

#### 5.1.1. Citizen Monitoring and Goals

#### 5.1.1.1 Mission

The mission of citizen monitoring is to produce environmental information which is needed to protect California's watersheds and aquatic resources. Citizen monitoring will also inform and engage the community in effective watershed stewardship.

### 5.1.1.2 Program Goals

- To provide an illustrated field guide for sampling and analysis performed by citizen monitors. This field guide will be patterned after the model provided by the Heal the Bay Stream Team Field Guide.
- To encourage and increase public involvement, and to standardize data quality from citizens in citizen monitoring programs. SCMI will provide training, guidance, field consultations, and quality assurance sessions open to all of the region's citizen monitoring organizations.
- To expand and coordinate seasonal water monitoring "snapshot" efforts. The existing citizen monitoring effort within Region 4 of the State Water Resources Control Board (SWRCB) would be restructured and expanded in order to assess and report water quality on the same day in all the region's watersheds, which include: Los Angeles River watershed, San Gabriel River watershed, Dominguez Channel watershed and Santa Monica Bay Water Management Area (WMA). Several of these snapshot efforts will be performed per year. At least two seasonal snapshots and one near the timing of the first flush will be planned.
- To assist groups in data entry and transmittal, thereby assisting the Los Angeles Regional Water Quality Control Board (LARWQCB) staff in their water quality assessment and Total Maximum Daily Load (TMDL) efforts. All credible data collected by participating citizen groups and lead agency, in all of the local watersheds will be entered on a Microsoft Excel database and transmitted to the LARWQCB via email.
- To increase public awareness and stewardship of our water resources, thereby altering behavior and unsustainable land use practices, and resulting in lower pollution levels over time.
- Recommend revisions to the Southern California Volunteer Monitoring Quality Assurance Project Plan (QAPP).

## 5.2. Intended Usage of Data

The data will be used by the Southern California Marine Institute for general watershed assessment purposes. Assessment of this data will be useful in providing information for watershed management and pollution prevention. The data will be made public for watershed education. It will also be made available to regulatory and resource management agencies to supplement their existing data collection efforts. One potential application of the data will be to provide information to the Regional and State Boards for their use, if they so choose, in Clean Water Act Section 305(b) reporting.

Data will be collected and shared with the State Water Resources Control Board, the LARWQCB, and upon request to other state, federal, and local agencies and organizations. The main database will be maintained at Southern California Marine Institute at 820 S. Seaside Ave., Terminal Island, CA 90731.

# 6. Project/Task Description

## 6.1. General Overview of Monitoring

The citizen monitoring organizations are assessing water quality in Los Angeles County and northern Orange County including, but not limited to, the following watersheds: Los Angeles River, San Gabriel River, Dominguez Channel, Ballona Creek, Malibu Creek, San Diego Creek, Bolsa Chica, Anaheim Bay, Santa Ana River, and Santa Monica Bay. Physical, chemical and biological parameters are measured, although not all groups are measuring all parameters. Table 6.1 summarizes the monitoring design that relates to each of the participating groups, including the physical, chemical and biological parameters to be measured, whether the samples will be analyzed by the monitoring group or sampled for later analysis by a professional lab, and the frequency of measurement.

Parameter	Heal the Bay, type & frequency	Santa Monica Bay Keeper, type & frequency	Surf-rider type & frequency	Algalita, type & frequency	CMA, type & frequency	DIVERS, type & frequency	Bolsa Chica, type & frequency
Temperature	F, M	F, X	F, M	F, X	F, M	F, M	F, W
Dissolved Oxygen	F, M	F, X	F, M	F, X	F, M	F, M	F, W
РН	F, M	F, X	F, M	F, X	F, M	F, M	F, W
Conductivity	F, M	F, X	F, M	F, X	F, M	F, M	F, W
(fresh water) or							
Salinity (marine)							
Turbidity/	F, M	F, X	F, M	F, X	F, M	F, M	N/A
Transparency							
Ammonia	L, M	L, X	L, M	L, X	L, M	L, M	L, S
Nitrate	L, M	L, X	L, M	L, X	L, M	L, M	L, S
Ortho-Phosphate	L, M	L, X	L, M	L, X	L, M	L, M	L, S
Bacteria	L, M	L, X	L, X, P	L, X, P	Р, Х	P, X	P, S
Odor and Visual	F, M	F, X	F, M	F, X	F, M	F, M	F, W
Observations							

### Table 6.1. Summary of Monitoring Design

#### **Codes for Table 6.1:**

**Type**: F: field analysis, L: in-house lab analysis, P: sample only, send to outside professional lab **Frequency**: W: weekly, M: monthly, S: seasonal, X: irregular **N/A**: parameter not sampled

All of the water quality data will be compared to the Regional Water Quality Control Board Basin Plan. For results that are not comparable to the Basin Plan we will review those data with our Technical Advisors.

This QAPP only addresses data quality objectives for the following parameters:

Weather and Sensory Observations Temperature Dissolved Oxygen pH Conductivity Salinity Turbidity/Transparency Ammonia (nitrogen) Nitrate (nitrogen) Ortho-Phosphate Total Coliform Bacteria *E. coli* Bacteria Enterococcus Bacteria Chemical, physical, and bacterial parameters will be monitored using protocols outlined in a Field Guide based on Heal the Bay Stream Team Field Guide-hereafter referred to as the Field Guide.

# 7. Data Quality Objectives

This section identifies how accurate, precise, complete, comparable, sensitive, and representative our measurements will be. These data quality objectives were derived by reviewing the QAPPs and performance of other citizen monitoring organizations (e.g. Chesapeake Bay, Texas Watch, Coyote Creek Riparian Station, Southern California Citizen Monitoring Steering Committee, Heal the Bay, Malibu Stream Team), by considering the specifications of the instruments and methods employed, and by considering the utility of the data. For purposes of this QAPP the data quality is considered adequate for the determination of general water quality conditions, with a potential application of the data to Section 305(b) reporting purposes.

Data Quality Objectives (DQOs) are summarized in Tables 7-1 to 7-4. Whenever possible the methods with the greatest sensitivity and lowest detection limit will be employed as the primary methods. Methods with lesser sensitivity and higher detection limits will be used for field confirmations or as back-up methods in the case that the primary methods are not available or functioning properly for a particular sampling event. Specific DQOs were not given for *in-situ* continuous monitoring devices. See Section 14 for quality control protocols to be followed when continuous monitoring devices are employed.

Parameter	Method/Range	Units	Detection Limit	Sensitivity *	Precision	Accuracy	Completeness
Temperature	Thermometer (-5 to 50)	°C	-5	0.5 ° C	$\pm$ 0.5 $^{\rm o}$ C	$\pm$ 0.5 $^{\rm o}$ C	80%
Dissolved	Electronic meter/probe	mg/l	0.1 mg/l	0.1 mg/l	± 10%	± 10%	80%
oxygen	Modified Winkler Titration	mg/l	0.2 mg/l	0.2 mg/l	± 10%	± 10%	80%
pH	pH meter	pH units	2.0	0.1 unit	$\pm 0.2$ units	$\pm 0.2$ units	80%
Conductivity	conductivity meter	µS/cm	10	10 µS/cm	± 10%	± 10%	80%
Salinity	Electronic meter/probe	ppt	0.1	0.1 ppt	<u>+</u> 10%	<u>+</u> 10%	80%
	Refractometer	ppt	1	1 ppt	<u>+</u> 1 ppt	<u>+</u> 1 ppt	80%
	Nephelometer	NTUs		0.1	± 10%	± 10%	80%
Turbidity/ Transparency	Secchi Disk	Secchi Depth	0.1 m	0.1 m	± 0.1 m	± 0.1 m	80%
Transparency	Dual tube optical	JTUs	5	5 JTUs	± 5 JTUs	N/A	80%

#### Table 7.1. Data Quality Objectives for Conventional Water Quality Parameters

NA: not applicable

\* Note: Some test kits vary in sensitivity over the range of detection. The specific range of readings is noted in parentheses.

	~ F F						
Parameter	Method/Range	Units	Detection	Sensitivity	Precision	Accuracy	Completeness
			Limit				
Ammonia	Nessler method	mg/l	0.05	0.01	±0.2 (<2.0)	±0.2 (<2.0)	80%
Nitrogen					±10% (>2)	±10% (>2)	
Nitrate	Cadmium	mg/l	0.05	0.01	±0.2 (<2.0)	±0.2 (<2.0)	80%
Nitrogen	reduction				±10% (>2)	±10% (>2)	
Ortho-	Ascorbic acid	mg/l	0.07	0.01	±0.2 (<2.0)	±0.2 (<2.0)	80%
Phosphate					±10% (>2)	±10% (>2)	

## Table 7.2. Data Quality Objectives for Nutrients Using Colorimeters or Spectrophotometers

# Table 7.3. Data Quality Objectives for Nutrients Using Comparators

Parameter	Method/range	Units	Detection Limit	Sensitivity*	Precision	Accuracy	Completeness
Ammonia Nitrogen	Salicylate method	mg/l	0.25	0.25 (0-0.5) 0.5 (0.5-1.0) 1.0 (1.0-2.0) 2.0 (2.0-4.0)		± 1.0	80%
Nitrate Nitrogen	Cadmium reduction	mg/l	1.0	1.0	± 1.0	± 1.0	80%
Ortho- Phosphate	Ascorbic acid	mg/l	0.2	0.2 (0-1.0) 0.5 (1.0-2.0)	± 0.5	± 1.0	80%

\* Note: Some test kits vary in sensitivity over the range of detection. The specific range of readings is noted in parentheses.

## **Table 7.4. Data Quality Objectives for Biological Parameters**

Parameter	Method/Range	Units	Detection Limit	Sensitivity	Precision	Accuracy	Completeness
Total Coliform Bacteria	Colilert-18	MPN/ 100ml	10	See IDEXX quantitray tables	Duplicates within 95% confidence limits	Positive standard within <sup>1</sup> / <sub>2</sub> of an order of magnitude	80%
<i>E. coli</i> Bacteria	Colilert-18	MPN/ 100ml	10	See IDEXX quantitray tables	Duplicates within 95% confidence limits	Positive standard within ½ of an order of magnitude	80%
Entero- coccus Bacteria	Enterolert	MPN/ 100ml	10	See IDEXX quantitray tables	Duplicates within 95% confidence limits	Positive standard within <sup>1</sup> / <sub>2</sub> of an order of magnitude	80%

## 7.1 Accuracy

#### 7.1.1. Chemical and Physical Parameters

Accuracy describes how close the measurement is to its true value. Accuracy is the measurement of a sample of known concentration and comparing the known value against the measured value. The accuracy of measurements and analyses will be checked by performing tests on standards and spiked samples at the quality control sessions <u>held twice a year</u>. A standard is a known concentration of a specific solution. Standards can be purchased from chemical or scientific supply companies. Standards can also be prepared by a professional partner, e.g. a commercial or research laboratory. The concentration of the standards, known to the volunteer leader, will be unknown to the monitors until after measurements are determined. The concentration of the standards should be within the mid-range of the equipment. The Data Quality Form: Accuracy, found in the Appendix, will be used to record accuracy and all QA data will be captured as per the Clean Water Team's (CWT) Data Quality Management (DQM) Guide.

#### 7.1.2. Biological Parameters

Accuracy for bacteria will be determined by analyzing a positive control sample twice annually. A positive control is similar to a standard, except that a specific discreet value is not assigned to the bacterial concentrations in the sample. This is because bacteria are living, reproducing organisms, with variable mortality rates. Instead of a specific value, an approximate target value of the bacterial concentration is assigned to the sample by the laboratory preparing the positive control sample. Split samples will be taken amongst the different groups and with certified labs.

## 7.2. Comparability

Comparability is the direct correlation of collected data to similar studies. Uniformity in data collection is essential for the utility of the data in the state database. Citizen monitoring groups will use the methods described in the following resource documents to ensure that their data can be compared to others:

- U.S. EPA's Volunteer Monitoring Manuals for Streams, Lakes or Estuaries,
- SWRCB Clean Water Team Compendium for Water Quality Monitoring and Assessment, and
- California's Department of Fish and Game's (CDFG) California Stream Bioassessment Protocol (CSBP) for Citizen Monitors.
- Heal the Bay's Malibu Creek Stream Team Pilot Project, Shattering the Myths of Volunteer Monitoring
- San Francisco Estuary Institute's Volunteer Monitoring Protocols.

Before modifying these methods, or developing alternative or additional methods, technical advisors will evaluate and review the effects of the potential modification. It will be important to address their concerns about data quality before proceeding with the monitoring program. Conformity of data is an essential part of citizen monitoring programs.

### 7.3. Completeness

Completeness is the fraction of planned data that must be collected. There are no statistical criteria that require minimal data. However, it is expected that 80% of all measurements could be taken when anticipated. This accounts for adverse weather conditions, safety concerns, and equipment problems.

Completeness is determined by comparing the number of measurements planned on being collected compared to the number of measurements actually collected, also deemed valid. An invalid measurement would be one that does not meet the sampling methods requirements and the data quality objectives. Completeness results will be checked quarterly.

This will allow for the identification and correction of problems. The Data Quality Completeness Form, found in Appendix B6, will be used to record completeness. It will alsocontribute to statewide data conformity and utility.

### 7.4. Precision

#### 7.4.1. Chemical and Physical Parameters

The precision objectives apply to duplicate and split samples taken as part of a QC session or as part of periodic in-field QC checks. Precision describes how well repeated measurements agree. The evaluation of precision described here relates to repeated measurements taken by either different volunteers on the same sample (at quality control sessions) or the same volunteer analyzing replicate samples (in the field). Sampling variability will not be covered in this section. The Data Quality Precision Form, found in Appendix B6, will be used to record precision.

#### 7.4.2. Biological Parameters

Precision for bacterial parameters will be determined by having the same analyst complete the procedure for laboratory duplicates of the same sample. At a minimum this should be done once per day, or run duplicates on a minimum of 5% of the samples if there are over 20 samples run per day. The results of the duplicates should be within the confidence limits supplied by the manufacturer.

#### 7.5. Representativeness

Representativeness describes how relevant the data are to the actual environmental condition. Problems can occur if:

- Samples are taken in a stream reach that does not describe the area of interest (e.g. a headwaters sample should not be taken downstream of a point source),
- Samples are taken in an unusual habitat type (e.g. a stagnant backwater instead of in the flowing portion of the creek),
- Samples are not analyzed or processed appropriately, causing conditions in the sample to change (e.g. water chemistry measurements are not taken immediately).

Representativeness will be ensured by processing the samples in accordance with Section 10, 11 and 12, by following the established methods, and by obtaining approval of this document.

### 7. 6. Method Detection Limit and Sensitivity

The Method Detection Limit is the lowest detectable concentration for the instrument, chemical procedure, or equipment. This is important because it can never be determined if a pollutant was not present, only that it was not detected. Sensitivity refers to the detectable differences in concentration for test instruments. Thus, the sensitivity is the number of decimal places that are useable and repeatable. Detection Limits and Sensitivities are noted in Tables 7.1. - 7.5.

# 8. Training Requirements

All citizen monitoring leaders must participate in a minimum of three days of hands-on training sessions on water quality monitoring conducted by the Clean Water Team of the State Water Resources Control Board or equivalent.

In addition to completion of the above described training course, the citizen monitoring leaders must participate in **<u>semi-annual</u>** Quality Control Sessions. These Quality Control Sessions will be supervised by Quality Control Trainers and will provide an opportunity for citizen monitors to check the accuracy and precision of their equipment and techniques. Quality Control Trainers are defined as water quality professionals from the U.S. Environmental Protection Agency, the State Water Resources Control Board, and the Regional Water Quality Control Boards. Additional qualified trainers may be recruited and designated by the above agencies from experienced citizen monitoring organizations, universities and colleges, commercial analytical laboratories, and other federal, state, and local agencies.

The volunteer leaders will bring their equipment to the Quality Control Session. Monitors will conduct duplicate tests on all analyses and meet the data quality objectives described in Section 7. If a monitor does not meet the objectives, the trainers will re-train and re-test the monitor. If there is insufficient time at the QC session to re-train and re-test monitors, the monitor will be scheduled for an additional training sessions as required. The volunteer leader will be encouraged to discontinue monitoring for the analysis of concern until training is completed.

The Quality Control Trainers (Trainers) will examine kits for all of components: date, condition, and supply of reagents, and whether the equipment is in good repair. The Trainers will check data quality by testing equipment against blind standards. The Trainers will also ensure that monitors are reading instruments and recording results correctly. Sampling and safety techniques will also be evaluated. The Trainer will make noted recommendations to the volunteers to improve sampling or safety procedures. The citizen monitoring leader is responsible for the follow-up reports. Certificates of completion will be provided once all corrective action has been completed or the data quality objective is obtained (see Appendix B-5). Refresher courses after initial training will be provided as needed, or when methodology changes.

# 9. Documentation and Records

All field results will be recorded at the time of collection using the field data sheets (see Appendix B-1). Data sheets will be reviewed for outliers and omissions before volunteers leave the sample site. Data sheets will be signed after review by the citizen monitoring leader. Data sheets will be stored in hard copy form at the location as specified in Section 5.2. Field data sheets are to be archived for three years from the time they were collected. If data entry is ever performed at another location, duplicate data sheets will be used, with the originals remaining at the headquarters site. Hard copies of all data as well as computer back-up disks are maintained at headquarters, and a second copy is to be kept at a different location.

# **10.** Sampling Process Design

### 10.1. Rationale for Selection of Sampling Sites

Sampling sites vary by group. The following criteria were evaluated when choosing sampling locations:

- access is safe,
- permission to cross private or posted public property is granted,
- sample can be taken in main river current or where homogeneous mixing of water occurs,
- sample is representative of the part of the water body of interest,
- location complements or supplements historical data, and
- location represents an area that possesses unique value for fish and wildlife or recreational use.

Reference sites are chosen upstream of any potential impact. A site chosen to reflect the impact of a particular discharge, tributary or land use is located downstream of the impact where the impact is completely integrated with the water, but upstream of any secondary discharge or disturbance.

Prior to final site selection, permission to access the water body was obtained from all property owners except publicly accessable water bodies. If access to the site becomes a problem, the citizen monitoring leader will select a new site following the site selection criteria identified in Section 10.1. Safety concerns are discussed in the field guide.

Sample sites will be reviewed by the leader before sending citizen monitors out to the site. The monitoring leader will document permission and terms obtained from landowners.

### **10.2.** Sample Design Logistics

Citizen monitors are instructed to work in teams of at least two people. If a scheduled team cannot conduct the sampling, the team captain is instructed to contact the citizen monitoring leader so that arrangements can be made for a substitute trained citizen monitor.

Safety measures will be discussed with all citizen monitors. No instream sampling will be conducted if there are small creek flood warnings or advisories, marine or small boat craft advisories, or storm advisories. It is the responsibility of the citizen monitoring organization to ensure the safety of their citizen monitor monitors. Safety issues are included in the Field Guide.

# 11. Sampling Method Requirements

The Field Guide describes the appropriate sampling procedure for collecting samples for water chemistry. Water sampling apparatus may include Van Dorn Samplers, Niskin Bottles, Kemmerer Tubes, LaMotte Oxygen Samplers. In those cases where glass bottles are required in Table 11.1, plastic samplers are allowed as long as the hold time in the sampling device is minimal before transfer to the glass sample bottle. Sampling devices and sample bottles (that are not pre-sterilized and do not contain preservatives/fixing agents) will be rinsed three times with sample water prior to collecting each sample. For sterile bottles, whirl-paks, and sample bottles which contain preservatives/fixing agents (e.g., acids, etc.) <u>never</u> rinse with sample water prior to collecting the sample. Also, never use a sample bottle containing preservatives/fixing agents for sampling; in these cases always use a sampling device to collect the sample prior to transferring the sample into the bottle.

Whenever possible, the collector will sample from a bridge so that the water body is not disturbed from wading. All samples are to be taken approximately in mid-stream, at least one inch below the surface. If it is necessary to wade into the water, the sample collector stands downstream of the sample, taking a sample upstream. If the collector disturbs sediment when wading, the collector will wait until the effect of disturbance is no longer present before taking the sample.

The following table describes the sampling equipment, sample holding container, sample preservation method and maximum holding time for each parameter.

Parameter	Sample Bottle	Preferred / Maximum Holding Times
	Conventional Parame	ters
Temperature	clear plastic bottle or sample directly	immediately
Dissolved oxygen	plastic bottle or sample directly	immediately / for wet chemistry fix per protocol instructions, continue analysis within 8 hr.
pН	plastic bottle or sample directly	immediately
conductivity	plastic bottle or sample directly	immediately / refrigerate up to 24 hours.
salinity	plastic bottle or sample directly	immediately
turbidity	plastic bottle or sample directly	immediately / store in dark for up to 24 hours.
	Nutrients	
Ammonia N	plastic bottle	immediately / up to 8 hours if the sample is acidified with sulfuric acid to less than 3.0 pH.
Nitrate N	plastic bottle	immediately / refrigerate in dark for up to 48 hours.
Ortho-Phosphate	plastic bottle	immediately / refrigerate in dark for up to 8 hours.
	<b>Biological Samples</b>	3
Bacteria	sterile plastic sampling bottle or whirl-pak	Refrigerate to 4 degrees C in the dark; delivered to the lab within 4 hours, start analysis within 6 hours.

**Table 11.1 Sampling Method Requirements** 

# 12. Sample Handling and Custody Procedures

### 12.1. Sample Handling

Identification information for each sample will be recorded on the field data sheets (see Appendix 2) when the sample is collected. Samples that are not processed immediately in the field will be labeled with the waterbody name, sample location, sample number, date and time of collection, sampler's name, and method of preservation.

### 12.2. Custody Procedures

The conventional water quality monitoring tests do not require specific custody procedures since they will, in most cases, be conducted immediately by the same person who performs the sampling. In certain circumstances (such as driving rain or extreme cold), samples will be taken to a nearby residence for analysis. Samples requiring chemical preservation will be fixed prior to transport.

When samples are transferred from one citizen monitor to another member of the same organization for analysis, or from the citizen monitoring group to an outside professional laboratory, then a Chain of Custody Form should be used. This form identifies the waterbody name, sample location, sample number, date and time of collection, sampler's name, and method used to preserve sample (if any). It also indicates the date and time of transfer, and the name and signature of the sampler and the sample recipient. In cases where the sample remains in the custody form. It is recommended that when a sample leaves the custody of the monitoring group, then the Chain of Custody Form used be the one provided by the outside professional laboratory. Similarly, when quality control checks are performed by a professional lab, their samples will be processed under their chain of custody procedures with their labels and documentation procedures.

### 12.3. Disposal

All analyzed samples or spent chemicals (except for waste from the nitrate/cadmium reduction test and the Nessler ammonia test) including used reagents, buffers or standards will be collected in a plastic bottle clearly marked "Waste" or "Poison". This waste material will be disposed of according to appropriate state and local laws and regulations. This will usually mean disposal into a drain connected to a sewage treatment plant.

Liquid waste from the cadmium reduction nitrate test will be kept separate and disposed of at a facility that is permitted to handle, transport, or dispose cadmium (Cd) waste. Liquid waste from the Nessler ammonia test (which contains mercury) will likewise be kept separate and disposed of at a facility that is permitted to handle, transport, or dispose mercury (Hg) waste. Waste from the salicylate ammonia test can be held in the regular waste container and disposed of as described in the previous paragraph.

Whenever possible, if waste includes reagents from the detergent test, these wastes will be poured down a drain underneath a flume hood.

# 13. Analytical Methods Requirements

Water chemistry is to be monitored using protocols outlined in the Field Guide. The methods were chosen based on the following criteria:

- capability of citizen monitors to use methods,
- provide data of known quality,
- ease of use,
- methods can be compared to professional methods in *Standard Methods*<sup>1</sup>.

If modifications of methods are needed, comparability will be determined by side-by-side comparisons with a US EPA or APHA Standard Method on no less than 50 samples. If the results meet the same precision and accuracy requirements as the approved method, the new method will be accepted.

<sup>&</sup>lt;sup>1</sup> Standard Methods for the Examination of Water and Wastewater is a comprehensive reference that covers all aspects of water and wastewater analysis techniques. A joint publication of the American Public Health Association, the American Water Works Association, and the Water Environment Federation, the most recent *Standard Methods*, 20th Edition was published in 1998.

Table 13.1 outlines the methods to be used, any modifications to those methods, and the appropriate reference to a standard method.

Parameter	Method	Modification	Reference (a)
Temperature	Thermometric	Alcohol-filled thermometer	2550 B.
remperature		marked in 0.5°C increments	
	Winkler Method,	Pre-packaged reagents, 20 ml	4500-O C.
Dissolved Oxygen	Azide Modification	sample size	
Dissolved Oxygen	Membrane Electrode	None	4500-O G.
	Colorimetric indigo carmine	Vacuum ampoules	ASTM D 888-87
pН	Electrometric	none	4500-Н В.
	Litmus indicator strips	Non-bleeding	Whatman Co.
Conductivity	Electrometric	none	2520 B.
Q = 1;; (	Electrometric	None	2520 B
Salinity	refractometer	None	
Turbidity	Nephelometric	None	2130 B
Transparency	Secchi disk	None	
	Phenate	Salicylate with Color	4500 – NH3 F.
Ammonia N		Comparator	
Ammonia N	Nessler or	prepackaged reagents,	4500 – NH3 C 18 <sup>th</sup>
	Phenate/Salicylate	colorimeter or	edition only (1992)
		spectrophotometer	
	Cadmium Reduction	Color Comparator	$4500 - NO_3^{-}E.$
Nitrate N	Cadmium Reduction	prepackaged reagents, colorimeter or spectrophotometer	4500 – NO <sub>3</sub> <sup>-</sup> E.
	Ascorbic acid	Color Comparator	4500 – P E.
Ortho-Phosphate	Ascorbic acid	prepackaged reagents, colorimeter or spectrophotometer	4500 – P E.
Total Coliform	Colilert-18	none	9223 B
Bacteria			
E. coli Bacteria	Colilert-18	none	9223 B
Enterococcus Bacteria	Enterolert	none	IDEXX Corp.

<b>Table 13.1</b>	Analytical	Methods for	Water	Quality H	Parameters
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<sup>(</sup>a) All of the above methods, with the exception of dissolved oxygen via indigo carmine, pH via non-bleeding indicator strips, turbidity via dual tube (JTUs), and Enterococcus bacteria are described in Standard Methods for the Examination of Water and Wastewater 20<sup>th</sup> Edition. American Public Health Association *et al.*, 1998.

# 14. Quality Control Requirements

Quality control samples will be taken to ensure valid data are collected. Depending on the parameter, quality control samples will consist of blanks (field, trip and laboratory), replicate samples, and split samples. The temperature of the sample transport container will also be measured. In addition, quality control sessions (a.k.a. intercalibration exercises) will be held twice a year to verify the proper working order of equipment, refresh citizen monitors in monitoring techniques and insure data quality objectives are met.

## 14.1. Cautions Regarding Test Procedures

#### 14.1.1. Winkler Method for Dissolved Oxygen

The Winkler method is not appropriate for highly alkaline waters.

Other citizen monitoring groups have noted problems with short shelf-life of the sodium thiosulfate reagent. Field measurements should be evaluated immediately to determine whether they are reasonable. The validity of the dissolved oxygen test will also be assured by taking these steps:

- Care is taken not to aerate water samples during collection,
- Water is added gently to the dissolved oxygen bottle,
- No air bubbles are present in the sample,
- The titration sample will be measured carefully with a graduated cylinder,
- The sample is swirled thoroughly after each drop of titrant,

If the endpoint is overrun, another 20 ml. of the sample will be titrated.

#### 14.1.2. Nutrients

The nitrate test measures nitrite as well as nitrate. Therefore the results for the nitrate test are actually mg/l Nitrite + Nitrate Nitrogen. When mixing nitrate reagents citizen monitors should take care not to agitate samples aggressively. Phosphate reagents (the LaMotte chemical test kits) have been shown to degrade well within their listed shelf life once opened.

#### 14.1.3. Urban Pollutants

The tests for detergent, chlorine, and phenol should not be conducted on saline waters. Suspended matter and algae may give false positive results for detergent. The low sensitivity of the copper test may preclude detecting copper as most falls out of solution forming copper carbonate.

## 14.2. Blanks, Replicates, Split Samples, and Standardization

<u>Field/Trip/Laboratory Blanks</u>: In our testing, a field blank will be distilled water exposed to the sampling environment at the sample site, a trip blank will be distilled water taken from the laboratory and into the field and returned to the lab unopened, and a lab blank will be distilled water put through the entire lab procedure and analysis. For all conventional water quality analyses, except temperature, dissolved oxygen and pH, field and trip blanks will be analyzed once daily. For nutrients using comparators, a field blank will be analyzed every sampling trip. Color can sometimes appear in these nutrient blanks, suggesting that the real samples may be overestimating the true nutrient concentration. When colorimeters or spectrophotometers are used at the group's facility for nutrient analysis, a laboratory reagent blank will be analyzed and recorded for each day of analysis. For urban pollutants, field and trip blanks will be run daily. For bacterial analysis performed at a group's facility, a laboratory blank will be performed for each sampling/analysis event.

<u>Instructions for Field, Trip, and Lab Blanks</u>: Distilled water is taken into the field or used in the laboratory and handled just like a sample. It will be poured into the sample container and then analyzed. When reagents are used in a test method, then the reagents are added to the distilled water and these types of blanks are referred to as reagent blanks. Field blanks are recorded on the field data sheet. For nutrients measured with comparators, results from the field reagent blanks should be "not detected". If nutrients are detected, corrective action will be taken to eliminate the problem. For nutrients measured with colorimeters, the lab reagent blanks should be less than 0.05 ppm and the specific value should be recorded and subtracted from the field sample result. For bacterial analysis, the reagents are added to distilled water (in the same manner as for a field sample) and that blank is then sealed in a quantitray and incubated along with the field samples. The blank should be below detection limits (i.e., no positive wells) at the end of the incubation period.

<u>Field Confirmations</u>: When a second method for measuring temperature, dissolved oxygen, pH, turbidity, and salinity is available in the field, then the monitors are encouraged to perform both measurements on a split sample at least once daily. Examples of this sort of redundant measurement would be:

- For temperature, the use of a thermistor (or electronic thermometer) (such as those that are built into dissolved oxygen meters) and an armored thermometer;
- For dissolved oxygen, the use of an oxygen meter and an indigo carmine colorimetric kit;
- For pH, a meter and a non-bleeding indicator strip,
- For turbidity, a nephelometer and a Secchi disk,
- For salinity, an electrode (such as those built into dissolved oxygen meters) and a refractometer.

This will serve to provide backup capability if the more sensitive electronic meters fail, and will provide additional confidence as to the quality of the data. The results of both measurements will be recorded along with the procedure used on the field data sheet. If both results are comparable, then the result produced using the method of greater sensitivity will be the one entered in the final data set by the data manager in consultation with the monitoring leader. If the two results are inconsistent, then the monitoring leader will note on the data sheet which of the results will be entered on the final data set by the data manager.

<u>Replicate Samples</u>: Replicate samples are two or more samples collected at the same time and place. When there are only two replicates then these are referred to as duplicates. For conventional water quality, nutrients, and urban pollutant analyses duplicate field samples will be taken once every 20 samples, or quarterly whichever comes first. Duplicate samples will be collected as soon as possible after the initial sample has been collected, and will be subjected to identical handling and analysis. For bacterial analysis, lab duplicates will be run at least once per sampling day, and when there are more than 20 samples run per day then there will be a minimum 5% of the samples analyzed in duplicate.

<u>Split Samples</u>: Twice a year, split spiked samples (standards) will be analyzed as part of the Quality Control Session. The split standard is one sample, containing a known concentration of an analyte, that is divided equally into two or more sample containers. Split standards will be analyzed by the citizen monitors, and sent to a professional laboratory (except for dissolved oxygen, temperature, and pH), before the maximum sample handling time is exceeded. Citizen monitors will analyze the split standard normally and will perform at least three analyses on that same sample. From these results accuracy and precision will be determined. The professional laboratory will analyze the sample using the method referenced in Table 13.1

For turbidity using the dual tube (JTU) method, split field samples will be analyzed as part of the Quality Control Session. The laboratory receiving the split sample will analyze it using the nephelometric method, even though these results are not strictly comparable to the visual JTU comparators. The results of turbidity using the two methods will be plotted to determine if there is a linear correlation. If this correlation is significant, then it will be used to estimate and compare results of the turbidity tubes with nephlometric results. The Technical Advisory Committee for all groups will use the product-moment correlation coefficient (r) to determine the adequacy of the correlation.

For bacteria, split field samples or split positive controls will be analyzed by the citizen monitoring group and an outside professional laboratory twice annually. In addition, at the quality control session different analysts from the citizen monitoring group(s) will each read a minimum of the three quantitrays and compare their results. These results should be within  $\pm$  one well for concentrations of less than 1000 MPN/100 ml, and within  $\pm$  two wells for concentrations of greater than 1000 MPN/100ml.

<u>Standardization of Instruments and Procedures:</u> At the Quality Assurance Sessions the temperature measurements will be standardized by comparing our thermometers to a National Institute of Standards and Technology (NIST)-certified or calibrated thermometer in ice water and ambient temperature water. All meters (pH, conductivity, oxygen) will be evaluated at the Quality Assurance Session using standards provided with the assistance of a professional laboratory and/or the technical advisors. For oxygen meters the standard will be distilled water saturated with oxygen. The Winkler kits for dissolved oxygen will be checked by standardizing the sodium thiosulfate solution in the test kit, and/or by comparing the entire kit to a saturated oxygen standard. Instructions for checking the sodium thiosulfate are included in the test kit. (Additional reagents and glassware must be purchased separately however.) If the result is unsatisfactory, as indicated in the instructions, the sodium thiosulfate and/or other reagent will be discarded and replaced with new reagents.

<u>Continuous Monitoring Devices:</u> Should continuous monitoring devices be used for any parameters then such devices must be calibrated and deployed according to the manufacturer's specifications and field confirmation will be performed using replicate sampling (for laboratory analysis) or standardized instruments. For example, there is the possibility of using *in-situ* continuous monitoring devices for flow or temperature measurements. Confirmations using a flow meter or a standardized field thermometer will be performed at the time of deploying and retrieving the device. This will serve to determine the accuracy of the continuous monitoring device.

Table 14.1 Summarizes the Quality Control Requirements.

Parameter	Blank	Duplicate Sample	Split Sample to lab	QC session				
Water quality								
Temperature	none	5% or a minimum of once a year	none	twice a year				
Dissolved oxygen	none	5% or a minimum of once a year	none	twice a year				
рН	none	5% or a minimum of once a year	none	twice a year				
Conductivity/Salinity	daily	5% or a minimum of once a year	twice a year	twice a year				
Turbidity/Transparency	daily	5% or a minimum of once a year	twice a year	twice a year				
		Nutrients (comparators)						
Ammonia	daily	5% or a minimum of once a year	twice a year	twice a year				
Nitrate	daily	5% or a minimum of once a year	twice a year	twice a year				
Ortho-Phosphate	daily	5% or a minimum of once a year	twice a year	twice a year				
	Nutr	ients (colorimeters or spectrophotometer	rs)					
Ammonia	daily	5% or a minimum of once a year	twice a year	twice a year				
Nitrate	daily	5% or a minimum of once a year	twice a year	twice a year				
Ortho-Phosphate	daily	5% or a minimum of once a year	twice a year	twice a year				
Biological Parameters								
Total Coliform and <i>E. coli</i> Bacteria	daily	5% or a minimum of once per day	twice a year	twice a year				
Enterococcus Bacteria	daily	5% or a minimum of once per day	twice a year	twice a year				

### Table 14.1 Summary of Quality Control Requirements

# 15. Instrument/Equipment Testing, Inspection, and Maintenance

## 15.1. Temperature

Before each use, thermometers are to be checked for breaks in the column. If a break is observed, the alcohol thermometer will be placed in hot water causing the alcohol to expand and fill the column. Verify accuracy by comparing with a calibrated or certified thermometer.

## 15.2. Dissolved oxygen

<u>Dissolved Oxygen Winkler Titration:</u> Before each use, bottles, droppers, and color comparators are checked to see if they are clean and in good working order. If the pH is high (alkaline), alternative testing methods should be used. Reagents are replaced annually according to manufacturer's recommendation.

<u>Dissolved Oxygen Meters</u>: Membranes and solutions should be replaced according to manufacturer's specifications, but at least quarterly. Membranes should be checked for bubbles after replacement. Before each use, D.O. meters are checked to see if they are clean and in good working order.

## 15.3. Conductivity and pH

Before each use, conductivity and pH meters are checked to see if they are clean and in good working order. Conductivity and pH meters are calibrated before each use. Conductivity standards and pH buffers are replaced at least annually. Conductivity standards are stored with the cap firmly in place and in a dry place kept away from extreme heat. Do not re-use pH or conductivity standards.

## 15.4. Turbidity and Transparency

<u>Secchi:</u> Meter measurements on the Secchi disk line should be checked for accuracy. Two measurements should be taken, one when the disk just disappears, and one when the disk just reappears.

<u>Dual Tube Turbidity (JTU's)</u>: Before each use, turbidity tubes are checked to ensure that they are clean. The turbidity standard will be replaced annually.

<u>Nephelometers:</u> Meters and tubes should be checked for cleanliness and proper operation. The tubes should not be smudged or scratched.

### 15.5. Nutrients and Urban Pollutants

Before each use, test kits are checked to ensure that droppers, sample containers, and color comparators are clean and in working condition. Colorimeter tubes should be checked to make sure they are clean and are not scratched. Reagents are replaced annually or according to manufacturer's instructions.

# 16. Instrument Calibration / Standardization and Frequency

Instruments will be calibrated and reagents checked against standards accordingly to the following schedule. Standards will be purchased from a chemical supply company or prepared by (or with the assistance of) a professional laboratory. Calibration records will be kept in the maintenance log at the headquarters location (described in Section 5.2.) where it can be easily accessed before and after equipment use. Calibrations that are performed by monitors in the field are recorded on the field data sheets, also archived at the headquarters. The frequency of calibration is described in Table 16.1.

Conventional Water Quality Parameters				
Equipment Type	Calibration Frequency	Standard or Calibration Instrument Used		
Temperature	Every 6 months	NIST calibrated or certified thermometer		
Dissolved Oxygen	Every 6 months	Check sodium thiosulfate and/or against a		
(Winkler)	-	saturated oxygen standard every 6 months.		
Dissolved Oxygen	Every sampling day	At a minimum, water saturated air, according		
meter		to manufacturer's instructions.		
рН	Every sampling day	pH 7.0 buffer and one other standard (4 or		
		10)		
conductivity	Every sampling day	Conductivity standard and distilled water		
salinity	Every sampling day	Check accuracy with distilled water.		
Transparency	Every 6 months	Check marked lengths on the line to a metric		
(Secchi disk)		tape measure.		
Turbidity meter	Every sampling day	For clear ambient conditions use an 1.0 NTU		
(nephelometer)		standard, for turbid conditions use an 10.0		
		NTU standard		
Dual Tube Turbidity	Every sampling day	Distilled water		

## **Table 16.1 Instrument Calibration and Frequency**

Nutrients (using comparators)			
Equipment type	Checked against Standard	Standard Used	
Ammonia	every 6 months or when reagents replaced	ammonia standard	
Nitrate	every 6 months or when reagents replaced	nitrate standard	
Ortho-Phosphate	every 6 months or when reagents replaced	ortho-phosphate standard	

Nutrients (using colorimeters or spectrophotometers)				
Equipment type	Checked against Standard	Standard Used		
Ammonia	every 6 months or when reagents replaced	ammonia standard		
Nitrate	every 6 months or when reagents replaced	nitrate standard		
Ortho-Phosphate	every 6 months or when reagents replaced	ortho-phosphate standard		

# 17. Inspection/Acceptance Requirements

Upon receipt, buffer solutions, standards, and reagents used in the field kits will be inspected by the citizen monitoring leader for leaks or broken seals, and to compare the age of each reagent to the manufacturer's recommended shelf-life. All other sampling equipment will be inspected for broken or missing parts, and will be tested to ensure proper operation.

Before usage, thermometers are inspected for breaks in the alcohol column. Breaks can be eliminated by heating (see Section 15.1). If not, they will be returned to the manufacturer.

Reagents are replaced before they exceed manufacturer's recommended shelf life. These shelf lives are typically one to two years. However, specific replacement dates can determined by providing the reagent lot number to the manufacturer. Reagent replacement dates are noted in the maintenance log.

# **18.** Data Acquisition Requirements

## **18.1.** Professional Analytical Data

Only certified analytical laboratories or academic laboratories (with approval of State and/or LARWQCB staff) will be used for quality assurance checks and analysis of field samples. The Technical advisory

Committee (TAC) or technical advisors will review these laboratories' data as well as the citizen monitors. They may also review the lab's own quality control data to ensure data validity.

## 18.2. Geographical Information/ Mapping

United States Geologic Survey (USGS) maps will be used to verify watershed boundaries and river courses. National Oceanographic and Atmospheric Administration (NOAA) navigation charts can be used for mapping marine sampling sites. Additional information on distribution of natural resources will be obtained from the National Park Service (NPS) and the California Department of Fish and Game (CDFG) Biodiversity database. Land use information will be obtained from local planning offices. When information is requested, the agency will be asked to provide appropriate megadata and any information on data limitations. This information will be maintained with the data files.

# **19.** Data Management

Field data sheets are checked and signed in the field by the citizen monitoring leader. The citizen monitoring leader will identify any results where holding times have been exceeded, sample identification information is incorrect, samples were inappropriately handled, or calibration information is missing or inadequate. Such data will be marked as unacceptable by the monitoring leader and will not be entered into the electronic data base.

As part of standard field protocols, any sample readings out of the expected range will be reported to the citizen monitoring leader. A second sample will be taken as soon as possible to verify the condition. If the data is invalid, then the data will be noted (flagged) on the data sheet. We will take further actions to trace the sources of error, and to correct those problems. If the error is a result of improper monitoring procedures, then we may re-train monitors until their performance is acceptable. It is the responsibility of the citizen monitoring leader to re-train citizen monitors until performance is acceptable.

Independent laboratories will report their results to the citizen monitoring leader. The leader will verify sample identification information, review the chain-of-custody forms, and identify the data appropriately in the database. These data are also reviewed by the technical advisors quarterly.

The data management coordinator will review the field sheets and enter the data deemed acceptable by the citizen monitoring leader and the technical advisors. Range checks and other QA/QC methods will be performed before accepting the data set and data analysis. Upon entering the data the data management coordinator will sign and archive the field data sheets. Data will be entered into a spreadsheet (MS Excel) or a database (MS Access) in a way that will be compatible with EPA's STORET<sup>2</sup> and the Regional WQCB's database guidelines. Following initial data entry the data coordinator will review electronic data, compare to the original data sheets and correct entry errors. After performing data checks, and ensuring that data quality objectives have been met, data analysis will be performed.

Raw data will be provided to the State WQCB and Regional WQCB in electronic form at least once every two years so that it can be included in the 305(b) report. Appropriate quality assurance information may be provided on request.

# 20. Assessment and Response Actions

Review of all field and data activities is the responsibility of the citizen monitoring leader, with the assistance of the technical advisory committee. Citizen monitors will be accompanied by the citizen monitoring leader, or a technical advisor on at least one of their first 5 sampling trips. If possible, citizen

<sup>&</sup>lt;sup>2</sup> EPA's STORET (short for STOrage and RETrieval) is a repository for water quality, biological, and physical data and is used by state environmental agencies, EPA and other federal agencies, universities, private citizens, and many others.

monitors in need of performance improvement will be retrained on-site. All citizen monitors must attend a refresher course offered by the citizen monitoring group. If errors in sampling technique are consistently identified, retraining may be scheduled more frequently.

# 21. Reports

Raw data screened by the Quality Assurance (QA) officer will be made available to data users per request. The citizen monitoring organization(s) will report their data to its (their) constituents only after QA/QC standards have been reviewed and approved by their technical advisors. Every effort will be made to submit data and/or a report to the State and/or LARWQCB staff in a fashion timely for their data uses, e.g. 305(b) reports.

# 22. Validation and Verification Methods

As part of standard field protocols, any sample readings out of the expected range will be reported to the citizen monitoring leader. A second sample will be taken as soon as possible to verify the condition. If the data is invalid, then the data will be noted (flagged) on the data sheet. We will take further actions to trace the sources of error, and to correct those problems. If the error is a result of improper monitoring procedures, then we may re-train monitors until their performance is acceptable. It is the responsibility of the citizen monitoring leader to re-train citizen monitors until performance is acceptable.

# APPENDICES

# Appendix A – Description of the Participating Organizations

## **1.0** Algalita Marine Research Non-Profit Foundation (AMRF)

AMRF's focus is on nearshore marine habitat and recreational waters affected by urban runoff. Currently, proliferation of marine dinoflagellates in Long Beach Harbor is causing a permanent red tide heretofore unknown in these waters. The relationship of urban runoff to red tide has been one focus of AMRF research since 1995. The possibility of toxic dinoflagellate proliferation is an important consideration in our research. Like the algae blooms that plague European seaports and those that kill Florida manatees, our permanent red tide may be yet another sign that humans are changing coastal environments in ways that could have serious consequences for wildlife and people.

#### **Technical** Advisors

Bill Yanko, LA County Sanitation Chuck Mitchell, Marine Biological Consultants Dr. Balchandra Bave, Consultech Walton Wright, professional environmental consultant

#### Group Leaders

Charles Moore, Algalita Marine Research Non-Profit Foundation

# 2.0 Bolsa Chica Conservancy

The Bolsa Chica Conservancy is a non-profit, non-political organization whose mission is to advocate the restoration and preservation of the Bolsa Chica Wetlands through public outreach, participation, education and leadership. We are a coalition of responsible community leaders from science, business, education, and government. In summary: *We restore. We educate. We advocate.* 

#### **Technical Advisors**

To be announced.

#### Group Leaders

Claire Grozinger, Bolsa Chica Conservancy Rick Wojciechowski, Bolsa Chica Conservancy

## 3.0 Cabrillo Marine Aquarium

Cabrillo Marine Aquarium interprets the marine life of southern California and is a facility of the City of Los Angeles Department of Recreation and Parks. Located by the Los Angeles Harbor, over 400,000 visitors come to the Aquarium annually and approximately 1 million visitors spend time at the surrounding coastal habitats of sandy beaches, rocky shore and salt marsh. The mission of the Aquarium is to provide education, research, conservation and recreation opportunities for the visitors. Over 400 volunteers assist the staff with interpretation during tours, classes and public programs. Additional volunteers assist with

animal husbandry and environmental monitoring including ongoing research surveys. Volunteers are trained to use field equipment in conducting surveys, to collect data and to help with maintaining a database. These results are shared with visitors to the aquarium and will be used to get a long term profile of the health of the surrounding habitats.

#### **Technical** Advisors

Technical advisor, Michael Shaadt, Cabrillo Aquarium

#### **Group Leaders**

Linda Chilton, Cabrillo Marine Aquarium

# 4.0 Divers Involved Voluntarily in Environmental Rehabilitation and Safety (D.I.V.E.R.S.)

D.I.V.E.R.S. focus is on near shore marine habitat and recreational waters affected by urban runoff. Our study proposes to identify characteristics of pollutants and effluent of city streets and storm drains with respect to their effect on the locale specific to Main Beach in Laguna Beach.

#### **Technical** Advisors

Technical advisors to be recruited.

#### Group Leaders

Dawn Marcova Martin Carreon

# 5.0 Heal the Bay

Heal the Bay's goal is to work with the California Coastal Conservancy to implement a pilot Malibu Creek Watershed Citizen Monitoring Program. The objective of the pilot program is to create the foundations for watershed "watchdogs" that will result in citizen stewardship of the watershed. Watchdogs, ultimately, are those who can singly or collectively, identify problems in the watershed. Heal the Bay believes that stewardship will lead to greater awareness of pollution problems, an eventual reduction of pollutant loads to the watershed, and greater protection of the Malibu Creek Watershed's beneficial uses-- habitat for fish and wildlife, and swimming, diving, and surfing. Heal the Bay would like to see as part of the overall goals of volunteer monitoring the formation of a standardized data format within watersheds such as the use of Geographic Information Systems to map and display results of the monitoring. Heal the Bay feels that a standardized format will make integrating and analyzing data easier and more useful to the public and decision makers.

#### **Technical Advisors**

Dr. Mark Gold, Executive Director, Heal the Bay, D.Env., Environmental Science and Engineering: UCLA 1994. M.A., Biology: UCLA, 1986. B.S., Biology: UCLA, 1984 Dr. Randal Orton, , D.Env., Environmental Science and Engineering: UCLA, 1992. Ph D. Population genetics: UCLA 1989. M.A., Marine Biology: San Francisco State University, 1983. B.S. Biology: UCLA, 1979.

#### Group Leaders

Lisa L. Dobbins, Program Director, Heal the Bay. Mark Abramson, Malibu Creek Watershed Volunteer Monitoring Project Coordinator, Heal the Bay.

## 6.0 Santa Monica BayKeeper

The primary mission of the Santa Monica BayKeeper is to continually survey the environmental health of the Santa Monica Bay, San Pedro Bay and adjacent coastal waters and watersheds; to alert the public to potential hazards and expose those who contribute in any way to the degradation of this ecosystem. Storm drain pollution contributes largely to the degradation of the Santa Monica Bay. The pollutants in the Santa Monica Bay come from thousands of different sources and from communities throughout Los Angeles County. The trash, used oil and other toxic materials that are dumped into gutters, or discarded on streets all over the county are eventually washed by rains into storm drains and flow directly into the Bay. Volunteer monitoring enables the BayKeeper to monitor each of the 350 storm drains that flow into the bay so that we may identify and stop pollution sources. In our most recent wet weather sampling we were able to identify a drain that had unusually high levels of particular contaminants. The BayKeeper traced the runoff to a point source and found that there was a facility whose runoff was washing directly into a gutter that flowed out where the sample was taken. After working with the polluter they have changed their facility drastically so as to eliminate the runoff.

#### **Technical** Advisors

Ladan Mohajerani, Santa Monica BayKeeper Jim Noblet, PhD, Santa Monica BayKeeper

#### Group Leaders

Kris Haddad, Santa Monica BayKeeper Terry Tamminen, Santa Monica BayKeeper Angie Bera, Santa Monica BayKeeper

# 7.0 Southern California Marine Institute (SCMI)

#### Technical Advisors

Rich Gossett, CRG Marine Labs Rick Pieper, Director, SCMI

#### **Group Leaders**

Kerry Flaherty For EMP: Carrie Wolfe, and participating teachers For Yacht Clubs: Tom Eskridge, Long Beach Yacht Club

# 8.0 Surfrider - Huntington/Seal Beach Chapter

The goals of the Surfrider monitoring program in Orange County are to identify pollutants or contaminants and pollution sources to Orange County beach waters.

#### Technical Advisors

Technical advisors will be recruited.

#### Group Leaders

Don Schulz

# 9.0 Surfrider – Long Beach Chapter

The goals of the Surfrider monitoring program in Long Beach are to identify pollutants or contaminants and pollution sources to Long Beach waters.

## **Technical** Advisors

Rich Gossett, CRG Marine Labs

#### Group Leaders

Emiko Kobiashie

# Appendix B – Data Sheets and Forms

- 1 Field Data Sheet
- 2 Volunteer Water Quality Monitoring Sign in Sheet

3 – Volunteer Monitors Inter-Calibration and QA Session Chemical and Physical Parameters

- 4 Citizen Monitor Training Evaluation Form
- 5 Certificate of Completion
- 6 Data Quality Forms (Accuracy, Precision, Completeness)

Appendix C – Map of Volunteer Sampling Stations