



Brake Pad Partnership Castro Valley Atmospheric Deposition Study

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ABSTRACT

This report summarizes the results of a study of atmospheric deposition of copper and other trace elements conducted in Castro Valley, California from March 2004 through February 2005. The primary objective of the study was to characterize the wet and dry deposition of selected trace metals and trace organic pollutants from the atmosphere to the Castro Valley Creek watershed, as part of a larger effort led by the Brake Pad Partnership examining the contribution of brake pad material to the release of copper in the environment and its potential to negatively impact environmental water quality. Precipitation and dry deposition samples were collected at four sites spread to measure deposition across a range of urban development and activity in the watershed. For several sampling events, ambient (gaseous) samples were also collected and analyzed for benzene.

Average dry deposition fluxes of copper at the sites in Castro Valley ranged from 12 to 31 $\mu\text{g}\cdot\text{m}^{-2}\cdot\text{d}^{-1}$. Deposition rates varied widely among sites and among sampling events within any site over the course of the study. Average concentrations of copper measured in precipitation or bulk samples ranged approximately 1.8 to 6 ppb ($\mu\text{g}/\text{L}$). Resultant average bulk and wet deposition fluxes in precipitation ranged 3.5 to 9 $\mu\text{g}\cdot\text{m}^{-2}\cdot\text{d}^{-1}$ for copper, with a large amount of variation among sampling events. The wet and dry deposition rate combined could therefore total around 40 $\mu\text{g}\cdot\text{m}^{-2}\cdot\text{d}^{-1}$, well above previous results for sites in the San Francisco Estuary, but comparable to rates measured in other urban areas in the United States..

Large temporal variability in measured deposition rates was found as expected. Both wet and dry deposition rates are influenced by conditions before and during each sampling period, which typically spanned two weeks for wet and bulk deposition samples, and two days for dry deposition. Dry deposition samples were also collected during weekend periods and show small and inconsistent differences between weekday and weekend deposition rates.

Significant differences in deposition at the lowest versus highest traffic sites were found, with differences among the three monitoring sites in the densely developed areas low in the watershed being less distinct. Deposition differences among high and low traffic sites may also in part be caused by differences in elevation, with the lowest traffic areas located high in the watershed, requiring substantial vertical transport for pollutants generated in the valley to be seen.

1.0 INTRODUCTION

Pollutants such as copper are emitted into the atmosphere either naturally by processes such as wind driven soil resuspension, or by human activities such as industrial discharges stationary sources such as furnaces and mobile sources like motor vehicle operation. The distance air pollutants are carried depends on multiple factors such as weather conditions, the nature and location of the source, and the particle size distribution of pollutants (or the aggregates they may form). Deposition of atmospheric pollutants to a watershed may occur through several processes, including rain scavenging of gases and particles, and dry deposition of particles. Because metals such as copper are largely not found in the gaseous phase, diffusive exchange and adsorption to surfaces are not significant deposition pathways of concern for these pollutants.

Previously, gross calculations of atmospheric deposition were made based on ambient air monitoring results of the California Air Resources Board (CARB) and the Bay Area Air Quality Management District (BAAQMD) (Kirschmann and Grovhoug 1996, Hauri 1998a, Hauri 1998b). In 1999-2000, the Regional Monitoring Program Atmospheric Deposition Pilot Study (Tsai, 2001) directly examined deposition in precipitation and onto surrogate surfaces at three sites around the estuary, and those methods are used in this study to evaluate deposition of copper and other trace elements in the Castro Valley Creek watershed.

This deposition monitoring was initiated and funded as part of a larger Brake Pad Partnership study examining the emission and transport of copper to the environment from vehicle brake pads. For this study, the San Francisco Estuary Institute (SFEI), coordinator of the previous RMP Atmospheric Deposition Pilot Study, conducted field sample collection and conducted the study in collaboration with Professor Thomas Holsen's Laboratory at the Clarkson University Department of Civil and Environmental Engineering, which provided advice, sampling materials and analytical support.

Methods used were largely identical to those of the RMP Atmospheric Deposition study, based on monitoring methods for trace metals in dry deposition (Holsen et al. 1993, Yi et al. 1997a, Yi et al. 1997b, Paode et al. 1998) and in precipitation (Vermette et al. 1995b) used by other researchers. The RMP study estimated contributions of pollutants to the estuary via tributaries using simplistic assumptions of runoff and retention and in watersheds. In this study we do not attempt to track pollutant transport beyond its deposition to the surface. Modeling of subsequent transport processes is addressed in the airshed modeling conducted by AER and watershed modeling by to be performed by EPA.

This report describes the methodology used in the Brake Pad Partnership Castro Valley Atmospheric Deposition Study (hereafter simply referred as this study) and presents the results obtained from monitoring in this study for copper and other selected trace elements. This report also presents results from more limited monitoring of gaseous benzene as a compound representative of vehicle exhaust gases. Sampling methods for ambient benzene were derived from those described in EPA Method TO-15 (EPA 1999), as specified by the contract analytical laboratory, Air Toxics, Ltd.

2.0 OBJECTIVES

Copper, a pollutant of concern in San Francisco Bay, is present in some formulations of brake pads. Copper from brake pads therefore likely contributes in part to copper loading in urban runoff to the San Francisco Bay. The Brake Pad Partnership (BPP), a consortium of manufacturers, environmental organizations, stormwater management agencies, and regulatory stakeholders have come together to implement a program to identify and prevent water quality impacts that may arise from the use of copper in automotive brake pads.

Current plans are to delist the Lower San Francisco Bay as a copper-impaired water body, but only on the condition that copper levels remain constant or decrease. However, copper content in brake pads has increased to meet safety regulations and consumer demand. Concerns of increased copper loadings to the Bay in urban runoff have therefore arisen, and this study, part of a larger Brake Pad Partnership effort, aims to better quantify any potential impacts that brake wear material will have on the environment to better mitigate or prevent any negative effects.

The BPP has selected the Castro Valley Creek watershed as an area for monitoring and modeling to be representative of mixed urban and open space land uses that may be found in the San Francisco Bay area. Information from these linked studies will be examined with the intention of identifying possibilities for extrapolating or extending data and models developed for this small watershed to other watersheds in the region.

The primary objectives of this study (within the larger BPP project) are to:

- ?? measure atmospheric deposition of copper to the watershed surface
- ?? determine the spatial and temporal variability of deposition
- ?? measure other trace elements and benzene to estimate or bound the contribution of vehicle versus other sources to copper to the watershed.

Characterizing and quantifying brake emissions, modeling of atmospheric and hydrologic transport, and water monitoring in Castro Valley Creek are other major elements of the larger BPP project. This air deposition monitoring will be used in evaluating the performance of the modeling efforts to verify that the developed models reasonably reflect the actual behavior and ultimate fate of brake pad material in the environment.

3.0 METHODOLOGY

Deposition of air pollutants to the Estuary and surrounding watersheds can occur by several processes, including rain scavenging of gases and particles, dry deposition of particles, deposition associated with condensation processes, and air-water exchange processes. Air-water exchange does not occur for the non-volatile trace metals addressed in this report, and deposition through condensation is not specifically targeted for quantitation in this study, although the methods employed may incidentally collect some metals deposited through this pathway. This study of atmospheric deposition consisted of sample collection during rain events for wet and/or bulk deposition (depending on the equipment available at the location and the weather conditions during the collection period) and collections during non-rainy periods for dry and/or bulk deposition.

The wet and bulk deposition portion of the project measured pollutant concentrations in rainfall, captured by specific precipitation like those used in the National Atmospheric Deposition Program Mercury Deposition Network (NADP/MDN) or with bulk deposition collectors for locations and times where use of the automated precipitation collectors was not possible or practical. This study employed a direct method for dry depositon monitoring, measuring pollutant concentration in particles deposited and trapped on a surrogate surface (a greased plate) during dry (non-rainy) periods. Other studies such as one recently conducted by SCCWRP and UCLA in Southern California (Lu et al, 2001) used an indirect method for measuring pollutants in ambient air and calculating deposition using relevant equations, and assumptions or measurements of particle size distributions and concentrations and meteorological parameters. The direct method using surrogate surface plates was chosen for this study to mirror the previous atmospheric monitoring study in this region. Ambient gaseous benzene was also measured in a limited number of samples using evacuated canisters following EPA Method TO-15.

An initial design for this study was included in the BPP Proposition 13 grant proposal to collect wet and dry atmospheric deposition samples from two sites in Castro Valley every two weeks over the course of a year. The study design was reviewed and refined subsequently by the BPP steering committee and Science Advisory Team (SAT). Components that evolved and were modified after the initial proposal over the course of the study included the suite of measured chemicals, monitoring duration and frequency, and sampling site selection. A brief description of the study scope and methods is presented below.

3.1 Monitoring Parameters

This study was designed to measure both dry deposition and wet deposition in precipitation (rain). Discussions among the BPP steering committee and SAT sought sought chemical indicators specific to brake wear material but none were identified. Antimony was identified as one metal enriched in vehicle sources, and measurements of elements common in soils such as iron, aluminum, and silica (Bradford, 1996) were included as possible indicators of non-vehicle sources. However, the latter elements are also common in automotive materials, so any estimates of “non-vehicle” contributions to copper deposition on the basis of these other compounds would represent likely represent overestimates or upper bounds of such sources.

Benzene, an aromatic hydrocarbon, constitutes approximately 1-2% of gasoline (CARB, 1997) and is emitted in vehicle exhaust from a combination of unburned fuel and as a by product of partial combustion of other fuel components. Benzene was added late in the study in September 2004 as an additional monitored pollutant and potential indicator of vehicle sources.

3.2 Monitoring Duration and Frequency

The aim of this study was to measure wet and dry deposition over the course of a year at various sites within the Castro Valley Creek watershed. Monitoring frequency and duration of wet deposition samples for most of the Study was set on a 14-day schedule, to include the same proportion of weekday and weekend days in most of the collected samples. Early in the study in spring 2004, some wet deposition collections were extended for one or two weeks during periods without precipitation, in an attempt to capture at least one rain event per deployment. Bulk deposition samplers were deployed in areas where automated wet deposition samplers could not be used, and later in the study during late spring, and early fall 2004, periods when precipitation was sporadic and the ability to collect wet samples would not be reliable. Wet or bulk deposition samples were taken beginning in March 2004 and ending February 2005.

Dry deposition samples were generally collected approximately twice a month beginning July 2004, but the samples were not distributed uniformly throughout each month. The two day exposure periods for the dry deposition samples needed to be scheduled during periods without rain, so samplers could not be deployed on a regular set schedule. Furthermore, with exposure periods lasting only 2 days, significant differences between weekend and weekday samples could perhaps be captured. However, because large changes in meteorological conditions that could occur over the course of a week or longer might also change deposition rates in collected samples for reasons unrelated to weekend or weekday differences in traffic and urban activity, all weekend samples were deployed immediately after weekday dry deposition samples were collected, to minimize the influence of changing weather or other longer term causes of variability. Dry deposition samples were collected until the end of the study in February 2005. However, only one dry sample was taken at each site per month for most of the wet season months in the study (Table 2).

3.3. Monitoring Stations

The Castro Valley Creek watershed encompasses most but not all of Castro Valley. Castro Valley is highly urbanized, with a mix of residential, commercial, light industrial, and undeveloped (open space) land uses. There are many congested roads within Castro Valley, with the highest volume roads being the Highway 580/238 corridor, Castro Valley Boulevard, and Redwood Road.

In the San Francisco Bay region, winds originate from the west and northwest during most of the year, (BAAQMD 1998). In the winter, stormy periods with moderate and strong winds (>5 m/sec) are interspersed with stagnant periods with very light winds (<1 m/sec) when air masses can flow from various directions. The SF Bay region has a "Mediterranean" climate with distinct wet (November to April) and dry seasons (Trujillo et al. 1991).

Where possible, especially for placing the wet deposition samplers, sampling locations were selected to comply with the siting criteria prescribed in the National Atmospheric Deposition Program (Bigelow 1984). Initial criteria for sampling locations in this Study were as follows:

- ?? accessible and secure
- ?? adequate AC power (for wet deposition samplers only)
- ?? within the Castro Valley Creek watershed
- ?? not in proximity to known major stationary sources or located up-wind (prevailing winds) of such sources

At the start of the study, two sampling sites were chosen using these criteria in order to accommodate the wet deposition samplers. Castro Valley Community Center (CVCC) is located on Lake Chabot Road, a four-lane road leading from downtown Castro Valley to Lake Chabot Park. The wet deposition sampler was placed on the rooftop on the western side of the building, approximately 30m east from the curb of Lake Chabot Road, and approximately 5m off the ground surface. Castro Valley Elementary (CVE) is a school on San Miguel Avenue, in a residential neighborhood and several hundred meters from Castro Valley Boulevard. The sampler at CVE was also placed on the roof, approximately 20 meters west of San Miguel Ave, and 10m above the ground surface.

Two other sampling sites were added later in the study in order to capture a wider range of traffic conditions based on their proximity or distance from major roads. These sites had not been previously considered, as they were not suitable for deployment of the wet deposition samplers. However, it was decided that bulk and dry deposition samples should be collected at these locations. The Redwood Professional Building is located on Redwood Road, at the end of an exit ramp from eastbound Highway 580. The samplers were placed on a pole attached to the roof of the building, about 8m above the ground, and 30m east of Redwood Road. The end of the sampler pole was also about 30m south of the edge of Highway 580, and 5m below the road surface; the highway is elevated in that section of Castro Valley to pass over Redwood Road. The last sampler was placed near Madison Reservoir, a small enclosed reservoir operated by East Bay Municipal Utility District located in the upper watershed around 800m above sea level, along one of the ridges above Castro Valley. The reservoir is adjoined by two houses in the Columbia housing development. The sampler was attached to a fence post, 3m above the ground surface, and about 30m west of Columbia Road, the main route into and out of the housing development, along a ridge about 10m higher than the road at that location.

3.4 Wet and Bulk Deposition

At sampling sites where continuous AC power was available, automatic precipitation collectors (Figure 2) designed for the National Atmospheric Deposition Program/Mercury Deposition Network (NADP/MDN) could be deployed to collect precipitation samples. The device has been described in detail and evaluated by other investigators (Vermette et al. 1995a). Samplers (Aerochem Metrics, Inc., Bushnell, FL) were modified by the Illinois State Water Survey for the previous RMP study. Up to two sampling bottle and funnel assemblies could be deployed simultaneously in each machine. A precipitation sensor activated the lid to expose the funnels during a rain

event. When the rain ceased, the sensor dried off, closing the lid. Bulk deposition samples were collected using identical bottle and funnel assemblies, but the assemblies were loaded into a passive continuously open chimney rather than an automated sampler. Procedures for the preparation of the bottles and funnels used in collecting wet and bulk samples are described below.

High-density polyethylene one-liter bottles and funnels with connecting adaptors were used to collect samples. Bottles and funnels were cleaned to reduce metal contamination. Bottles and funnels were cleaned sequentially with detergent, nitric acid, and HCl, with multiple rinses of ultrapure water between each solution. After the last rinse and soak with ultrapure water, sample bottles were emptied, capped, and double bagged for shipment. Because the connectors used to attach the funnel and sample bottle did not have direct contact with the precipitation samples, they were subject to slightly less stringent cleaning procedures. Connectors were washed in detergent, soaked in HCl, rinsed with distilled water, dried, and bagged for shipment.

Materials for wet and bulk sample collection were handled in the field using methods adapted from EPA 1669. The empty sample bottle was uncapped at the monitoring site, and a connector and funnel were attached. The assembly was checked for tightness and placed in the precipitation collector or passive sampler chimney. Site and sample conditions were noted on FOFs. At the end of the collection period, the funnel and connector were removed, and the sample bottle recapped and bagged for shipment to the laboratory.

Precipitation samples were weighed and acidified with trace-metal grade nitric acid to an acid concentration of 0.2%. The samples were equilibrated for more than 48 hours before being analyzed. For samples containing less than 10 mL of precipitation, 20.0 mL of reagent water was added, and the sample was acidified to 0.2%.

Before analysis, concentrated HNO₃ was added to 10 mL of the sample until it contained 2% nitric acid. The sample was then digested at 85°C for 2 hours. The digested samples were then analyzed by ICP-MS. Blanks and calibration standards were measured periodically with samples in accordance with the EPA Methods and the project QAPP.

3.5 Dry Deposition Methods

Various techniques have been used in an effort to make reliable estimates of dry deposition flux of pollutants. There are two general approaches: 1) direct measurement of deposition fluxes using surrogate surfaces (direct method), and 2) calculated deposition fluxes using airborne concentration data and modeled deposition velocity (indirect method). With direct methods, geometry of the surface, roughness, and collection substrate all influence the wind profile and local turbulence above the sampling surface, which in part controls the deposition flux. Surrogate surfaces are generally designed to minimize turbulence in the flow field so that minimum fluxes are measured. In the indirect modeling approach, particle size distribution is a critical parameter influencing deposition velocity and the resulting deposition fluxes.

Large particles are responsible for a large proportion of the deposition flux, but they may account for only a small fraction of the total airborne mass (Holsen & Noll 1992, Holsen et al. 1993, Shahin et al. 2000) due to large increases in deposition velocity

with small increases in particle diameter over 2 μm . Shahin et al. (2000) asserted that large particles are not sampled efficiently by most samplers and are therefore underrepresented in most indirect flux estimates. This often results in modeling estimates often lower than deposition flux measurements made using surrogate surfaces. Due to these limitations described above, a direct method, employing surrogate surface plates, had been chosen for use in the RMP Atmospheric Deposition Pilot Study to collect dry deposition samples of trace metals.

To generate data most easily comparable to this previous monitoring effort in the SF Bay region, in this study of Castro Valley atmospheric deposition we also elected to use the same direct surrogate surface method. The sampler "Egret I" (Figure 3) used in this study and the previous RMP study was an adaptation of the prototype developed previously by Professor Holsen, (Holsen et al. 1991). The original sampling device, with greased Mylar films mounted on a knife-edge plate, has been described by Yi and associates (Yi et al. 1997a, Yi et al. 1997b), and evaluated by other investigators (Holsen et al. 1993, Pirrone et al. 1995, Paode et al. 1998). A modified sampler with a larger surface area developed and manufactured for the previous RMP study was also used here. The total exposed surface area was 500 cm^2 . Plastic materials were used to construct the frame and various parts the sampler. In addition, transparent plastics were used whenever possible to reduce thermal microclimates induced from blackbody radiation.

A cartridge plate bearing a greased Mylar film was pointed into the wind with a leading knife-edge to provide a laminar or non-turbulent flow of air over the surface of the film. The L-Apiezon grease applied to the Mylar film provided a sticky surface to capture particles, and prevent particle bounce and loss (Noll et al. 1990). L-Apezion grease is largely free of the elements being monitored.

At the analytical laboratory, Mylar film was cleaned sequentially with methanol, ~2% nitric acid, and deionized distilled water. After washing, the Mylar film was air-dried and mounted in a plastic cartridge plate. Approximately 150 mg of L-Apiezon grease was then applied to each film, using low heat to aid in spreading the grease. Each cartridge plate was placed into a plastic box that was in turn enclosed in two plastic bags by the laboratory. The sampling plates were then shipped to SFEI.

When installing or retrieving samples, ultra-clean field techniques adapted from EPA Method 1669 "Sampling Ambient Water for Trace Metals at EPA Water Quality Criteria Levels" (USEPA 1996) were practiced to minimize contamination from field operations. The Egret I sampler could hold up to two wind-vanes that pivot according to the wind direction (Figure 3). However, generally only one vane was mounted in the sampler at a time, with the exception of collection periods when field replicates were collected. A Mylar film on a surrogate surface cartridge plate prepared by the laboratory was removed from its shipping container and mounted and secured to the wind-vane with plastic clips.

Each greased Mylar film was exposed 48 hours for collection. Field site and sample conditions were recorded on a field observation form (FOF). Duplicate samples as well as trip blank samples were collected for a subset of sampling events (approximately 1 for each 10 field samples). Trip blanks were transported to field sites but were not removed from their containers while in the field.

After samples were retrieved and returned to the laboratory, the greased Mylar film was rinsed with 50ml hexane to recover the collected material. The Mylar film was rinsed a second (and often a third time) with hexane to ensure that all material was recovered, with the rinses collected for separate analysis. Sample rinses were dried under vacuum, then microwave digested individually in 10 mL of 10% ultra-pure nitric acid. Samples were subsequently analyzed using inductively coupled plasma-mass spectrometer (ICP-MS) following EPA Method 200.8. Urban particulate matter (UPM) standard reference material (SRM) from NIST was analyzed along with samples to evaluate method recovery and quantitation accuracy.

3.6 Gaseous Trace Organics

Benzene in ambient air was collected and analyzed using EPA Method TO-15, which employs an initially evacuated stainless-steel canister (at subatmospheric pressure). A sample of ambient air is drawn through a coarse filter to exclude particles which would jam the canister valve or contaminate the canister interior. A mass flow controller regulates the rate and duration of sampling into the pre-evacuated canister. For this study ambient air samples were collected concurrently with dry deposition samples (48 hour integrated samples). After the air sample was collected, the canister valve was closed, an identification tag was attached to the canister, a chain-of-custody (COC) form completed, and the canister was shipped to the laboratory for analysis.

Each sample canister after being initially evacuated at the analytical laboratory was individually proofed to ensure that the valve operated properly and that no contaminating gases remain in the canister. A mass flow controller (with attached vacuum gauge) was also calibrated to ensure that the proper flow rate was set for the planned deployment period. The field team attached the flow controller to the canister and checked for tightness of the entire assembly by checking for loss of vacuum while the sample assembly was still capped and presumably closed to the atmosphere. The initial vacuum of the canister was recorded on the FOF. If all the equipment functioned properly to that point, the valve was opened for sample collection and left at the site for the collection period. On completion of sampling, the field team noted the final vacuum of the canister and closed the canister valve. The flow controller was removed from the canister, and the equipment returned to the laboratory for analysis.

For analysis of volatile organics in ambient air samples, water vapor is reduced in the gas stream (as needed if applicable), and the VOCs are then concentrated by collection in a cryogenically-cooled trap. The cryogen is then removed and the temperature of the trap is raised. The VOCs originally collected in the trap are revolatilized, separated on a GC column, then detected by one or more detectors for identification and quantitation. In this study the analytical laboratory used gas chromatography with a mass spectrometer operated using selective ion monitoring mode (GC-MS-SIM)

3.7 Quality Assurance and Quality Control

Overall sample handling techniques used in the study closely followed the general clean technique principles as described in USEPA Method 1669 (USEPA 1996). Equipment and material that had direct contact with samples were rigorously cleaned (see above). Additionally, clean sample bottles and funnels were double bagged. Surrogate surface plates with greased Mylar films were placed in polyethylene boxes before being

enclosed in clean plastic bags. Powder-free polyethylene gloves were worn when handling samples.

3.7.1 Wet and Bulk Deposition

For wet and bulk deposition, the majority of samples were analyzed at Clarkson University in Professor Thomas Holsen's laboratory. For samples collected during the first two wet sampling events, a commercial laboratory (Caltest) analyzed wet deposition samples for copper only. A summary of QA/QC results for the parameters reported are presented in Table 3. The copper reporting limit of the commercial laboratory was 0.2 µg/L (ppb), which was comparable to the method detection limit (MDL) for copper reported by the Holsen laboratory of 0.23 ppb. MDLs for other analyzed elements, estimated as three times the standard deviations on replicates of laboratory blanks, are also shown in Table 3. Data presented in this report are not blank-corrected although laboratory method blanks are used in calibrating the instrument.

System blanks were periodically collected to assess any contamination resulting from field operations. A system blank was generated by rinsing a small volume (20ml) of ultra-pure water through the sample funnel and collecting it in the sample bottle to determine the amount of contamination of the sampling train, similar to the process used at the end of collecting a bulk deposition sample. Table 3 also presents concentrations of analytes found in system blanks. Many of the trace elements were detected in the system blanks at concentrations above the MDL, particularly for elements typically abundant in both natural and anthropogenic sources (Mg, Al, Si, Fe, Zn). However, because the system blank volumes are small, the amounts per sample of even the most abundant contaminants (Fe and Mg) in system blanks averaged ~1 µg, well below the total amounts of these elements typically found in field samples, which typically have larger volumes.

3.7.2 Dry Deposition

Summary results of quality assurance and quality control (QA/QC) measurements for dry deposition samples are presented in Table 4. One blank result for copper was over 4 standard deviations from the mean of the remaining blank results and was discarded as an outlier. Method detection limits (MDL) were defined as three times the standard deviation of blanks and are shown in Table 4 (in units of µg per sample). Like wet deposition samples, results for dry deposition samples in this report were also not blank-corrected.

Recoveries and RSDs on replicate analyses of the UPM reference material on average met the DQOs established for the Pilot Study (Table 4). Recoveries on trace elements on average deviated from the certified value by 10% or less, with the exception of Ni and Al. The average recovery error for Ni was 18%, and the average for Al was 26%, slightly over the target DQO. Precision on replicate analyses of the reference material was generally quite good, ranging 1-22% RSD.

3.7.3 Ambient Gas Collection

Benzene sampling QA/QC consisted of blank proofs on the collection canisters, and surrogates spiked to the canister prior to analysis. All canisters were proofed by the laboratory individually prior to shipment for field sampling. Benzene was not detected in any canisters. On arrival, canisters were tested for tightness (< -29 in Hg vacuum) with a

vacuum gauge supplied by the laboratory. After deployment, canisters were checked for remaining vacuum. Some canisters that reached ambient pressure (due to small leaks in connectors, misadjusted flow controllers, or any other reasons) at the end of sampling were not analyzed, as the time of sampling completion could not be known and some loss of gases from the canister could have occurred. At the lab prior to analysis, canisters were spiked with labeled recovery surrogates. Recoveries of all surrogates were within control limits of the lab.

4.0 RESULTS AND DISCUSSION

This section presents results of the monitoring and loading estimates of the selected trace metals from the atmosphere to sampling sites in the Castro Valley Creek Watershed. These estimates are further compared with estimates of trace metal loading from atmospheric deposition previously found in the SF Bay areas in the RMP Atmospheric Deposition Pilot Study, and results from studies in other regions.

4.1 Wet Deposition

A total of 49 precipitation samples were collected and analyzed from the two to four stations monitored over the course of the study, including field replicates and bulk/wet splits (Table 1). Sample volumes ranged from approximately 20 mL to nearly 800 mL for wet and bulk samples collected. Smaller sample funnels were deployed starting in 2005 to reduce the threat of overfilling bottles during a series of severe winter storms, so collected volumes of wet and bulk samples are proportional only within each of the two sampling periods (before and after December 29, 2004). Furthermore, all bulk samples contain approximately 20mL of added water used to rinse the collection funnel, so collected volumes for bulk samples do not represent only precipitation. A number of samples were not successfully collected over the course of the study, indicated by “MF” (malfunction) in Table 1. Malfunctions in sample collection have included failure of the automated sampler lid to open due to jamming or sensor failure, funnels becoming disconnected from bottles, and unauthorized removal of a bulk sampler (by an unknown party) in one case.

Concentrations of the measured trace elements in wet and bulk samples varied widely among sampling events at each site and among the four sites. Average concentrations measured by the laboratory in each collected sample are presented in the appendix in Table A-1. Average concentrations of trace elements measured in samples for the different sites are summarized in Table 5. For bulk samples in particular, metal concentrations in the collected water would be expected to vary widely, as some bulk samples would represent 14-day periods that were entirely or nearly dry, with all dry deposition in the collection funnel to that point being washed down and collected in a small volume of water rinsing the funnel at the end of the collection period. In contrast, collections from extended rainy periods would have a large volume of water effectively diluting any dry material that had deposited in the funnel.

One sample from Madison (October 20-Nov3) had an unusually high concentration of copper (~22 ppb) despite occurring during a period with a moderate amount of rain. On reanalysis by the laboratory, the high measured concentration was reconfirmed. However, for nearly all of the samples collected, a number of the trace

elements correlated extremely well; for example copper concentrations correlate extremely well with zinc, but the one sample from Madison in this period did not, with a ratio of Cu/Zn well outside the usual range (Figure 4). This value was therefore flagged as an outlier and was not included in calculations of flux rates or in statistical analyses.

Fluxes for wet and bulk deposition rates ($\mu\text{g}\cdot\text{m}^{-2}\cdot\text{d}^{-1}$) at the various sampling sites were calculated by using equation (1) shown below:

$$F = C*V/A/EP \quad (1)$$

Where

- F = Deposition flux rate
- C = Concentration of trace elements in the collected sample
- V = Volume of sample collected, including water for funnel rinse
- A= Cross sectional surface area of the collection funnel
- EP = The exposure period

The surface area of collection funnels was calculated from their diameters at their widest points, and the other parameters were measured or recorded for each sample collected. Deposition fluxes for all the wet and bulk deposition samples are determined using this calculation, and tabulated in the appendix in Table A-2. Although for a number of measured elements, a large number of concentration results measured in wet and bulk samples were below the MDL (defined as 3*stdev of replicate blank analyses determined at the laboratory), it was decided to use the concentrations reported at less than MDL to calculate fluxes, as measures commonly used to assign values to non-detect records (e.g. assigning 0, half the MDL, or the MDL) can potentially introduce biases to the data set.

Similar to concentrations, flux rates in bulk and wet samples also varied widely among sites and between collection events within any particular site. Despite the variability, when the data are aggregated for each site and type of collection, some patterns emerge. Table 6 shows averages and standard deviations of calculated flux rates for all wet and bulk collection events during the study.

As shown in Table 6, for many of the trace elements, flux rates at Redwood averaged higher than at other sites. Mean copper deposition rates for Redwood were significantly higher ($p < 0.05$) than at the other sites using both parametric (Student's T-test, Tukey HSD) and nonparametric (Kruskal Wallis) statistical tests. Other elements for which deposition was significantly higher for Redwood than at other sites include Fe, Zn, Sr, Sb, and Ba. Parametric and nonparametric analyses generally agreed on significance of differences, although the Tukey HSD test indicated a significant difference for Pb, whereas the Kruskal-Wallis did not.

On three occasions, bulk samplers were deployed at one or both of the wet deposition locations in order to collect both sample types side by side. On two of those events, either one or both of the samplers malfunctioned, and a sample could not be collected. However, two bulk samples (one at CVE and one at CVCC) were collected on events where the precipitation sampler appeared to be functioning properly. For the CVE sample (collected 29-Dec-2004 to 12-Jan-2005), the mass of water collected differed only

by approximately the amount of water used for the collection rinse (651-633g = ~20mL). However, for the CVCC sample (collected during 9-Feb to 23-Feb-2005), the sample volumes collected differed by over 150mL, even though the automated sampler appeared to be functioning properly at the time the sample was collected. One possibility is that power to the roof outlet had been turned off for part of the collection period.

As was expected, wet deposition collected in these samples was lower than bulk deposition in the samples from the same period at the same sites. Despite the collected volumes being virtually identical, nearly all deposition rates for the bulk collection bottle at CVE 29-Dec to 12-Jan were higher than for the corresponding wet deposition sample collected. Similarly, the pair of samples from CVCC 9-Feb to 23-Feb had greater deposition rates for nearly all trace metals in the bulk sample, although the larger difference in collected volumes may have also contributed to the differences in observed deposition.

All other bulk and wet deposition samples were collected on different events at the various sites, and biases may occur because the bulk of bulk samples were collected from wet deposition sites during periods where rain was expected to be sporadic. However, the patterns seen are consistent with the expectation that for any given site and event; bulk samples would contain at the least all the elements in precipitation collected in wet deposition samples, plus any additional deposition occurring during periods when it was not raining.

4.2 Dry Deposition

A total of 47 dry deposition samples were collected at up to the four monitoring stations over the course of the study. For some of the sampling events, duplicate samples were collected from one of the stations. The exposure duration was 48 hours for all collection events.

Similar to the wet and bulk deposition monitoring, the dry deposition rate ($\mu\text{g}\cdot\text{m}^{-2}\cdot\text{day}^{-1}$) at any given sampling site can be calculated using the same general formula:

$$F = C \cdot V / A / EP \quad (1)$$

Where here

- F = Deposition flux rate
- C = Concentration of trace element in the analyzed extract
- V = Volume of final extract
- A = Exposed area of the surrogate surface collector
- EP = The exposure period

Dry deposition fluxes for individual samples, including field replicates, are tabulated in appendix Table A-3. Dry deposition fluxes varied among sampling events at all the sampling locations and average dry deposition rates are summarized for the various sampling sites in Table 7.

Similar to trends seen in the wet and bulk deposition data, for nearly all elements, deposition at Redwood was significantly higher (for Ti, V, Mn, Fe, Co, Cu, Zn, Sr, Ba, Pb, using both Tukey HSD and Kruskal Wallis tests) than at least one other site.

Generally Redwood was significantly different from at least Madison, but for some elements Redwood was higher than all other sites. Similarly, dry deposition at Madison was nearly always the lowest, with significant differences from one or more other sites. CVCC and CVE generally fell in between, significantly different from Redwood or Madison for some elements but not others.

Unlike results found previously in the SF Bay area (Tsai et al 2001), deposition fluxes were not significantly lower during the wet season, although a contributing factor may have been the small sample number for any given site (a maximum of five wet season sampling events, and eight dry season events). Similarly, comparisons of weekend versus weekday samples taken on three occasions showed no significant differences.

Average dry deposition flux rates (converted to an annual basis) were higher than reported in the previous RMP atmospheric deposition study (Table 8). Dry deposition fluxes in Castro Valley fell within the middle and upper range of those observed in urban areas and around other lakes and bays in the USA (Holsen et al. 1993, Wu et al. 1994, Golomb et al. 1997, Paode et al. 1998, Sweet et al. 1998, Zufall et al. 1998, Shahin et al. 2000). Results are most comparable to other studies using similar methods, as direct methods such as surrogate surface plates may yield measurements much higher than indirect methods with dichotomous air samplers. This discrepancy due to different sampling methods is apparent in Table 8, as indirect methods generally reported lower deposition rates.

Concentrations of copper detected in precipitation samples collected from Castro Valley were similar to those in a previous study of the San Francisco Estuary and those found in the Great Lakes area by Sweet et al. (1998) (Table 9). In Table 9, both wet and bulk results for this study are aggregated to derive averages. Deposition fluxes of trace metals to Castro Valley estimated in this study were also similar but higher than those in the RMP pilot study and within the range of the fluxes reported for the Great Lakes (Sweet et al. 1998), Massachusetts Bay (Golomb et al. 1997), and Chesapeake Bay (Scudlark et al. 1994).

Dry deposition fluxes around Lake Michigan from December 1993 to October 1995, using an automated dry deposition sampler with knife-edge surrogate surfaces, were approximately 10, 7, 0.2, and 1.0 $\mu\text{g}\cdot\text{m}^{-2}\cdot\text{day}^{-1}$ (corresponding to 3,650, 2,555, 73, and 365 $\mu\text{g}\cdot\text{m}^{-2}\cdot\text{yr}^{-1}$) for Cu, Ni, Cd, and Cr, respectively (Shahin et al. 2000). Indirect estimates made using dichotomous sampler during 1993 and 1994 and an estimated overall deposition velocity around Lake Michigan as one of the IADN monitoring stations yielded lower dry deposition fluxes of 1,300, 320, 380, 130 $\mu\text{g}\cdot\text{m}^{-2}\cdot\text{yr}^{-1}$ for copper, nickel, cadmium, and chromium respectively (Sweet et al. 1998). Golomb et al. (1997) used both direct and indirect methods to measure the dry deposition of trace metals to the Massachusetts Bay, and found higher deposition flux of chromium by over a factor of two for the direct method versus an indirect method.

Although both this study and the previous RMP study in SF Bay both used the same sampler design for direct measurements of dry deposition, the much higher Cu deposition fluxes measured in Castro Valley (e.g. $\sim 20 \mu\text{g}\cdot\text{m}^{-2}\cdot\text{d}^{-1}$ or $\sim 7000 \mu\text{g}\cdot\text{m}^{-2}\cdot\text{yr}^{-1}$) compared to bay sites are not surprising. That study located monitoring sites as near to

the bay as possible while avoiding sources such as roads as much as practicable, whereas this study aimed to capture the maximum impact of traffic and urban activities.

4.3 Benzene Monitoring

Results of Benzene monitoring are presented in Table 10. Ambient benzene concentrations ranged 0.10 to 0.48 ppbv. For all sampling events, benzene concentrations at Redwood were higher than those at Madison. One pair of samples taken concurrently at Redwood (Nov 19-21) averaged 0.34 ± 0.05 ppbv, whereas a sample from Madison taken at the same time (a weekend period) measured 0.14 ppbv. A sample taken at Madison on the prior weekdays (Nov 17-19) measured 0.25 ppbv, suggesting a weekend versus weekday difference in ambient benzene. However, considering the variability of field replicate samples taken, and assuming that variability at Madison would be similar (± 0.05 ppbv), the weekend/weekday difference would not be statistically significant ($p > 0.05$) in large part due to the small sample number. When all the sampling events are considered together, the difference between Redwood and Madison is significant. Ambient benzene concentrations are moderately correlated to copper deposition rates ($R^2 = 0.49$, Figure 7).

4.4 Sources of Copper and Other Elements

Examining the ratio of Cu to Fe or other elements found in environmental samples is one method of bounding or scaling of the origin of their sources. In this study, the Fe/Cu ratio was approximately 100 in the dry deposition samples. This is in contrast to a ratio of 10-20 in the precipitation samples, and 30 in bulk deposition samples. All these are in a range similar to a ratio of around 60 for the NIST UPM (used in this study as an SRM for analyzing samples). Because the ratio of Fe/Cu in soils is approximately 1400 (Bradford, 1996), differing ratios of copper enhancement in samples may reflect smaller and larger contributions of iron from soil sources diluting copper from brake pads and other urban pollution sources.

If we assume that all the iron in an average dry deposition sample (Fe/Cu of 100) originates from soil sources with (Fe/Cu approximately 1000), then at most (i.e., assuming all other copper in the sample originated from a pure copper source), we can expect that at most 10% of the copper in the sample originated from the soil. Taking a ratio of the sample Fe/Cu to a potential source Fe/Cu (e.g. 100/1000 in this case) rapidly indicates the maximum contribution of the potential source to the Cu in a sample, up to the point where the sample and source ratios are identical (i.e. in that case, all Cu in a sample could then have come from the source, as the ratios are identical).

In the RMP Atmospheric Deposition Pilot Study, the Cu/Ni ratio was explored as a possible indicator of differing pollutant sources or transport mechanisms. The average in this study is similar to that reported by CARB in ambient air monitoring, with average Cu/Ni ratio of about 7:1. The Cu/Ni ratio found in CARB's ambient air samples varied with locations, ranging from 4:1 in San Jose to 11:1 in Concord. However, because the detection limit in this study is higher than in the RMP study, determining ratios with more abundant elements (such as Fe) is preferable, as Cu/Ni ratios would vary widely, especially for measurements of Ni near its detection limit.

4.5 Comparison to Other Loading Pathways

In the previous RMP atmospheric deposition study, one objective was to quantify the contribution of atmospheric deposition relative to other sources and pathways of pollutants (Table 11). Loading estimates via atmospheric deposition in Table 11 were based on measurements of deposition flux at the sites in that previous study. Although many of the assumptions used in that exercise were quite crude, it served as a screening level evaluation, highlighting first-order estimates for various sources and pathways, and revealing the large uncertainties associated with many of these estimates.

One goal of this BPP project is to combine monitoring and modeling to develop our best integrated understanding of pollutants (i.e. particularly copper Cu) from their emissions, to transport, and fate in the environment. Other members of the project team will be engaged in measuring and modeling those components of the study of the environmental fate of brake pad wear material. Therefore, for this portion of the study, refining the previous calculated watershed loadings of pollutants and their transport to the Bay is not an objective, as those questions will be better addressed by the detailed modeling components of the study.

However, we will note that bulk deposition rates measured in Castro Valley for this study for copper are at least 50% higher than the combined wet and dry copper flux estimates found previously for San Francisco Estuary sites in the RMP pilot study. Although those measurements are more appropriate for determining direct deposition to the water surface given the closer proximity of those monitoring sites to the bay, for estimates of pollutant deposition and loading rates in the watersheds, this study likely provides more appropriate data. Furthermore, if the average dry deposition rates found for these sites in Castro Valley are presumed to be representative of those in other watersheds around the Bay (about 4 times higher than those in the RMP study), then indirect atmospheric deposition could account for up to a quarter of the stormwater inputs of copper estimated in Table 11.

One open question is whether surfaces in the watershed behave more like the surrogate surface plates, which prevent resuspension once particles are deposited, or more like the bulk collectors, on which materials can be continually deposited and resuspended to a degree. Watersheds are likely to have both types of surfaces, but the surfaces that are most likely to prevent wind resuspension such as vegetated soils are also most likely to retard surface runoff of deposited particles, whereas smooth impervious surfaces that will allow rapid surface runoff of particles are also likely to yield particles to wind resuspension.

5.0 CONCLUSIONS

Various trace elements were measured in collected precipitation and from surrogate surface plates deployed at various locations in the Castro Valley Creek watershed. Copper, and other pollutants of interest were measured in samples, and similar to results from a previous study in San Francisco Bay, concentrations found in samples varied widely among sites and among collection events for any site. Average wet, bulk, and dry deposition rates fell within the range seen for other urban areas in the United States, with average deposition rates exceeding those measured previously for sites located nearer the bay.

Wet or bulk deposition rates were significantly different among the sites for some trace elements, with the site at Redwood Road under Highway 580 receiving the most deposition. However, for many of the trace elements, the Redwood site was higher but not significantly above the other sites in the lower watershed, Castro Valley Community Center and CVE. The site at Madison Reservoir, high in the watershed, generally received the least deposition, as vehicle traffic and other activities are generally restricted to residents of the housing development located there.

Dry deposition fluxes were also found to be significantly different among the various sites, again with the Redwood Road site generally showing the highest flux, Madison showing the lowest as expected, and the other two sites falling in between. Dry deposition rates measured for the surrogate plates were much higher than previously found for sites in SF Bay. Sampling site conditions span the range of those that may be found in many Bay area communities, from heavily traveled highway corridors and commercial districts, to quiet residential neighborhoods. Deposition rates (whether bulk, wet, or dry) measured here likely better represents the typical conditions seen in regional watersheds for the modeling and estimating atmospheric pollutant loads.

A bulk deposition sampler was collocated with one of the automated precipitation samplers on two of the sampling events, as expected measured additional deposition not captured the wet sampler. On one occasion, the differences in collected precipitation were negligible, so it is likely that the added deposition was from dry material deposited in the collector during dry periods during the sampling. In the other sampling event, the bulk collector captured a greater volume of fluid, suggesting shielding of the collection funnel during part of the collection period (e.g. moisture sensor not being sufficiently sensitive for some small rain events, power interruption, temporary jamming).

However, even the highest deposition rates measured for bulk samplers remain below those seen for the greased surrogate surface plates (dry deposition collectors). Unlike the grease on the surrogate surface plate, which prevents resuspension and loss of deposited material, bulk collector surfaces do not bind the deposited particles and thus allow resuspension losses and lower measured net flux. Despite the complications of modeling and interpreting measured results for bulk samplers versus more idealized collectors such as surrogate plates and wet-only precipitation collectors, their simplicity, small size, and low expense allow monitoring at sites that would be otherwise difficult to utilize.

For a limited number of dry deposition collection events near the end of the study, canister collectors were deployed to capture ambient gases for measurement of benzene. As expected, the ambient concentrations at the Redwood Road site were higher than for Madison Reservoir. However, one sample taken at CVE indicates concentrations similar to those at Redwood, within the analytical variability for a pair of samples collected simultaneously at Redwood. The benzene data further confirms our expectations from observations that traffic volume and thus benzene and other automotive pollutants are likely to be higher at Redwood than at the upper watershed site at Madison Reservoir.

The data collected in this study show that deposition rates of trace element pollutants such as copper, zinc, and other metals associated with vehicle traffic and other urban activities would be found higher than at sites in a previous SF Bay region study

that intentionally located monitoring sites away from such sources. The data suggest average copper deposition rates four to five times those previously measured, although results are likely skewed by location of one of the sites virtually below one of the region's busiest highways. However, data from less impacted but still highly urbanized sites within Castro Valley indicate deposition rates still well above those in the previous regional study. The range of sites monitored in Castro Valley greatly improves our knowledge of local atmospheric deposition for the modeling of pollutant transport within this watershed, as well as providing more appropriate data for extending modeling efforts and atmospheric pollutant load estimates to other urban watersheds surrounding San Francisco Bay.

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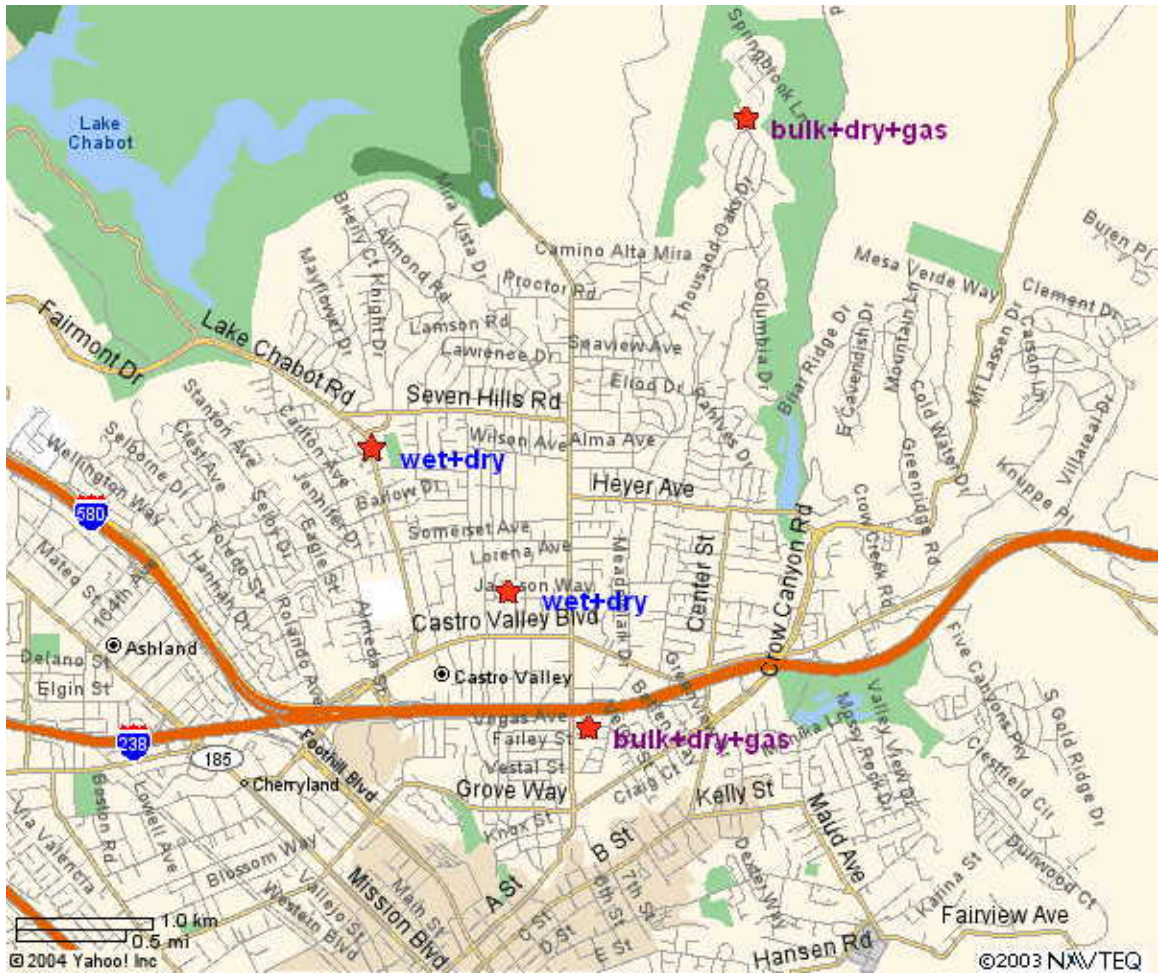


Figure 1. Sampling Sites in Castro Valley
(left to right) Castro Valley Community Center, CV Elementary, Redwood Professional Building, Madison Reservoir



Figure 2. Wet Deposition Sampling Device



Figure 3. Dry Deposition Sampling Device (Egret I)

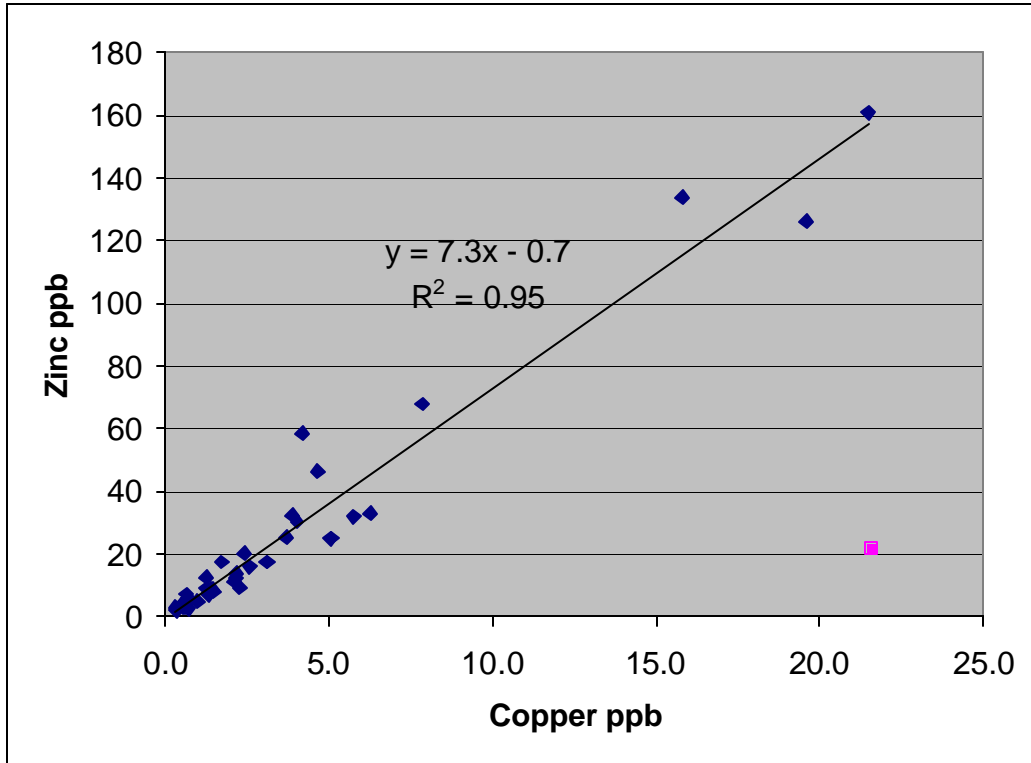


Figure 4. Copper and Zinc Correlation in Wet & Bulk Samples
One likely outlier from Madison (Oct20-Nov3) in contrasting color

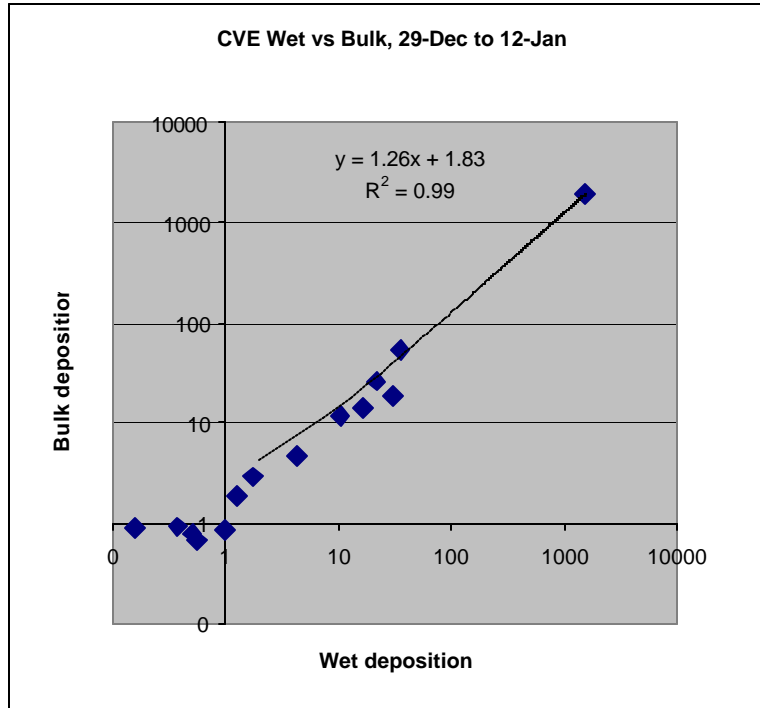


Figure 5. Collocated Wet versus Bulk Samples CV Elementary
 Wet sample deposition rates were lower for most trace elements despite similar volumes

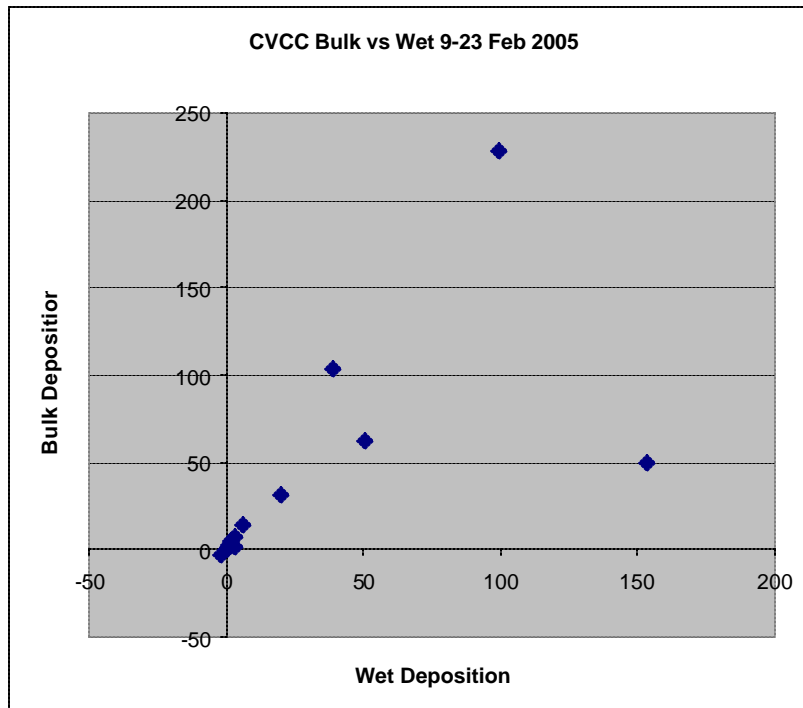


Figure 6. Collocated Wet vs Bulk Samples CV Community Center
 Like in previous figure (CVE), most elements in bulk sample > wet

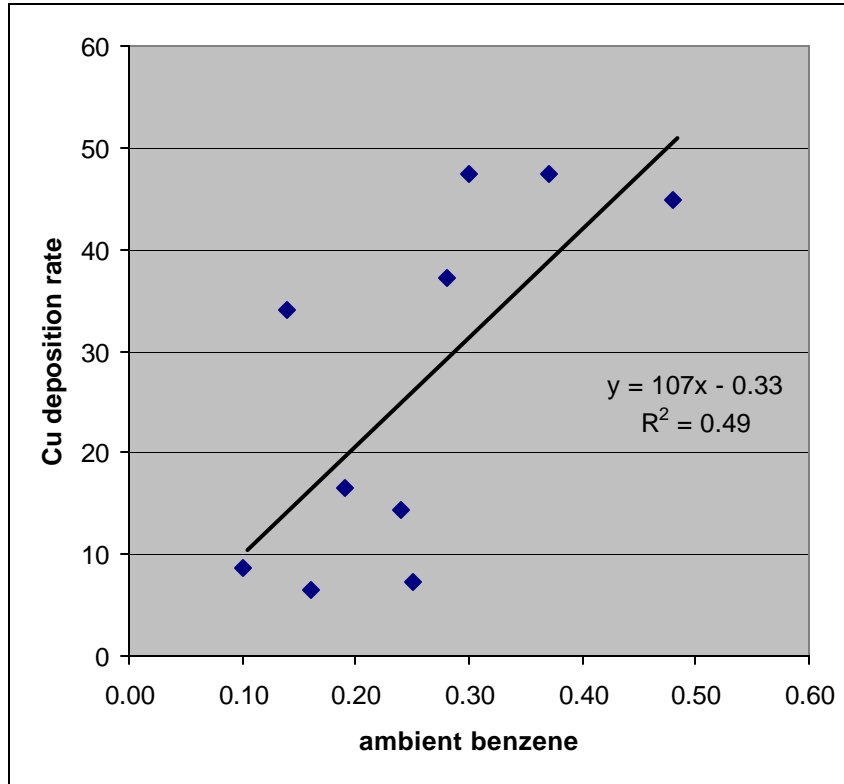


Figure 7. Relationship of Cu Dry Deposition Rate to Ambient Benzene Measurements

Table 1. Wet and Bulk Samples Collected

Start	end	wet	CVCC	CVE	Madison	Redwood	bulk	CVCC	CVE	Madison	Redwood
3/17/2004	3/31/2004		1	1							
3/31/2004	4/21/2004		1	1							
4/21/2004	5/26/2004		NR	NR							
5/26/2004	6/9/2004		MF	1							
6/9/2004	7/7/2004							1	1	1	
10/6/2004	10/20/2004							1	1	1	1
10/20/2004	11/3/2004							1	1	1	1
11/3/2004	11/17/2004							1	1	1	1
11/17/2004	12/3/2004		1	1						MF	1
12/3/2004	12/17/2004		MF	MF				1	1	2	1
12/17/2004	12/29/2004		1	1						1	1
12/29/2004	1/12/2005		MF	1				1	1	2	1
1/12/2005	1/26/2005										
1/26/2005	2/9/2005		1	1						1	1
2/9/2005	2/23/2005		1	2				1		2	1
	total		6	9				7	6	12	9

MF = malfunction of sampler or sampling train

NR = no rain on wet collection

Table 2. Dry and Gaseous (Benzene) Samples Collected

Start	end	dry	CVCC	CVE	Madison	Redwood	gas	CVCC	CVE	Madison	Redwood
7/7/2004	7/9/2004		1		1	1					
7/28/2004	7/30/2004		1		1	1					
8/18/2004	8/20/2004		1		1	1					
8/20/2004	8/22/2004		1		1	1					
8/31/2004	9/2/2004		2		1	1					
9/15/2004	9/17/2004		2		1	1					
9/29/2004	10/1/2004		2		1	1					
10/1/2004	10/3/2004		2		1	1					
11/17/2004	11/19/2004		1	1	1	1				1	
11/19/2004	11/21/2004		1	1	MF	1				1	2
12/15/2004	12/17/2004		1	1	1	1				1	1
1/19/2005	1/21/2005		1	1	1	1				1	MF
2/9/2005	2/11/2005		1	1	1	1			1	1	1
	total		17	5	12	13			1	5	4

MF = malfunction of sampler or sampling train

Table 3. QA/QC for Wet and Bulk Deposition Samples (in ppb)

	Method blank stdev	MDL	System blank average	Average RSD
24Mg	0.083	0.25	57	17%
27Al	0.1	0.3	19	19%
28Si	1.454	4.36	7.7	22%
47Ti	0.209	0.63	0.22	32%
51V	0.26	0.78	0.077	9%
52Cr	0.067	0.20	0.35	16%
55Mn	0.016	0.048	0.85	9%
56Fe	0.15	0.45	54.7	17%
59Co	0.006	0.018	-0.014	26%
60Ni	0.287	0.86	-0.21	13%
65Cu	0.076	0.23	0.77	14%
66Zn	0.095	0.29	8.19	13%
88Sr	0.006	0.018	0.07	8%
111Cd	0.006	0.018	0.021	12%
121Sb	0.002	0.006	-0.054	20%
137Ba	0.022	0.066	0.21	9%
208Pb	0.005	0.015	0.33	29%

MDL - Method detection limit

RSD – Relative Standard Deviation of lab replicate analyses, for FS averaging at least 3xMDL

Table 4. Quality Assurance and Quality Control Parameters for Dry Deposition Samples (in ug/sample)

	Method & trip blank avg	Method & trip blank stdev	MDL	RSD on SRM	SRM %ERR avg
24Mg	0.67	0.48	1.43		
27Al	1.53	1.34	4.03	4%	26
28Si	4.38	3.15	9.46		
47Ti	0.08	0.08	0.24		
51V	0.01	0.03	0.09		
52Cr	0.41	0.28	0.85		
55Mn	0.13	0.32	0.95		
56Fe	1.65	1.37	4.11	4%	10
59Co	0.01	0.03	0.09		
60Ni	-0.02	0.13	0.38	22%	19
65Cu	0.18	0.61	1.84	6%	6
66Zn	0.88	0.81	2.43	1%	8
88Sr	0.02	0.04	0.13		
111Cd	0.02	0.04	0.11	2%	8
121Sb	0.004	0.03	0.08	2%	9
137Ba	0.07	0.12	0.36		
208Pb	0.09	0.14	0.41	2%	7

MDL – Method detection limit

RSD – Relative standard deviation

SRM – Standard reference material NIST UPM (1648)

%ERR – % difference from certified value

Table 5. Average Wet and Bulk Sample Concentrations (ppb) in Castro Valley

	CVCC bulk		CVCC wet		CVE bulk		CVE wet		Madison bulk		Redwood bulk	
	av	sd	av	sd	av	sd	av	sd	av	sd	av	sd
24Mg	326	± 471	91	± 110	361	± 441	186	± 173	155	± 165	175	± 148
27Al	189	± 346	31	± 15	183	± 240	105	± 165	43	± 78	101	± 99
28Si	197	± 376	17	± 13	208	± 254	58	± 121	46	± 101	112	± 97
47Ti	1.7	± 1.5	0.4	± 0.3	2.1	± 1.3	2.9	± 5.2	1.1	± 1.8	2.0	± 1.2
51V	0.7	± 1.1	0.1	± 0.1	0.7	± 0.8	0.3	± 0.3	0.3	± 0.5	0.4	± 0.3
52Cr	0.5	± 0.6	0.2	± 0.1	0.5	± 0.4	0.6	± 0.9	0.2	± 0.2	0.4	± 0.3
55Mn	11.4	± 19.3	1.2	± 0.7	9.9	± 12.9	2.3	± 2.3	4.9	± 11.9	6.6	± 5.9
56Fe	195	± 310	23	± 15	194	± 217	19	± 13	41	± 58	186	± 186
59Co	0.2	± 0.4	0.0	± 0.0	0.2	± 0.3	0.1	± 0.1	0.1	± 0.2	0.1	± 0.1
60Ni	0.9	± 1.4	0.2	± 0.6	1.0	± 0.8	0.3	± 0.9	0.3	± 0.8	0.6	± 0.8
65Cu	5.6	± 8.2	1.8	± 1.3	5.9	± 7.0	1.9	± 1.1	3.0	± 5.9	4.9	± 4.2
66Zn	43	± 76	10	± 8	36	± 45	13	± 9	23	± 44	37	± 37
88Sr	4.9	± 7.6	0.9	± 0.7	4.9	± 5.8	2.3	± 1.5	2.2	± 4.2	6.1	± 6.9
111Cd	0.1	± 0.1	0.0	± 0.0	0.1	± 0.1	0.0	± 0.1	0.0	± 0.1	0.0	± 0.1
121Sb	0.2	± 0.2	0.1	± 0.1	0.4	± 0.5	0.3	± 0.3	0.1	± 0.1	0.4	± 0.2
137Ba	9.7	± 15.8	1.9	± 0.9	9.2	± 11.2	2.6	± 1.8	3.3	± 5.6	11.1	± 9.5
208Pb	3.5	± 6.0	0.6	± 0.4	3.1	± 4.4	0.7	± 0.4	1.2	± 2.4	1.9	± 1.8

Table 6. Average Wet and Bulk Deposition Rates ($\mu\text{g}\cdot\text{m}^{-2}\cdot\text{d}^{-1}$) of Castro Valley Sites

	CVCC bulk avg sd		CVCC wet avg sd		CVE bulk avg Sd		CVE wet avg sd		Madison bulk avg sd		Redwood bulk avg sd	
24Mg	603	± 104	545	± 100	671	± 284	658	± 484	489	± 642	425	± 406
27Al	186	± 50	143	± 51	223	± 53	185	± 31	59	± 66	174	± 107
28Si	175	± 24	108	± 16	275	± 33	267	± 19	58	± 87	204	± 140
47Ti	3.4	± 0.7	3.1	± 0.2	4.4	± 0.8	3.5	± 0.5	1.8	± 1.7	4.0	± 2.4
51V	0.9	± 0.2	0.6	± 0.1	1.1	± 0.2	0.6	± 0.1	0.5	± 0.3	0.7	± 0.4
52Cr	0.9	± 0.4	0.3	± 0.4	0.9	± 0.4	0.5	± 0.2	0.6	± 0.3	0.9	± 0.4
55Mn	14.4	± 1.6	14.1	± 0.4	13.8	± 1.9	17.3	± 0.9	5.4	± 8.8	12.4	± 8.6
56Fe	236	± 31	165	± 9	271	± 31	224	± 25	77	± 81	321	± 158
59Co	0.2	± 0.0	0.3	± 0.1	0.2	± 0.0	0.2	± 0.1	0.0	± 0.2	0.2	± 0.2
60Ni	1.1	± 0.4	2.1	± 1.1	1.8	± 0.7	1.1	± 2.1	-0.1	± 2.4	1.0	± 2.5
65Cu	8.5	± 2.1	4.2	± 1.4	7.9	± 2.5	4.8	± 2.3	3.6	± 1.9	9.0	± 3.0
66Zn	50	± 13	22	± 4	46	± 19	31	± 15	25	± 11	64	± 22
88Sr	7.4	± 1.1	5.1	± 0.5	7.9	± 3.1	6.1	± 3.1	3.9	± 3.9	12.2	± 9.5
111Cd	0.1	± 0.0	0.1	± 0.0	0.1	± 0.0	0.1	± 0.1	0.1	± 0.1	0.1	± 0.1
121Sb	0.5	± 0.1	0.3	± 0.1	0.6	± 0.3	0.3	± 0.4	0.2	± 0.3	0.7	± 0.2
137Ba	13.7	± 2.6	6.5	± 1.7	13.4	± 5.6	9.0	± 5.6	6.4	± 5.7	21.1	± 6.1
208Pb	4.1	± 1.0	3.5	± 1.1	3.3	± 1.5	2.6	± 2.2	1.5	± 0.8	3.5	± 1.4

Table 7. Average Dry Deposition Rates ($\mu\text{g}\cdot\text{m}^{-2}\cdot\text{d}^{-1}$) to Castro Valley Sites

	CVCC dry avg sd		CVE dry avg sd		Madison dry avg sd		Redwood dry avg Sd	
24Mg	593	± 322	397	± 433	212	± 161	1275	± 1405
27Al	891	± 459	788	± 741	307	± 232	2636	± 3964
28Si	123	± 56	113	± 14	91	± 34	138	± 44
47Ti	41.4	± 17.5	38.6	± 31.0	18.0	± 8.1	93.5	± 70.5
51V	3.0	± 1.4	2.7	± 2.3	1.3	± 0.6	6.2	± 4.8
52Cr	8.7	± 2.9	11.8	± 3.3	7.2	± 3.1	12.5	± 5.5
55Mn	30.7	± 13.7	33.3	± 26.1	15.9	± 7.1	63.3	± 42.1
56Fe	1568	± 834	1181	± 1054	579	± 437	4541	± 4160
59Co	0.8	± 0.3	1.1	± 0.7	0.3	± 0.2	1.4	± 1.0
60Ni	4.0	± 2.0	8.2	± 8.6	1.5	± 1.8	6.1	± 3.5
65Cu	18.5	± 5.4	16.3	± 7.6	11.8	± 9.6	30.8	± 10.1
66Zn	81	± 24	79	± 22	34	± 12	174	± 60
88Sr	8.0	± 3.1	7.2	± 3.8	3.2	± 1.6	25.9	± 13.4
111Cd	0.2	± 0.3	0.6	± 0.2	0.4	± 0.5	0.3	± 0.3
121Sb	0.5	± 0.3	0.7	± 0.2	0.3	± 0.2	0.8	± 0.3
137Ba	21.0	± 7.4	18.9	± 9.9	9.2	± 3.3	52.1	± 21.5
208Pb	10.2	± 4.3	8.4	± 5.3	4.8	± 2.3	15.0	± 5.9

Table 8. Estimated Dry Deposition Flux of Trace Metals Reported in the Literature

Sampling Site	Sampling Method	Cu	Nil	Cd	Cr	Citation
Urban Chicago ²	D	21900	NA	NA	NA	Paode et al. 1998
Lake Michigan ²	D	3650	NA	NA	NA	Paode et al. 1998
South Haven, MI ²	D	2555	NA	NA	NA	Paode et al. 1998
Lake Superior ³	I	2400	570	380	130	Sweet et al. 1998
Lake Michigan ³	I	1300	320	380	130	Sweet et al. 1998
Lake Erie ³	I	3300	460	400	1000	Sweet et al. 1998
So. Lake Michigan ⁴	D	2000	NA	NA	NA	Zufall et al. 1998
Massachusetts Bay ⁵	D	2000	930	130	1200	Golomb et al. 1997
Massachusetts Bay ⁶	I	NA	NA	NA	470	Golomb et al. 1997
Chesapeake Bay ⁷	I	290-810	330-910	11-32	110-300	Wu et al. 1994
Lake Michigan	D	3650	2550	75	360	Shahin et al. 2000
SF Estuary ⁸	D	1100?730	600?350	22?15	1300?900	Tsai et al. 2001
Castro Valley	D	7150±3900	1530±1460	120±140	3500±1600	This study

¹ All data are presented in $\mu\text{g}\cdot\text{m}^{-2}\cdot\text{yr}^{-1}$.

² Average of three weekly composite samples taken in May 1994, July 1994, and January 1995.

³ Average of monthly composite samples taken between summer 1993 and summer 1994.

⁴ Average of four day measurements with two samples/day in July 1994.

⁵ Average of bi-weekly composite samples collected at two sites from September 1992 to September 1993.

⁶ Average of alternative bi-weekly composite samples collected at one site from September 1992 to September 1993.

⁷ Range of low and high estimates from weekly composite samples collected between June 1990 and July 1991.

⁸ Values represent mean \pm standard deviation from all bi-weekly two 24-hr samples collected at three sites from August 1999 to August 2000.

D – Direct method collecting particles deposited directly on surrogate surface plates.

I – Indirect method collecting ambient air concentration and using modeling estimate.

NA – Not analyzed.

Table 9. Comparison of Trace Metal Concentrations and Deposition Fluxes Reported in the Literature to Results from this Study.

	Cu	Ni	Cd	Cr	Citation
Concentration in Precipitation ($\mu\text{g/L}$)					
Lake Superior	0.9 \pm 0.5	0.3 \pm 0.3	0.1 \pm 0.1	0.1 \pm 0.1	Sweet et al. 1998
Lake Michigan	0.8 \pm 0.1	0.4 \pm 0.4	0.1 \pm 0.3	0.1 \pm 0.2	Sweet et al. 1998
Lake Erie	0.9 \pm 0.1	0.3 \pm 0.2	0.1 \pm 0.1	<0.1	Sweet et al. 1998
San Francisco Estuary*	1.2	0.4	0.1	0.2	Tsai et al. 2001
Castro Valley	3.5	0.51 (<MDL)	0.043	0.41	This study
Wet Deposition Flux ($\mu\text{g}\cdot\text{m}^{-2}\cdot\text{d}^{-1}$)					
Lake Superior	1.92	0.63	0.21	0.21	Sweet et al. 1998
Lake Michigan	1.56	0.79	0.20	0.20	Sweet et al. 1998
Lake Erie	2.33	0.77	0.26	0.17	Sweet et al. 1998
Massachusetts Bay	1.37	1.70	0.38	4.11	Golomb et al. 1997
Chesapeake Bay	0.71	0.70	0.13	0.24	Scudlark et al. 1994
San Francisco Estuary	1.73	0.63	0.16	0.33	Tsai et al. 2001
Castro Valley	2.34	ND	0.02	0.40	This study
Total (Dry+Wet) Deposition Flux ($\mu\text{g}\cdot\text{m}^{-2}\cdot\text{d}^{-1}$)					
Southern Quebec, Canada	3.26	1.62	0.74	0.63	Gelinas et al. 2000
Lake Superior	8.49	2.19	1.25	0.57	Sweet et al. 1998
Lake Michigan	5.12	1.67	1.24	0.55	Sweet et al. 1998
Lake Erie	11.37	2.03	1.35	2.91	Sweet et al. 1998
Massachusetts Bay	6.85	4.11	0.74	7.40	Golomb et al. 1997
Commencement Bay, WA	20-149	8.4-49	NA	4.0-17	Crecelius 1991
San Francisco Estuary	4.66	2.25	0.22	3.84	Tsai et al. 2001
Castro Valley (bulk deposition)	6.9	0.76	0.068	0.78	This study

* Volume-weighted average concentration

NA – Not analyzed

Table 10. Ambient Benzene Concentrations in Castro Valley

site	start date	end date	conc ppbv	conc ug/m3
Madison	11/17/2004	11/19/2004	0.25	0.81
Madison	11/19/2004	11/21/2004	0.14	0.46
Madison	12/15/2004	12/17/2004	0.19	0.6
Madison	1/19/2005	1/21/2005	0.16	0.53
Madison	2/9/2005	2/11/2005	0.10	0.32
Redwood A	11/19/2004	11/21/2004	0.30	0.97
Redwood B	11/19/2004	11/21/2004	0.37	1.2
Redwood	12/15/2004	12/17/2004	0.48	1.5
Redwood	1/19/2005	1/21/2005	MF	
Redwood	2/9/2005	2/11/2005	0.28	0.88
CVE	2/9/2005	2/11/2005	0.24	0.75

Table 11. Comparison of Trace Metal Loading to the San Francisco Estuary from Atmospheric Deposition and Other Sources/Pathways (kg/year)

Sources/Pathways	Copper	Nickel	Cadmium	Chromium	Citation
Total Atmospheric Deposition	5,900	2,800	280	4,800	Tsai et al. 2001
Direct Atmospheric Deposition (dry + wet)	1,900	930	90	1,600	Tsai et al. 2001
Indirect Atmospheric Deposition ¹	4,000 (6%) ²	1,900 (4%) ²	190 (8%) ²	3,200 (8%) ²	Tsai et al. 2001
Effluent Discharges ³	6900- 7300	5600- 6100	98-330	1500-2000	Davis et al. 2000
Stormwater Runoff from San Francisco Bay Region ⁴	66,000 (36,000 - 150,000)	49,000 (27,000 - 78,000)	2,300 (1,300 - 3,700)	40,000 (22,000 - 64,000)	Davis et al. 2000
Stormwater Runoff from Central Valley Region ^{4,5}	270,000	410,000	1,600	550,000	Davis et al. 2000

¹ Indirect atmospheric deposition is derived from runoff and tributaries that are atmospheric in origin.

² Number in the parenthesis represents percentage of the loading from stormwater runoff in the San Francisco Bay Region.

³ Data is extrapolated from the 1998 monitoring data from effluent dischargers that represented approximately 85% of the total discharges; lower number assumes zero value for non-detectable samples, and upper number uses value of detection limit for non-detectable samples.

⁴ Include loading that is atmospheric in origin

⁵ Crude estimates, according to Davis et al. (2000)

APPENDIX Data Tables

Table A-1 Wet and Bulk Sample Concentrations (ppb)

Site	start date	end date	Field Rep	g water	area m2	bw/d	24Mg	27Al	28Si	47Ti	51V	52Cr	55Mn	56Fe	59Co	60Ni	65Cu	66Zn	88Sr	111Cd	121Sb	137Ba	208Pb
CVCC	17-Mar-04	31-Mar-04		205	0.01227	wet											4.1						
CVCC	31-Mar-04	21-Apr-04		37	0.01227	wet											1.7						
CVCC	09-Jun-04	07-Jul-04		79	0.01227	bulk	1384	964	1044	3.50	3.10	1.91	54.3	882.2	1.093	3.94	23.9	213	21.9	0.251	0.63	45.3	16.79
CVCC	06-Oct-04	20-Oct-04		650	0.01227	bulk	193	113	78	1.98	0.44	0.36	11.9	133.3	0.214	0.64	4.6	25	4.6	0.022	0.24	7.3	2.96
CVCC	20-Oct-04	03-Nov-04		460	0.01227	bulk	166	105	101	2.54	0.65	0.39	4.4	140.5	0.115	0.99	3.5	22	2.5	0.031	0.28	4.7	1.99
CVCC	03-Nov-04	17-Nov-04		316	0.01227	bulk	180	114	126	3.05	0.76	0.48	4.7	164.5	0.135	0.59	3.4	21	2.6	0.039	0.29	4.9	2.13
CVCC	17-Nov-04	03-Dec-04		147	0.01227	wet	45	39	39	0.78	0.24	0.37	2.5	44.7	0.094		2.0	21	1.3	0.019	0.15	3.0	0.81
CVCC	03-Dec-04	17-Dec-04		MF 378	0.01227	wet	61	23	26	0.53	0.14	0.14	1.1	28.4	0.043	0.33	1.9	6	0.8	0.007	0.08	1.4	0.43
CVCC	03-Dec-04	17-Dec-04		755	0.01227	bulk	88	12	16	0.21	0.03	0.14	3.1	18.2	0.019	0.18	1.6	12	1.0	0.041	0.07	2.0	0.30
CVCC	17-Dec-04	29-Dec-04		399	0.01227	wet	26	15	5	0.28	0.03	0.23	0.7	12.0	0.011		0.6	5	0.4	0.013	0.05	0.8	0.17
CVCC	17-Dec-04	29-Dec-04		399	0.01227	wet	26	14	6	0.22	0.04	0.00	0.7	11.1	0.004	0.00	0.5	5	0.3	0.004	0.03	0.7	0.17
CVCC	29-Dec-04	12-Jan-05		MF 210	0.00581	wet	104	61	11	0.04	0.08	0.11	0.5	6.8	0.007	0.14	0.8	4	0.8	0.026	0.22	3.0	0.18
CVCC	29-Dec-04	12-Jan-05		612	0.00581	bulk	237	7	7	0.18	0.05	0.11	0.5	10.8	0.003	0.40	0.8	4	1.4	0.012	0.11	1.8	0.19
CVCC	12-Jan-05	26-Jan-05		39	0.00581	wet	94	36	10	0.53	0.16	0.43	1.7	28.3	0.024	1.29	4.2	24	1.0	0.087	0.19	2.3	1.25
CVCC	26-Jan-05	09-Feb-05		70	0.00581	wet	353	26	31	0.94	0.33	0.36	1.7	41.0	0.015	0.19	1.9	11	2.4	0.006	0.08	2.5	0.91
CVCC	09-Feb-05	23-Feb-05		359	0.00581	wet	22	35	12	0.21	0.06	0.25	0.4	8.9	0.023	0.46	0.8	5	0.3	0.004	0.00	1.4	0.75
CVCC	09-Feb-05	23-Feb-05		526	0.00581	bulk	35	8	10	0.14	0.08	0.15	0.8	16.0	0.006	0.49	1.2	5	0.4	0.008	0.04	2.1	0.29
CVE	17-Mar-04	31-Mar-04		216	0.01227	wet											1.4						
CVE	31-Mar-04	21-Apr-04		26	0.01227	wet											2.6						
CVE	26-May-04	09-Jun-04		88	0.01227	wet	181	41	44	0.53	0.34	0.18	4.0	32.6	0.076	0.18	1.7	17	2.6	0.046	0.29	2.4	0.90
CVE	09-Jun-04	07-Jul-04		38	0.01227	bulk	1255	659	707	3.13	2.17	1.22	34.9	615.4	0.764	2.48	19.6	126	16.5	0.240	1.29	31.6	11.99
CVE	06-Oct-04	20-Oct-04		690	0.01227	bulk	207	137	191	2.28	0.46	0.40	12.1	160.6	0.155	0.75	3.7	25	4.6	0.024	0.22	7.4	1.97
CVE	20-Oct-04	03-Nov-04		420	0.01227	bulk	182	127	142	2.99	0.68	0.41	5.0	162.0	0.105	1.24	5.1	25	2.7	0.036	0.32	5.9	1.81
CVE	03-Nov-04	17-Nov-04		253	0.01227	bulk	185	129	153	3.03	0.70	0.42	5.0	168.8	0.106	1.02	5.0	25	2.7	0.033	0.26	5.9	1.80
CVE	17-Nov-04	03-Dec-04		159	0.01227	wet	75	16	14	0.34	0.18	0.25	1.5	17.4	0.262		1.2	12	1.3	0.019	0.31	1.8	0.33
CVE	03-Dec-04	17-Dec-04		MF 333	0.01227	wet	333	344		11.97	0.86	2.25	6.2		0.224	1.88	3.1	17	4.2	0.023	0.97	2.7	0.93
CVE	03-Dec-04	17-Dec-04		20	0.01227	wet	399	477	379	13.23	0.87	2.48	6.3		0.239		3.1	17	4.2	0.034	0.92	2.7	0.93
CVE	03-Dec-04	17-Dec-04		747	0.01227	bulk	95	38	50	1.02	0.19	0.27	1.9	52.2	0.057	0.24	1.4	9	1.2	0.052	0.10	2.3	0.63
CVE	17-Dec-04	29-Dec-04		433	0.01227	wet	48	30	9	0.21	0.05	0.11	0.9	12.7	0.008	0.21	0.5	4	1.8	0.012	0.04	0.9	0.19
CVE	29-Dec-04	12-Jan-05		633	0.00581	wet	197	5	4	0.13	0.06	0.07	0.2	0.4	0.014	0.16	0.5	3	1.3	0.002	0.02	2.1	0.05
CVE	29-Dec-04	12-Jan-05		651	0.00581	bulk	241	7	2	0.11	0.10	0.08	0.4	5.3	0.002	0.33	0.6	3	1.5	0.002	0.11	1.8	0.11
CVE	12-Jan-05	26-Jan-05		53	0.00581	wet	82	39	11	0.48	0.22	0.34	1.5	25.9	0.031	1.17	3.9	32	2.3	0.288	0.28	2.6	0.89
CVE	26-Jan-05	09-Feb-05		73	0.00581	wet	508	61	36	1.52	0.44	0.41	1.5	38.3	0.004	0.06	2.1	12	4.1	0.062	0.26	7.5	0.93
CVE	09-Feb-05	23-Feb-05	A	422	0.00581	wet	17	17	12	0.16	0.06	0.12	0.5	11.0	0.016	0.71	1.3	7	0.6	0.007	0.12	2.4	1.31
CVE	09-Feb-05	23-Feb-05	B	436	0.00581	wet	17	18	11	0.17	0.03	0.15	0.6	13.4	0.020	0.69	1.2	9	0.3	0.009	0.22	1.4	0.63
Madison	09-Jun-04	07-Jul-04		38	0.01227	bulk	621	285	374	6.78	1.93	0.91	43.9	218.5	0.543	2.38	21.5	161	15.9	0.181	0.40	21.7	9.20
Madison	06-Oct-04	20-Oct-04		680	0.01227	bulk	250	64	86	1.50	0.28	0.20	8.7	76.9	0.133	0.63	2.2	9	2.6	0.040	0.07	3.8	0.70
Madison	20-Oct-04	03-Nov-04		380	0.01227	bulk	83	25	15	0.78	0.48	0.19	1.6	29.6	0.033	0.26	OL	22	0.9	0.018	0.30	1.5	0.89
Madison	03-Nov-04	17-Nov-04		245	0.01227	bulk	154	40	32	1.38	0.73	0.26	1.7	46.8	0.025	0.20	1.4	9	1.3	0.026	0.10	1.9	0.97
Madison	03-Dec-04	17-Dec-04	A	745	0.01227	bulk	72	24	19	1.05	0.06	0.28	1.0	43.1	0.013	0.40	0.7	5	0.7	0.014	0.07	0.9	0.31
Madison	03-Dec-04	17-Dec-04	B	775	0.01227	bulk	60	11	7	0.41	0.10	0.12	0.8	17.5	0.004	0.14	0.6	7	0.7	0.046	0.06	0.8	0.26
Madison	17-Dec-04	29-Dec-04		435	0.01227	bulk	21	8	6	0.28	0.04	0.08	0.9	12.3	0.004	0.12	1.0	5	0.3	0.118	0.04	0.9	0.22
Madison	29-Dec-04	12-Jan-05	A	706	0.00581	bulk	187	3	4	0.07	0.04	0.06	0.3	3.8	0.005	0.10	0.5	3	1.1	0.000	0.05	1.1	0.09
Madison	29-Dec-04	12-Jan-05	B	744	0.00581	bulk	218	5	4	0.17	0.04	0.09	0.4	6.5	0.004	0.16	0.3	2	1.3	0.011	0.08	2.2	0.10
Madison	12-Jan-05	26-Jan-05		58	0.00581	bulk	73	31	23	1.34	0.30	0.34	2.2	47.0	0.035	1.15	4.2	58	1.4	0.069	0.22	3.0	1.45
Madison	26-Jan-05	09-Feb-05		142	0.00581	bulk	253	24	24	0.62	0.36	0.24	1.7	22.8	0.024	0.21	2.1	11	2.2	0.008	0.07	2.7	1.31
Madison	09-Feb-05	23-Feb-05	A	539	0.00581	bulk	10		3	0.08	0.07	0.12	0.2	4.2	0.020	0.81	0.3	3	0.1	0.015	0.03	0.5	0.11
Madison	09-Feb-05	23-Feb-05	B	564	0.00581	bulk	11	1	4	0.08	0.05	0.14	0.3	5.7	0.027	0.76	0.7	2	0.2	0.008	0.01	1.5	0.45
Redwood	06-Oct-04	20-Oct-04		620	0.01227	bulk	178	121	165	2.79	0.45	0.49	10.2	196.7	0.155	1.47	4.0	30	5.3	0.018	0.28	9.4	1.78
Redwood	20-Oct-04	03-Nov-04		370	0.01227	bulk	166	96	99	2.74	0.63	0.44	5.6	185.8	0.101	0.83	5.8	32	4.5	0.030	0.35	11.1	2.05
Redwood	03-Nov-04	17-Nov-04		222	0.01227	bulk	175	100	115	3.20	0.70	0.46	5.7	202.0	0.109	0.36	6.3	33	4.4	0.033	0.59	11.5	2.06
Redwood	17-Nov-04	03-Dec-04		159	0.01227	bulk	99	98	153	2.56	0.34	0.57	8.4	235.0	0.140		4.6	46	4.2	0.131	0.37	12.0	2.71
Redwood	03-Dec-04	17-Dec-04		661	0.01227	bulk	89	43	44	1.00	0.13	0.21	2.9	79.3	0.032	0.21	2.2	14	2.3	0.014	0.18	5.1	0.80
Redwood	03-Dec-04	17-Dec-04		672	0.01227	bulk	95	45	57	1.35	0.20	0.46	3.3	90.4	0.056		2.6	16	2.6	0.038	0.20	5.6	0.88
Redwood	17-Dec-04	29-Dec-04		486	0.01227	bulk	53	46	55	1.21	0.16	0.23	3.0	84.4	0.047	0.52	2.4	20	1.9	0.014	0.28	5.7	0.82
Redwood	29-Dec-04	12-Jan-05		630	0.00581	bulk	193	9	10	0.26	0.09	0.10	0.7	18.6	0.004	0.24	0.7	6	4.8	0.014	0.11		

Table A-2 Wet and Bulk Sample Deposition Rates ($\mu\text{g}\cdot\text{m}^{-2}\cdot\text{d}^{-1}$)

Site	start date	end date	Field Rep	g water	area m2	bww/d	24Mg	27Al	28Si	47Ti	51V	52Cr	55Mn	56Fe	59Co	60Ni	65Cu	66Zn	88Sr	111Cd	121Sb	137Ba	208Pb
CVCC	17-Mar-04	31-Mar-04		205	0.012272	wet											4.89						
CVCC	31-Mar-04	21-Apr-04		37	0.012272	wet											0.24						
CVCC	09-Jun-04	07-Jul-04		79	0.012272	bulk	318	222	240.1	0.80	0.71	0.44	12.49	203	0.25	0.91	5.50	49.1	5.04	0.06	0.15	10.42	3.86
CVCC	06-Oct-04	20-Oct-04		650	0.012272	bulk	730	429	295.1	7.47	1.56	1.37	45.19	504	0.81	2.44	17.40	93.2	17.54	0.08	0.92	27.70	11.18
CVCC	20-Oct-04	03-Nov-04		460	0.012272	bulk	445	280	269.9	6.81	1.75	1.05	11.88	376	0.31	2.64	9.42	59.6	6.58	0.08	0.75	12.57	5.32
CVCC	03-Nov-04	17-Nov-04		316	0.012272	bulk	331	210	231.7	5.60	1.40	0.88	8.68	303	0.25	1.08	6.27	39.5	4.76	0.07	0.54	9.09	3.91
CVCC	17-Nov-04	03-Dec-04		147	0.012272	wet	34	29	29.1	0.59	0.18	0.28	1.84	33	0.07		1.50	15.4	0.96	0.01	0.11	2.21	0.61
CVCC	03-Dec-04	17-Dec-04		MF 378	0.012272	wet																	
CVCC	03-Dec-04	17-Dec-04		755	0.012272	bulk	387	52	72.0	0.91	0.15	0.60	13.70	80	0.08	0.78	7.03	51.0	4.48	0.18	0.33	8.78	1.33
CVCC	17-Dec-04	29-Dec-04		399	0.012272	wet	70	39	13.5	0.76	0.06	0.62	2.00	32	0.03		1.58	12.7	1.00	0.03	0.14	2.08	0.45
CVCC	17-Dec-04	29-Dec-04		399	0.012272	wet	71	39	17.1	0.58	0.10	0.00	1.84	30	-0.01	0.01	1.30	12.4	0.87	0.00	0.09	1.97	0.46
CVCC	29-Dec-04	12-Jan-05		MF 210	0.005809	wet																	
CVCC	29-Dec-04	12-Jan-05		612	0.005809	bulk	1786	56	50.4	1.38	0.40	0.85	3.83	82	0.02	3.00	5.82	28.7	10.39	0.09	0.82	13.69	1.45
CVCC	12-Jan-05	26-Jan-05		39	0.005809	wet	45	17	4.8	0.25	0.08	0.21	0.79	13	0.01	0.61	1.98	11.2	0.46	0.04	0.09	1.09	0.59
CVCC	26-Jan-05	09-Feb-05		70	0.005809	wet	303	22	26.5	0.81	0.28	0.31	1.49	35	0.01	-0.16	1.62	9.2	2.07	0.01	0.07	2.16	0.78
CVCC	09-Feb-05	23-Feb-05		359	0.005809	wet	99	153	50.9	0.91	0.27	1.11	1.88	39	-0.10	-2.04	3.38	20.0	1.20	0.00	-0.01	5.99	3.30
CVCC	09-Feb-05	23-Feb-05		526	0.005809	bulk	228	50	62.6	0.90	0.52	0.94	5.22	104	-0.04	-3.18	7.81	31.9	2.74	-0.05	0.28	13.69	1.85
CVE	17-Mar-04	31-Mar-04		216	0.012272	wet											1.76						
CVE	31-Mar-04	21-Apr-04		26	0.012272	wet											0.26						
CVE	26-May-04	09-Jun-04		88	0.012272	wet	92	21	22.6	0.27	0.17	0.09	2.03	17	0.04	0.09	0.87	8.9	1.33	0.02	0.15	1.23	0.46
CVE	09-Jun-04	07-Jul-04		38	0.012272	bulk	137	72	77.2	0.34	0.24	0.13	3.81	67	0.08	0.27	2.14	13.8	1.80	0.03	0.14	3.44	1.31
CVE	06-Oct-04	20-Oct-04		690	0.012272	bulk	830	550	765.6	9.16	1.83	1.59	48.47	645	0.62	3.00	14.87	101.5	18.41	0.10	0.90	29.81	7.90
CVE	20-Oct-04	03-Nov-04		420	0.012272	bulk	445	311	346.1	7.31	1.67	1.00	12.19	396	0.26	3.04	12.38	61.3	6.58	0.09	0.77	14.51	4.43
CVE	03-Nov-04	17-Nov-04		253	0.012272	bulk	273	189	226.0	4.47	1.03	0.62	7.42	249	0.16	1.50	7.40	36.7	3.95	0.05	0.39	8.74	2.64
CVE	17-Nov-04	03-Dec-04		159	0.012272	wet	61	13	11.2	0.27	0.15	0.20	1.18	14	0.21		1.01	10.1	1.05	0.01	0.25	1.43	0.27
CVE	03-Dec-04	17-Dec-04		MF	0.012272	wet																	
CVE	03-Dec-04	17-Dec-04		20	0.012272	wet	47	56	44.2	1.54	0.10	0.29	0.73				0.36	2.0	0.49	0.00	0.11	0.31	0.11
CVE	03-Dec-04	17-Dec-04		747	0.012272	bulk	414	163	219.1	4.42	0.84	1.18	8.09	227	0.25	1.06	5.99	38.0	5.28	0.22	0.45	9.99	2.72
CVE	17-Dec-04	29-Dec-04		433	0.012272	wet	140	87	26.7	0.62	0.14	0.31	2.72	37	0.02	0.63	1.49	11.7	5.17	0.03	0.12	2.66	0.56
CVE	29-Dec-04	12-Jan-05		633	0.005809	wet	1531	35	30.5	0.98	0.50	0.56	1.76	-3	-0.11	1.26	4.22	21.4	10.30	-0.01	0.16	16.35	0.37
CVE	29-Dec-04	12-Jan-05		651	0.005809	bulk	1927	53	18.9	0.85	0.79	0.67	2.93	42	-0.02	1.85	4.63	26.1	11.64	0.02	0.89	14.08	0.92
CVE	12-Jan-05	26-Jan-05		53	0.005809	wet	54	25	7.2	0.32	0.14	0.22	0.97	17	0.02	0.77	2.53	21.1	1.50	0.19	0.18	1.73	0.58
CVE	26-Jan-05	09-Feb-05		73	0.005809	wet	455	55	31.9	1.36	0.39	0.37	1.36	34	0.00	-0.05	1.91	10.9	3.69	0.06	0.23	6.68	0.83
CVE	09-Feb-05	23-Feb-05	A	422	0.005809	wet	86	88	60.6	0.84	0.33	0.63	2.77	57	-0.08	-3.66	6.82	36.5	2.85	-0.03	0.62	12.41	6.78
CVE	09-Feb-05	23-Feb-05	B	436	0.005809	wet	90	95	58.6	0.92	0.17	0.79	3.35	72	-0.11	-3.68	6.64	47.1	1.74	-0.05	1.18	7.46	3.36
Madison	09-Jun-04	07-Jul-04		38	0.012272	bulk	68	31	40.8	0.74	0.21	0.10	4.79	24	0.06	0.26	2.35	17.6	1.74	0.02	0.04	2.37	1.00
Madison	06-Oct-04	20-Oct-04		680	0.012272	bulk	989	255	340.8	5.95	1.12	0.78	34.38	304	0.53	2.48	8.89	36.5	10.26	0.16	0.27	15.11	2.78
Madison	20-Oct-04	03-Nov-04		380	0.012272	bulk	184	54	33.8	1.72	1.07	0.42	3.48	65	0.07	0.57	47.73	48.3	1.90	0.04	0.67	3.21	1.97
Madison	03-Nov-04	17-Nov-04		245	0.012272	bulk	220	58	45.3	1.97	1.04	0.37	2.37	67	0.04	0.28	2.04	12.4	1.89	0.04	0.15	2.73	1.39
Madison	03-Dec-04	17-Dec-04	A	745	0.012272	bulk	312	104	83.0	4.55	0.27	1.22	4.46	187	0.06	1.73	3.06	21.3	3.15	0.06	0.28	3.77	1.34
Madison	03-Dec-04	17-Dec-04	B	775	0.012272	bulk	270	50	33.3	1.83	0.43	0.55	3.55	79	0.02	0.62	2.89	32.6	3.11	0.21	0.25	3.49	1.17
Madison	17-Dec-04	29-Dec-04		435	0.012272	bulk	61	24	17.8	0.82	0.12	0.24	2.51	36	0.00	0.34	2.89	13.9	1.00	0.35	0.11	2.75	0.64
Madison	29-Dec-04	12-Jan-05	A	706	0.005809	bulk	1621	23	12.0	0.64	0.36	0.54	2.36	33	-0.05	0.87	4.05	29.5	9.60	0.00	0.47	9.17	0.77
Madison	29-Dec-04	12-Jan-05	B	744	0.005809	bulk	1993	44	35.1	1.52	0.38	0.84	3.89	59	-0.03	1.44	3.20	17.9	11.68	0.10	0.74	20.26	0.94
Madison	12-Jan-05	26-Jan-05		58	0.005809	bulk	52	22	16.3	0.96	0.22	0.24	1.58	34	0.02	0.83	3.01	41.9	0.99	0.05	0.16	2.19	1.04
Madison	26-Jan-05	09-Feb-05		142	0.005809	bulk	441	42	42.3	1.09	0.62	0.41	2.91	40	0.04	0.37	3.69	19.3	3.76	0.01	0.12	4.71	2.29
Madison	09-Feb-05	23-Feb-05	A	539	0.005809	bulk	66		19.5	0.53	0.44	0.77	1.62	28	-0.19	-5.35	1.94	18.7	0.99	-0.10	-0.21	3.59	0.72
Madison	09-Feb-05	23-Feb-05	B	564	0.005809	bulk	80	7	29.4	0.52	0.38	0.96	1.96	39	-0.18	-5.25	4.72	16.8	1.20	-0.06	-0.10	10.12	3.10
Redwood	06-Oct-04	20-Oct-04		620	0.012272	bulk	644	437	593.7	10.08	1.61	1.75	36.98	710	0.56	5.29	14.50	109.5	19.18	0.06	1.01	33.98	6.43
Redwood	20-Oct-04	03-Nov-04		370	0.012272	bulk	358	206	212.3	5.90	1.37	0.95	12.03	400	0.22	1.79	12.38	69.2	9.67	0.06	0.75	23.85	4.40
Redwood	03-Nov-04	17-Nov-04		222	0.012272	bulk	226	129	148.1	4.13	0.91	0.60	7.40	261	0.14	0.46	8.08	42.6	5.75	0.04	0.76	14.83	2.67
Redwood	17-Nov-04	03-Dec-04		159	0.012272	bulk	80	79	123.5	2.07	0.28	0.46	6.82	190	0.11		3.76	37.4	3.39	0.11	0.30	9.75	2.19
Redwood	03-Dec-04	17-Dec-04		661	0.012272	bulk	344	166	168.7	3.83	0.51	0.80	11.20	305	0.12	0.80	8.34	53.4	8.97	0.05	0.69	19.46	3.09
Redwood	03-Dec-04	17-Dec-04		672	0.012272	bulk	372	177	222.0	5.29	0.77	1.81	13.01	353	0.22		10.00	61.8	10.31	0.15	0.80	21.77	3.45
Redwood	17-Dec-04	29-Dec-04		486	0.012272	bulk	174	152	181.3	4.00	0.53	0.76	9.90	278	0.16	1.71	7.91	66.2	6.14	0.05	0.93	18.72	2.70
Redwood	29-Dec-04	12-Jan-05		630	0.005809	bulk	1496	71	77.3	2.01	0.71	0.79	5.60	144	0.01	1.83	5.34	44.1	36.99	0.01	0.86	24.25	4.98
Redwood	12-Jan-05	26-Jan-05		51	0.005809	bulk	192	219	214.7	2.56	0.53	0.66	13.28	431	0.21	1.36	9.87	83.6	15.94	0.10	0.57	22.06	4.12
Redwood	26-Jan-05	09-Feb-05		100	0.005809	bulk	675	236	221.3	2.59	0.75	0.76	12.76	297	0.19	0.32	9.61	83.0	12.66	0.07	0.68	24.69	3.35
Redwood	09-Feb-05	23-Feb-05		524	0.005809	bulk	110	43	75.8	2.04													

Table A-3 Dry Deposition Rates ($\mu\text{g}\cdot\text{m}^{-2}\cdot\text{d}^{-1}$)

site	start date	end date	field rep	g water	area m2	bw/d	24Mg	27Al	28Si	47Ti	51V	52Cr	55Mn	56Fe	59Co	60Ni	65Cu	66Zn	88Sr	111Cd	121Sb	137Ba	208Pb
CVCC	07-Jul-04	09-Jul-04		10	0.05	DRY	999	1429	130	43.0	2.77	7.13	26.63	2638	0.53	5.60	15.8	85.5	8.44	0.09	0.74	21.7	8.55
CVCC	28-Jul-04	30-Jul-04		10	0.05	DRY	641	991	115	48.9	3.70	7.97	32.85	1976	0.69	2.73	21.5	89.8	9.47	0.09	0.38	26.4	13.22
CVCC	18-Aug-04	20-Aug-04		10	0.05	DRY	669	1087	108	40.4	3.50	7.54	32.57	2238	0.78	4.07	21.5	97.6	9.40	0.09	0.56	21.7	16.24
CVCC	20-Aug-04	22-Aug-04		10	0.05	DRY	508	873	109	32.9	3.14	7.79	26.32	1693	0.61	2.08	18.6	86.2	6.70	0.08	0.36	20.0	14.86
CVCC	31-Aug-04	02-Sep-04	A	10	0.05	DRY	487	775	118	43.2	3.00	7.84	28.71	1361	0.64	1.92	16.1	74.9	8.81	0.01	0.28	19.9	10.77
CVCC	31-Aug-04	02-Sep-04	B	10	0.05	DRY	572	834	119	40.3	2.95	7.81	32.13	1659	0.73	2.74	19.7	81.3	9.02	0.05	0.35	21.1	14.14
CVCC	15-Sep-04	17-Sep-04	A	10	0.05	DRY	1327	1985	191	83.8	6.33	13.17	59.86	3390	1.43	6.79	24.6	130.1	14.70	0.11	0.63	38.3	15.34
CVCC	15-Sep-04	17-Sep-04	B	10	0.05	DRY	1237	1751	121	77.8	6.07	12.22	62.20	3151	1.44	7.18	31.6	132.3	16.06	0.25	0.43	37.9	18.82
CVCC	29-Sep-04	01-Oct-04	A	10	0.05	DRY	537	747	102	38.5	2.37	7.14	23.57	1330	0.84	1.48	19.8	68.2	7.33	0.01	1.00	18.7	6.43
CVCC	29-Sep-04	01-Oct-04	B	10	0.05	DRY	607	843	69	39.1	2.61	4.56	24.30	1450	1.00	3.34	15.3	75.6	7.37	0.07	0.28	19.0	6.80
CVCC	01-Oct-04	03-Oct-04	A	10	0.05	DRY	239	448	77	21.9	1.52	6.12	13.92	801	0.38	1.46	10.1	43.9	5.60	0.00	0.20	13.2	5.91
CVCC	01-Oct-04	03-Oct-04	B	10	0.05	DRY	282	500	88	23.1	1.65	5.77	14.85	790	0.44	8.41	11.2	43.6	5.87	0.02	0.56	14.4	5.96
CVCC	17-Nov-04	19-Nov-04		10	0.05	DRY	313	734	260	44.3	2.80	8.97	22.96	968	1.07	3.60	17.6	80.2	6.48	0.61	0.95	19.4	8.94
CVCC	19-Nov-04	21-Nov-04		10	0.05	DRY	746	1307	97	59.5	4.38	12.83	56.20	2040	1.33	6.34	16.0	103.8	8.87	0.69	0.70	28.9	13.98
CVCC	15-Dec-04	17-Dec-04		10	0.05	DRY	245	426	109	22.1	1.50	12.45	24.27	683	0.65	3.33	28.1	69.5	3.70	0.57	0.51	14.1	4.77
CVCC	19-Jan-05	21-Jan-05		10	0.05	DRY	178	325	103	20.8	1.87	11.71	26.35	630	0.69	3.98	16.7	58.9	4.29	0.58	0.57	13.2	6.57
CVCC	09-Feb-05	11-Feb-05		10	0.05	DRY	756	289	86	22.8	1.49	11.45	27.71	571	0.69	3.51	14.9	57.7	5.70	0.63	0.45	12.7	6.79
CVE	17-Nov-04	19-Nov-04		10	0.05	DRY	232	516	120	31.1	2.66	8.51	17.65	829	1.12	2.61	13.5	71.9	5.91	0.87	0.99	16.9	7.45
CVE	19-Nov-04	21-Nov-04		10	0.05	DRY	1171	2110	126	93.7	6.70	17.05	79.57	3058	2.30	8.60	29.7	115.4	13.31	0.85	0.66	36.3	17.30
CVE	15-Dec-04	17-Dec-04		10	0.05	DRY	212	503	116	24.6	1.52	12.22	25.89	740	0.82	22.93	12.5	81.6	8.24	0.55	0.64	15.9	5.71
CVE	19-Jan-05	21-Jan-05		10	0.05	DRY	195	429	90	21.2	1.50	11.80	23.46	714	0.62	3.94	11.3	66.8	4.61	0.47	0.60	13.6	8.27
CVE	09-Feb-05	11-Feb-05		10	0.05	DRY	176	381	111	22.5	1.35	9.45	19.89	564	0.52	2.96	14.3	58.1	3.72	0.33	0.38	11.8	3.40
Madison	07-Jul-04	09-Jul-04		10	0.05	DRY	607	901	80	24.5	1.40	5.39	15.97	1693	0.26	0.59	6.2	32.5	5.33	0.00	0.21	11.0	4.47
Madison	28-Jul-04	30-Jul-04		10	0.05	DRY	267	485	167	32.2	2.20	6.75	21.05	887	0.34	1.77	13.1	54.5	5.91	0.05	0.40	13.9	9.37
Madison	18-Aug-04	20-Aug-04		10	0.05	DRY	263	429	161	24.7	2.05	6.52	21.51	879	0.34	0.52	10.6	42.6	5.57	0.03	0.25	13.3	8.11
Madison	20-Aug-04	22-Aug-04		10	0.05	DRY	243	351	74	21.3	1.94	5.90	23.90	793	0.28	1.03	9.5	38.9	3.90	-0.01	0.41	13.4	6.59
Madison	31-Aug-04	02-Sep-04		10	0.05	DRY	219	224	77	12.7	0.99	4.22	11.42	466	0.16	-0.26	5.1	26.5	2.68	0.01	0.15	7.1	3.70
Madison	15-Sep-04	17-Sep-04		10	0.05	DRY	484	648	93	32.6	2.14	6.66	24.09	1137	0.48	1.05	8.7	37.7	4.96	0.05	0.21	12.9	7.09
Madison	29-Sep-04	01-Oct-04		10	0.05	DRY	261	313	67	20.6	1.14	6.10	11.89	573	0.19	-0.92	6.3	29.4	3.15	0.00	-0.05	9.1	2.66
Madison	01-Oct-04	03-Oct-04		10	0.05	DRY	159	186	66	13.3	1.20	4.77	7.37	389	0.14	0.42	4.9	13.7	3.04	-0.04	-0.03	7.2	3.26
Madison	17-Nov-04	19-Nov-04		10	0.05	DRY	71	107	93	12.8	1.27	7.40	5.49	175	0.90	1.12	7.2	19.6	1.88	0.89	0.78	5.2	3.27
Madison	19-Nov-04	21-Nov-04		10	0.05	DRY	133	251	75	18.9	1.04	10.13	22.41	418	0.47	4.46	34.1	50.7	2.86	1.52	0.34	10.1	5.81
Madison	15-Dec-04	17-Dec-04		10	0.05	DRY	36	57	96	10.7	0.58	10.95	17.42	161	0.46	4.21	16.5	28.3	1.44	0.58	0.39	7.0	3.07
Madison	19-Jan-05	21-Jan-05		10	0.05	DRY	38	75	93	7.0	0.52	11.28	13.64	129	0.36	2.87	6.4	32.9	1.34	0.44	0.25	5.9	2.77
Madison	09-Feb-05	11-Feb-05		10	0.05	DRY	51	96	76	7.6	0.41	7.53	9.15	95	0.22	0.94	8.7	21.2	1.06	0.27	0.06	4.1	1.93
Redwood	07-Jul-04	09-Jul-04		10	0.05	DRY	5767	15695	161	208.2	17.17	13.92	137.56	16136	3.54	8.98	35.6	156.9	32.66	0.14	0.30	61.0	17.60
Redwood	28-Jul-04	30-Jul-04		10	0.05	DRY	704	1304	172	73.3	4.12	9.32	39.79	2726	0.89	3.93	21.9	146.6	14.79	0.11	0.82	40.6	11.66
Redwood	18-Aug-04	20-Aug-04		10	0.05	DRY	962	1523	178	69.8	4.86	9.98	50.26	3649	1.17	5.80	31.8	192.9	36.44	0.14	0.93	48.6	17.98
Redwood	20-Aug-04	22-Aug-04		10	0.05	DRY	489	847	193	47.3	3.36	9.34	27.24	1954	0.58	4.54	25.9	133.0	20.97	0.10	0.91	31.2	14.40
Redwood	31-Aug-04	02-Sep-04		10	0.05	DRY	456	694	96	38.4	2.47	7.46	25.95	1493	0.63	1.97	17.9	101.5	10.73	0.03	0.46	24.3	8.47
Redwood	15-Sep-04	17-Sep-04		10	0.05	DRY	994	1415	89	65.3	4.66	10.02	57.41	3088	1.19	4.59	21.5	154.8	19.61	0.08	0.66	49.4	11.66
Redwood	29-Sep-04	01-Oct-04		10	0.05	DRY	563	829	108	45.5	2.59	7.47	24.41	1616	0.50	0.94	15.3	90.0	9.10	0.01	0.72	26.3	5.77
Redwood	01-Oct-04	03-Oct-04		10	0.05	DRY	316	517	81	28.5	1.87	6.26	18.24	1120	0.38	2.43	24.5	86.5	7.98	0.06	0.66	26.0	10.25
Redwood	17-Nov-04	19-Nov-04		10	0.05	DRY	1022	1978	179	125.7	7.25	16.97	84.05	5092	1.98	7.70	42.3	255.6	24.66	0.85	1.39	75.4	18.33
Redwood	19-Nov-04	21-Nov-04		10	0.05	DRY	2622	5736	167	283.0	16.80	24.75	161.16	11199	3.34	14.11	47.4	265.6	42.09	0.68	0.80	86.8	21.45
Redwood	15-Dec-04	17-Dec-04		10	0.05	DRY	1020	1845	151	85.3	5.85	19.23	72.76	4609	1.65	9.26	44.9	223.7	35.35	0.91	1.11	78.3	14.61
Redwood	19-Jan-05	21-Jan-05		10	0.05	DRY	1163	1624	95	95.6	6.26	19.36	78.67	4296	1.73	8.93	32.7	240.5	51.21	0.66	1.11	80.3	28.50
Redwood	09-Feb-05	11-Feb-05		10	0.05	DRY	807	1378	83	73.9	4.89	11.31	58.08	2955	1.33	5.92	37.2	194.9	20.07	0.43	0.64	53.3	11.58