

CHARACTERIZATION OF THE PRESENCE AND
SOURCES OF DIAZINON IN THE
CASTRO VALLEY CREEK WATERSHED

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EXECUTIVE SUMMARY

Diazinon, an organophosphate insecticide, has been detected in creeks throughout the Bay Area. Previous toxicity studies showed that storm runoff in Castro Valley Creek was frequently toxic to the aquatic test organism *Ceriodaphnia dubia*, and that diazinon was the most likely cause of this toxicity (Hansen, 1995). Since its watershed is fairly representative of urban land use patterns in the East Bay and the stream is also part of long-term stream monitoring studies, Castro Valley Creek was selected as the site for a more detailed study of diazinon in surface runoff.

The main purpose of this study was to characterize the temporal and spatial patterns of occurrence of diazinon in the Castro Valley Creek watershed. Runoff at the discharge point for the entire watershed was sampled during multiple storm events to record both seasonal and within-event variations in diazinon concentration. Analysis of these data suggested how seasonal and hydrologic factors may affect diazinon levels, and provided the basis for estimates of concentrations in unsampled runoff and of the total mass of diazinon discharged during the 1995-96 water year. Repeated sampling during and after storm events showed how long diazinon concentrations remained elevated after storms. Since a pollutant's toxicity to aquatic organisms depends on both the concentration and the duration of exposure, the persistence of diazinon in creeks is an important consideration in assessing the ecological impact of diazinon.

Sampling at multiple locations was used to evaluate the spatial distribution of diazinon within the watershed. Results from five major subcatchments indicated that sources are not restricted to particular areas, and samples from street gutters showed that at a smaller scale individual diazinon sources may be very localized. A few street gutter samples contained very high concentrations of diazinon. Data from street gutter samples were used to form a rough estimate of the number of sources in the watershed. Three selected residences were given experimental applications of diazinon to see if subsequent runoff could contain high concentrations of diazinon. Although limited in scope, the test results suggest that residential users applying diazinon in accordance with label directions may contribute significantly to the diazinon in runoff in Castro Valley Creek.

This report is part of a coordinated effort to increase available information about urban pesticide toxicity in surface waters of the Bay Area and to develop a control strategy. Related reports summarize the current problems and questions related to diazinon (Katznelson and Mumley, 1997), discuss its use and formulation (Scanlin and Cooper, 1997), and propose a framework for control strategies to reduce diazinon levels in creeks (Scanlin and Gosselin, 1997).

ES-1 DIAZINON IN CASTRO VALLEY CREEK

Temporal variation in diazinon levels were studied at the lower end of the watershed at station S3, where flow meters provide continuous records of flow rates and discharge volumes. Discrete samples were taken at fixed flow intervals by an automatic sampler during selected storm events in the 1995-96 and 1996-97 rainy seasons. Samples were analyzed for diazinon using ELISA (enzyme linked immuno-sorbent assay), which permits rapid testing of large numbers of samples.

Diazinon was detected in all of the sampled events. Event mean concentrations (EMCs) during 1995-96 ranged from 180 to 820 nanograms per liter (ng/l; equivalent to parts per trillion). High EMCs of diazinon were observed during fall and spring months, corresponding to periods when estimated application rates were high. High EMCs were inversely correlated with flow volumes during the two weeks preceding sampled events; this suggests that diazinon accumulates on watershed surfaces during intervals of dry weather and is then washed into the creek by subsequent storms. These relationships with season and antecedent flow appeared to be stronger for the 1995-96 season than for 1996-97. Average EMCs for 1996-97 events were somewhat lower than for 1995-96, which may be related to year-to-year differences in application rate and weather patterns.

The temporal variation of diazinon concentrations within storms followed two main patterns: concentrations either rose to a high initial peak and then gradually declined, or remained at a more constant, intermediate level throughout the storm. Low antecedent flow appeared to be associated with events showing high early peaks, possibly because more diazinon had accumulated, or because recently applied diazinon was easily washed off during the early part of the storm. Diazinon concentrations decreased gradually after storms, taking about 2 days to decrease 50% from the end-of-storm levels. Although individual storms were usually relatively short, closely spaced events or high within-storm EMC's can maintain diazinon concentrations over 150 ng/l for extended periods of up to a week. Previous toxicity tests have shown that this level of exposure can cause toxicity to *Ceriodaphnia dubia* test organisms during that time. Animals in the creek may thus receive toxic exposures to diazinon several times during the rainy season, which could have an impact on aquatic life.

EMCs and the discharged volumes of water were used to calculate the total mass of diazinon discharged in each sampled event. Mass discharge increased with increasing event flow volume, even though dilution may have reduced the average concentration. The seasonal and hydrologic patterns observed in sampled events were used to estimate EMCs and mass discharge for unsampled events. Non-storm or dry weather flows, defined as flows less than 5 cubic feet per second (cfs), were assumed to have lower diazinon concentrations than storm flows. The total amount of diazinon discharged from Castro Valley Creek during the 1995-96 water year was estimated at about 600 grams, or about 0.3% of the estimated annual usage in the watershed. About 90% of this amount was discharged during periods of storm flow (greater than 5 cubic feet per second), while the remainder was discharged during dry weather and non-storm flows.

ES-2 DISTRIBUTION OF SOURCES WITHIN THE CASTRO VALLEY CREEK WATERSHED

Grab samples were taken by hand at different locations within the watershed to evaluate spatial patterns of diazinon occurrence. During different storm events, sampling was performed at either the subcatchment level or more locally in residential neighborhoods. Analysis of these samples showed that diazinon sources are widely distributed throughout the watershed.

Runoff was sampled at half-hour intervals during the storm of April 1, 1996 at five subcatchments that comprised 78% of the watershed area. Diazinon concentrations ranged from 50 to 1250 ng/l in individual samples and from 200 to 690 ng/l when averaged for each subcatchment. Mean subcatchment concentration was higher in subcatchments with less open space.

Grab samples taken from 45 street gutters in residential sections of Subcatchments 2 and 3 on May 16, 1996 were more variable, with concentrations ranging from non-detect (<30 ng/l) to 79,000 ng/l. Most sample concentrations were much lower, and the median sample concentration (80 ng/l) was similar to levels previously observed in rainfall samples. Higher diazinon levels were not restricted to particular neighborhoods, and adjacent gutters which drained independent sets of residential properties often had very different concentrations.

Some street gutters contain high diazinon levels on more than one occasion, suggesting that there may be some consistent sources. However, average street gutter concentrations were more similar to the concentrations observed for S3 and the subcatchments. An experimental calculation was made to estimate the number of sources in the watershed. If each street gutter sample with a diazinon concentration greater than 100 to 400 ng/l contained one source property in its drainage area, an estimated two to four percent of the properties in the total area sampled could be considered to be sources. (Extrapolation of these percentages suggests that a total of 300-600 households in the watershed may be sources during spring storm events.)

ES-3 RESIDENTIAL RUNOFF STUDY

Since outdoor residential application appears to be a major use of diazinon in urban areas (Scanlin and Cooper, 1997), an attempt was made to verify whether application at recommended rates could produce the high concentrations observed in street gutter samples. Test applications were made during February 1997 at two residences in Castro Valley and one in a similar neighborhood in Oakland. Liquid concentrate (25% diazinon) was mixed and applied to outdoor paved areas in a quantity two-thirds of the amount recommended for control of ants. Grab samples were taken of runoff from roof, patio or driveway areas at each site during subsequent rainfall events.

On-site runoff on the sprayed properties contained diazinon concentrations of up to 1,200,000 ng/l several days after application. Patio and driveway runoff concentrations were higher than those in roof drains. Diazinon was still detected on-site seven weeks after application, and rainfall samples taken at one sprayed property after application showed elevated concentrations of diazinon up to 1300 ng/l. Rainfall during this period was insufficient to produce enough runoff for sampling in street gutters near these properties; however, these results indicate that outdoor residential use of liquid diazinon in accordance with the label directions is a possible source of the diazinon observed in the creek and storm drains. Alternative formulations such as granules were not tested and may contribute an unknown amount of diazinon to runoff.

1.0 INTRODUCTION

1.1 BACKGROUND

Diazinon is an organophosphate insecticide. It is widely used for both agricultural and urban pest control. In urban areas, it is primarily used for the outdoor control of ants, fleas, spiders and grubs. The estimated annual per capita urban use of diazinon in Alameda County (CA) is 0.02 lbs. (9g) of active ingredient (Scanlin and Cooper, 1997).

This widespread use of diazinon may be having a negative impact on water quality in urban creeks. Toxicity tests conducted on storm water samples from throughout Alameda County have shown toxicity to a standard laboratory test organism, *Ceriodaphnia dubia* (WCC, 1991, 1994, 1995, 1996a). A toxicity identification evaluation (TIE) determined that diazinon was the primary cause of this toxicity (Hansen, 1995). The presence of diazinon in storm water is not limited to Alameda County. Diazinon was consistently detected in storm water samples collected from residential watersheds throughout the San Francisco Bay area and the Central Valley of California, often at concentrations that cause toxicity to test organisms (Katznelson and Mumley, 1997).

Concern about the potential impact of diazinon in Bay Area waters led to the formation of the Urban Pesticide Committee by representatives of diverse agencies and organizations (Katznelson and Mumley, 1997). Several related reports arising from efforts by the committee discuss different aspects of diazinon use (Scanlin and Cooper, 1997), policy issues and current knowledge (Katznelson and Mumley, 1997) and proposals for a control strategy (Scanlin and Gosselin, 1997). This report provides information about diazinon in a typical urban watershed as a complement and support to the other publications.

The objectives of this study were (1) to characterize the temporal variations of diazinon concentrations in Castro Valley Creek, (2) to track the sources of diazinon in the watershed, and (3) to determine if the application of diazinon in accordance with label directions could be contributing to the observed toxic concentrations.

1.2 WATERSHED DESCRIPTION

The Castro Valley Creek Watershed, a sub-watershed of the San Lorenzo Creek drainage, covers an area of 5.5 square miles in west central Alameda County (Figure 1.2.1). It has been selected for ongoing water quality monitoring by Alameda County because it contains a representative mix of residential and commercial land uses (WCC 1995, 1996a). Previous sampling has shown consistently high levels of both diazinon and toxicity in runoff samples from Castro Valley Creek and suggested that a number of hydrologic factors may affect diazinon concentrations.

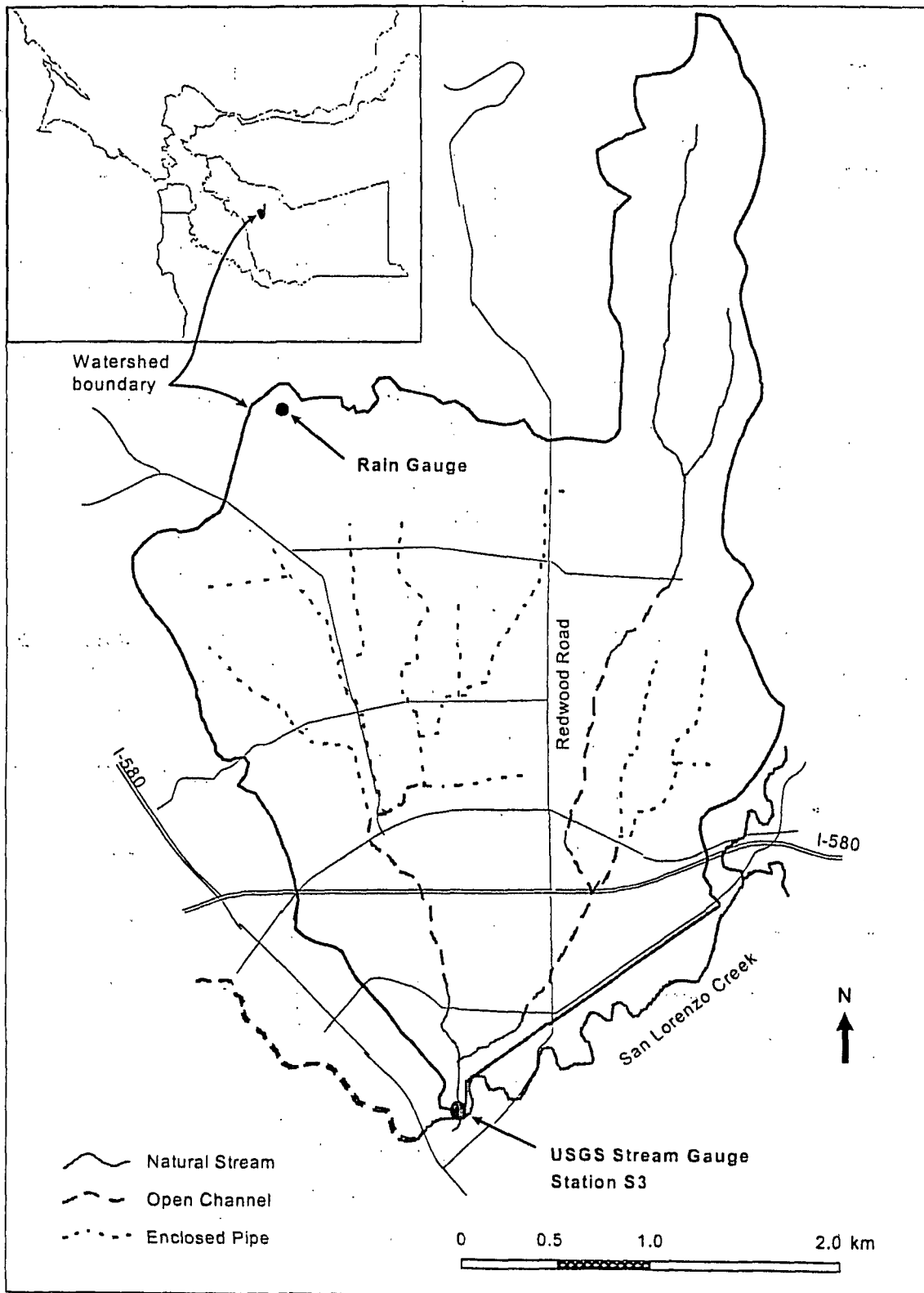


Figure 1.2.1 Location Map of Castro Valley Creek Watershed

The population in the area is approximately 35,000. The area is predominantly low density residential development (50%), with some open space (35%), and commercial development (15%). There are 11,600 assessed parcels with single family residential units and 3,200 with multiple family units. Mean household income is approximately \$50,000 (US Census Data, 1990, Sandy Rivera, Alameda County Planning Department, pers. comm.).

Mean annual rainfall in the watershed is about 21 inches, with over 95% occurring between October 1 and May 31. The soils are predominantly clays and silty clays. The downstream portion of the creek is still in a relatively natural state but most of the upper portions have been channelized or enclosed (Figure 1.2.1). The elevation ranges from 200 to 500 feet (65 to 160 meters) above mean sea level.

Castro Valley Creek flows year round. Summertime flows average about 0.2 cubic feet per second (cfs). The peak flow of record was 1,350 cfs during a storm event on January 23, 1983. The annual flow for 1995-96 water year (October 1 to September 30) was approximately 120 million cubic feet.

2.0 DIAZINON IN CASTRO VALLEY CREEK

2.1 OBJECTIVES AND METHODS

This portion of the study examined the occurrence of diazinon in combined runoff from all parts of the watershed, which is discharged at sampling station S3 (figure 1.2.1). The primary objective was to characterize temporal variations in diazinon over the course of a storm season. The relationship of these temporal variations with hydrologic (flow-related) variables was also examined and the results used to estimate the total mass of diazinon that was discharged during the year. The sampling data were also used to study how diazinon levels varied within and after each storm event.

All of the samples discussed in this section of the report were collected at station S3 near the mouth of Castro Valley Creek (Figure 1.2.1), which is the site of United States Geological Survey (USGS) gaging station number 1181008. Storm runoff samples were collected using an ISCO 3700 sampling device and an ISCO 3230 flow meter; which are housed in the USGS gaging station. The flow meter provided continuous stream level measurements that were converted to flow discharge (cubic feet/second) using the USGS rating curve for the site. These level or discharge observations were automatically recorded at two-minute intervals during most storm events and at ten-minute intervals throughout the rainy season. USGS discharge data (USGS, 1996) were used to to determine flow volumes at other times of the year. Flow meter readings were used by the sampler to control the discharge volume (cubic feet) represented by each water sample collected. The sampler could be set to keep separate discrete samples for different segments of a storm event or to combine flow-weighted subsamples into a single composite sample. Supplementary "grab" samples were collected manually during periods of non-storm flow (< 5 cfs), using 20 ml borosilicate glass vials that were rinsed in creek water immediately before sampling.

All samples were analyzed for diazinon using an enzyme linked immuno-sorbent assay (ELISA) method. Some samples were also sent to a laboratory for confirmation. (See Appendix A for additional description of sampling protocol and QA/QC procedures.) The event mean concentration (EMC) of diazinon for each sampled event was obtained either through analysis of a composite sample or through a calculation based on analysis of discrete samples (see Appendix B).

2.2 SEASONAL VARIATION IN DIAZINON CONCENTRATION

To examine if there were any seasonal trends in diazinon levels, flow-weighted samples were taken for twelve storm events throughout the 1995-96 rainy season (October - May). Diazinon was found above the detection level of 30 ng/l in all samples; EMCs for the sampled events ranged from a low of 180 nanograms/liter

(ng/l, or ppt) to a high of 820 ng/l (Table 2.2.1). EMCs were highest in December and May, as shown in Figure 2.2.1, which also plots the seasonal hydrograph (flow discharge as a function of time) at S3 from December 1995 through May 1996. There was no significant rainfall for the season prior to December 1995.

Table 2.2.1 Hydrologic (Stream Flow) Parameters and Mean Diazinon Concentrations for Sampled Events at Station S3 in Castro Valley Creek During the 1995-1996 Storm Season.

Event Number	Start Date	Start Time for Sampling	Sampling Duration (hr:min)	Flow Volume (cubic feet x 1,000)	Diazinon Event Mean Concen. (ng/l)	Flow Volume Previous 2 weeks (cf x 1,000)	Avg. Temp. Previous 30 Days (°F) (1)
1	12/4/95	2:30	28:00	147	820	225	59.5
2	12/10/95	17:50	10:50	1,746	700	565	59.5
3	12/11/95	18:40	13:00	6,063	300	6,240	59.4
4	12/29/95	12:10	3:40	745	660	2,100	54.5
5	1/16/96	5:50	3:00	4,739	320	460	52.9
6	1/18/96	14:40	13:00	3,075	250	9,190	53.1
7	1/20/96	22:20	8:10	2,688	200	13,030	53.1
8	2/3/96	20:50	12:20	2,732	190	24,360	51.8
9	3/4/96	8:10	14:00	3,300	180	20,780	56.1
10	4/1/96	9:50	6:20	2,893	290	1,250	57.5
11	4/17/96	16:00	6:00	1,713	450	4,451	58.4
12	5/15/96	12:00	12:00	4,700	700	720	63.7

(1) Source: NOAA daily temperature data for Newark, CA

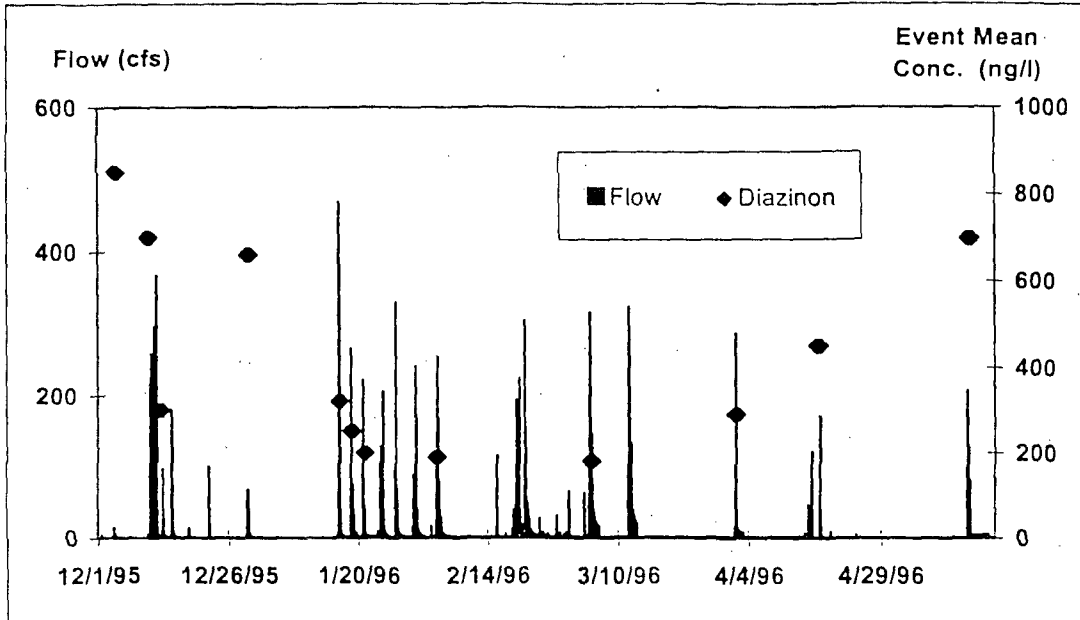


Figure 2.2.1 Event Mean Concentrations of Diazinon for Sampled Events and Stream Flow (Discharge) at Station S3 Between December 1, 1995 and May 21, 1996

Diazinon concentrations appear to vary seasonally, declining in the winter and increasing again in spring. Sampling during previous years also found that higher concentrations occurred during fall and spring (WCC, 1994, 1995, 1996a). This variation in diazinon levels parallels the seasonal pattern of use: the amount of diazinon applied in urban areas is lowest in winter and rises again in March, with the heaviest monthly applications in summer and early fall (Department of Pesticide Regulation data, from Scanlin and Cooper, 1997). Correlation analysis showed that for 1995-96 events EMC was highly correlated ($p < 0.05$) with mean air temperature for the previous month (Table 2.2.2). Average temperatures generally vary seasonally and are also associated with greater insect activity, which may stimulate application of pesticides.

Some of the variation in diazinon is probably also due to hydrologic factors. EMCs had a strong negative correlation with two-week antecedent flow (the total volume of water discharged during two weeks prior to a given storm event). In other words, less rainfall prior to an event was associated with higher EMCs, possibly through accumulation of diazinon on the watershed area between storms. Similar strong negative correlations between diazinon levels and antecedent flow or antecedent rainfall have also been identified in previous monitoring data from Castro Valley Creek (WCC, 1996a).

Table 2.2.2 Pearson Correlation Coefficients for Diazinon with Seasonal and Hydrologic Parameters at Station S3 During 1995-1996

	<i>EMC</i>	<i>Antecedent Flow Volume</i>	<i>Prev. 30 Day Temp</i>	<i>Event Flow Volume</i>	<i>Event Mass Discharge</i>
EMC Diazinon	1.00				
2 Week Antecedent Flow Volume	-0.70	1.00			
Previous 30 Day Average Temperature	0.64	-0.54	1.00		
Event Flow Volume	-0.47	0.10	0.10	1.00	
Event Mass Discharge	0.20	-0.35	0.59	0.70	1.00

Bold indicates $p < 0.05$

A similar correlation analysis performed on the ten events sampled in 1996-97 showed no significant relationships between EMC and the other variables (i.e. correlation coefficients were all lower, with $p > 0.05$). This difference may have been partly due to the smaller samples size. (EMCs and hydrologic variables for 1996-97 events are given in Appendix B.) When the pooled 1995-97 data were analyzed, the correlation relationships of EMC to temperature and antecedent flow were similar to the 1995-96 results shown in Table 2.2.2. Spearman correlation analysis, which uses relative ranking of sample values for each variable instead of the values themselves, showed similar or higher coefficients for these relationships on both the 1995-96 and 1995-97 datasets ($p < 0.01$); good agreement between the Pearson and Spearman coefficients suggests that the data do not present any major conflicts with the statistical assumptions necessary for the Pearson analysis.

Larger storms were somewhat associated with lower EMCs: there was a weak negative correlation between EMC and the event flow volume, which was only significant ($p < 0.05$) for Spearman correlations on the two-year dataset. Part of this difference is probably due to year-to-year variation, since the 1995-96 events had a lower average event flow volume and higher average EMC than the 1996-97 events ($p \leq 0.05$, t-test). The 1996-97 storm season also began and ended earlier, with the sampled period starting on October 29 and ending January 22, so that the lack of sampled events during spring could have affected the average EMC (see-May 1997 data in Table 3.2.2).

However, when the total amount of diazinon per storm event was calculated (see Section 2.3 below), it continued to increase as event flow volume increased (Pearson and Spearman correlation coefficients ≥ 0.70 , $p < 0.01$). This suggests that additional diazinon was washed into runoff during heavy rainfall, but that it was more diluted.

These results suggest that diazinon concentrations in Castro Valley Creek were affected by both seasonal use and hydrologic factors. Multiple regression analyses were performed on the 1995-96 data to examine how combinations of variables were related to EMCs. The adjusted value of R^2 , the proportion of variation in EMC that was explained by the regression equation, was 0.44 when only antecedent flow volume was used as the independent variable, but increased for the combinations of event flow volume with log-transformed antecedent flow (adj. $R^2 = 0.65$) or event flow with previous temperature (adj. $R^2 = 0.66$). Based on these results, the timing of storm events, antecedent flow and event flow volume were all considered in developing the estimate for total mass discharge of diazinon in 1995-96 (Section 2.3).

Relationships derived from a single storm season may be a poor predictor for EMCs during other years. Figure 2.2.2 shows EMC values for 1995-96 and 1996-97 plotted as a function of antecedent flow volume. Linear and log-linear curves that are plotted in Figure 2.2.2 to show the trend of the 1995-96 data do not fit the lower EMCs found in the 1996-97 data. Year-to-year variations in use and rainfall patterns could make it difficult to detect trends in diazinon levels measured over several seasons.

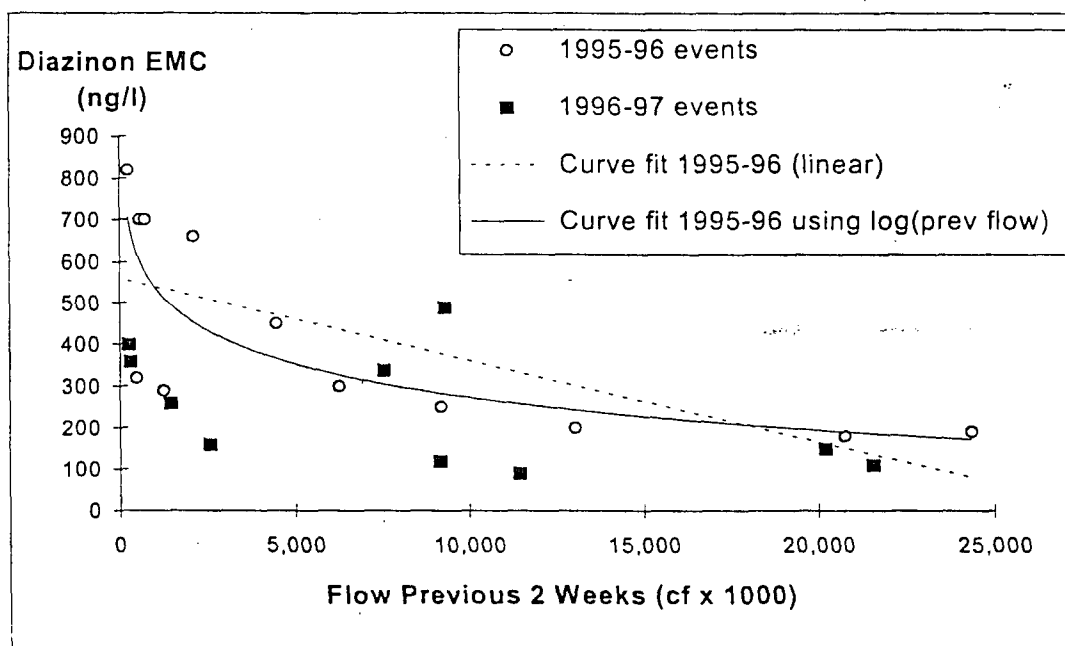


Figure 2.2.2 Diazinon Event Mean Concentration and 2-Week Antecedent Flow Volume for Two Seasons at Station S3, with Predicted Relationships Based on 1995-1996 Data

2.3 MASS DISCHARGE

An estimate of the total mass of diazinon discharged from Castro Valley Creek during the period from October 1, 1995 through September 30, 1996 was made by combining estimates for different components of the total discharge volume (Table 2.3.1). First, the EMC and event flow volume from each of the twelve sampled events were multiplied to calculate the amount discharged in the sampled storm events. Then, estimated EMCs were used to calculate the amount discharged in unsampled storm flow. For the purpose of this report, storm flow was defined as having a discharge rate greater than 5 cfs. Periods when flow was less than 5 cfs were categorized as either non-storm flow between storms, or dry weather flows outside of the rainy season. EMCs for these two categories were estimated from occasional samples taken during periods of low flow (see Section 2.5, WCC 1997b).

The total flow volume from December 1995 through May 1996 was approximately 116 million cubic feet (mcf) of which 84.2 mcf was storm flow (Figure 2.3.1). The twelve sampled events accounted for 44% of the storm flow volume. Figure 2.3.2 shows the seasonal hydrograph and the mass of diazinon discharged for each sampled storm event. The per event mass of diazinon discharged during the sampled events ranged from a low of 3.4 grams to a high of 93.2 grams (Table 2.3.2). Mass discharge was highly correlated (0.70) with total flow volume for a given event and also less strongly with season (represented by previous temperature in Table 2.2.2). The estimated total mass discharge of diazinon for these sampled events was 354 grams (12 oz.).

Table 2.3.1 Estimated Mass Discharge of Diazinon from the Castro Valley Creek Watershed from October 1, 1995 to September 30, 1996

	Flow Volume (mcf)	Average Concentration (ng/l)	Mass (grams)
Sampled Events	35	361	354
Unsampled Events	44	287	358
Non-storm Flow (Dec-May)	38	60	64
Dry Weather Flow (Oct, Nov, Jun-Sep)	3	60	5
Totals	120		781

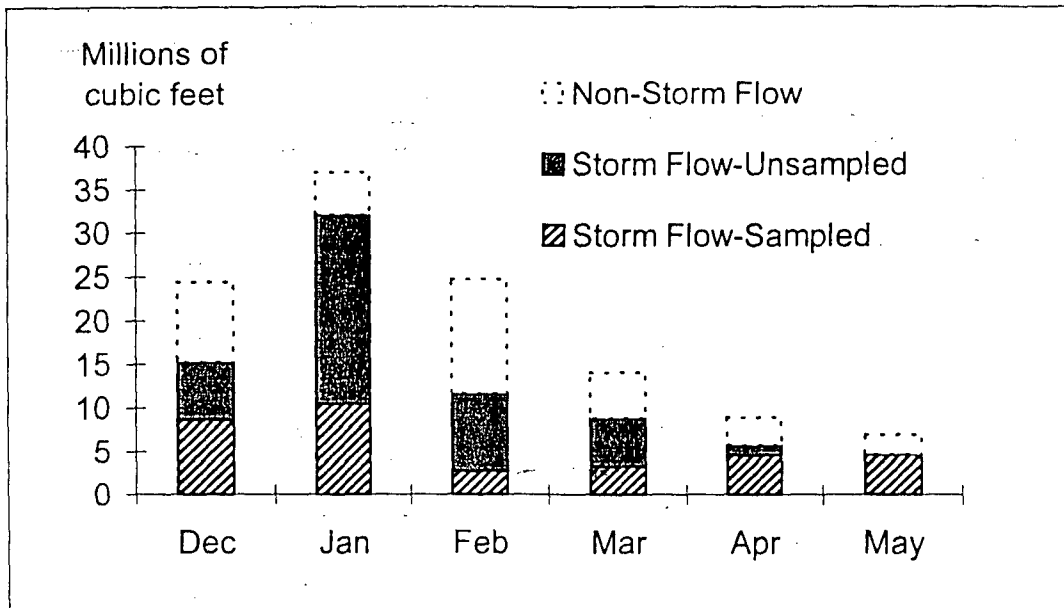


Figure 2.3.1 Monthly Flow at Station S3 between December 1, 1995 and May 21, 1996

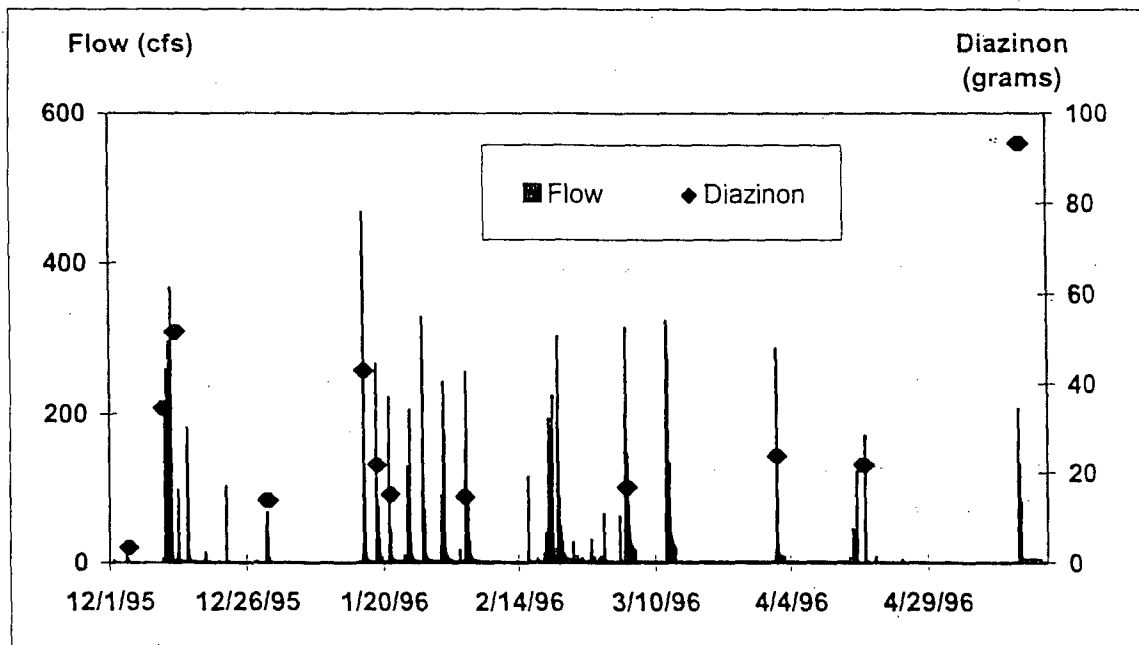


Figure 2.3.2 Estimated Diazinon Mass Discharge for Sampled Events and Stream Flow at Station S3 during 1995-1996 Storm Season

Table 2.3.2 Hydrologic Parameters and Mass Discharge of Diazinon for Sampled Events at Station S3 During 1995-1996

Event Number	Start Date	Sampling Duration (hr:min)	Flow Volume (cubic feet x 1,000)	Diazinon Event Mean Concn. (ng/l)	Diazinon Estimated Mass (grams)	Flow Volume Previous 2 weeks (cf x 1,000)
1	12/4/95	28:00	147	820	3.4	225
2	12/10/95	10:50	1,746	700	34.6	565
3	12/11/95	13:00	6,063	300	51.5	6,240
4	12/29/95	3:40	745	660	13.9	2,100
5	1/16/96	3:00	4,739	320	42.9	460
6	1/18/96	13:00	3,075	250	21.8	9,190
7	1/20/96	8:10	2,688	200	15.2	13,030
8	2/3/96	12:20	2,732	190	14.7	24,360
9	3/4/96	14:00	3,300	180	16.8	20,780
10	4/1/96	6:20	2,893	290	23.8	1,250
11	4/17/96	6:00	1,713	450	21.8	4,451
12	5/15/96	12:00	4,700	700	93.2	720
Totals			34,541		354	

Table 2.3.3 lists the event flow volume, estimated EMC, and calculated mass discharge for the 15 storm events that were not sampled. Most of the unsampled storm flow volume occurred between January 20th and March 11th. EMCs for most of the storm flow during this period were estimated at 200 ng/l based on sampled events during this period. Higher values were assumed for the storm events on February 15th and April 16th because these were small storms with low antecedent flow volumes (see Section 2.2). EMCs for the unsampled December events were estimated at 400 ng/l because of the higher EMCs during sampled December events. The average non-storm and dry weather concentration was estimated at 60 ng/l based on a limited number of samples taken (see Tables 2.5.1 and 2.5.2, WCC 1997). All mass discharge estimates were obtained by multiplying the estimated concentration by the flow volume.

Combining the mass discharge of diazinon from the 12 sampled storm events with the mass discharge from the 15 events that were not sampled, together with the diazinon in non-storm flow, resulted in an estimated mass discharge of approximately 800 grams (26 oz.). Assuming an estimated annual usage of 0.02 ~~oz.~~^{lb.} (9g) per person (Scanlin and Cooper, 1997), the total mass discharged in Castro Valley Creek was approximately 0.3 percent of the total mass applied in the watershed.

**Table 2.3.3 Stream Flow Volume and Estimated Diazinon for
Unsamped Events at Station S3 During the 1995-1996
Storm Season**

Start Date	Start Time	Event Duration (hr:min)	Flow Volume (cf x 1,000)	Assumed Diazinon Concentration (ng/l)	Estimated Diazinon Mass (grams)
12/11/95	4:40	14:00	4,182	400	47.4
12/13/95	9:50	9:00	1,697	400	19.2
12/15/95	4:12	8:44	3,012	400	34.1
12/18/95	11:10	3:42	181	400	2.1
12/22/95	6:18	3:44	850	400	9.6
12/29/95	16:00	4:00	839	400	9.5
1/16/96	8:58	11:02	3,248	280	25.8
1/18/96	13:00	1:40	595	300	5.1
1/24/96	7:38	23:04	5,048	200	28.6
1/27/96	5:10	16:52	5,308	200	30.1
1/30/96	15:20	28:40	7,441	200	42.1
2/4/96	9:10	20:50	4,558	160	20.7
2/15/96	16:52	4:20	985	600	16.7
2/18/96	16:30	89:30	2,234	200	12.7
2/23/96	23:20	2:40	274	200	1.6
2/27/96	5:30	2:40	298	200	1.7
2/28/96	17:40	10:00	251	200	1.4
2/29/96	15:00	3:00	376	180	1.9
3/3/96	12:00	6:00	347	180	1.8
3/11/96	23:00	20:00	5,281	200	29.9
4/16/96	5:00	5:00	1,138	500	16.1
4/19/96	15:50	0:50	44	200	0.2
Totals			44,005		358

2.4 WITHIN STORM VARIATION

To examine how diazinon concentrations varied during storm events, discrete samples were taken throughout nine of the sampled events at station S3. Discrete concentrations ranged from 118 to 1,832 ng/l. Figure 2.4.1 shows that the variation in diazinon concentrations appeared to follow one of two main patterns during an event. During Events 2 and 5 (Figure 2.4.1 (a) and (c)) the peak concentration occurred early and was followed by a marked decline, with peak concentrations more than 3 times the concentration at the end of the storm. During other storms such as Events 3 and 10 (Figure 2.4.1 (b) and (d)), diazinon concentrations remained relatively consistent. The high peak concentrations observed in events 2 and 5 appeared to be associated with low antecedent flow volume (Table 2.2.1). This relationship could reflect an accumulation of diazinon between storms that was readily washed off at the start of the next rainfall. However, if accumulation alone was responsible for initial peaks, they would be expected to occur only in events with high EMCs. Since different parts of the watershed contribute to different portions of the hydrograph (Baker and Richards, 1990), differences in spatial variation of diazinon could also produce different temporal patterns at Station S3 for different events.

Table 2.5.1 Diazinon Concentrations in Non-Storm Flow (<5 cfs) at Station S3 during 1995-1996 and 1996-1997 Storm Seasons

Sample Date	Time	Diazinon Concentration (ng/l)	Hours after End of Storm Flow	% of Initial Post-Storm Concentration
12/4/95	12:00	760	0	100
12/5/95	6:30	570	18.5	75
4/16/96	9:00	640	0	100
4/16/96	16:20	610	7.3	95
4/17/96	5:40	250	20.7	40
5/16/96	1:00	400	0	100
5/17/96	2:00	210	26	52
10/29/96	13:00	460	0	100
10/30/96	8:00	420	19	91
10/31/96	23:00	190	58	41
11/1/96	15:30	110	74.5	24

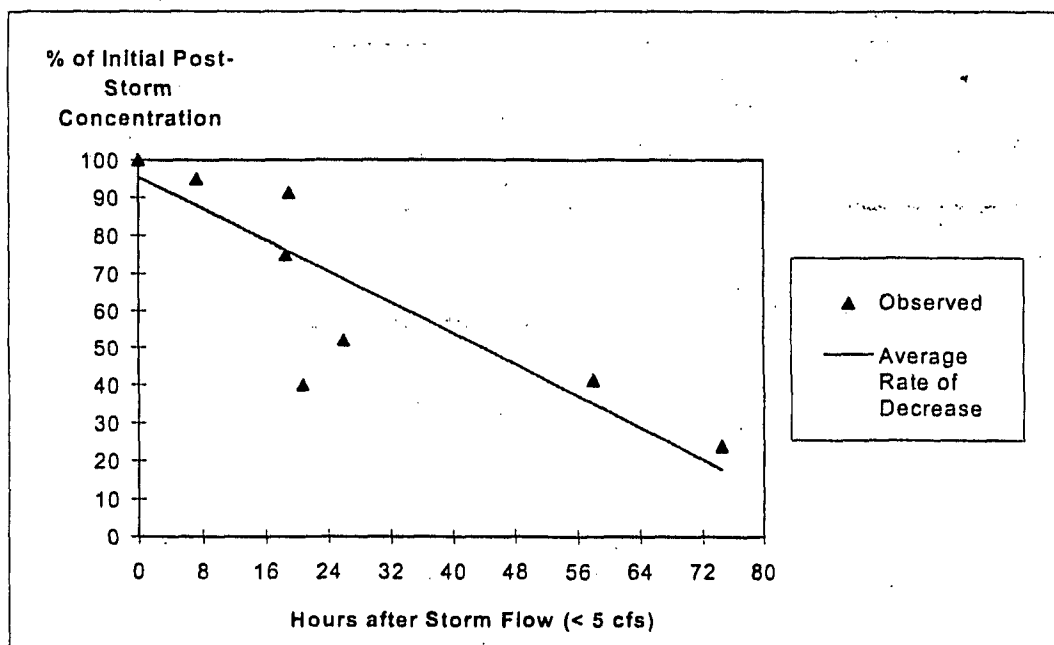
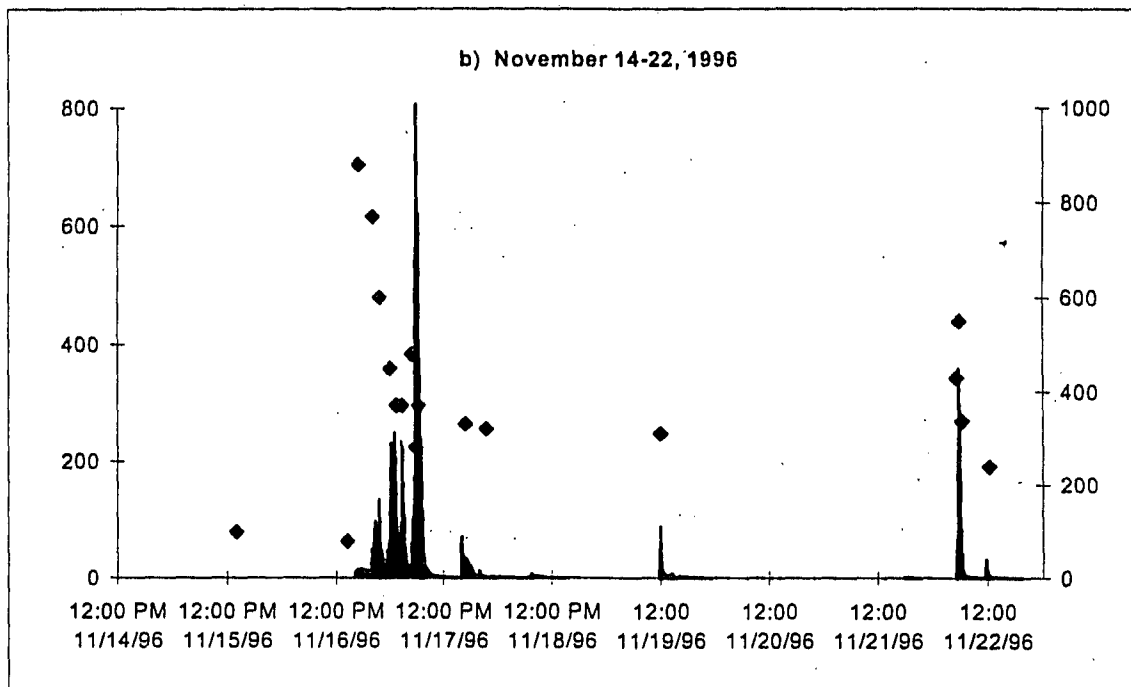
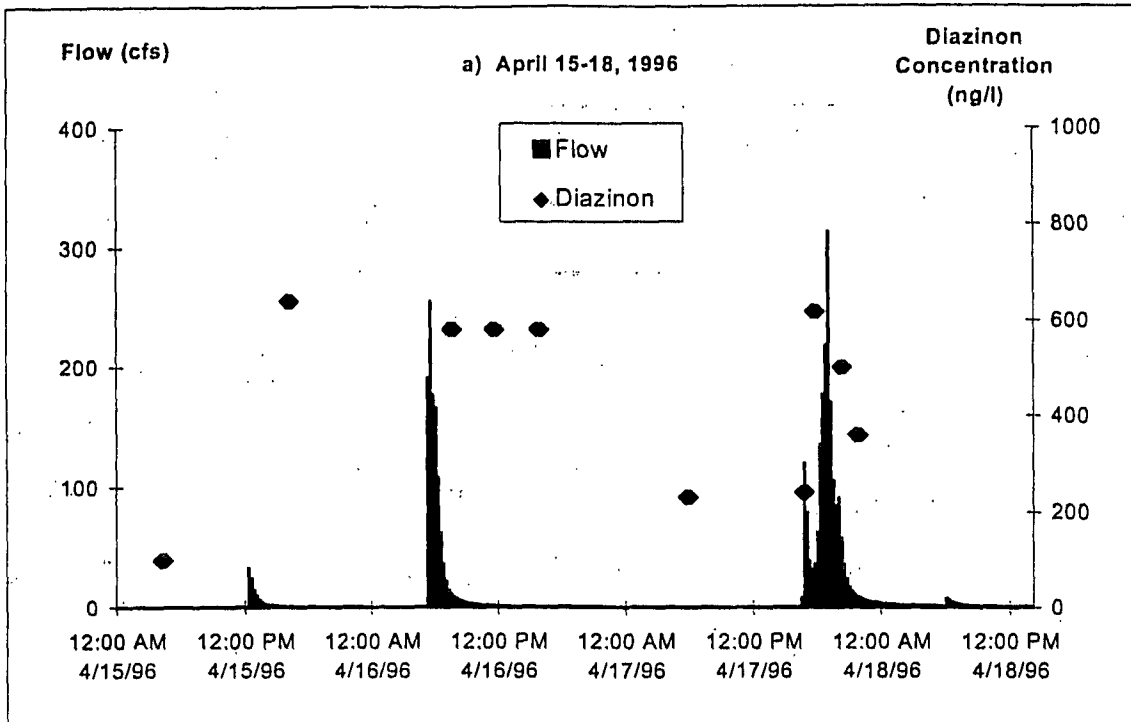


Figure 2.5.2 Proportion of Diazinon Concentration Remaining in Runoff at Varying Intervals After End of Storm Flows for Selected Storm Events at Station S3

The size of the watershed and the pattern of rainfall also affect in-stream concentrations. Since Castro Valley is a small urbanized watershed, storm flow recedes quickly following a rain event (WCC, 1995). Duration of storm flow during 1995-96 ranged from 50 minutes on April 19, 1996 to 28 hours on December 11, 1995. As seen above, concentrations declined when the storm flow receded. In a larger watershed, peak concentrations would tend to be lower while median concentrations would tend to be higher (Baker and Richards, 1990).

Previous toxicity testing of Castro Valley Creek runoff samples has shown that diazinon concentrations of 300 to 500 ng/l are lethal to *Ceriodaphnia dubia* at exposure times of 48 hours (LT_{50} , median time to lethality; Katznelson and Mumley, 1997). Thus, post-storm concentrations during December 5-6, 1995 could have remained high enough to kill 50% of *C. dubia* populations in the 48 hour period during and after the storm. Also, as can be seen in Figure 2.5.3, frequent storm events can cause concentrations to remain elevated for an extended period of time. Since diazinon concentrations of 150-300 ng/l are lethal to *C. dubia* when exposure times (LT_{50}) are 4 to 7 days (Hansen 1995, Katznelson and Mumley, 1997; see also Bailey et al., 1997), periods of intermittent storm events could also cause mortality to animals living in the creek.

Grab samples taken at S3 during longer intervals of dry weather occasionally contained diazinon concentrations as high as 220 ng/l (Table 2.5.2). During May 1997, concentrations at S3 remained above 150 ng/l long enough to be toxic to *C. dubia*. The dry weather sampling summarized in Table 2.5.2 was performed during months when storms may be expected in most years, but always took place after at least 21 days without significant rainfall in the watershed. The factors affecting dry weather concentrations of diazinon are outside the scope of this report; see WCC (1997b) for discussion.



Note: Horizontal (time) and vertical flow scales vary

Figure 2.5.3 Diazinon Concentration and Flow at Station S3 for Selected Periods of Storm Flow (<5 cfs) and Non-Storm Flow in 1996

Table 2.5.2 Diazinon Concentrations in Dry Weather (Non-Storm) Flows at Station S3 During 1996-97

Sample Date	Time	Diazinon Concentration (ng/l)
10/4/96	10:20	80
10/6/96	N.A.	75
10/8/96	12:00	50
2/3/97	10:10	35
2/21/97	10:00	80
5/15/97	10:10	220
5/17/97	10:00	200
5/19/97	11:00	170
5/21/97	10:00	180

Source: WCC (1997b)

N.A.: Sampling time not available

3.0 DISTRIBUTION OF DIAZINON SOURCES IN THE CASTRO VALLEY CREEK WATERSHED

3.1 OBJECTIVES AND METHODS

The purpose of this portion of the study was to gather data on the geographic distribution of diazinon sources in the watershed. A major objective was to determine if the sources were distributed throughout the watershed or if there were localized "hot spots." All of the samples discussed in this section of the report were "grab" samples which were collected in 20 ml glass vials. Samples were analyzed using ELISA.

3.2 SUBCATCHMENT CONCENTRATIONS

The first step in determining the distribution of diazinon sources was to collect samples from five subcatchments within the watershed (Figure 3.2.1). These subcatchments drain 78% of the watershed. Samples were not collected from the other portions of the watershed because there were no discrete discharge points from other large areas.

During a storm event on April 1, 1996, grab samples were collected every half hour at the discharge points of the subcatchments. Diazinon was detected in all of the samples collected, in concentrations ranging from 51 ng/l to 1248 ng/l. The mean subcatchment concentrations ranged from 201 ng/l to 675 ng/l (Table 3.2.1). Diazinon levels in discrete samples at S3 during this event are also shown for reference, but since there is no hydrograph data for the subcatchments, subcatchment results are not directly comparable with those from station S3.

The mean subcatchment concentrations were all similar in magnitude to the EMCs measured at S3. The area-weighted average of all subcatchments, 390 ng/l, was similar to the watershed EMC of 290 ng/l at S3. These results suggest that the sources of diazinon were distributed throughout the watershed. Much of the difference in concentrations appears to be related to differences in land use. Subcatchments with higher percentages of open space (such as parks or undeveloped land) had lower concentrations of diazinon than those with little open space. This suggests that application of diazinon is more associated with developed areas. Runoff samples from other watersheds with a higher proportion of open space than the Castro Valley watershed also tend to contain lower diazinon concentrations (WCC, 1994, 1995; Hansen 1995). Subcatchment 5, with the highest proportion of commercial development, had the second highest mean diazinon concentration. (For purposes of this report, "commercial" land use includes multi-family residential areas).

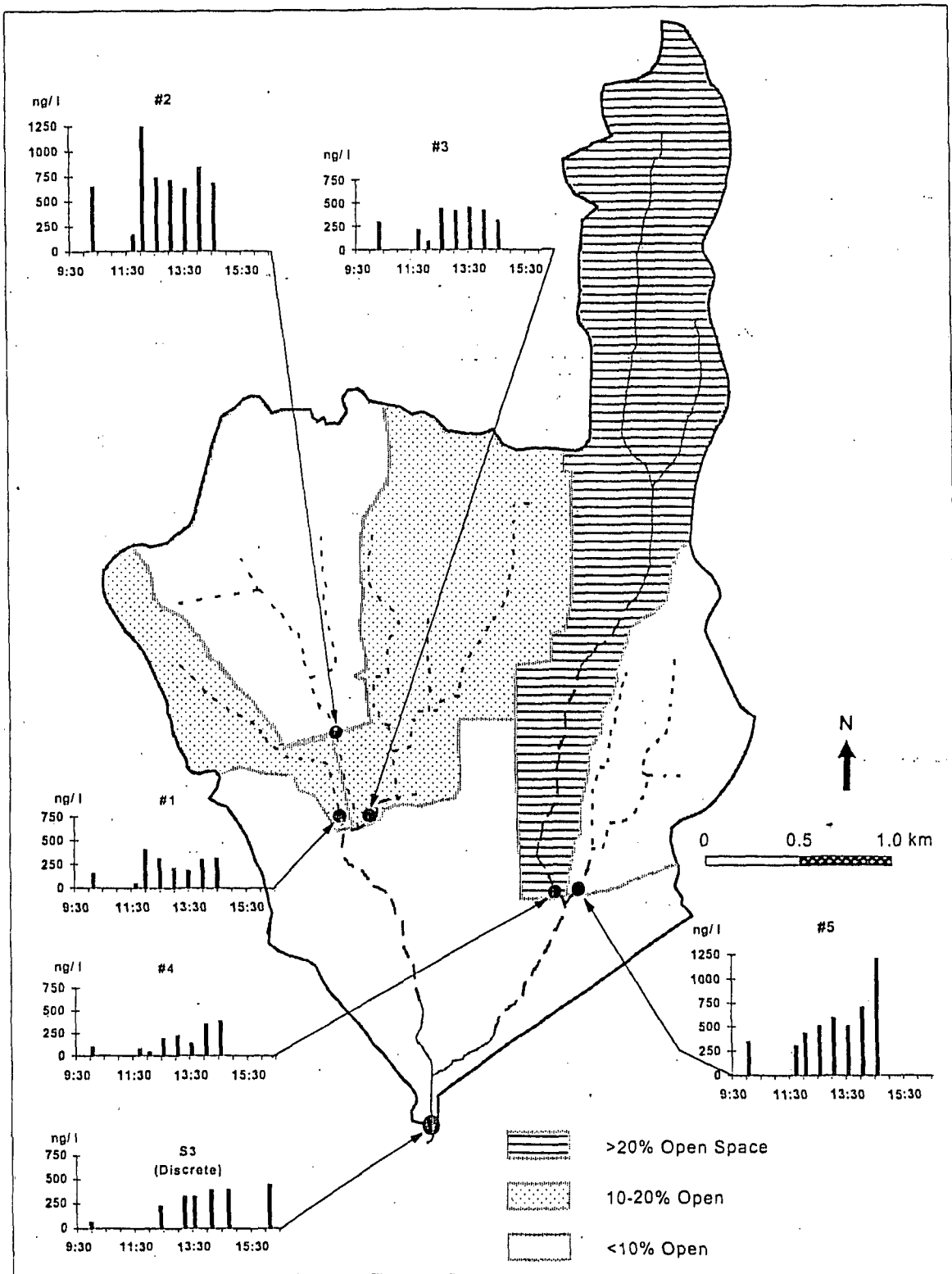


Figure 3.2.1 Diazinon Concentrations in Storm Runoff from Selected Subcatchments of the Castro Valley Creek Watershed on April 1, 1996

Table 3.2.1 Mean Diazinon Concentrations in Runoff from Subcatchments of the Castro Valley Creek Watershed on April 1, 1996

Subcatchment Number	Area (acres)	Land Use Percentage			% Pervious Area	Mean Diazinon Concentration (ng/l)
		Commercial	Open	Residential		
1	214	15	15	70	61	246
2	536	4	9	87	62	675
3	678	19	10	71	53	343
4	908	4	52	44	80	201
5	391	37	3	60	41	595

Source for subcatchment data: Alameda County Flood Control District

Table 3.2.2 show that diazinon concentrations varied among all subcatchments that were sampled during extended intervals (greater than 3 weeks) of dry weather within the 1996-97 storm season. Subcatchment 4, which had the lowest average concentrations during the May 1996 storm event, had a high concentration of 3400 ng/l in one dry weather sample in May 1997. All of the subcatchments sampled in dry weather tended to have higher diazinon concentrations in the spring. While the conditions affecting dry weather concentrations of diazinon are outside the scope of this report (see WCC 1997b for details and discussion), these results also suggest that diazinon sources are found in all subcatchments.

Table 3.2.2 Diazinon Concentrations in Dry Weather (Non-Storm) Flow from Selected Subcatchments of the Castro Valley Creek Watershed During 1996-97

Month	Date	Diazinon (ng/l) in Subcatchment:			
		1	2	3	4
October 1996	4	50	228	107	326
	6	51	38	87	135
	8	36	39	62	116
	11	ND	50	ND	57
February 1997	3	14	19	12	16
	21	ND	228	107	ND
	23	42	59	133	128
	25	ND	ND	53	47
May 1997	15	323	50	ND	ND
	17	662	2959	188	31
	19	113	ND	48	3400
	21	106	48	69	285

Source: WCC (1997b)

ND: non-detect (< 30 ng/l)

3.3 STREET GUTTER CONCENTRATIONS,

To gather data on the distribution of sources within the subcatchments, samples were collected from 45 randomly selected street gutters during a storm event on May 15, 1996. All of the samples were collected in residential areas within Subcatchments 2 and 3 (Figure 3.3.1; see also Appendix C). Each sample represented a separate local drainage area, usually consisting of between 5 and 15 residential properties.

Diazinon concentrations in street gutters were highly variable, ranging from less than 30 ng/l to over 70,000 ng/l (Figure 3.3.2). The median sample concentration was 80 ng/l, a level which has previously been found in rainfall samples from Castro Valley. The mean sample concentration was 3,900 ng/l for all samples, or 450 ng/l with the two highest concentrations excluded. Since the two samples with concentrations above 10,000 ng/l both came from small drainage areas (about 5 properties each; Jim Scanlin, pers. obs.), the high concentrations may have been due to a smaller dilution factor for diazinon coming from a single property. Therefore the area-weighted average concentration for all street gutter samples was probably less than 3,900 ng/l. For comparison, the average concentration at the discharge point for the watershed (Station S3) during this event was about 700 ng/l (Figure 2.2.1).

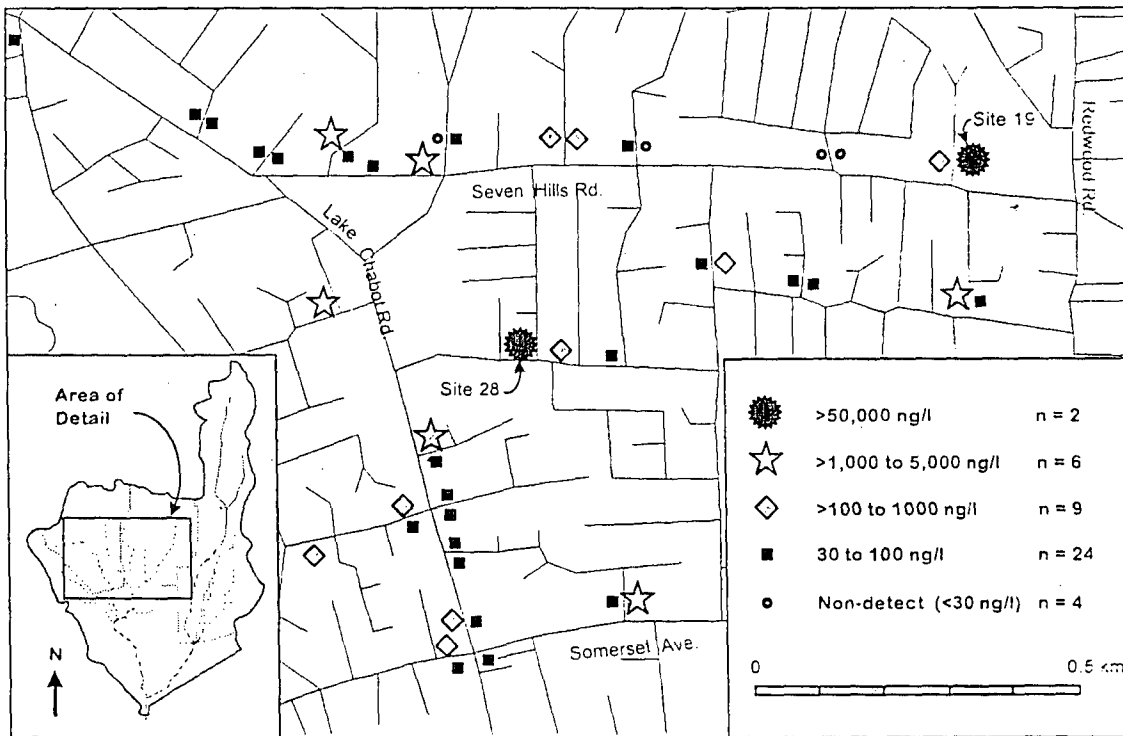


Figure 3.3.1 Diazinon Concentrations in Storm Water Runoff Collected in Street Gutters in Castro Valley on May 16, 1996

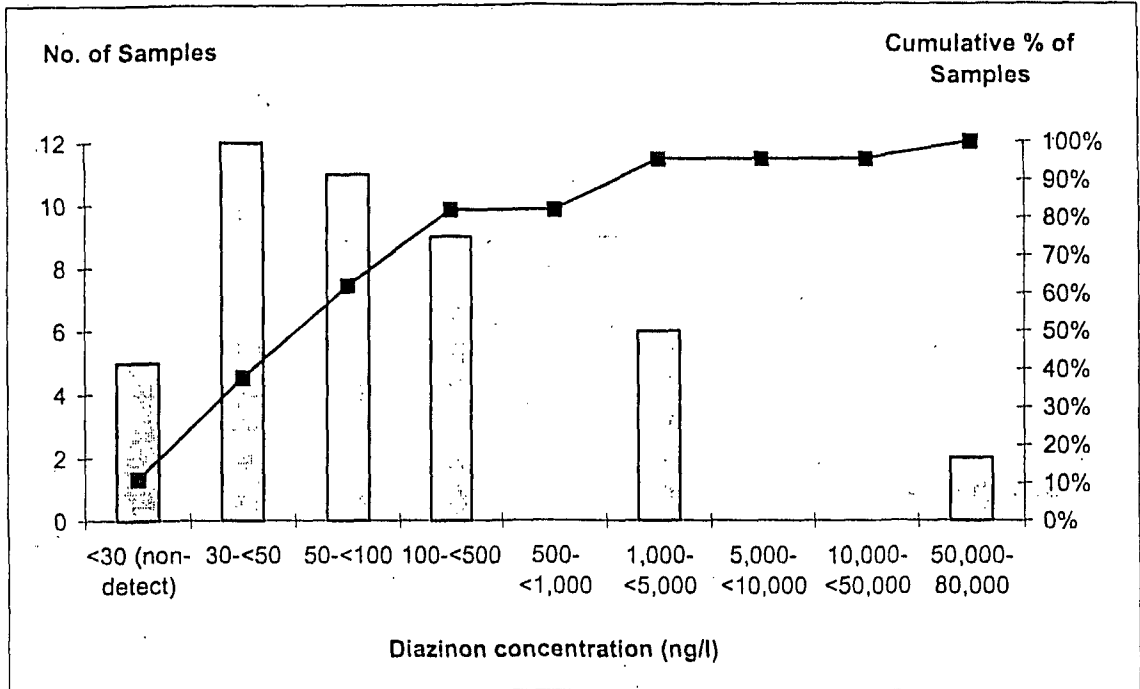


Figure 3.3.2 Distribution of Diazinon Concentrations in Selected Street Gutters in Castro Valley on May 15, 1996

Based on these results, an experimental calculation was used to form a rough estimate of 300-600 sources of diazinon in the entire watershed during the sampled event (See Appendix C for assumptions and details of the estimation). A basic assumption used for estimation was that a street gutter sample with a diazinon concentration above an arbitrary threshold level of 100-400 ng/l was considered a source. It was further assumed that when a sample met this criterion, the individual source was a single property within the area drained by that street gutter. Using an average estimate of 10 properties per street gutter sample, 2 to 4 percent of the 450 properties in the sampled areas were sources. Since most of the properties in Subcatchments 2 and 3 were single-family homes, this was roughly equivalent to 2 to 4 percent of the households. Extrapolation of these percentages to the 14,800 properties in the entire watershed suggests that 300 to 600 of these were diazinon sources for the sample event.

To determine whether the two sites with the highest concentrations were consistent "hot spots", these sites were sampled again during the fall of 1996. Samples were taken at the beginning and end of a rainfall event that occurred on November 16-17. Street gutter concentrations at Site 19 were moderately high in November but much lower than in May (Table 3.3.1). Site 28 had a very high concentration of 39,000 ng/l at the beginning of the November storm. These results suggest that there may be some consistent sources of diazinon in the watershed. Both of these "hot spots" were located in Subcatchment 3,

which had intermediate levels of diazinon during the sampling on April 1, 1996. Street gutter samples from a similar residential area draining to nearby San Leandro Creek showed similar consistency, with high concentrations persisting at some sites during several storm events (WCC, 1997a).

Table 3.3.1 Diazinon Concentrations in Selected Castro Valley Street Gutters Sampled in May and November 1996

Site	Sample Date	Sample Time	Cumulative Event Rainfall	Diazinon Concentration (ng/l)
19	5/15/96	13:54	0.1"	71,000
19	11/16/96	18:40	0.1"	314
19	11/17/96	8:00	0.4"	562
28	5/15/96	14:21	0.1"	79,000
28	11/16/96	18:50	0.1"	37,000
28	11/17/96	8:00	0.4"	2,000

4.0 RESIDENTIAL RUNOFF STUDY

4.1 OBJECTIVES AND METHODS

The purpose of this portion of the study was to determine if the use of diazinon in accordance with label directions could result in the observed diazinon concentrations in Castro Valley Creek.

Diazinon was applied to two residential properties in Castro Valley and one residential property in Oakland (located approximately 14 km northwest of the Castro Valley watershed). The diazinon was applied at the Castro Valley sites between 14:00 and 15:00 PST on February 26, 1997. The diazinon was applied at the Oakland site between 15:00 and 15:45 PST on February 28, 1997. All of the properties receiving test applications of diazinon were single-family residences on approximately one-eighth acre lots. Foundation perimeter length was approximately 120-150 feet (35-45 m) for all three residences.

The amount of diazinon applied at each property was two-thirds of the quantity recommended on the product label for the control of ants. At each site, four ounces of liquid concentrate diazinon (25% active ingredient) was mixed with two gallons of water in a hand held sprayer. (The directions called for six ounces in three gallons of water.) The solution was sprayed in a strip approximately 3 feet (1m) wide around the foundation of each house, and on the edges and in the cracks of paved surfaces of driveways, walkways and patios.

Grab samples of runoff from roofs, patios and driveways were taken at each site during subsequent rainfall events. (See Appendix D for a detailed description of sampling locations, and Appendix E for rainfall data.) None of these events produced enough runoff to permit sampling from street gutters beyond the edges of the properties. Samples were collected using 20 ml glass vials and were analyzed using ELISA.

At the Oakland site, rainfall samples were also collected several days after application. The rainfall samples were collected in a glass 9" x 13" baking pan. The pan was soaked in a 10% bleach solution then washed with soap and water and rinsed with tap water. The pan was placed on the railing of a deck 20 feet above the ground. The rain that accumulated in the pan was poured into a glass vial for cold storage prior to analysis. A similar sample was collected at the same time at an unsprayed Oakland residence located 6 km away.

4.2 RESULTS AND DISCUSSION

Diazinon was detected in all samples, which were taken up to seven weeks after application. Roof runoff samples ranged from 57 ng/l to 17,000 ng/l (Table 4.2.1). Patio and driveway runoff samples ranged from 735 ng/l to 1,200,000 ng/l. Concentrations were highest immediately after application and decreased over time. The highest on-site concentration was 15 times the highest concentration observed in street gutter samples. Since street gutter samples from Sites 19 and 28 each represented small areas of no more than five residences (James Scanlin, pers. obs.), a 33 percent dilution of the on-site sample could produce concentrations similar to the highest street gutter samples.

Table 4.2.1 Diazinon Concentrations in Storm Water Runoff at Residential Properties Following Test Spray Application on February 26-28, 1997

Site	Sample Date	Sample Time	Days Since Application	Cumulative Rainfall	Concentration ng/l
Oak. Roof Drain 1	3/2/97	2:40	1.5	0.03"	17,000
Oak. Roof Drain 1	3/2/97	7:30	1.7	0.19"	1,100
Oak. Roof Drain 1	3/15/97	19:30	15	0.24"	1,250
Oak. Roof Drain 1	3/16/97	8:55	16	0.24"	350
Oak. Roof Drain 1	3/30/97	20:05	30	0.51"	250
Oak. Roof Drain 1	4/18/97	5:30	48	0.52"	590
Oak. Roof Drain 1	4/18/97	20:00	48	0.91"	57
Oak. Drain 1 + Patio	3/2/97	2:40	1.5	0.03"	150,000
Oak. Drain 1 + Patio	3/3/97	7:30	1.7	0.19"	40,000
Oak. Drain 1 + Patio	4/18/97	20:00	48	0.91"	735
Oak. Roof Drain 2	3/16/97	10:30	16	0.26"	3,800
Oak. Roof Drain 3	3/16/97	10:30	16	0.26"	3,300
Oak. Driveway	3/16/97	10:30	16	0.26"	110,000
Oak. Patio	3/16/97	10:30	16	0.26"	880,000
CV1 Patio	3/3/97	8:30	5	0.08"	1,200,000
CV1 Patio	4/18/97	18:00	50	0.44"	3,500
CV1 Patio	4/19/97	NA	51		16,000
CV1 Driveway	4/18/97	18:00	50	0.44"	6,000
CV1 Roof Drain	4/18/97	18:00	50	0.44"	50
CV1 Roof Drain	4/19/97	NA	51		340
CV2 Patio 1	4/18/97	18:00	50	0.44"	110,000
CV2 Patio 2	4/18/97	18:00	50	0.44"	1,400
CV2 Driveway	4/18/97	18:00	50	0.44"	91,000
CV2 Roof Drain	4/18/97	18:00	50	0.44"	220

Diazinon concentrations in rainfall at the sprayed site ranged from 240 to 1,300 ng/l (Table 4.2.2). This was higher than the concentration found in many street gutter samples. In contrast, rainfall samples taken at the Oakland site prior to the application of diazinon were consistently below 100 ng/l, as was the sample from the unsprayed site. Diazinon in rainfall near treated properties may thus be a pathway for diazinon to enter storm drain runoff.

Recommended rates of application for lawns and garden areas differ from those for paved areas (six fluid ounces of liquid concentrate per 1000 square feet of lawn). A test application was also made on a lawn at a third residence in Castro Valley, but no runoff was produced from the lawn during the sampling period. This study was not able to evaluate the likelihood of applications to lawn and garden areas contributing to diazinon in runoff. Other formulations such as granules are commonly used on lawns (Scanlin and Cooper, 1997), but their potential contribution to diazinon is unknown.

Table 4.2.2 Diazinon Concentrations in Oakland Rainfall Samples, March 1997

Site and Sample Number	Sample start		Sample end		Cumulative Event Rainfall	Diazinon Concentration (ng/l)
	Date	Time	Date	Time		
Test 1	3/1/97	18:00	3/2/97	9:00	0.23"	1,300
Test 2	3/15/97	19:00	3/15/97	19:30	0.01"	240
Test 3	3/15/97	19:35	3/16/97	10:00	0.02"	930
Unsprayed	3/1/97	20:00	3/2/97	8:30	0.23"	60

4.3 CONCLUSIONS

Preliminary results indicate that use of liquid concentrate formulations of diazinon in accordance with label directions can not be ruled out as a source of the observed toxic concentrations in Castro Valley watershed. This has important implications for design of strategies to reduce diazinon concentrations in runoff (Scanlin and Gosselin, 1997).

5.0 SUMMARY AND CONCLUSIONS

Diazinon was found in storm runoff throughout the 1995-96 and 1996-97 storm seasons, in all parts of the Castro Valley Creek watershed. This study tracked temporal variations in diazinon concentrations for the aggregate runoff from the watershed, and also evaluated spatial variation at both intermediate (subcatchment) and small (residential neighborhood) scales.

Event mean concentrations for sampled 1995-96 storm events in Castro Valley Creek ranged from 180 to 820 ng/l, with peak concentrations of diazinon reaching 1800 ng/l for short periods. Low antecedent flow during the 2 weeks preceding an event was a major factor in predicting EMC, but seasonal patterns of diazinon application probably increased diazinon levels during events in fall and spring. Variations in EMC may also have been caused by differences in the hydrographs for individual events, or by year-to-year differences in diazinon application.

The total amount of diazinon discharged from Castro Valley Creek during the 1995-96 water year was estimated at about 600 grams, or about 0.3 percent of the estimated annual usage in the watershed. Approximately 90% of the mass discharge occurred during storm flows greater than 5 cfs.

Higher flows appeared to reduce EMCs, but increased the mass discharge for individual events. This suggests that some diazinon is readily washed off during small storm events, while a larger quantity is present in the watershed and enters runoff during heavier rainfall.

Diazinon concentrations over 300 ng/l sometimes persisted in Castro Valley Creek for periods of 48 hours after storm flow ended. These exposures were sufficient to be toxic to aquatic animals, and confirm the suggestions of previous studies that diazinon probably causes toxicity in Castro Valley Creek. Chronic toxic exposure to diazinon, caused by persistence of concentrations above 150 ng/l for periods up to a week long, may also have an impact on aquatic life in Castro Valley Creek.

The sources of diazinon are widespread, occurring in all subcatchments of the watershed. During the storm of April 1, 1996, runoff from more densely developed subcatchments generally contained higher average concentrations than runoff from subcatchments with more open space. Samples from street gutters were also geographically variable; a few street gutter sampling sites had diazinon levels over 2000 ng/l on different dates, but average street gutter concentrations were more similar to the concentrations observed for Station S3 and the subcatchments.

Based on street gutter results and various sets of assumptions about the number of sources within each street gutter drainage area, the estimated number of sources in the watershed is 300 to 600, or up to 4 percent of the households. Some sources may be consistent, producing high street gutter concentrations during more than one storm event.

Test applications of diazinon on exterior paving and foundation perimeter areas produced maximum runoff concentrations of 1,200,000 ng/l on the residential properties that were sprayed. This was greater than the maximum observed concentrations in street gutter samples, and over 300 times the average. These results suggest that applications of liquid diazinon according to the label directions could be responsible for much of the diazinon entering storm runoff. The potential runoff contribution of other formulations of diazinon or of applications to lawn and garden areas is unknown.

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Appendix A . ELISA Analyses for Diazinon: Quality Assurance

A.1.0 ELISA--General Description

This appendix focuses on the use of enzyme linked immunosorbent assay (ELISA) kits specific for diazinon. Most of the discussion is reproduced from WCC (1996b). ELISA techniques for other environmental contaminants, e.g., PCBs, PAHs, etc., will be referred to briefly in the following text where applicable. For in-depth understanding of the general features of immunoassays and, specifically, enzyme linked immunosorbent assays (ELISA), the reader may consult publications by ELISA kit manufacturers (Millipore, Ohmicron, or Ensys product lines) available at the ACFCWCD Water Resources Laboratory (Hayward) or by request from the manufacturers themselves.

A.1.1 WHY USE ELISA

Analytical laboratory techniques for detection of diazinon and other pesticides at low concentrations are expensive and require lengthy turnaround times. If the cost of analysis can be reduced, pesticide concentrations can be determined for more samples and the extent of spatial and temporal contamination can be better characterized. If results can be obtained more rapidly, control strategies can be implemented in a timely fashion. The development of enzyme linked immunosorbent assays (ELISAs) provides a rapid and inexpensive alternative to the traditional analytical methods used to determine pesticide concentrations.

A.1.2 SPECIFICITY OF THE IMMUNE RESPONSE

The immune response of mammals to foreign materials introduced into their bodies utilizes both cellular mechanism (for example, appearance of "killer T-cells" that recognize and kill viruses and bacteria), and humoral response (e.g., production of soluble antibody proteins that circulate in the blood). Scientists have been "raising" antibodies for decades by injecting inactivated pathogenic bacteria into laboratory or farm animals. These inactivated organisms are foreign antigens, i.e., materials that the animal has not been exposed to before and which elicit an immune response in the injected animal. Some very important human vaccines were produced that way.

Specific antibodies have been raised for research and analytical work as well. Small organic molecules such as diazinon are not good antigens and need to be attached to a bigger "carrier" in order for the immune system to respond and start producing antibodies against them. A batch of antibodies raised against a specific compound xxx is often called "anti-xxx antibody" (e.g., anti-diazinon antibody).

Polyclonal antibodies are isolated from the serum of immunized animals and consist of a mixture of antibody proteins that recognize different areas in the antigen molecule. Monoclonal antibodies are raised in tissue-cultures of antibody-producing cells that have been derived (cloned) from one cell which was isolated very early in the immune response process, when each cell "learns" to produce one type of antibody protein. Naturally, monoclonal antibodies are more specific in their recognition ability because they "see" only one component of the antigen molecule and will not recognize other components which different molecules may share with the target antigen.

Antibodies work by binding their antigens and disposing of them or making them harmless in other ways. Specificity is essential, because the body contains a host of functional molecules that should not be bound. This feature is the cornerstone of immunoassays: we are using antibodies to bind specific substances in a mixture of non-target molecules. The antigen-antibody complex is relatively stable and can be isolated from the mixture by various methods. The most common method is to immobilize the antibodies on a solid surface, expose them to the mixture and allow them to bind our target analyte, and then wash off the rest of the mixture. The diazinon ELISA uses anti-diazinon antibodies attached to the inside surfaces of plastic microwells to "pull" diazinon molecules out of a complex mixture of substances. An indirect way, utilizing the principle of competition with labeled diazinon, allows for quantification of the diazinon pulled from the mixture.

A.1.3 THE PRINCIPAL STEPS IN ELISA

In the first step, the unknown water sample suspected of containing diazinon is added to a microwell together with a known quantity of enzyme-labeled diazinon (a conjugate of diazinon and horseradish peroxidase). The ambient (free) diazinon and the added enzyme-labeled diazinon compete for the limited number of binding sites on the antibodies, and eventually bind in proportion to their relative abundance (concentration) during the incubation period.

In the second step, all unbound molecules are removed by rinsing the well thoroughly, while the bound analytes remain attached.

In the third step, a substrate for the enzyme is added at non-limiting concentration. The enzyme transforms the substrate into a colored molecule at a rate that depends on the number of enzyme molecules in this well. After a fixed period of time, the enzyme reaction is stopped by addition of acid.

In the fourth step, the color intensity is quantified by measuring the amount of light absorbed by the color molecules present in the solution in the well. If the intensity of color is high it means that there was a lot of enzyme in the well, which means that the conjugate "won" the competition, which means that there was very little or no diazinon in our ambient sample. On the other hand, if the intensity of color is low, we know that we had many molecules of diazinon in our sample.

In the fifth step, the amount of diazinon in the sample is calculated. Each ELISA "run" uses enough microwells (24 or 36) to allow comparison of color produced in wells that had environmental samples (of unknown diazinon concentrations) with color produced in wells with known concentrations of free diazinon and in wells without diazinon at all. Three concentrations of free diazinon are used for calibration of each run, and the concentrations of diazinon in all the ambient samples tested in the run are calculated from that calibration curve.

A.1.4 LABORATORY PROCEDURES

The diazinon ELISA reagent kits sold commercially include five essential components: 1) diazinon stock solution; 2) anti-diazinon antibodies (attached to wells or tubes); 3) enzyme-labeled diazinon (a conjugate of diazinon and horseradish peroxidase); 4) substrate for the enzyme, and 5) acid to stop the enzymatic reaction. For this study, EnviroGard diazinon plate ELISA kits intended for water testing were purchased from

EnSys Environmental Products (now Strategic Diagnostics Inc., Newark DE). All tests of water samples were performed in the ACFCWCD Water Resources Laboratory. The kits were used according to the manufacturer's instructions, except that the concentrations used for diazinon calibrators were altered slightly, from 30, 100 and 500 nanogram per liter (ng/l) to 25, 100 and 400 (ng/l). All measurements were made using adjustable-volume micropipettors; all dilutions were made with HPLC grade water (Fisher).

Diazinon calibrators were prepared in water by serial dilutions of the stock solution (component 1). Water samples obtained from streams or street drains were initially tested without dilution or at 25% dilution; if the initial test showed concentrations greater than the highest calibrator concentration, another test was made with a larger dilution factor. Samples and calibrators were introduced into the antibody coated wells (component 2), and immediately mixed with the diazinon-enzyme conjugate (component 3). There are 12 wells in one plastic strip.

Diazinon molecules in samples or calibrators were allowed to compete with conjugated diazinon for one hour (A1.3, first step), all the unbound molecules were washed away in cold tap water (second step), a substrate (component 4) was added and allowed to react with the enzyme for 30 minutes (third step) until the reaction was stopped with acid (component 5), and the intensity of the color in each well was recorded by measuring the absorbance of light at 450 nm (fourth step).

All data were recorded carefully throughout the entire procedure. Data for weights, volumes, dilution factors, and absorbance were subsequently entered into a spreadsheet that has been developed for this purpose (fifth step). The spreadsheet incorporates the calibrator data into a built in curve and calculates the diazinon concentration in nanogram per liter for water samples.

A.2.0 SOURCES OF UNCERTAINTY AND ERROR

Several sources of uncertainty were identified concerning the nature of diazinon in water samples, separate experiment to evaluate the effects of holding time in different containers prior to ELISA testing. Several sources of analytical error are also outlined below.

A.2.1 Sources of Uncertainty

Representativeness of the sample: Discrete or composite samples taken by the stream sampler represented known volumes of runoff water associated with specific portions of the storm hydrograph. However, samples from storm drain lines or street gutters could represent different parts of the hydrograph depending on local rainfall and drainage conditions.

Holding temperature: Samples were kept refrigerated after collection and prior to the ELISA assay. Occasionally the cooling ice in the stream sampler melted before samples could be removed, resulting in higher temperatures and potential loss of part of the diazinon.

Holding time: Samples were typically tested within 1 to 3 days. Repeated sampling from spiked samples held at 0-4°C for up to 14 days showed that none of the original diazinon was lost during the first four days, but up to 30% was lost after 14 days (Figure A.2.1).

Holding container: Compositing and subsampling of the spiked sample followed the

same procedures used in preparing composite samples. The sample was composited in a 20 liter glass carboy and pumped into containers with a peristaltic pump. Three replicates were filled for five types of containers: one liter polyethylene bottles used to collect discrete samples (PL), one liter glass bottles (GL), 20 ml glass scintillation vials (GS), 125 ml wide mouth glass jars with Teflon-lined screw caps (WGM) and 10 l plastic collapsible cubitainers (PC). Results were similar for all container types (Figure A.2.1).

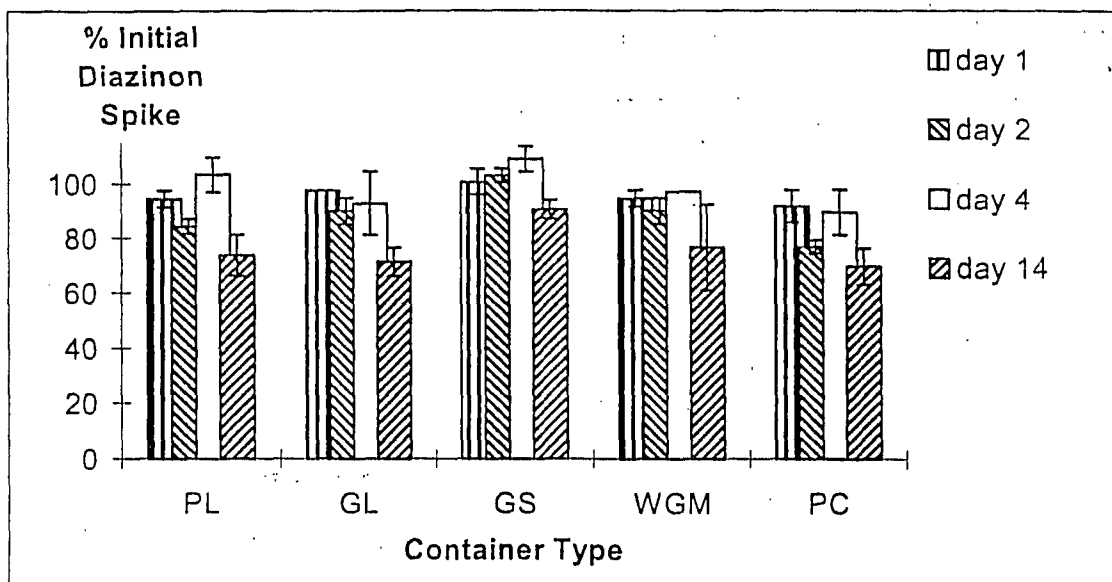


Figure A.2.1 Percentage of Original Diazinon Spike Remaining in Test Containers After Different Holding Times

Subsample representativeness: Sample containers were agitated and allowed to settle for a few minutes before aliquots were removed for assay. However, the variation among replicates of the spiked samples suggest that mixing may have been less complete in larger sample containers than in the scintillation vials. In samples with large amounts of suspended sediment, these particles may have "scavenged" some diazinon from the water, resulting in a lower concentration in the aliquots than in the sample at the time of collection.

A2.2 Sources of Error

Pipetting: Dispensing small volumes of water or samples using air displacement pipettors is a source of error. We pipetted at least 100 microliters (μ l) when air displacement pipettors were employed, and used positive displacement pipetting for smaller volumes.

Incubation time: Differences among wells are inevitable because filling the wells with samples takes part of the incubation period. To minimize the difference in incubation time, we restricted the number of wells used in a single run, put replicate calibration standards both at the beginning and the end of the run, and used the average for the calibration curve.

Absorbance reading: For the first 22 ELISA runs made during the 1995-96 runoff season, the absorbance (color density) was measured by diluting the contents of

each well in 500 µl of water in a test tube and reading it against a water blank in a photospectrometer (EnSys, now SDI, Newark DE). Samples were read in order of increasing density, or decreasing diazinon concentration. For the last 17 runs absorbances were read directly from the microwells using a strip reader (Millipore, now SDI, Newark DE). The photospectrometer method does not permit rereading of the sample and introduces an additional dilution step. It also takes more time; repeat readings made using the strip reader show that there is some loss of color during the 10-20 minutes required to read one run on the photospectrometer.

A.3.0 PERFORMANCE EVALUATION AND QA/QC RESULTS

ELISA methodology differs from laboratory analytical methods in several aspects. Table A.3.1 summarizes some of these inherent differences.

These inherent differences dictate different ways to assess the performance of the methods and to evaluate the quality and reliability of the data. However, the two methodologies share many quality assurance elements.

TABLE A.3.1 COMPARISON OF ELISA METHODOLOGY WITH CONVENTIONAL ANALYTICAL TECHNIQUES

Aspect	ELISA	Analytical method
Extraction (Water)	None	Liquid-Liquid extraction
Separation from other analytes	Specific antibody (does not separate all the analyte molecules)	Gas chromatography or Liquid chromatography
Detection	Competition for specific antibody followed by color production and absorbance measurements	Mass spectroscopy or nitrogen-phosphorous detectors
Method blank	Highest output reading	Lowest output reading

A.3.1 ELISA-Specific QA/QC Issues

Detection limit

In contrast to analytical methods, the ELISA method has no blank value (because zero diazinon results in highest color production) and the detection limit is defined by what is considered as a significant difference in absorbance between zero diazinon and the lowest concentrations of free diazinon in the calibration curve. For the work performed at the

concentrations of free diazinon in the calibration curve. For the work performed at the ACFCWCD laboratory in Hayward, a detection limit of 30 ng/l in water was considered reliable.

Cross reactivity

The specificity of an antibody is measured as cross-reactivity, or the degree to which the antibody will bind with non-target compounds. With the exception of some affinity for pirimiphos-ethyl and pirimiphos-methyl, the antibody in the diazinon kit used for this study is very specific to diazinon. Chlorpyrifos, a closely related organophosphorous insecticide, is not detected by this ELISA diazinon kit at concentrations of up to 1000 parts per billion (ppb).

Interference

Substances that affect the antibodies in a non-specific way (e.g. sediment or oil molecules that may stick to the surface of the wells and cover the antibody) will hinder attachment of diazinon conjugate even in the absence of free diazinon. This will cause a false positive result.

Confirmation requirements

ELISA reagent kits have been approved by EPA as screening tools generally used in a semi-quantitative mode (e.g., analyte concentration is greater than 1 ppm and lower than 10 ppm). In the regulatory context, ELISA results need confirmation by conventional analytical techniques for a certain percentage of samples (5-20%). Although our study was not within regulatory context, several composite samples were analyzed both by ELISA and by a sensitive LC/MS method (Quanterra Analytical, Sacramento), and a good agreement was found (see "accuracy" below).

A.3.2 General QA/QC Elements

Precision

The precision of a measurement is an expression of the degree of reproducibility of results. It can be determined by evaluating the variability among laboratory replicates and by analyzing duplicate samples. The percent coefficient of variation (%CV) calculated for each set of replicates or duplicates is a measure of precision. The % CV is the standard deviation of the diazinon concentration in replicates, divided by the mean concentration and multiplied by 100.

The precision of the diazinon ELISA kits used in this study was evaluated in two ways: "true" replicates (solution from the same sample dilution is introduced into two wells), and "dilution" replicates (solution from two different dilutions (e.g., 25% and 6.25%) of the same sample are introduced into two wells). Calibrators mixed from stock solution are "true" replicates. The kit manufacturer recommends that replicate calibrators placed in adjacent wells have a maximum %CV of 15%. Due to assay drift (see below), calibrators placed at the beginning and end of the run would be expected to vary somewhat more. Out of 35 ELISA runs containing two sets of calibrators, 29 had less than 15%CV for all four calibrator pairs; the mean %CV of the 140 calibrator pairs was 14.5%, slightly less than 15%CV (the standard deviation, or SD, was 11.5%).

Table A.3.2 summarizes the %CV results for 73 samples that had two or more replicates falling within the range of the calibration curve. While the average %CV was less than 15%, a few replicates were much more variable. There was no difference in variability between "true" and "dilution" replicates, or for within-run replicates as opposed to replicates tested in different runs. These results show that the ELISA methodology gave consistent results throughout the study period.

Table A.3.2 Variation in Diazinon ELISA Results for Replicate Runoff Samples Taken in Castro Valley and San Leandro Watersheds during 1995-1996 Storm Season

		all replicates	"true" replicates	"dilution" replicates	within same ELISA run	multiple runs
No. of replicated samples		73	16	57	53	20
Using average calibrators	mean %CV	11.68	13.81	11.08	12.57	9.32
	SD of %CV	9.89	10.96	9.58	10.23	8.70
	maximum %CV	50.46	31.03	50.46	50.46	31.03
Using different calibrator sets	mean %CV	9.56	9.46	9.59	9.83	8.84
	SD of %CV	6.97	6.81	7.07	6.98	7.07
	maximum %CV	28.99	22.32	28.99	28.99	22.32

Three of the runs showed a systematic shift in calibrator absorbances between the beginning and end calibrator sets, suggesting assay drift was occurring due to differences in incubation times within the runs. Sample values were recalculated for these runs, with beginning wells calculated from the beginning calibrator set only, the end wells from the end calibrators, and the middle wells from the average of both calibrator sets. After this recalculation, which affected seven replicate sets, the maximum %CV was reduced, but there were still some high values up to 29%CV that could be due to a variety of error sources.

Accuracy

Accuracy is the nearness of a measurement to its true value. Accuracy of the analysis is evaluated through the use of standard analyte solutions. In the ELISA runs, the diazinon stock solution provided in the ELISA reagent kit was used both for calibration and for reference to the "true" concentration. Stock dilutions from different kits and different manufacturing lots were tested together and produced similar results.

The samples analyzed by Quanterra for confirmation provide an independent measure of accuracy, because the laboratory uses a different standard analyte solution. Table A.3.3 summarizes the diazinon data obtained by the analytical laboratory and by ELISA for the same composite samples. Here, the relative percent difference (RPD) is the difference between the laboratory and ELISA results divided by the average of the two results and multiplied by 100. Unlike the %CV, the RPD may have either a negative or positive

value, depending which of the two results was used first; this is useful in detecting a trend of bias caused by the method. The average %CV and the RPD are greater than that used in the laboratory's quality assurance program, and also greater than for replicates within ELISA runs. However, the 40% average of the absolute values for RPD is reasonable agreement for tests from different laboratories using different analysis methods.

Table A.3.3 Comparison of ELISA Results and Laboratory Analysis for Diazinon in Composite Runoff Samples

sample date	Diazinon concentration, ng/l		Holding time, days		RPD	%CV
	Lab	ELISA	Lab	ELISA		
12/11/95	320	276	14	2	14.6	10.3
1/16/96	205	386	23	1	-61.1	43.2
1/16/96	220	367	23	1	-50.0	35.4
1/19/96	170	254	20	2	-39.6	28.0
1/18/96	140	240	21	2	-52.8	37.3
3/4/96	280	156	14	1	56.9	40.2
3/4/96	120	89	14	1	29.4	20.8
4/1/96	280	342	16	2	-19.9	14.1
4/1/96	370	267	16	2	32.3	22.8
Average:					-10.0	28.0
Average of absolute RPD:					39.6	

Laboratory samples were typically held for longer periods of time before analysis. Figure A.3.1 plots the RPD against the difference in holding time for the two types of analysis; the RPD shows a bias that is positive for samples held less than three weeks but negative when laboratory holding times were longer. Laboratory samples that were tested more than three weeks after sample collection showed consistently lower results than the ELISA tests that were performed within 48 hours, as would be expected from the results of holding time tests (Section A.2.1). However, laboratory analyses made between 2 and 3 weeks after collection produced consistently higher results than the ELISA. Based on these results, laboratory tests that were made within the same time frame as the ELISA would probably show higher RPDs for all samples.

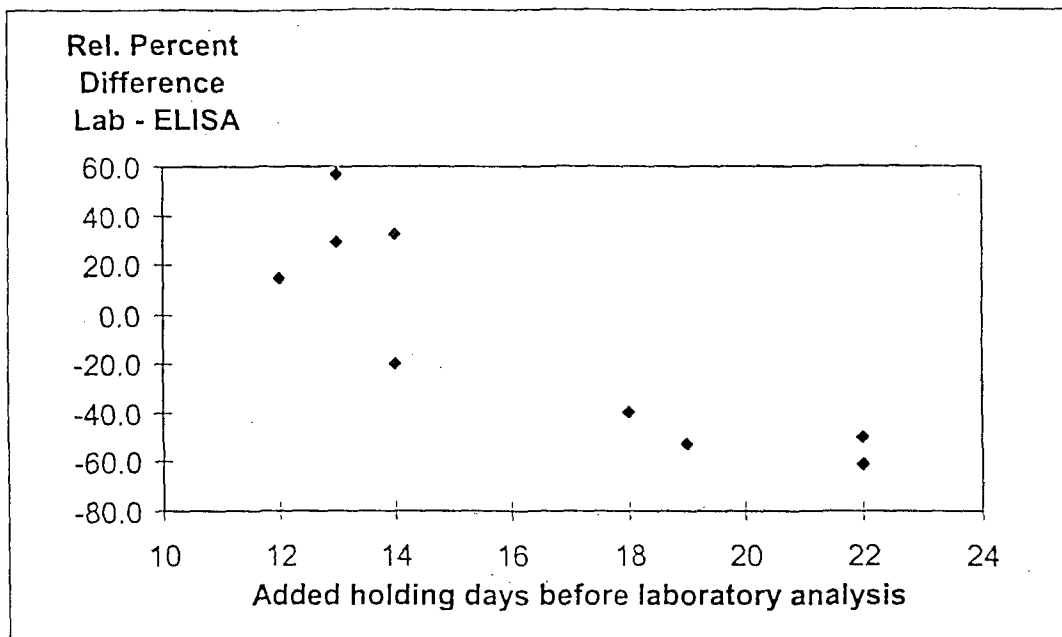


Figure A.3.1 Relative Percent Difference for ELISA Results and Laboratory Analyses Made After Different Sample Holding Times

Appendix B. Sample Results and Hydrological Data From Station S3

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- Table B.2 Sample Data and Diazinon Event Mean Concentration Calculations for Station S3 During 1996-97 Storm Season**
- Table B.3 Flow Summaries and Calculations for 1996-97 Storm Season**
- Table B.4 Diazinon in Storm Runoff Samples Taken from Castro Valley During 1994-95 Storm Season**
- Table B.5 Pesticide Test Results for Composites Samples Taken at Station S3 on 10/29/96 and 12/11/96**

Table B.1 Sample Data and Diazinon Event Mean Concentration Calculations for Station S3 During 1995-96 Storm Season

1995-96 Station S3 SAMPLE ID (1)	Sample date	time	Initial Sample (ppt)	Notes (2)	Beginning Calibrator (3)	End Calibrator (3)	Multiplier Factor (4)	Event Mean Concentration Used
cv56d1#1(25%)	12/4/95	237	758		585			
cv56d1#2(25%)	12/4/95	249	874		689		2	
cv56d1#4(25%)	12/4/95	316	1008		811		2	
cv56d1#6(6.25%)	12/4/95	353	844			1192		
cv56d1#6(25%)	12/4/95	353	1339		1124		2	
cv56d1#8(6.25%)	12/4/95	455	635			927		
cv56d1#8(25%)	12/4/95	455	874		689		2	
cv56d1#9(6.25%)	12/4/95	543	551			817		
cv56d1#9(25%)	12/4/95	543	874		689			
cv56d1#10(6.25%)	12/4/95	658	414			635		
cv56d1#10(25%)	12/4/95	658	874		689	1051	2	
cv56d1#12(25%)	12/4/95	816	874		689	1051	2	
cv56d1#14(25%)	12/4/95	941	874		689	1051	2	
cv56d1#16(25%)	12/4/95	1120	874		689	1051	2	
cv56d1#17(25%)	12/4/95	1212	758			927	2	
cv56d1#20(25%)	12/5/95	625	571			720	2	
Discrete #1	12/4/95	2:37-6:23	34/50 = 823					820
cv56d2#1(12.5%)	12/10/95	1754	241			303		
cv56d2#1(50%)	12/10/95	1754	241		191		1	
cv56d2#2(50%)	12/10/95	1943	1106	r, e-	877		1	
cv56d2#3(12.5%)	12/11/95	236	1459			1838	1	
cv56d2#3(25%)- retest12/29	12/11/95	236	782	h18				
cv56d2#4(50%)	12/11/95	241	1106	r, e-	877		1	
cv56d2#5(50%)	12/11/95	245	838	r, e-	665		3	
cv56d2#8(12.5%)	12/11/95	258	838			1056		
cv56d2#8(25%)- retest12/29	12/11/95	258	516	h18				
cv56d2#8(50%)	12/11/95	258	729		579		5	
cv56d2#15(12.5%)	12/11/95	330	635			800	4	
cv56d2#15(50%)	12/11/95	330	481		382	606	4	
cv56d2#18(50%)	12/11/95	346	553		439	696	4	
cv56d2#23(12.5%)	12/11/95	431	635			800		
cv56d2#23(50%)	12/11/95	431	419		332	528	3	
Discrete #2	12/10/95	17:54- 4:31(12/1 1)	16,870/24 = 703					700
cv56C#1(50%)	12/11/95	18:48- 7:73(12/1 2)	276		219	348		
Quanterra LC/MS	12/11/95	18:48- 7:73(12/1 2)	320	14 days				
Composite #1	12/11/95	18:48- 7:73(12/1 2)						300

1995-96 Station S3 SAMPLE ID (1)	Sample date	time	Initial Sample (ppt)	Notes (2)	Beginning Calibrator (3)	End Calibrator (3)	Multiplier Factor (4)	Event Mean Concentration Used
cv56d3#3(25%)	12/29/95	1218	681			800		
cv56d3#3(25%)	12/29/95	1218	1031		877			
cv56d3#4(25%)	12/29/95	1234	1031		877			
cv56d3#5(25%)	12/29/95	1242	898		764		2	
cv56d3#7(25%)	12/29/95	1258	898		764			
cv56d3#8(25%)	12/29/95	1308	898		764		2	
cv56d3#10(25%)	12/29/95	1328	681		579		2	
cv56d3#12(25%)	12/29/95	1345	898		764	1056	2	
cv56d3#14(25%)	12/29/95	1348	592		504	696	2	
cv56d3#17(25%)	12/29/95	1520	516		439	606	2	
cv56d3#19(25%)	12/29/95	1528	516		439	606	2	
cv56d3#21(25%)	12/29/95	1535	449			528	2	
cv56d3#24(25%)	12/29/95	1549	449			528	2	
Discrete #3	12/29/95	12:18- 15:49	13,916/21=663					660
cv56d4#1(20%)	1/16/96	609	771				1	
cv56d4#3(20%)	1/16/96	626	324				3	
cv56d4#7(20%)	1/16/96	656	193				3	
cv56d4#9(20%)	1/16/96	713	193				4	
cv56d4#13(20%)	1/16/96	742	193				4	
cv56d4#17(20%)	1/16/96	803	193				5	
cv56d4#23(20%)	1/16/96	847	229				3	
Discrete #4	1/16/96	5:50 - 8:35						250
Composite #2	1/16/96	5:50 - 8:35	386					390
Quanterra LC/MS	1/16/96	5:50 - 8:35	190	22 days				
Quanterra Dup.	1/16/96	5:50 - 8:35	220	22days				320
cv56d5#2(20%)	1/18/96	1509	379	sc				
cv56d5#6(20%)	1/18/96	1702	218	sc				
cv56d5#8(20%)	1/18/96	1832	144	sc				
S3C38(25%)	1/18/96	14:40- 3:40(1/19)	267					
S3C38(100%)	1/18/96	14:40- 3:40(1/19)	240					
Composite #3	1/18/96	14:40-3:40(1/19)						250
Quanterra LC/MS	1/18/96	14:40-3:40(1/19)		1709 days				

1995-96 Station S3 SAMPLE ID (1)	Sample date	time	Initial Sample (ppt)	Notes (2)	Beginning Calibrator (3)	End Calibrator (3)	Multiplier Factor (4)	Event Mean Concentration Used
cv56g6S3(25%)	1/20/96	~14:00	67					
cv56d6#1(25%)		2222	131					
cv56d6#3(25%)	1/20/96	2235	201					
cv56d6#4(25%)	1/20/96	2240	267					
cv56d6#6(25%)	1/20/96	2258	267					
cv56d6#8(25%)	1/20/96	2322	201					
cv56d6#10(25%)	1/21/96	7	201					
cv56d6#12(25%)	1/21/96	135	232					
cv56d6#13(25%)	1/21/96	208	175					
cv56d6#15(25%)	1/21/96	605	151					
cv56d6#16(100%)	1/21/96	624	118					
cv56d6#16(25%)	1/21/96	624	131					
Discrete #6	1/20/96	22:20-6:38	205					200

1995-96 Station S3 SAMPLE ID (1)	Sample date	time	Initial Sample (ppt)	Notes (2)	Beginning Calibrator (3)	End Calibrator (3)	Multiplier Factor (4)	Event Mean Concentration Used
cv56d7#4(25%)	2/3/96	2100	98		81	115		
cv56d7#4	2/3/96	2100	105		95			
cv56d7#5(25%)	2/4/96	410	420	r, e+	380	459		
cv56d7#5	2/4/96	410	292		279		1	
cv56d7#7(25%)	2/4/96	425	314			348	1	
cv56d7#7	2/4/96	425	252		239		1	
cv56d7#9(25%)	2/4/96	504	175			200		
cv56d7#9	2/4/96	504	163		151		2	
cv56d7#13(25%)	2/4/96	619	175			200		
cv56d7#13	2/4/96	619	163		151		1	
cv56d7#15(25%)	2/4/96	632	175			200		
cv56d7#15	2/4/96	632	188		176		2	
cv56d7#20(25%)	2/4/96	709	113			132		
cv56d7#20	2/4/96	709	105		95	115	2	
cv56d7#24(25%)	2/4/96	902	131			152		
cv56d7#24	2/4/96	902	121		111	132	1	
Discrete #7	2/4/96	4:10-9:02	2041/11=185.5					190
cv56g8S3(25%)	2/15/96	1655	539		471			
cv56g8S3(6.25%)	2/15/96	1655	497		434	570		

cv56g9S3#1 (25%)	3/4/96	1400	228					
cv56g9S3#2 (25%)	3/4/96	1406	228					
cv56g9S3c39	3/4/96	8:10- 22:10	185					
cv56g9S3c39 (25%)	3/4/96	8:10- 22:10	127					
Quanterra LC/MS	3/4/96	8:10- 22:10	280	14 days				
Composite #4	3/4/96	8:10- 22:10						180

CV56D10#1	4/1/96	956	73		54	82	1	
CV56D10#3 (25%)	4/1/96	1220	235		202	269	1	
CV56D10#4 (25%)	4/1/96	1232	235		202	269	1	
CV56D10#7 (25%)	4/1/96	1310	310			351	1	
CV56D10#9 (25%)	4/1/96	1332	310			351	2	
CV56D10#10 (25%)	4/1/96	1338	270			307	1	
CV56D10#14 (25%)	4/1/96	1407	356			400	1	
CV56D10#16 (25%)	4/1/96	1440	356			400	1	
CV56D10#19 (25%)	4/1/96	1608	409			456	1	
Discrete #10		9:50- 16:10						310
CV56g10S3gMar31	4/1/96	-14:00	44		42			
CV56g10S3C	4/1/96	8:50- 17:00	288		289			290
CV56g10S3C (25%)	4/1/96	8:50- 17:00	396		382			
Quanterra LC/MS	4/1/96	8:50- 17:00	280	16 days				

1995-96 Station S3 SAMPLE ID (1)	Sample date	time	Initial Sample (ppt)	Notes (2)	Beginning Calibrator (3)	End Calibrator (3)	Multiplier Factor (4)	Event Mean Concentration Used
CV56D11#1 (6.25%)	4/15/96	1723	538		481	600		
CV56D11#1 (25%)	4/15/96	1723	681		628	738	1	
CV56D11#2 (6.25%)	4/16/96	904	473			528		
CV56D11#2 (25%)	4/16/96	904	642			696	1	
CV56D11#3 (6.25%)	4/16/96	1619	647		550			
CV56D11#3 (25%)	4/16/96	1619	692		609		1	
CV56D11#4 (25%)	4/17/96	543	190		246	221		
CV56D11#4 (25%)	4/17/96	543	286		221			
CV56D11#4	4/17/96	543	284		254		1	
CV56D11#5 (25%)	4/17/96	1645	232		119			
CV56D11#5	4/17/96	1645	252		224	282	1	
CV56D11#6 (25%)	4/17/96	1654	511		447	581	1	
CV56D11#6	4/17/96	1654	390	rc, e-	350	432		
CV56D11#8 (25%)	4/17/96	1717	617		542	699	1	
CV56D11#10 (25%)	4/17/96	1810	468			533	1	
CV56D11#10	4/17/96	1810	396	rc, e-		438		
CV56D11#13 (25%)	4/17/96	1831	433			494	1	
CV56D11#13	4/17/96	1831	382	rc, e-		423		
CV56D11#1B (25%)	4/17/96	1950	403				0.5	
CV56D11#1B (25%)	4/17/96	1950	507				0.5	
CV56D11#2B (25%)	4/17/96	2006	511				1	
CV56D11#3B (25%)	4/17/96	2024	365				1	
CV56D11#4B (25%)	4/17/96	2119	360				1	
CV56D11#6B	4/18/96		271				1	
CV56D11#6B (25%)	4/18/96		257					
Estimated EMC			450	*Estimated				450

cv56d12# 1 (25%)	5/15/96		957		909			
cv56d12# 2 (25%)	5/15/96	12:35	898			947	1	
cv56d12# 2 (25%)	5/15/96	12:35	992		941			
cv56d12# 3 (25%)	5/15/96	12:38	1307		1235		1	
cv56d12# 4 (25%)	5/15/96	12:41	1189		1125		2	
cv56d12# 7 (25%)	5/15/96	12:49	1099		1041		3	
cv56d12# 9 (25%)	5/15/96	12:53	1175		1112		2	
cv56d12# 11 (25%)	5/15/96	12:57	1069		1013	1129	2	
cv56d12# 15 (25%)	5/15/96	13:04	938		891	989	4	
cv56d12# 18 (25%)	5/15/96	13:09	758			797	4	
cv56d12# 21 (25%)	5/15/96	13:15	801			843	2	
cv56d12# 24 (25%)	5/15/96	13:22	805			846	2	
cv56d12# 1B (25%)	5/16/96	15:06	881				5	
cv56d12# 2B (25%)	5/16/96	15:26	753				2.5	
cv56d12# 3B (25%)	5/16/96	15:53	785				2.5	
cv56d12# 4B (25%)	5/16/96	16:16	759				2.5	
cv56d12# 5B (25%)	5/16/96	16:42	704				2.5	
cv56d12# 6B (25%)	5/16/96	18:00	532				2.5	
cv56d12# 6B (25%)	5/16/96	18:00	656				2.5	
cv56d12# 7B (25%)	5/16/96	18:17	565				2.5	
cv56d12# 8B (25%)	5/16/96	18:34	556				2.5	
cv56d12# 9B (25%)	5/16/96	18:50	559				2.5	
cv56d12# 10B (25%)	5/16/96	20:04	552				10	
cv56d12# 11B (25%)	5/16/96	22:55	413				10	
cv56d12# 12B (25%)	5/16/96	0:44	402				10	
CV56d12# 13B	5/17/96	1:57	305	c1	270			
CV56d12# 13B (25%)	5/17/96	1:57	256	c1	209			
Discrete # 12								700

Footnotes to Table B.1

- (1) Sample type codes: cv56dxx discrete, flow-weighted sample
 cv56cxx composited from discrete samples
 cv56gxx grab sample
- (2) Codes in "Notes": Estimate Peak flow was not sampled. Concentrations estimated from samples surrounding peak.
 d below detection limit = 25 ppt
 r above range of standard curve = 400 ppt (derived)
 rc %BO beyond range for lowest of 400 calibrators
 e- probably underestimates diazinon concentration
 e+ probably overestimates diazinon concentration
 c1,c2,c3 one or more calibrators exceed 15%CV between replicates
 sc single set of calibrators in this run--no replicates
 cx first set of calibrators from previous run, not on same strips
 cy last set of calibrators from following run, not on same strips
 h18 holding time = 18 days before this run, some diazinon has broken down
 ~time time of sampling is approximate
- (3) Beginning/End Calibrator: Alternative values for ELISA results using single set of calibrators closest to each sample within the run, instead of average of both sets of calibrators
- (4) Multiplier Factor: Concentrations of analyzed samples assigned to adjacent unanalyzed samples

Notes on interpretation of ELISA results:

- 1 15%CV = maximum suggested allowance for 2 replicate samples, equivalent to:

ppt	at 25 ppt
ppt	at 100 ppt
ppt	at 250 ppt
ppt	at 400 ppt

When the difference between replicates is less than the above, either or both may be reasonable estimates.
 Multiply these numbers by dilution factor for approx. range of variation in the concentration of original sample
- 2 Linear standard curve approximates actual absorbance curve, but tends to underestimate concentration at 25 and 400 ppt, and overestimate at 100 ppt. This bias may be 10-15% of the sample value, depending on run.
 "e-" and "e+" notes are shown when the best estimate in bold face is probably biased as noted.
 "rc" note indicates actual sample concentration is probably over 400 ppt
- 3 "d" or "r" values have been deleted unless no other value is available

Table B.2 Sample Data and Diazinon Event Mean Concentration Calculations for Station S3 During 1996-97 Storm Season

1996-97station S3 SAMPLE ID (1)	Sample date	time	Initial Sample (ppt)	Notes (2)	Beginning Calibrator (3)	End Calibrator (3)	Multiplier Factor (4)	Event Mean Concentration Used
cv67d1#1 (25%)	10/29/96	305	113				1	
cv67d1#2 (25%)	10/29/96	316	336					
cv67d1#2 (25%)	10/29/96	316	246					
cv67d1#2 (25%)	10/29/96	316	256				1	
cv67d1#3 (25%)	10/29/96	327	381				1	
cv67d1#4 (25%)	10/29/96	334	435					
cv67d1#4 (25%)	10/29/96	334	466				1	
cv67d1#4 (25%)	10/29/96	334	472					
cv67d1#4 (25%)	10/29/96	334	485					
cv67d1#5 (25%)	10/29/96	339	429					
cv67d1#5 (25%)	10/29/96	339	460				1	
cv67d1#5 (25%)	10/29/96	339	541	c1				
cv67d1#6 (25%)	10/29/96	343	300				1	
cv67d1#7 (25%)	10/29/96	347	370				1	
cv67d1#8 (25%)	10/29/96	351	372				1	
cv67d1#9 (25%)	10/29/96	355	379				0.5	
cv67d1#9 (25%)	10/29/96	355	429				0.5	
cv67d1#10 (25%)	10/29/96	359	501	c1				
cv67d1#10 (25%)	10/29/96	359	512				1	
cv67d1#10 (25%)	10/29/96	359	528					
cv67d1#11 (25%)	10/29/96	403	457	sc				
cv67d1#12 (25%)	10/29/96	408	415					
cv67d1#12 (25%)	10/29/96	408	442				1	
cv67d1#12 (25%)	10/29/96	408	493					
cv67d1#12 (25%)	10/29/96	408	529					
cv67d1#13 (25%)	10/29/96	414	455					
cv67d1#13 (25%)	10/29/96	414	485				1	
cv67d1#13 (25%)	10/29/96	414	543					
cv67d1#13 (6.25%)	10/29/96	414	465					
cv67d1#13 (6.25%)	10/29/96	414	485					
cv67d1#13 (6.25%)	10/29/96	414	503					
cv67d1#14 (25%)	10/29/96	422	353					
cv67d1#14 (25%)	10/29/96	422	396					
cv67d1#14 (25%)	10/29/96	422	424				1	
cv67d1#14 (25%)	10/29/96	422	470					
cv67d1#15 (25%)	10/29/96	435	379				1	
cv67d1#16 (25%)	10/29/96	454	336				0.5	
cv67d1#16 (25%)	10/29/96	454	492				0.5	
cv67d1#17 (25%)	10/29/96	510	472					
cv67d1#17 (25%)	10/29/96	510	495				1	
cv67d1#17 (25%)	10/29/96	510	540	sc				
cv67d1#18 (25%)	10/29/96	521	194					
cv67d1#18 (25%)	10/29/96	521	342					
cv67d1#18 (25%)	10/29/96	521	365				1	
cv67d1#18 (25%)	10/29/96	521	416					
cv67d1#18 (25%)	10/29/96	521	509					

1996-97station S3 SAMPLE ID (1)	Sample date	time	Initial Sample (ppt)	Notes (2)	Beginning Calibrator (3)	End Calibrator (3)	Multiplier Factor (4)	Event Mean Concentration Used
cv67d1#20 (25%)	10/29/96	538	290				1	
cv67d1#21 (25%)	10/29/96	549	306				1	
cv67d1#22 (25%)	10/29/96	610	216				1	
cv67d1#23 (25%)	10/29/96	635	431	c1			1	
cv67d1#24 (25%)	10/29/96	653	279				1	
Discrete #1							375	
cv67d1#1B (25%)	10/29/96	1251	460					
cv67d1#2B	10/30/96	751	358					
cv67d1#2B (25%)	10/30/96	751	479					
cv67d1#3B	10/31/96	2257	206					
cv67d1#3B	10/31/96	2257	175					
cv67c1S3A (25%)	10/29/96		293					
cv67c1S3A (25%)	10/29/96		313				0.333	
cv67c1S3A (25%)	10/29/96		326					
cv67c1S3B (25%)	10/29/96		306					
cv67c1S3B (25%)	10/29/96		359				0.333	
cv67c1S3B (25%)	10/29/96		553					
cv67c1S3C (25%)	10/29/96		396					
cv67c1S3C (25%)	10/29/96		405				0.333	
cv67c1S3C (25%)	10/29/96		411					
Event 1 composite							359	360
cv67d1Grab	11/1/96	1530	87					
cv67d1Grab	11/1/96	1530	131					
cv67d1Grab (25%)	11/1/96	1530	104					
cv67d2#1 (25%)	11/16/96	2010	770				1	
cv67d2#1 (25%)	11/16/96	2010	934					
cv67d2#2 (25%)	11/16/96	2101	818				1	
cv67d2#3 (25%)	11/16/96	2149	532				0.5	
cv67d2#3 (25%)	11/16/96	2149	608				0.5	
cv67d2#3 (25%)	11/16/96	2149	706					
cv67d2#3 (25%)	11/16/96	2149	812					
cv67d2#5 (25%)	11/17/96	12	413				1	
cv67d2#5 (25%)	11/17/96	12	496				1	
cv67d2#7 (25%)	11/17/96	50	392				1	
cv67d2#7 (25%)	11/17/96	50	409				1	
cv67d2#9 (25%)	11/17/96	126	364				1	
cv67d2#9 (25%)	11/17/96	126	374				1	
cv67d2#11 (25%)	11/17/96	244	368				1	
cv67d2#11 (25%)	11/17/96	244	375				2	
cv67d2#11 (25%)	11/17/96	244	513					
cv67d2#11 (25%)	11/17/96	244	614					

1996-97 station S3 SAMPLE ID (1)	Sample date	time	Initial Sample (ppt)	Notes (2)	Beginning Calibrator (3)	End Calibrator (3)	Multiplier Factor (4)	Event Mean Concentration Used
cv67d2#14 (25%)	11/17/96	501	402				1	
cv67d2#16 (25%)	11/17/96	550	274				1	
cv67d2#16 (25%)	11/17/96	550	292				1	
cv67d2#18 (25%)	11/17/96	601	216				1	
cv67d2#18 (25%)	11/17/96	601	245				1	
cv67d2#20	11/17/96	611	332				2	
cv67d2#22	11/17/96	626	370				2	
Discrete #2							406	
cv67C2S3 (25%)	11/17/96		380					
cv67C2S3 (25%)	11/17/96		473	c1				
Event 2 Composite							380	400
cv67d2#1B	11/17/96	1650	331				1	
cv67d2#2B	11/17/96	2035	320				1	
cv67d2#3B	11/19/96	1203	323				0.5	
cv67d2#3B	11/19/96	1203	302				0.5	
cv67d2#4B (25%)	11/22/96	517	414				0.5	
cv67d2#4B (25%)	11/22/96	517	456				0.5	
cv67d2#5B (25%)	11/22/96	546	529				0.5	
cv67d2#5B (25%)	11/22/96	546	595				0.5	
cv67d2#6B (25%)	11/22/96	557	204				0.5	
cv67d2#6B (25%)	11/22/96	557	229				0.5	
cv67d2#7B (25%)	11/22/96	613	336				1	
cv67d2#8B	11/22/96	1215	271					
cv67d2#8B (25%)	11/22/96	1215	206					
cv67d2#8B (25%)	11/22/96	1215	234				1	
Discrete Event 2A							344	340
cv67d2gS3-1	11/15/96	1400	90					
cv67d2gS3-1	11/15/96	1400	111					
cv67d2gS3-2	11/16/96	1515	78					
cv67d2gS3-2	11/16/96	1515	93					
cv67d2gS3-3 (25%)	11/16/96	1700	876					
cv67d2gS3-4	11/22/96	1235	45					
cv67d2gS3-5	11/25/96	1245	54					
cv67d2gS3-6	12/2/96	1300	211					
cv67d2#9B	12/1/96	410	327					
cv67d2#9B	12/1/96	410	295					
cv67d2#10B	12/3/96	1848	380					
cv67d2#10B	12/3/96	1848	437	r				

1996-97station S3 SAMPLE ID (1)	Sample date	time	Initial Sample (ppt)	Notes (2)	Beginning Calibrator (3)	End Calibrator (3)	Multiplier Factor (4)	Event Mean Concentration Used
cv67d3#1	12/4/96	2209	441	r				
cv67d3#1 (25%)	12/4/96	2209	351				3	
cv67d3#5 (25%)	12/5/96	4	305				1	
cv67d3#5 (25%)	12/5/96	4	261				1	
cv67d3#7 (25%)	12/5/96	109	292				1	
cv67d3#7 (25%)	12/5/96	109	196	sc			1	
cv67d3#9 (25%)	12/5/96	231	217	sc			2	
cv67d3#11 (25%)	12/5/96	316	208				3.5	
cv67d3#15 (25%)	12/5/96	420	362				4	
cv67d3#19 (25%)	12/5/96	556	163				3.5	
cv67d3#22 (25%)	12/5/96	750	228				2	

Discrete #3

261 260

cv67d3#1B	12/6/96	2	252					
cv67d3gS3-1	12/4/96	1340	228	sc				
cv67d3gS3-1	12/4/96	1340	75					
cv67d3gS3-1 (25%)	12/4/96	1340	83	sc,d				
cv67d3gS3-2	12/6/96	1440	153	sc				
cv67d3gS3-2 (25%)	12/6/96	1440	117	sc				
cv67d4#2 (25%)	12/6/96	15:44	249		197			
cv67d4#2 (25%)	12/6/96	15:44	131			168		

cv67d4#3 (25%)	12/9/96	8:25	213		166		2	
cv67d4#3 (25%)	12/9/96	8:25	134			171		
cv67d4#7 (25%)	12/10/96	4:34	245		193			
cv67d4#7 (25%)	12/10/96	4:34	136			174	4	
cv67d4#11 (25%)	12/10/96	6:02	165		127		3	
cv67d4#15 (25%)	12/10/96	7:04	172		132		2.5	
cv67d4#15 (25%)	12/10/96	7:04	141			179	2.5	
cv67d4#19 (25%)	12/10/96	8:39	121		91			
cv67d4#19 (25%)	12/10/96	8:39	118			152	4	
cv67d4#22 (25%)	12/10/96	9:57	113			145	2	
cv67d4#23	12/10/96	10:56	137		114		1	
cv67d4#23 (25%)	12/10/96	10:56	127		96			
cv67d4#23 (25%)	12/10/96	10:56	161			204		

Discrete #4

123 120

cv67c4S3 (25%) 12/10/96 8:05 167

Event 4 Composite (includes Dec 6 sample)

170

1996-97station S3 SAMPLE ID (1)	Sample date	time	Initial Sample (ppt)	Notes (2)	Beginning Calibrator (3)	End Calibrator (3)	Multiplier Factor (4)	Event Mean Concentration Used
cv67c5S3	12/23/96		380	c1,sc- 400,rc				
cv67c5S3 (25%)	12/23/96		560	c1,sc-400				
cv67c5S3 (25%)	12/23/96		488					
Event 5 composite								490
cv67c6S3-1	12/30/96		100					
cv67c6S3-1 (25%)	12/30/96		123					
cv67c6S3-1 (25%)	12/30/96		106					
cv67c6S3-1	12/30/96		141					
cv67c6S3-1	12/30/96		149					
Event 6 Composite								150
cv67c6S3-2	12/31/96		79					
cv67c6S3-2	12/31/96		155					
cv67c6S3-2 (25%)	12/31/96		110					
cv67c6S3-2 (25%)	12/31/96		114					
cv67c6S3-3	1/1/97		67					
cv67c6S3-3	1/1/97		110					
cv67c6S3-3	1/1/97		117					
cv67c6S3-3 (25%)	1/1/97		97	d				
cv67c6S3-3 (25%)	1/1/97		98	d				
Event 6A composite								110
cv67g6S3	1/2/97	13:45	60					
cv67g6S3	1/2/97	13:45	68					
cv67g6S3 (25%)	1/2/97	13:45	47	d				
CV67C7 S3-1 (100%)	1/23/97	-	157					
CV67C7 S3-1 (25%)	1/23/97		115					
Event 7 Composite								160
CV67C7 S3-2 (100%)	1/26/97		88					
Event 7A Composite								90

Footnotes to Table B.2

- (1) Sample type codes: cv56dxx discrete, flow-weighted sample
 cv56cxx composited from discrete samples
 cv56gxx grab sample
- (2) Codes in "Notes": Estimate Peak flow was not sampled. Concentrations estimated from samples surrounding peak.
 d below detection limit = 25 ppt
 r above range of standard curve = 400 ppt (derived)
 rc %BO beyond range for lowest of 400 calibrators
 e- probably underestimates diazinon concentration
 e+ probably overestimates diazinon concentration
- c1,c2,c3 one or more calibrators exceed 15%CV between replicates
 sc single set of calibrators in this run--no replicates
 cx first set of calibrators from previous run, not on same strips
 cy last set of calibrators from following run, not on same strips
- h18 holding time = 18 days before this run, some diazinon has broken down
 -time time of sampling is approximate
- (3) Beginning/End Calibrator: Alternative values for ELISA results using single set of calibrators closest to each sample within the run, instead of average of both sets of calibrators
- (4) Multiplier Factor: Concentrations of analyzed samples assigned to adjacent unanalyzed samples

Notes on interpretation of ELISA results:

- 1 15%CV = maximum suggested allowance for 2 replicate samples, equivalent to:-

ppt	at 25 ppt
ppt	at 100 ppt
ppt	at 250 ppt
ppt	at 400 ppt

When the difference between replicates is less than the above, either or both may be reasonable estimates.
 Multiply these numbers by dilution factor for approx. range of variation in the concentration of original sample
- 2 Linear standard curve approximates actual absorbance curve, but tends to underestimate concentration at 25 and 400 ppt, and overestimate at 100 ppt. This bias may be 10-15% of the sample value, depending on run.
 "e-" and "e+" notes are shown when the best estimate in bold face is probably biased as noted.
 "rc" note indicates actual sample concentration is probably over 400 ppt
- 3 "d" or "r" values have been deleted unless no other value is available

Table B.3 Hydrologic (Stream Flow) Parameters and Mean Diazinon Concentrations for Sampled Events at Station S3 in Castro Valley Creek During the 1996-1997 Storm Season.

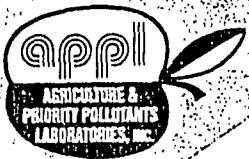
Event Number	Start Date	Start Time for Sampling	Sampling Duration (hr:min)	Flow Volume (cubic feet x 1,000)	Diazinon Event Mean Concn. (ng/l)	Flow Volume Previous 2 weeks (cf x 1,000)	Avg. Temp. Previous 30 Days (°F) (1)
1	10/29/96	2:50	4:00	524	360	282	60.9
2	11/16/96	16:10	14:20	5,209	400	248	55.2
2A	11/22/96	5:00	7:10	920	340	7,521	55.3
3	12/4/96	20:20	11:30	6,879	260	1,431	54.1
4	12/6/96	12:00 (2)	27:40 (2)	6,345 (3)	120	9,152	54.3
5	12/21/96	1:30 (2)	5:50 (2)	10,920 (3)	490	9,304	53.1
6	12/29/96	10:30 (2)	20:45 (2)	1,006	150	20,199	52.2
6A	1/1/97	2:40 (2)	13:50 (2)	9,581	110	21,568	53.1
7	1/21/97	12:00 (2)	21:50 (2)	6,207	160	2540	49.7
7A	1/21/97	12:00 (2)	32:10 (2)	9,977	90	11,437	49.9

Notes:

- (1) Source: NOAA daily temperature data for Oakland Airport
- (2) estimated starting time and duration of sampling for composited samples
- (3) estimated from flow data at 10 minute intervals instead of 2 minutes

Table B.4 Diazinon in Storm Runoff Samples Taken from Castro Valley During 1994-95 Storm Season

SAMPLE ID	Sampling date	Sampling time	Diazinon (ppt)
cv45c1S3	11/5/94	1320	533
cv45c2S3	11/25/94		460
cv45c3S3	1/3/95	1030	186
cv45c4S3	1/8/95	2300	92
cv45g1Bee	2/28/95	1225	3260
cv45g1Con	2/28/95	1245	2499
cv45g1Hey	2/28/95	1300	1081
cv45g1CVcrk-Knox	2/28/95	1210	100
cv45g1S3	2/28/95	1200	65
cv45g2CVcrk-Knox	3/9/95	1105	205
cv45g2S3	3/9/95	1050	190
cv45g3Con	4/7/95	1230	556
cv45g3Chabcrk-Knox	4/7/95	1215	285
cv45g3Hey	4/7/95	1245	395
cv45g3CVcrk-Knox	4/7/95	1210	377
cv45g3Seav	4/7/95	1300	105
cv45g3Grab1	4/7/95	1125	<30
cv45g3Grab2	4/7/95	1135	99
cv45g3Grab3	4/7/95	1145	535
cv45g3Grab4	4/7/95	1155	207
cv45g3Grab5	4/7/95	1205	173
cv45g3Grab6	4/7/95	1250	397
cv45g4Con	4/15/95	1420	272
cv45g4CVBvd	4/15/95	1520	292
cv45g4Hey	4/15/95	1510	1352
cv45g4Seav	4/15/95	1500	291
cv45d1#1	4/15/95	1410	33
cv45d1#2	4/15/95	1430	<30
cv45d1#3	4/15/95	1440	<30
cv45d1#4	4/15/95	1450	58
cv45d1#5	4/15/95	1510	236
cv45d1#6	4/15/95	1530	289
cv45d1#7	4/15/95	1550	233
cv45d1#8	4/15/95	1600	252
cv45d1#9	4/15/95	1620	266
cv45d1#10	4/15/95	1640	284
cv45d1#11	4/15/95	1750	319



Woodward-Clyde Consultants
500 12th Street, Suite 100
Oakland, CA 94607-4014
Attn: Revital Katznelson

Sample Date: 12/10/96
Report Date: 12/27/96

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Sample I.D. No: 941202NC/C9943
S3
APPL Sample No: R24179-45962W

Date Received: 12/12/96
Date Extracted: 12/13/96

Method 8140 Results (OP Pesticides):

<u>Compound</u>	<u>Concentration $\mu\text{g/L}$</u>	<u>Quantitation Limit $\mu\text{g/L}$</u>
Azinphosmethyl	ND*	1.0
Bolstar	ND	0.10
Chlorpyrifos	0.03 J	0.05
Coumaphos	ND	0.10
Def	ND	0.10
Demeton-s	ND	0.20
Diazinon	0.16	0.05
Dichlorvos	ND	0.20
Dimethoate	ND	0.10
Disulfoton	ND	0.10
EPN	ND	0.10
EPTC	ND	0.10
Ethion	ND	0.10
Ethoprop	ND	0.10
Fensulfothion	ND	0.50
Fenthion	ND	0.10
Malathion	ND	0.10
Merphos	ND	0.10
Mevinphos	ND	0.70
Naled	ND	0.50
Parathion, ethyl	ND	0.10
Parathion, methyl	ND	0.10
Phorate	ND	0.10
Prowl	ND	0.10
Ronnel	ND	0.10
Stirophos	ND	0.10
Tokuthion	ND	0.10
Trichloronate	ND	0.10
Trifluralin	ND	0.10

Non Target

Prometon	0.12	0.10
Simazine	0.69	0.50

J = Estimated value, below quantitation limit.

ND = None Detected

Tested By Paula Young
Checked By Misha [Signature]

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500 12th Street, Suite 100
Oakland, CA 94607-4014
Attn: Revital Katznelson

Sample Date: 12/10/96
Report Date: 12/27/96

Page 2 of 2

Sample I.D. No: 941202NC/C9943
S3
APPL Sample No: R24179-45962W

Date Received: 12/12/96
Date Extracted: 12/13/96

Method 8321 Results (Carbamates):

<u>Compound</u>	<u>Concentration $\mu\text{g/L}$</u>	<u>Quantitation Limit $\mu\text{g/L}$</u>
Aminocarb	ND*	0.4
Aldicarb	ND	0.4
Barban	ND	3.5
Benomyl (Carbendazim)	ND	0.4
Bromacil	ND	0.4
Carbaryl	ND	0.07
Carbofuran	ND	0.07
Chloroprotham	ND	3.5
Chloroxuron	ND	0.4
Diuron	1.4	0.4
Fenuron	ND	0.4
Fluometuron	ND	0.4
Linuron	ND	0.4
Methomyl	ND	0.07
Methiocarb	ND	0.4
Mexacarbate	ND	3.5
Monuron	ND	0.4
Neburon	ND	0.4
Oxamyl	ND	0.4
Propachlor	ND	3.5
Protham	ND	3.5
Propoxur	ND	0.4
Siduron	ND	0.4
Tebuthiuron	ND	0.4

ND = None Detected

Tested By *Brad Anderson*⁷⁴
Checked By *Lance Cass*



Woodward-Clyde Consultants
500 12th Street, Suite 100
Oakland, CA 94607-4014
Attn: Revital Katznelson

Sample Date: NA
Report Date: 12/27/96

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Sample I.D. No: 941202NC/C9943
Blank for samples
taken 12/10/96

Date Received: NA

APPL Sample No: R24179-961213W

Date Extracted: 12/13/96

Method 8140 Results (OP Pesticides):

<u>Compound</u>	<u>Concentration $\mu\text{g/L}$</u>	<u>Quantitation Limit $\mu\text{g/L}$</u>
Azinphosmethyl	ND*	1.0
Bolstar	ND	0.10
Chlorpyrifos	ND	0.05
Coumaphos	ND	0.10
Def	ND	0.10
Demeton-s	ND	0.20
Diazinon	ND	0.05
Dichlorvos	ND	0.20
Dimethoate	ND	0.10
Disulfoton	ND	0.10
EPN	ND	0.10
EPTC	ND	0.10
Ethion	ND	0.10
Ethoprop	ND	0.10
Fensulfothion	ND	0.50
Fenthion	ND	0.10
Malathion	ND	0.10
Merphos	ND	0.10
Mevinphos	ND	0.70
Naled	ND	0.50
Parathion, ethyl	ND	0.10
Parathion, methyl	ND	0.10
Phorate	ND	0.10
Prowl	ND	0.10
Ronnel	ND	0.10
Stirophos	ND	0.10
Tokuthion	ND	0.10
Trichloronate	ND	0.10
Trifluralin	ND	0.10
Non Target		
Prometon	ND	0.10
Simazine	ND	0.50

* ND = None Detected

Checked By Samuel Cooper

Woodward-Clyde Consultants
500 12th Street, Suite 100
Oakland, CA 94607-4014
Attn: Revital Katznelson

Sample Date: NA
Report Date: 12/27/96

Page 3 of 4

Sample I.D. No: 941202NC/C9943
Blank for samples
taken 12/10/96
APPL Sample No: R24179-961213W

Date Received: NA
Date Extracted: 12/13/96

Method 8321 Results (Carbamates):

<u>Compound</u>	<u>Concentration $\mu\text{g/L}$</u>	<u>Quantitation Limit $\mu\text{g/L}$</u>
Aminocarb	ND*	0.4
Aldicarb	ND	0.4
Barban	ND	3.5
Benomyl (Carbendazim)	ND	0.4
Bromacil	ND	0.4
Carbaryl	ND	0.07
Carbofuran	ND	0.07
Chloroprotham	ND	3.5
Chloroxuron	ND	0.4
Diuron	ND	0.4
Fenuron	ND	0.4
Fluometuron	ND	0.4
Linuron	ND	0.4
Methomyl	ND	0.07
Methiocarb	ND	0.4
Mexacarbate	ND	3.5
Monuron	ND	0.4
Neburon	ND	0.4
Oxamyl	ND	0.4
Propachlor	ND	3.5
Protham	ND	3.5
Propoxur	ND	0.4
Siduron	ND	0.4
Tebuthiuron	ND	0.4

* ND = None Detected

Checked By Lance Coxe

Woodward-Clyde Consultants
500 12th Street, Suite 100
Oakland, CA 94607-4014
Attn: Revital Katznelson

Report Date: 12/27/96

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Project ID No: 941202NC/C9943

APPL Spike ID: R24179 45962W 961213WA

Concentration Units: $\mu\text{g/L}$

SPIKES

Method	Analysis	Date	Amt in Sample	Amt Spiked	Results	Percent Recovery	RPD
8140	Diazinon	12/13/96	0.00	0.333	0.284	85.3	18
8140	Ethion	12/13/96	0.00	0.333	0.346	104	16

APPL Spike ID: R24179 45962W 961213WB

Method	Analysis	Date	Amt in Sample	Amt Spiked	Results	Percent Recovery	RPD
8140	Diazinon	12/13/96	0.00	0.333	0.237	71.2	18
8140	Ethion	12/13/96	0.00	0.333	0.294	88.3	16

Comments:

Checked By Jamie Coax

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500 12th Street, Suite 100
Oakland, CA 94607-4014
Attn: Revital Katznelson

Report Date: 12/27/96

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Project ID No: 941202NC/C9943

APPL Spike ID: R24179 45962W 961213WA

Concentration Units: $\mu\text{g/L}$

SPIKES

Method	Analysis	Date	Amt in Sample	Amt Spiked	Results	Percent Recovery	RPD
8321	Methomyl	12/13/96	0.00	0.267	0.197	73.9	1.01
8321	Carbofuran	12/13/96	0.00	0.267	0.204	76.5	6.18
8321	Linuron	12/13/96	0.00	0.533	0.439	82.3	0.46

APPL Spike ID: R24179 45962W 961213WB

Method	Analysis	Date	Amt in Sample	Amt Spiked	Results	Percent Recovery	RPD
8321	Methomyl	12/13/96	0.00	0.267	0.199	74.6	1.01
8321	Carbofuran	12/13/96	0.00	0.267	0.217	81.4	6.18
8321	Linuron	12/13/96	0.00	0.533	0.437	81.9	0.46

Comments:

Checked By Lamar Coxe

Woodward-Clyde Consultants

500 12th Street, Suite 100, Oakland, CA 94607-4014
(510) 893-3600

Chain of Custody Record

Appl Lab

PROJECT NO. <i>941020 NC/C9943</i>			ANALYSES					Number of Containers	REMARKS (Sample preservation, handling procedures, etc.)		
SAMPLERS: (Signature)			Sample Matrix (Soil, Water, Air)	EPA Method	EPA Method	EPA Method	EPA Method				
DATE	TIME	SAMPLE NUMBER									
<i>12-10</i>	<i>11:00</i>	<i>S3</i>	<i>W</i>	<i>X</i>	<i>X</i>			<i>3</i>	<p><i>OPs and Carbamates at the lowest DL available</i></p> <p><i>Please send invoice and report to R. Katznelson, WCC (address above) (510) 874 3048</i></p>		
TOTAL NUMBER OF CONTAINERS								<i>3</i>			
RELINQUISHED BY (Signature) <i>[Signature]</i>	DATE/TIME <i>12-10-96 11:00</i>	RECEIVED BY (Signature) <i>Fed Express</i>	RELINQUISHED BY (Signature) <i>Fed Express</i>	DATE/TIME	RECEIVED BY (Signature)	METHOD OF SHIPMENT:		SHIPPED BY (Signature)	COURIER (Signature)	RECEIVED FOR LAB BY (Signature) <i>[Signature]</i>	DATE/TIME <i>12/11/96 11:15</i>



Woodward-Clyde Consultants
500 12th Street, Suite 100
Oakland, CA 94607-4014
Attn: R. Katznelson

Sample Date: 10/29/96
Report Date: 11/19/96

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Sample ID No : 941020NA/C9943
CV67C153

Date Received: 10/31/96
Date Extracted: 11/01/96

APPL Sample No: R23944-44765

Results of OCl Pesticides Water Analysis by EPA Method 8080

Compound	Units	Concentration	Quantitation
			Limit
Aldrin	µg/L	0.014 J	0.025
a-BHC	µg/L	ND	0.025
b-BHC	µg/L	ND	0.025
d-BHC	µg/L	0.011 J	0.025
γ-BHC (Lindane)	µg/L	ND	0.025
Chlordane	µg/L	0.017 J	0.025
Dieldrin	µg/L	ND	0.025
Endosulfan I	µg/L	ND	0.025
Endosulfan II	µg/L	ND	0.025
Endosulfan sulfate	µg/L	ND	0.025
Endrin	µg/L	ND	0.025
Endrin aldehyde	µg/L	ND	0.025
Endrin ketone	µg/L	ND	0.025
Heptachlor	µg/L	ND	0.025
Heptachlor epoxide	µg/L	ND	0.025
4,4'-DDE	µg/L	0.022 J	0.025
4,4'-DDT	µg/L	0.032	0.025
4,4'-TDE/DDD	µg/L	0.014 J	0.025
Methoxychlor	µg/L	ND	0.025
Toxaphene	µg/L	ND	1.00

AMENDED

J = Estimated value below quantitation limit
ND = None Detected

Tested by: Paula Young

Checked by: Mick De

Woodward-Clyde Consultants
500 12th Street, Suite 100
Oakland, CA 94607-4014
Attn: R. Katznelson

Sample Date: 10/29/96
Report Date: 11/19/96

Page 2 of 3

Sample ID No : 941020NA/C9943
CV67C153
APPL Sample No: R23944-44765

Date Received: 10/31/96
Date Extracted: 11/01/96

Results of OP Pesticides Water Analysis by EPA Method 8140

Compound	Units	Concentration	Quantitation Limit
Azinphosmethyl	µg/L	ND	1.0
Bolstar	µg/L	ND	0.10
Chlorpyrifos	µg/L	0.13	0.05
Coumaphos	µg/L	ND	0.20
Def	µg/L	ND	0.10
Demeton-s	µg/L	ND	0.20
Diazinon	µg/L	0.45	0.05
Dichlorvos	µg/L	ND	0.20
Dimethoate	µg/L	ND	0.10
Disulfoton	µg/L	ND	0.10
EPN	µg/L	ND	0.10
Ethion	µg/L	ND	0.10
Ethoprop	µg/L	ND	0.10
Fensulfothion	µg/L	ND	0.50
Fenthion	µg/L	ND	0.10
Malathion	µg/L	0.11	0.10
Merphos	µg/L	ND	0.10
Mevinphos	µg/L	ND	0.70
Naled	µg/L	ND	0.50
Parathion, ethyl	µg/L	ND	0.10
Parathion, methyl	µg/L	ND	0.10
Phorate	µg/L	ND	0.10
Prowl	µg/L	ND	0.10
Ronnel	µg/L	ND	0.10
Stirophos	µg/L	ND	0.10
Tokuthion	µg/L	ND	0.10
Trichloronate	µg/L	ND	0.10
Trifluralin	µg/L	ND	0.10

ND = None Detected

Tested by: Paula Young
Checked by: Mike Day

Woodward-Clyde Consultants
500 12th Street, Suite 100
Oakland, CA 94607-4014
Attn: R. Katznelson

Sample Date: 10/29/96
Report Date: 11/19/96

Page 3 of 3

Sample ID No : 941020NA/C9943
CV67C153
APPL Sample No: R23944-44765

Date Received: 10/31/96
Date Extracted: 11/05/96

Results of Water Analysis by EPA Method 8321

Compound	Units	Concentration	Quantitation Limit
Aminocarb	µg/L	ND	0.4
Aldicarb	µg/L	ND	0.4
Barban	µg/L	ND	3.5
Benomyl (Carbendazim)	µg/L	ND	0.4
Bromacil	µg/L	ND	0.4
Carbaryl	µg/L	0.09	0.07
Carbofuran	µg/L	ND	0.07
Chloroprotham	µg/L	ND	3.5
Chloroxuron	µg/L	ND	0.4
Diuron	µg/L	1.0	0.4
Fenuron	µg/L	ND	0.4
Fluometuron	µg/L	ND	0.4
Linuron	µg/L	ND	1
Methomyl	µg/L	ND	0.07
Methiocarb	µg/L	ND	0.4
Mexacarbate	µg/L	ND	3.5
Monuron	µg/L	ND	0.4
Neburon	µg/L	ND	0.4
Oxamyl	µg/L	ND	0.4
Propachlor	µg/L	ND	3.5
Propham	µg/L	ND	3.5
Propoxur	µg/L	ND	0.4
Siduron	µg/L	ND	0.4
Tebuthiuron	µg/L	ND	0.4

ND = None Detected

Tested by: Brad Anderson
Checked by: Mike Ray



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500 12th Street, Suite 100
Oakland, CA 94607-4014
Attn: R. Katznelson

Sample Date: NA
Report Date: 11/19/96

Sample I.D. No: **BLANK** Associated with
sample group: R23944
APPL Sample No: R23944-961111W BLK

Date Received: NA
Date Extracted: 11/01/96

Results of OCl Pesticides Water Analysis by EPA Method 8080

Compound	Units	Concentration	Quantitation Limit
Aldrin	µg/L	ND	0.01
a-BHC	µg/L	ND	0.01
b-BHC	µg/L	ND	0.01
d-BHC	µg/L	ND	0.01
g-BHC (Lindane)	µg/L	ND	0.01
Chlordane	µg/L	ND	0.01
Dieldrin	µg/L	ND	0.01
Endosulfan I	µg/L	ND	0.01
Endosulfan II	µg/L	ND	0.01
Endosulfan sulfate	µg/L	ND	0.01
Endrin	µg/L	ND	0.01
Endrin aldehyde	µg/L	ND	0.01
Endrin ketone	µg/L	ND	0.01
Heptachlor	µg/L	ND	0.01
Heptachlor epoxide	µg/L	ND	0.01
4,4'-DDE	µg/L	ND	0.01
4,4'-DDT	µg/L	ND	0.01
4,4'-TDE/DDD	µg/L	ND	0.01
Methoxychlor	µg/L	ND	0.01
Toxaphene	µg/L	ND	0.20

AMENDED

ND = None Detected

Checked by: Samuel Cooper



4203 West Swift ▼ Fresno, California 93722 ▼ Phone 209.275-2175 ▼ Fax 209.275-4422

Woodward-Clyde Consultants
500 12th Street, Suite 100
Oakland, CA 94607-4014
Attn: R. Katznelson

Sample Date: NA
Report Date: 11/18/96

Sample I.D. No: **BLANK** Associated with
sample group: R23944
APPL Sample No: R23944-961101W BLK

Date Received: NA
Date Extracted: 11/01/96

Results of OP Pesticides Water Analysis by EPA Method 8140

Compound	Units	Concentration	Quantitation Limit
Azinphosmethyl	µg/L	ND	1.0
Bolstar	µg/L	ND	0.10
Chlorpyrifos	µg/L	ND	0.05
Coumaphos	µg/L	ND	0.10
Def	µg/L	ND	0.10
Demeton-s	µg/L	ND	0.20
Diazinon	µg/L	ND	0.05
Dichlorvos	µg/L	ND	0.20
Dimethoate	µg/L	ND	0.10
Diphenamid	µg/L	ND	0.10
Disulfoton	µg/L	ND	0.10
Ethion	µg/L	ND	0.10
Ethoprop	µg/L	ND	0.10
Fensulfothion	µg/L	ND	0.20
Fenthion	µg/L	ND	0.10
Malathion	µg/L	ND	0.40
Merphos	µg/L	ND	0.10
Methyl trithion	µg/L	ND	0.20
Mevinphos	µg/L	ND	0.70
Naled	µg/L	ND	0.50
Parathion, ethyl	µg/L	ND	0.10
Parathion, methyl	µg/L	ND	0.10
Phorate	µg/L	ND	0.10
Phosalone	µg/L	ND	0.10
Prometon	µg/L	ND	0.10
Prowl	µg/L	ND	0.10
Ronnel	µg/L	ND	0.10
Simazine	µg/L	ND	0.50
Trichloronate	µg/L	ND	0.10
Trifluralin	µg/L	ND	0.10

ND = None Detected

Checked by: James Coore



Woodward-Clyde Consultants
500 12th Street, Suite 100
Oakland, CA 94607-4014
Attn: R. Katznelson

Sample Date: NA
Report Date: 11/19/96

Sample I.D. No: **BLANK** Associated with
sample group: R23944
APPL Sample No: R23944-961105W BLK1

Date Received: NA
Date Extracted: 11/05/96

Results of Water Analysis by EPA Method 8321

Compound	Units	Concentration	Quantitation Limit
Aminocarb	µg/L	ND	0.4
Aldicarb	µg/L	ND	0.4
Barban	µg/L	ND	3.5
Benomyl (Carbendazim)	µg/L	ND	0.4
Bromacil	µg/L	ND	0.4
Carbaryl	µg/L	ND	0.07
Carbofuran	µg/L	ND	0.07
Chloroprotham	µg/L	ND	3.5
Chloroxuron	µg/L	ND	0.4
Diuron	µg/L	ND	0.4
Fenuron	µg/L	ND	0.4
Fluometuron	µg/L	ND	0.4
Linuron	µg/L	ND	0.07
Methomyl	µg/L	ND	0.07
Methiocarb	µg/L	ND	0.4
Mexacarbate	µg/L	ND	3.5
Monuron	µg/L	ND	0.4
Neburon	µg/L	ND	0.4
Oxamyl	µg/L	ND	0.4
Propachlor	µg/L	ND	0.4
Protham	µg/L	ND	3.5
Propoxur	µg/L	ND	3.5
Siduron	µg/L	ND	0.4
Tebuthiuron	µg/L	ND	0.4

ND = None Detected

Checked by: James Coops

APPL Sample: 44765W

Method

Extraction
Date

Analysis
Date

Spike Report: MS / MSD

8080

11/1/96

11/18/96

Analyte	CAS Number	Units	Amount in Sample	Amount Added	Recovery MS	Recovery MSD	Percent MS	Percent MSD	FLAG	RPD	Recovery Limit	RPD Limit
Aldrin	309-00-2	µg/L	0.0181	0.0267	0.0283	0.0266	38.2%	31.8%	6.2%	30-99	31	
4,4'-DDT	50-29-3	µg/L	ND	0.0667	0.0581	0.0538	87.1%	80.7%	7.7%	47-118	18	
Dieldrin	60-57-1	µg/L	ND	0.0667	0.0433	0.0423	64.9%	63.4%	2.3%	45-122	14	
Endrin	72-20-8	µg/L	ND	0.0667	0.0585	0.0490	87.7%	73.5%	18%	61-133	19	
Heptachlor	76-44-8	µg/L	0.0556	0.0267	0.0656	0.0673	37.5%	43.8%	2.6%	28-125	26	
Lindane	58-89-9	µg/L	ND	0.0267	0.0180	0.0192	67.4%	71.9%	6.5%	49-132	16	

ABBREVIATIONS & FLAGS

ND None Detected

NA Not Applicable

NE Not Established

◆ Recovery is outside of the acceptable percentage range

APPL Sample: 44765W

Spike Report: MS / MSD

Method
8140

Extraction
Date
11/1/96

Analysis
Date
11/7/96

Analyte	CAS Number	Units	Amount in Sample	Amount Added	Recovery MS	Recovery MSD	Percent MS	Percent FLAG MSD	Percent FLAG MSD	RPD FLAG	Recovery Limit	RPD Limit
Disulfoton	298-04-4	µg/L	ND	0.333	0.245	0.259	73.6%	77.8%	5.6%		47-117	22
Ethion	563-12-2	µg/L	ND	0.333	0.319	0.312	95.8%	93.7%	2.2%		65-134	20
Parathion, methyl	298-00-0	µg/L	ND	0.333	0.249	0.261	74.8%	78.4%	4.7%		55-164	24
Phorate	298-02-2	µg/L	ND	0.333	0.151	0.153	45.3%	45.9%	1.3%		22-96	24
Stirophos	22248-79-9	µg/L	ND	0.333	0.289	0.261	86.8%	78.4%	10%		68-128	25

ABBREVIATIONS & FLAGS

ND None Detected
NA Not Applicable

NE Not Established
◆ Recovery is outside of the acceptable percentage range

APPL Sample: 961105W

Spike Report: LCS

Method
8321

Extraction
Date
11/5/96

Analysis
Date
11/5/96

Analyte	CAS Number	Units	Amount in Sample	Amount Added	Recovery LCS	Percent LCS	FLAG	Recovery Limit
Carbaryl	63-25-2	µg/L	ND	0.267	0.257	96.3%		40-131
Carbofuran	1563-66-2	µg/L	ND	0.267	0.218	81.6%		44-128
Methomyl	16752-77-5	µg/L	ND	0.267	0.223	83.5%		37-113

ABBREVIATIONS & FLAGS

ND None Detected

NA Not Applicable

NE Not Established

◆ Recovery is outside of the acceptable percentage range

APPL Sample: 44756W

Spike Report: MS / MSD

Method
8321

Extraction
Date
11/5/96

Analysis
Date
11/5/96

Analyte	CAS Number	Units	Amount in Sample	Amount Added	Recovery MS	Recovery MSD	Percent MS	Percent FLAG MSD	Percent FLAG MSD	RPD FLAG	Recovery Limit	RPD Limit
Carbaryl	63-25-2	µg/L	ND	0.267	0.245	0.215	91.8%	80.5%		13%	40-131	30
Carbofuran	1563-66-2	µg/L	ND	0.267	0.141	0.090	52.8%	33.7% ♦		44% ♦	44-128	30
Methomyl	16752-77-5	µg/L	ND	0.267	0.267	0.221	100%	82.8%		19%	37-113	30

ABBREVIATIONS & FLAGS

ND None Detected

NA Not Applicable

NE Not Established

♦ Recovery is outside of the acceptable percentage range

Woodward-Clyde Consultants

500 12th Street, Suite 100, Oakland, CA 94607-4014
(510) 893-3600

Chain of Custody Record

Appl Lab

PROJECT NO. <i>941020NA/C9943</i>			ANALYSES				Number of Containers	REMARKS (Sample preservation, handling procedures, etc.)
SAMPLERS: (Signature) <i>Arleen Ferguson (FOR JIM SCANLON)</i>			Sample Matrix (Soil, Water, Air)	EPA Method	EPA Method	EPA Method		
DATE	TIME	SAMPLE NUMBER						
<i>10/29</i>	<i>compost</i>	<i>CY67CL53</i>	<i>water</i>	<i>X</i>	<i>X</i>	<i>X</i>	<i>4</i>	<p><i>OCs, OPs and Carbamates at the lowest DL available</i></p> <p><i>Please send invoice and report to R. Katznelson, WCC (address above) (510) 874 3048</i></p>
RELINQUISHED BY (Signature)						TOTAL NUMBER OF CONTAINERS		
RELINQUISHED BY (Signature) <i>Arleen Ferguson</i>		DATE/TIME <i>10/30/96</i>	RECEIVED BY (Signature) <i>Fed Express</i>		RELINQUISHED BY (Signature) <i>Fed Express</i>	DATE/TIME	RECEIVED BY (Signature)	
METHOD OF SHIPMENT: <i>FED EX</i>			SHIPPED BY (Signature)	COURIER (Signature)	RECEIVED FOR LAB BY (Signature) <i>R. Katznelson</i>	DATE/TIME <i>10/30/96</i>		

Chilled & stored

Appendix C. Estimation of Number of Sources in Castro Valley Watershed

A basic assumption used for estimation of the number of sources was that each street gutter sample with a diazinon concentration elevated above an arbitrary threshold level represented an individual diazinon source. It was further assumed that only one property within the area drained by a street gutter sample was responsible for the elevated concentration. If a threshold level of 400-500 ng/l was selected, only 8 of the 45 samples contained sources; if the threshold was 100 ng/l, then 17 samples contained sources (Table C.1). These figures were used to bracket the minimum and maximum ranges of the estimate.

Taking an average estimate of 10 properties per street gutter sample, 450 properties were represented in the total area sampled. This estimate was verified by a rough visual estimation of the proportion of Subcatchments 2 and 3 that was covered by the street gutter sampling (Figure C.1), combined with census figures for the number of households in these subcatchments (Table C.2). The 8 to 17 assumed sources then represented between two and four percent of the properties in the sampled area. Since most of the properties in Subcatchments 2 and 3 were single-family homes ("residential" in Table 3.2.1), this was roughly equivalent to 2-4 percent of households.

Application of these percentages to the 14,800 residential units in the entire watershed results in an estimate of 300-600 sources of diazinon in the watershed. About one-fourth of the residential units represent multiple-family dwellings (which are lumped under "commercial" land use in Table 3.2.1) than the sampled area. Patterns of diazinon use and application may be somewhat different for households living in multiple unit dwellings, and it is possible that on an area basis these areas contribute more diazinon to runoff (See Section 3.2). However, it was assumed here that the percentage of these households associated with diazinon in runoff would be the same as for single-family residential areas.

Table C.1 Diazinon in Street Gutters in Castro Valley on May 15, 1996

Site	Sample time	Diazinon (ng/l)	Assumed No. of Sources Per Sample	
			Maximum (100 ng/l)	Minimum (500 ng/l)
28	14:21	79,000	1	1
19	13:54	71,000	1	1
10	13:27	4,100	1	1
36	14:45	3,700	1	1
29	14:24	2,700	1	1
21	14:00	2,400	1	1
42	15:03	1,500	1	1
7	13:18	1,400	1	1
2	13:03	400	1	
3	13:06	400	1	
43	15:06	400	1	
12	13:33	310	1	
18	13:51	260	1	
24	14:09	180	1	
27	14:18	130	1	
13	13:36	120	1	
45	15:12	110	1	
6	13:15	80		
30	14:27	80		
1	13:00	80		
4	13:09	80		
25	14:12	80		
31	14:30	80		
35	14:42	70		
39	14:54	60		
32	14:33	60		
34	14:39	60		
20	13:57	50		
22	14:03	40		
23	14:06	40		
26	14:15	40		
33	14:36	40		
38	14:51	40		
41	15:00	30		
5	13:12	30		
8	13:21	30		
15	13:42	30		
37	14:48	30		
40	14:57	30		
44	15:09	30		
9	13:24	ND		
11	13:30	ND		
14	13:39	ND		
17	13:48	ND		
16	13:45	ND		
Totals			17	8

Table C.2 Separate estimate of number of households in sampled area

(Approximated by census tracts)	Maximum			Minimum		
	Zone		Total	Zone		Total
	2	3		2	3	
% of subcatchment area sampled (Figure C.1)	.15	10		10	5	
Households per subcatchment	2600	4380	6980	2600	4380	6980
Households in sampled area	390	438	828	260	219	479

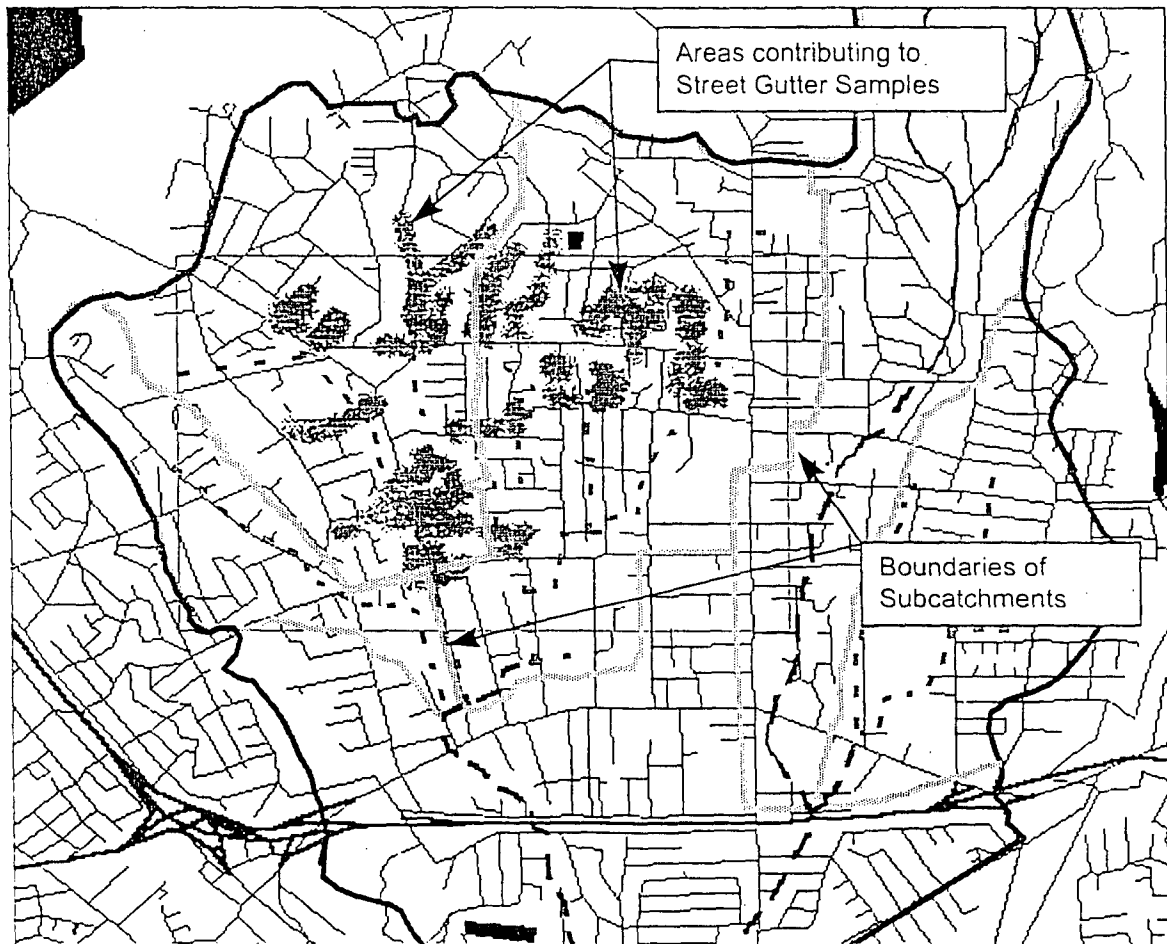


Figure C.1 Estimated Area Drained by Street Gutters Sampled in Castro Valley on May 15, 1996

Appendix D. Runoff Study Sampling Locations

Site	Sampling Location	Description of Sampling Location
Oakland	Roof Drain	Downspout from roof gutter
Oakland	Drain + Patio	Runoff from roof downspout after running across 6 ft (1.8m) of patio
Oakland	Driveway	Puddle at bottom of 40 ft. (12m) long driveway
Oakland	Patio	Puddle on paved patio
Castro Valley 1	Patio	Puddle on paved patio
Castro Valley 1	Driveway	Runoff from base of 20 ft. (6m) long driveway
Castro Valley 1	Roof Drain	Downspout from roof gutter
Castro Valley 2	Patio	Puddle on paved patio
Castro Valley 2	Driveway	Puddle at bottom of 20 ft. (6m) long driveway
Castro Valley 2	Roof Drain	Downspout from roof gutter

Appendix E. 1995-1997 Castro Valley Rainfall Data

Rainfall Data from Gauge at Arcadian Reservoir (Castro Valley) for 1995-97
Storm Season

Rainfall Data from Gauge at Arcadian Reservoir (Castro Valley) for 1996-97
Storm Season

Rainfall Data from Gauge in Backyard near Redwood Road and SR 13
("Andreasgauge", Oakland) from 2/3/97 to 4/24/97

Rain Gauge Location: Arcadian Reservoir, Centre Valley

MONTH	DAY	YEAR	HOUR	INCHES	MONTHLY TOTAL
11	1	95	3	0.04	0.86
12	1	95	18	0.04	
12	1	95	19	0.04	
12	3	95	20	0.04	
12	3	95	24	0.04	
12	4	95	2	0.04	
12	4	95	3	0.07	
12	4	95	5	0.04	
12	4	95	7	0.04	
12	4	95	8	0.04	
12	4	95	10	0.04	
12	4	95	12	0.04	
12	7	95	3	0.04	
12	8	95	7	0.04	
12	10	95	18	0.04	
12	10	95	24	0.04	
12	11	95	3	0.31	
12	11	95	4	0.20	
12	11	95	5	0.12	
12	11	95	6	0.12	
12	11	95	7	0.04	
12	11	95	8	0.19	
12	11	95	9	0.28	
12	11	95	10	0.24	
12	11	95	11	0.07	
12	11	95	12	0.04	
12	11	95	15	0.04	
12	11	95	16	0.04	
12	11	95	17	0.08	
12	11	95	19	0.20	
12	11	95	20	0.27	
12	11	95	21	0.24	
12	11	95	22	0.24	
12	11	95	23	0.07	
12	12	95	5	0.04	
12	12	95	6	0.04	
12	12	95	7	0.32	
12	12	95	8	0.27	
12	12	95	11	0.04	
12	12	95	12	0.04	
12	13	95	9	0.04	
12	13	95	14	0.24	
12	13	95	16	0.16	
12	15	95	4	0.07	
12	15	95	5	0.20	
12	15	95	7	0.16	
12	15	95	8	0.16	
12	15	95	9	0.04	
12	18	95	7	0.03	
12	18	95	10	0.04	
12	18	95	11	0.04	
12	18	95	12	0.04	
12	18	95	13	0.08	
12	18	95	14	0.04	
12	22	95	6	0.08	
12	22	95	7	0.16	
12	22	95	8	0.08	
12	22	95	9	0.04	
12	27	95	15	0.03	
12	27	95	12	0.12	
12	29	95	13	0.12	
12	29	95	14	0.04	
12	29	95	15	0.12	
12	29	95	16	0.12	
12	29	95	17	0.08	
12	29	95	18	0.07	
12	29	95	19	0.04	
12	30	95	2	0.04	
1	15	96	23	0.04	6.63
1	15	96	24	0.04	
1	16	96	5	0.04	
1	16	96	6	0.20	
1	16	96	7	0.43	
1	16	96	8	0.47	
1	16	96	9	0.40	
1	16	96	10	0.12	
1	16	96	11	0.03	
1	16	96	14	0.08	
1	18	96	14	0.16	
1	18	96	15	0.24	
1	18	96	16	0.15	
1	18	96	17	0.04	
1	18	96	21	0.04	
1	19	96	1	0.04	
1	19	96	2	0.04	
1	19	96	3	0.04	
1	19	96	5	0.04	
1	19	96	6	0.04	
1	19	96	11	0.04	
1	20	96	21	0.04	
1	20	96	22	0.04	
1	20	96	23	0.39	
1	20	96	24	0.04	
1	21	96	2	0.12	
1	21	96	6	0.08	
1	23	96	17	0.04	
1	23	96	21	0.04	
1	24	96	6	0.04	
1	24	96	7	0.04	
1	24	96	8	0.04	
1	24	96	9	0.11	
1	24	96	10	0.20	
1	24	96	12	0.08	
1	24	96	13	0.04	
1	24	96	15	0.08	
1	24	96	20	0.12	
1	24	96	21	0.15	
1	24	96	22	0.04	
1	24	96	23	0.04	

MONTH	DAY	YEAR	HOUR	INCHES	MONTHLY TOTAL
1	27	95	5	0.08	6.88
1	27	95	6	0.20	
1	27	95	7	0.23	
1	27	95	8	0.24	
1	27	95	10	0.04	
1	27	95	16	0.04	
1	27	95	17	0.04	
1	30	95	15	0.08	
1	30	95	16	0.04	
1	30	95	18	0.11	
1	30	95	19	0.12	
1	30	95	20	0.08	
1	30	95	21	0.04	
1	30	95	22	0.04	
1	30	95	24	0.16	
1	31	95	1	0.08	
1	31	95	2	0.04	
1	31	95	4	0.13	
1	31	95	5	0.24	
1	31	95	6	0.08	
1	31	95	7	0.04	
1	31	95	8	0.12	
1	31	95	9	0.16	
2	3	96	4	0.08	6.88
2	3	96	5	0.04	
2	3	96	6	0.04	
2	3	96	22	0.04	
2	4	96	4	0.08	
2	4	96	5	0.12	
2	4	96	6	0.08	
2	4	96	7	0.23	
2	4	96	9	0.32	
2	4	96	10	0.11	
2	4	96	11	0.08	
2	4	96	13	0.08	
2	4	96	14	0.04	
2	4	96	19	0.04	
2	4	96	20	0.08	
2	4	96	23	0.04	
2	5	96	2	0.04	
2	7	96	6	0.04	
2	15	96	15	0.04	
2	15	96	17	0.08	
2	15	96	18	0.03	
2	15	96	19	0.24	
2	15	96	20	0.08	
2	17	96	22	0.04	
2	18	96	16	0.04	
2	18	96	17	0.04	
2	18	96	19	0.04	
2	18	96	21	0.04	
2	18	96	24	0.15	
2	19	96	1	0.08	
2	19	96	12	0.12	
2	19	96	13	0.24	
2	19	96	14	0.08	
2	19	96	15	0.07	
2	19	96	17	0.20	
2	20	96	4	0.24	
2	20	96	5	0.12	
2	21	96	1	0.07	
2	21	96	2	0.08	
2	21	96	3	0.40	
2	21	96	4	0.15	
2	21	96	5	0.20	
2	21	96	6	0.12	
2	21	96	7	0.04	
2	21	96	10	0.08	
2	22	96	22	0.31	
2	23	96	23	0.08	
2	23	96	24	0.04	
2	24	96	18	0.04	
2	24	96	19	0.04	
2	25	96	16	0.04	
2	27	96	5	0.04	
2	27	96	6	0.08	
2	27	96	7	0.04	
2	27	96	20	0.08	
2	28	96	17	0.03	
2	28	96	23	0.04	
2	28	96	24	0.04	
2	29	96	15	0.08	
2	29	96	16	0.16	
3	3	96	13	0.12	3.52
3	3	96	14	0.04	
3	4	96	8	0.11	
3	4	96	12	0.08	
3	4	96	13	0.12	
3	4	96	14	0.12	
3	4	96	15	0.20	
3	4	96	20	0.04	
3	4	96	23	0.04	
3	5	96	2	0.03	
3	10	96	7	0.04	
3	10	96	9	0.08	
3	11	96	2	0.04	
3	11	96	9	0.04	
3	11	96	10	0.12	
3	11	96	21	0.04	
3	11	96	23	0.08	
3	11	96	24	0.12	
3	12	96	1	0.23	
3	12	96	2	0.12	
3	12	96	3	0.24	
3	12	96	4	0.15	
3	12	96	5	0.04	
3	12	96	6	0.12	
3	12	96	7	0.12	

MONTH	DAY	YEAR	HOUR	INCHES	MONTHLY TOTAL
3	12	96	13	0.12	2.67
3	12	96	14	0.08	
3	12	96	15	0.04	
3	28	96	24	0.11	
3	31	96	24	0.04	
4	2	96	14	0.39	
4	15	96	12	0.04	
4	16	96	5	0.16	
4	16	96	6	0.20	
4	17	96	16	0.04	
4	17	96	17	0.08	
4	17	96	18	0.11	
4	17	96	19	0.24	
4	17	96	20	0.04	
4	18	96	5	0.08	
4	19	96	15	0.04	
4	19	96	16	0.04	
4	19	96	17	0.04	
4	24	96	10	0.04	
5	15	96	12	0.04	1.15
5	15	96	13	0.31	
5	15	96	14	0.20	
5	15	96	15	0.11	
5	15	96	16	0.12	
5	15	96	17	0.04	
5	15	96	18	0.12	
5	15	96	19	0.20	
5	15	96	20	0.04	
5	15	96	21	0.04	
5	15	96	22	0.07	
5	15	96	23	0.08	
5	15	96	24	0.03	
5	16	96	1	0.12	
5	16	96	11	0.04	
5	17	96	4	0.04	
5	17	96	21	0.04	
5	21	96	10	0.04	
5	21	96	11	0.04	
5	21	96	14	0.08	
5	21	96	15	0.04	

GRAND TOTAL 25.91

CUMMULATIVE TOTAL DUE TO COMPUTER LOCKUP

Arcadian Reservoir 1996-97

Rain Gauge Location: Arcadian Reservoir, Castro Valley
 Sensor 1932 Cumulative rainfall 1996-97

DATE	TIME	INCHES	DATE	TIME	INCHES	DATE	TIME	INCHES	DATE	TIME	INCHES	DATE	TIME	INCHES
12/31/96	22.54	12.05	12/21/96	11:34	9.88	12/5/96	23:26	6.38	11/20/96	13:22	3.54	11/7/96	20:25	0.67
12/31/96	17:21	12.01	12/21/96	11:05	9.84	12/5/96	19:19	6.34	11/20/96	4:51	3.54	11/7/96	8:13	0.67
12/31/96	5:09	12.01	12/21/96	10:36	9.8	12/5/96	8:31	6.34	11/20/96	1:10	3.5	11/6/96	20:01	0.67
12/30/96	19:50	12.01	12/21/96	10:14	9.76	12/5/96	7:07	6.3	11/19/96	13:45	3.5	11/6/96	7:49	0.67
12/30/96	19:38	11.97	12/21/96	9:39	9.72	12/5/96	6:19	6.3	11/19/96	12:58	3.46	11/5/96	19:37	0.67
12/30/96	18:57	11.93	12/21/96	9:09	9.69	12/5/96	5:33	6.26	11/19/96	12:21	3.46	11/5/96	7:25	0.67
12/30/96	4:45	11.93	12/21/96	8:47	9.65	12/5/96	5:23	6.22	11/19/96	11:33	3.43	11/4/96	19:13	0.67
12/29/96	18:33	11.89	12/21/96	8:29	9.61	12/5/96	5:14	6.18	11/19/96	11:23	3.39	11/4/96	7:01	0.67
12/29/96	15:07	11.89	12/21/96	8:07	9.57	12/5/96	5:07	6.14	11/19/96	10:43	3.31	11/3/96	18:49	0.67
12/29/96	13:39	11.85	12/21/96	7:24	9.49	12/5/96	4:59	6.1	11/19/96	10:46	3.27	11/3/96	8:38	0.67
12/29/96	12:54	11.81	12/21/96	6:55	9.45	12/5/96	4:32	6.06	11/18/96	12:34	3.27	11/2/96	18:26	0.67
12/29/96	12:48	11.77	12/21/96	6:39	9.41	12/5/96	3:55	6.02	11/18/96	6:57	3.27	11/2/96	6:14	0.67
12/29/96	12:38	11.73	12/21/96	6:25	9.37	12/5/96	3:39	5.98	11/18/96	6:00	3.23	11/1/96	5:50	0.67
12/29/96	12:23	11.69	12/21/96	6:06	9.33	12/5/96	3:34	5.94	11/18/96	0:22	3.19	10/31/96	17:38	0.67
12/29/96	12:08	11.65	12/21/96	5:54	9.29	12/5/96	3:25	5.91	11/17/96	19:30	3.19	10/30/96	17:14	0.67
12/29/96	9:48	11.61	12/21/96	5:33	9.25	12/5/96	3:17	5.87	11/17/96	18:23	3.15	10/30/96	9:55	0.67
12/29/96	4:22	11.57	12/21/96	5:13	9.21	12/5/96	3:10	5.83	11/17/96	16:46	3.11	10/30/96	6:09	0.63
12/28/96	16:10	11.57	12/21/96	4:53	9.17	12/5/96	3:00	5.79	11/17/96	16:25	3.07	10/30/96	5:02	0.59
12/28/96	3:58	11.57	12/21/96	4:33	9.13	12/5/96	2:54	5.75	11/17/96	15:46	3.03	10/29/96	16:51	0.59
12/27/96	19:50	11.57	12/21/96	4:14	9.09	12/5/96	2:41	5.71	11/17/96	15:31	2.99	10/29/96	10:15	0.59
12/27/96	19:45	11.54	12/21/96	3:50	9.06	12/5/96	2:26	5.67	11/17/96	15:11	2.95	10/29/96	8:06	0.55
12/27/96	15:48	11.5	12/21/96	3:22	9.02	12/5/96	2:12	5.63	11/17/96	12:11	2.91	10/29/96	7:22	0.51
12/27/96	14:23	11.5	12/21/96	3:14	8.98	12/5/96	2:05	5.59	11/17/96	6:48	2.91	10/29/96	7:14	0.47
12/27/96	3:34	11.46	12/21/96	3:10	8.94	12/5/96	1:54	5.55	11/17/96	6:34	2.87	10/29/96	7:08	0.43
12/28/96	23:57	11.46	12/21/96	3:08	8.9	12/5/96	1:47	5.51	11/17/96	6:19	2.83	10/29/96	6:04	0.39
12/28/96	22:05	11.42	12/21/96	3:07	8.85	12/5/96	1:36	5.47	11/17/96	6:15	2.8	10/29/96	5:30	0.35
12/28/96	21:59	11.38	12/21/96	3:06	8.82	12/5/96	1:16	5.43	11/17/96	6:12	2.76	10/29/96	4:44	0.31
12/28/96	21:53	11.34	12/21/96	3:03	8.78	12/5/96	0:32	5.39	11/17/96	6:09	2.72	10/29/96	4:39	0.28
12/28/96	21:47	11.3	12/21/96	3:01	8.74	12/4/96	23:53	5.31	11/17/96	6:05	2.68	10/29/96	3:55	0.28
12/28/96	21:44	11.28	12/21/96	2:49	8.66	12/4/96	23:40	5.28	11/17/96	6:00	2.64	10/29/96	3:29	0.24
12/28/96	21:42	11.22	12/21/96	2:41	8.62	12/4/96	23:29	5.24	11/17/96	5:54	2.6	10/29/96	3:13	0.2
12/28/96	21:41	11.18	12/21/96	2:31	8.58	12/4/96	23:22	5.2	11/17/96	5:49	2.56	10/29/96	2:58	0.16
12/28/96	21:39	11.14	12/21/96	2:20	8.54	12/4/96	23:13	5.16	11/17/96	5:40	2.52	10/29/96	2:39	0.12
12/28/96	21:37	11.1	12/21/96	2:08	8.5	12/4/96	22:54	5.12	11/17/96	5:36	2.48	10/29/96	2:17	0.08
12/28/96	18:53	11.06	12/21/96	1:57	8.46	12/4/96	22:38	5.08	11/17/96	5:33	2.44	10/28/96	16:26	0.04
12/28/96	17:54	11.02	12/21/96	1:46	8.43	12/4/96	22:28	5.04	11/17/96	5:28	2.4	10/15/96	17:29	25.91
12/28/96	17:08	10.98	12/21/96	1:20	8.35	12/4/96	22:16	5	11/17/96	5:25	2.36	10/15/96	5:17	25.91
12/28/96	16:38	10.94	12/21/96	1:11	8.31	12/4/96	22:08	4.96	11/17/96	5:19	2.32	10/14/96	4:54	25.91
12/28/96	15:22	10.91	12/21/96	1:04	8.31	12/4/96	21:57	4.92	11/17/96	4:57	2.28	10/8/96	13:55	25.91
12/28/96	9:12	10.91	12/21/96	0:39	8.27	12/4/96	21:51	4.88	11/17/96	4:29	2.24	9/29/96	23:21	25.91
12/28/96	8:00	10.87	12/20/96	12:59	8.23	12/4/96	21:40	4.84	11/17/96	4:14	2.2	9/28/96	10:45	25.91
12/28/96	3:10	10.83	12/20/96	0:48	8.23	12/4/96	21:27	4.76	11/17/96	3:41	2.17	9/21/96	20:10	25.91
12/25/96	14:58	10.83	12/19/96	12:36	8.23	12/4/96	21:17	4.72	11/17/96	2:49	2.13	9/21/96	7:59	25.91
12/25/96	2:46	10.83	12/19/96	0:24	8.23	12/4/96	20:59	4.69	11/17/96	2:30	2.09	9/18/96	18:59	25.91
12/24/96	14:35	10.83	12/18/96	12:12	8.23	12/4/96	20:26	4.65	11/17/96	2:24	2.05	8/21/96	7:54	25.91
12/24/96	2:23	10.83	12/18/96	0:00	8.23	12/4/96	19:52	4.61	11/17/96	2:20	2.01	8/19/96	7:07	25.91
12/23/96	14:11	10.83	12/17/96	11:48	8.23	12/4/96	18:55	4.57	11/17/96	2:06	1.97	8/17/96	18:31	25.91
12/23/96	1:59	10.83	12/16/96	23:37	8.23	12/4/96	18:54	4.57	11/17/96	1:55	1.93	8/16/96	18:07	25.91
12/22/96	13:47	10.83	12/16/96	11:25	8.23	12/4/96	18:31	4.53	11/17/96	1:03	1.85	8/13/96	16:56	25.91
12/22/96	11:54	10.83	12/15/96	23:13	8.23	12/4/96	18:31	4.53	11/17/96	0:44	1.81	8/10/96	3:37	25.91
12/22/96	1:35	10.79	12/15/96	11:01	8.23	12/3/96	18:07	4.53	11/17/96	0:30	1.77	8/9/96	15:25	25.91
12/21/96	23:07	10.79	12/14/96	22:49	8.23	12/2/96	18:07	4.53	11/17/96	0:21	1.73	8/9/96	3:13	25.91
12/21/96	21:34	10.75	12/13/96	22:25	8.23	12/2/96	17:43	4.53	11/17/96	0:11	1.69	8/8/96	15:01	25.91
12/21/96	21:11	10.71	12/13/96	10:13	8.23	12/1/96	17:43	4.53	11/17/96	0:04	1.65	8/8/96	2:49	25.91
12/21/96	20:47	10.67	12/12/96	22:40	8.23	12/1/96	5:32	4.53	11/16/96	23:59	1.61	8/7/96	14:37	25.91
12/21/96	20:14	10.63	12/12/96	22:30	8.19	12/1/96	4:16	4.53	11/16/96	23:57	1.61	8/7/96	2:26	25.91
12/21/96	19:28	10.59	12/12/96	22:24	8.15	12/1/96	3:53	4.49	11/16/96	23:54	1.57	8/6/96	14:14	25.91
12/21/96	18:14	10.55	12/12/96	21:52	8.11	12/1/96	3:36	4.45	11/16/96	23:40	1.54	8/6/96	2:02	25.91
12/21/96	17:43	10.51	12/12/96	9:50	8.07	12/1/96	3:24	4.41	11/16/96	23:28	1.5	8/5/96	13:50	25.91
12/21/96	16:41	10.43	12/12/96	6:08	8.07	12/1/96	2:59	4.37	11/16/96	23:12	1.46	8/4/96	13:26	25.91
12/21/96	16:26	10.39	12/12/96	5:16	8.03	12/1/96	2:40	4.33	11/16/96	22:39	1.38	8/4/96	1:14	25.91
12/21/96	16:07	10.35	12/12/96	4:48	7.99	12/1/96	2:28	4.29	11/16/96	21:53	1.34	8/3/96	13:02	25.91
12/21/96	15:51	10.31	12/12/96	1:10	7.95	12/1/96	2:14	4.25	11/16/96	21:31	1.3	8/3/96	0:50	25.91
12/21/96	15:23	10.28	12/12/96	0:05	7.91	12/1/96	0:33	4.21	11/16/96	21:18	1.26	8/2/96	12:39	25.91
12/21/96	14:58	10.24	12/11/96	23:59	7.87	11/30/96	17:20	4.17	11/16/96	21:08	1.22	8/2/96	0:27	25.91
12/21/96	14:40	10.2	12/11/96	23:55	7.83	11/30/96	5:08	4.17	11/16/96	20:50	1.18	8/1/96	0:03	25.91
12/21/96	14:28	10.16	12/11/96	21:38	7.8	11/29/96	16:56	4.17	11/16/96	20:33	1.14	7/31/96	11:51	25.91
12/21/96	14:07	10.12	12/10/96	4:16	6.93	11/29/96	4:44	4.17	11/16/96	20:18	1.1	7/30/96	11:27	25.91
12/21/96	13:49	10.08	12/10/96	4:09	6.89	11/28/96	7:42	4.17	11/16/96	20:04	1.06	7/29/96	23:15	25.91
12/21/96	13:26	10.04	12/10/96	3:51	6.81	11/28/96	4:20	4.13	11/16/96	19:49	1.02	7/29/96	11:03	25.91
12/21/96	13:23	10	12/10/96	3:34	6.77	11/27/96	16:08	4.13	11/16/96	19:34	0.98	7/28/96	10:40	25.91
12/21/96	12:54	10	12/10/96	3:17	6.73	11/27/96	3:56	4.13	11/16/96	19:09	0.94	7/27/96	22:28	25.91
12/21/96	12:26	9.96	12/10/96	2:41	6.65	11/26/96	15:45	4.13	11/16/96	18:32	0.91	7/27/96	10:16	25.91
			12/10/96	2:22	6.61	11/26/96	3:33	4.13	11/16/96	17:50	0.87	7/26/96	22:04	25.91
			12/9/96	20:54	6.57	11/25/96	15:21	4.13	11/16/96	17:10	0.83	7/26/96	9:52	25.91

Rain Gauge Location: near Redwood Road x SR 13, Oakland

cumulative rainfall, inches

Recorder Status
 Type: 674L-8K Event Capacity: 2035 Recorder ID: TEST
 Time at Recorder: 04/24/97 12:30:14 Last Update: 12/18/95 12:18:17
 Signal process: Not Applicable
 Types of Events Recorded: Lo==>Hi
 Total time LOW: 00:04:15 Total time HI: 492 days 19:44:4

Number of Events Recorded: 2036

Rainfall per Event: 0.01"

cumulative rainfall, inches

rain sample	spray test	Time of event	Type of Event	cumulative rainfall, inches	spray test	Time of event	Type of Event
		2/3/97 19:23:01	Pulse Low to HIGH			4/18/97 14:53:27	Pulse Low to HIGH
		2/3/97 21:50:12	Pulse Low to HIGH			4/18/97 15:07:22	Pulse Low to HIGH
		2/3/97 21:52:23	Pulse Low to HIGH			4/18/97 15:22:29	Pulse Low to HIGH
		2/3/97 21:55:36	Pulse Low to HIGH			4/18/97 15:33:58	Pulse Low to HIGH
		2/3/97 22:20:06	Low to HIGH			4/18/97 15:41:32	Pulse Low to HIGH
		2/3/97 23:34:35	Pulse Low to HIGH			4/18/97 15:45:30	Pulse Low to HIGH
		2/4/97 7:00:28	Pulse Low to HIGH			4/18/97 15:51:32	Pulse Low to HIGH
		2/4/97 18:23:29	Pulse Low to HIGH			4/18/97 15:57:47	Pulse Low to HIGH
		2/8/97 8:45:10	Pulse Low to HIGH			4/18/97 16:03:38	Pulse Low to HIGH
		2/17/97 1:32:51	Pulse Low to HIGH			4/18/97 16:09:49	Pulse Low to HIGH
		2/17/97 1:35:20	Pulse Low to HIGH			4/18/97 16:16:55	Pulse Low to HIGH
		2/17/97 1:35:57	Pulse Low to HIGH			4/18/97 16:27:23	Pulse Low to HIGH
		2/17/97 1:36:15	Pulse Low to HIGH			4/18/97 16:31:17	Pulse Low to HIGH
		2/17/97 1:36:27	Pulse Low to HIGH			4/18/97 16:33:58	Pulse Low to HIGH
		2/17/97 1:36:48	Pulse Low to HIGH			4/18/97 16:37:57	Pulse Low to HIGH
		2/17/97 1:37:18	Pulse Low to HIGH			4/18/97 16:51:06	Pulse Low to HIGH
		2/17/97 1:37:52	Pulse Low to HIGH			4/18/97 17:01:26	Pulse Low to HIGH
		2/17/97 1:38:25	Pulse Low to HIGH			4/18/97 17:05:16	Pulse Low to HIGH
		2/17/97 1:39:27	Pulse Low to HIGH			4/18/97 17:17:06	Pulse Low to HIGH
		2/17/97 1:41:04	Pulse Low to HIGH			4/18/97 17:22:30	Pulse Low to HIGH
		2/17/97 1:43:08	Pulse Low to HIGH			4/18/97 17:30:21	Pulse Low to HIGH
		2/17/97 1:44:03	Pulse Low to HIGH			4/18/97 17:38:23	Pulse Low to HIGH
		2/17/97 1:52:10	Low to HIGH			4/18/97 17:52:39	Pulse Low to HIGH
0.00		2/19/97 12:48:44	Pulse Low to HIGH			4/18/97 18:12:25	Pulse Low to HIGH
		3/2/97 1:54:56	Pulse Low to HIGH			4/18/97 18:27:38	Pulse Low to HIGH
		3/2/97 2:04:51	Pulse Low to HIGH			4/18/97 18:42:53	Pulse Low to HIGH
0.03		3/2/97 2:16:59	Pulse Low to HIGH	0.52		4/18/97 18:53:39	Pulse Low to HIGH
		3/2/97 3:24:09	Pulse Low to HIGH			4/18/97 18:55:34	Pulse Low to HIGH
		3/2/97 4:29:36	Pulse Low to HIGH			4/18/97 18:57:29	Pulse Low to HIGH
		3/2/97 5:15:33	Pulse Low to HIGH			4/18/97 18:59:22	Pulse Low to HIGH
		3/2/97 5:39:28	Pulse Low to HIGH			4/18/97 19:00:41	Pulse Low to HIGH
		3/2/97 5:43:18	Pulse Low to HIGH			4/18/97 19:02:06	Pulse Low to HIGH
		3/2/97 5:47:52	Pulse Low to HIGH			4/18/97 19:03:34	Pulse Low to HIGH
		3/2/97 5:51:56	Pulse Low to HIGH			4/18/97 19:04:16	Pulse Low to HIGH
		3/2/97 5:55:35	Pulse Low to HIGH			4/18/97 19:21:12	Pulse Low to HIGH
		3/2/97 5:57:58	Pulse Low to HIGH			4/18/97 19:51:18	Pulse Low to HIGH
		3/2/97 6:01:39	Pulse Low to HIGH			4/18/97 19:52:32	Pulse Low to HIGH
		3/2/97 6:06:30	Pulse Low to HIGH			4/18/97 19:55:27	Pulse Low to HIGH
		3/2/97 6:11:48	Low to HIGH			4/18/97 19:59:12	Pulse Low to HIGH
		3/2/97 6:16:43	Pulse Low to HIGH			4/18/97 20:03:00	Pulse Low to HIGH
		3/2/97 6:17:39	Pulse Low to HIGH			4/18/97 20:06:02	Pulse Low to HIGH
		3/2/97 6:21:01	Pulse Low to HIGH			4/18/97 20:12:27	Pulse Low to HIGH
0.19		3/2/97 6:34:31	Pulse Low to HIGH			4/18/97 20:21:08	Pulse Low to HIGH
		3/3/97 5:31:55	Pulse Low to HIGH			4/18/97 20:27:42	Pulse Low to HIGH
		3/3/97 5:54:01	Pulse Low to HIGH			4/18/97 20:35:11	Pulse Low to HIGH
		3/3/97 5:56:36	Pulse Low to HIGH			4/18/97 20:36:25	Pulse Low to HIGH
		3/3/97 7:45:25	Pulse Low to HIGH			4/18/97 20:37:18	Pulse Low to HIGH
0.24		3/15/97 19:04:37	Pulse Low to HIGH			4/18/97 20:38:04	Pulse Low to HIGH
		3/16/97 9:27:46	Pulse Low to HIGH			4/18/97 20:39:55	Pulse Low to HIGH
		3/16/97 9:44:20	Pulse Low to HIGH			4/18/97 20:42:31	Low to HIGH
		3/16/97 13:08:42	Pulse Low to HIGH			4/18/97 20:42:31	Low to HIGH
		3/16/97 13:12:05	Pulse Low to HIGH			4/18/97 20:55:08	Pulse Low to HIGH
		3/16/97 13:16:34	Pulse Low to HIGH			4/18/97 20:56:34	Pulse Low to HIGH
		3/16/97 13:20:02	Pulse Low to HIGH			4/18/97 20:57:08	Pulse Low to HIGH
		3/16/97 13:22:54	Pulse Low to HIGH			4/18/97 20:57:49	Pulse Low to HIGH
		3/16/97 13:25:54	Pulse Low to HIGH			4/18/97 20:58:52	Pulse Low to HIGH
		3/16/97 13:29:11	Low to HIGH			4/18/97 21:00:05	Pulse Low to HIGH
		3/16/97 13:31:37	Pulse Low to HIGH			4/18/97 21:05:46	Low to HIGH
		3/16/97 13:36:48	Pulse Low to HIGH			4/18/97 21:07:22	Pulse Low to HIGH
		3/16/97 17:16:41	Pulse Low to HIGH			4/18/97 21:08:11	Pulse Low to HIGH
		3/16/97 17:21:19	Pulse Low to HIGH			4/18/97 21:10:00	Pulse Low to HIGH
		3/16/97 17:22:50	Pulse Low to HIGH			4/18/97 21:13:09	Pulse Low to HIGH
		3/16/97 17:23:32	Pulse Low to HIGH			4/18/97 21:14:29	Pulse Low to HIGH
		3/16/97 17:25:34	Pulse Low to HIGH			4/18/97 21:16:36	Pulse Low to HIGH
		3/16/97 17:27:45	Pulse Low to HIGH			4/18/97 21:18:56	Pulse Low to HIGH
		3/16/97 17:28:28	Pulse Low to HIGH			4/18/97 21:25:07	Pulse Low to HIGH
		3/16/97 17:33:49	Pulse Low to HIGH			4/18/97 21:35:23	Pulse Low to HIGH
		3/16/97 19:54:12	Pulse Low to HIGH			4/18/97 21:39:42	Pulse Low to HIGH
		3/16/97 20:01:08	Pulse Low to HIGH			4/18/97 21:44:36	Low to HIGH
		3/16/97 21:45:02	Pulse Low to HIGH			4/18/97 21:48:01	Low to HIGH
		3/16/97 21:46:24	Pulse Low to HIGH			4/18/97 21:53:00	Low to HIGH
		3/16/97 22:14:37	Pulse Low to HIGH			4/18/97 21:59:11	Pulse Low to HIGH
		3/16/97 22:26:56	Pulse Low to HIGH			4/18/97 22:09:34	Pulse Low to HIGH
		3/16/97 22:33:43	Pulse Low to HIGH			4/18/97 22:15:18	Pulse Low to HIGH
0.51		3/21/97 17:35:15	Pulse Low to HIGH			4/18/97 22:25:24	Pulse Low to HIGH
0.52		3/31/97 11:47:18	Pulse Low to HIGH			4/18/97 22:33:32	Pulse Low to HIGH
						4/18/97 22:36:23	Pulse Low to HIGH
						4/18/97 22:39:18	Pulse Low to HIGH
						4/18/97 22:41:36	Pulse Low to HIGH
						4/18/97 22:44:30	Pulse Low to HIGH
						4/18/97 22:49:35	Pulse Low to HIGH
						4/18/97 22:53:12	Low to HIGH
						4/18/97 22:55:23	Pulse Low to HIGH
						4/18/97 22:56:56	Pulse Low to HIGH
						4/18/97 23:00:23	Pulse Low to HIGH
						4/18/97 23:09:12	Pulse Low to HIGH
						4/18/97 23:18:53	Pulse Low to HIGH
						4/19/97 0:24:35	Pulse Low to HIGH
						4/20/97 14:12:15	Pulse Low to HIGH
						4/22/97 21:10:54	Pulse Low to HIGH
						4/24/97 6:51:40	Pulse Low to HIGH