

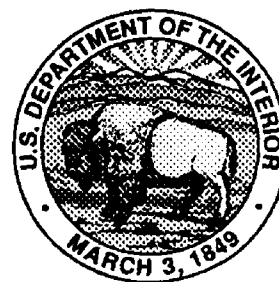
Selected Chemical Characteristics and Acute Toxicity of Urban Stormwater, Streamflow, and Bed Material, Maricopa County, Arizona

By THOMAS J. LOPES and KENNETH D. FOSSUM

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CONVERSION FACTORS

Multiply	By	To obtain
inch (in.)	25.4	millimeter
inch (in.)	2.540	centimeter
foot (ft)	0.3048	meter
mile (mi)	1.609	kilometer
square mile (mi ²)	2.590	square kilometer
cubic foot per second (ft ³ /s)	0.02832	cubic meter per second
pound (lb)	0.907	megagram

In this report, temperature is reported in degrees Fahrenheit (°F), which can be converted to degrees Celsius (°C) by the following equation:

$$^{\circ}\text{C} = (^{\circ}\text{F} - 32)/1.8$$

ABBREVIATED WATER-QUALITY UNITS

Chemical concentrations are given only in metric units. Chemical concentration in water is given in milligrams per liter (mg/L) or micrograms per liter (µg/L). Milligrams per liter is a unit expressing the solute mass per unit volume (liter) of water. One thousand micrograms per liter is equivalent to 1 milligram per liter. For concentrations less than 7,000 milligrams per liter, the numerical value is about the same as for concentrations in parts per million (ppm). Specific conductance is given in microsiemens per centimeter (µS/cm) at 25°C.

VERTICAL DATUM

Sea level: In this report, “sea level” refers to the National Geodetic Vertical Datum of 1929—A geodetic datum derived from a general adjustment of the first-order level nets of the United States and Canada, formerly called “Sea Level Datum of 1929.”

Selected Chemical Characteristics and Acute Toxicity of Urban Stormwater, Streamflow, and Bed Material, Maricopa County, Arizona

By Thomas J. Lopes and Kenneth D. Fossum

Abstract

The chemistry and toxicity of urban stormwater, streamflow, and bed material in the Phoenix, Arizona, area were characterized to determine if urban stormwater could degrade the quality of streams. Toxic phases of stormwater (oil and grease, suspended solids, dissolved metals, and dissolved organics) were identified to aid water-quality managers minimize the sources of toxicants. Acute aquatic toxicity tests were done using the water flea *Ceriodaphnia dubia* and fathead minnows (*Pimephales promelas*), and acute sediment toxicity tests were done using the amphipod *Hyaella azteca*. Statistical analyses also were used to determine the effect of urbanization on the quality of water and bed material and to identify toxic constituents.

Statistical analyses indicated that urban stormwater could degrade the quality of streamflow with oil and grease, pesticides, dissolved trace metals, and ammonia, and that ammonia, lead, cadmium, and zinc are released by urban activities and accumulate in bed material. Ammonia may be from fertilizers, fecal matter, and other sources. Lead probably is from vehicles that use leaded gasoline. Cadmium and zinc could be from particulate metal in oil, brake pads, and other sources.

Samples of the initial runoff from urban drainage basins appeared to be more toxic than flow-weighted composite samples, and stormwater was more harmful to fathead minnows than to *Ceriodaphnia dubia*. Streamflow samples from the Salt River were not toxic to either species, which indicates that urban stormwater could degrade the quality of the Salt River. The enhanced mortality rate of fathead minnows exposed to urban stormwater from most urban drainage basins indicated that the toxicants were more detrimental to fish than to insects and could be present in stormwater throughout the Phoenix area. The most toxic stormwater samples were collected from the drainage basins with residential and commercial land use, and the toxicity probably was due to surfactants and (or) other constituents leached from asphalt and resealant. Results of toxicity identification evaluations indicated that the toxicity of stormwater mostly was due to organic constituents; dissolved zinc and copper also appeared to contribute to stormwater toxicity. Statistical comparisons of chemical data to toxicity data indicated that organophosphate pesticides were not the toxic constituents, and the toxicity generally was due to organic constituents that were not analyzed.

The most toxic bed-material samples were collected from a drainage basin with undeveloped land use. In these bed-material samples, mortality rates were significantly higher than in samples from ephemeral channels. Comparisons between the toxicity of bed-material samples from undeveloped and urban drainage basins and between urban drainage basins and ephemeral channels showed no significant difference. In urban drainage basins, bed-material samples collected from areas where stormwater accumulates appeared to be more toxic than samples collected from areas where stormwater does not accumulate.

For bed-material samples from the undeveloped drainage basin, mortality rates strongly correlated with recoverable concentrations of zinc and moderately correlated with recoverable concentrations of copper. The high mortality rate probably was due to naturally occurring trace metals. For bed-material samples from urban drainage basins, mortality rates significantly correlated with recoverable concentrations of cadmium and zinc, which resulted from urban activities. The bioavailability of trace metals in bed material appeared to be controlled by the adsorption properties of organic carbon, iron, and manganese. Organochlorine pesticides were detected in most bed-material samples; however, mortality rates were poorly correlated with pesticide concentrations.

INTRODUCTION

Protecting ecosystems that are influenced by human activities has become a primary goal in managing public lands. Some land-use activities release toxic chemicals that could inhibit the survival, growth, and reproduction of fauna and flora in aquatic ecosystems. Thousands of chemicals are used in urban activities, and some of these chemicals accumulate on impervious surfaces and in sediments. Storms flush chemicals and sediments from streets, parking lots, and rooftops. These flushes could produce surges of toxic stormwater that are discharged into streams and lakes. Most urban stormwater in the Phoenix area is routed into drainage channels, which are tributary to ephemeral streams that include the Gila, Salt, New, and Agua Fria Rivers. Accumulation of sediments in depositional environments, such as lakes and reservoirs, could adversely affect benthic organisms because many chemicals in urban stormwater are associated with the solid phase (Lopes and others, 1995).

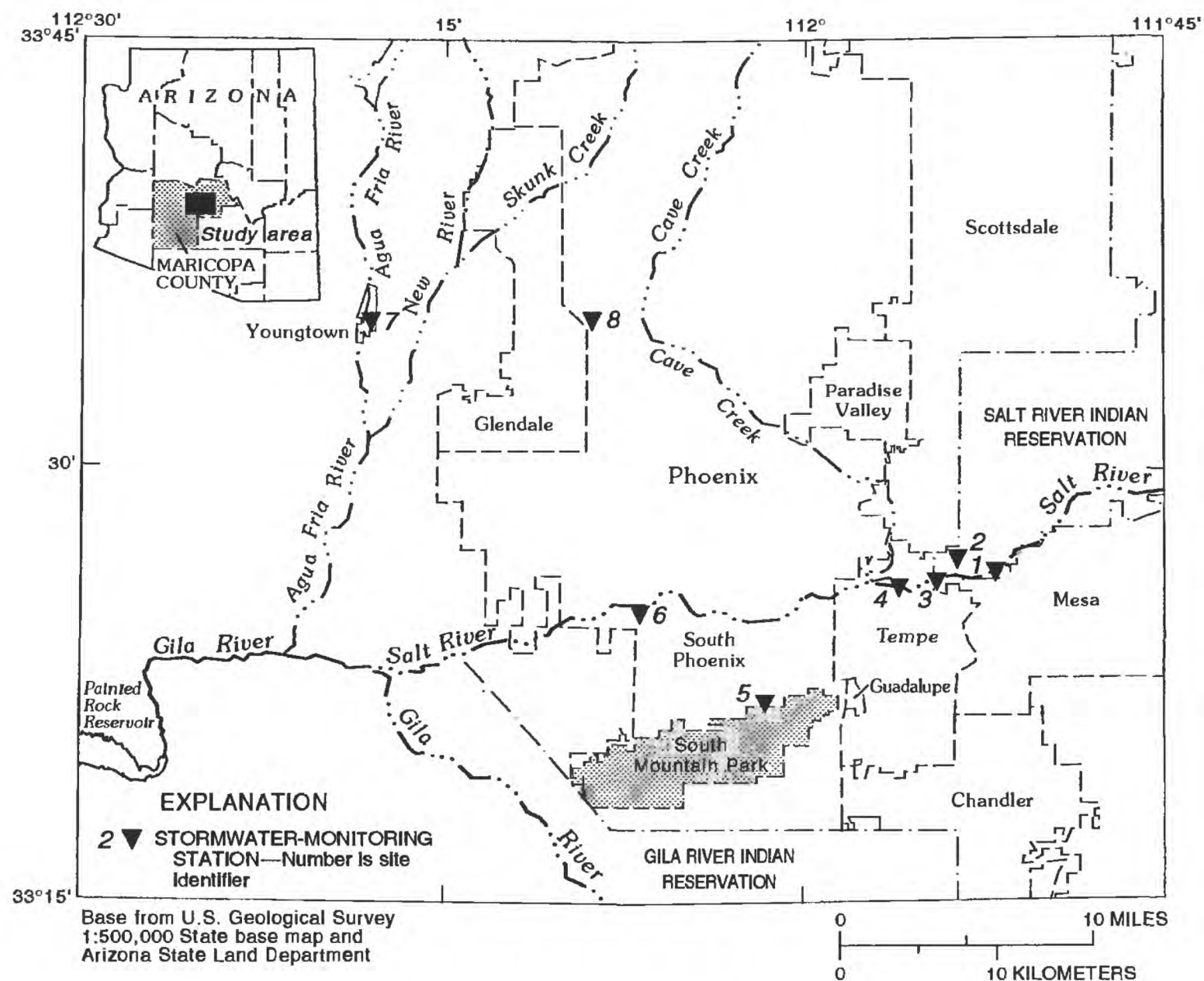
Analyzing stormwater for a myriad of constituents is not an efficient way to determine whether stormwater degrades the quality of receiving streams. Toxicity testing is an alternative technique that screens stormwater for toxic constituents by measuring the adverse effect of stormwater on selected species. Acute toxicity tests are used to determine if solutions and sediment are toxic to selected species during short-duration (between 2 to 10 days) exposures. Acute toxicity testing of urban stormwater and bed material measures the effect of land use on the quality of receiving streams and identifying the toxic phases of stormwater can help water-quality managers

implement strategies to minimize the sources of toxicants.

In 1992, the U.S. Geological Survey (USGS) in cooperation with the Arizona Department of Environmental Quality began a study of the quality of urban stormwater. The objectives of this study were to (1) characterize the chemistry and acute toxicity of stormwater from drainage basins with urban and undeveloped land use and of streamflow from the Salt River (fig. 1), (2) identify the phases of stormwater (oil and grease, suspended solids, dissolved trace metals, and dissolved organic compounds) that are causing the toxic effect, and (3) characterize the chemistry and acute toxicity of bed material from drainage basins with urban and undeveloped land use and ephemeral streams that receive urban runoff. The USGS monitored stormwater in Phoenix, Tempe, and unincorporated Maricopa County, and estimated constituent loads for Phoenix and the surrounding municipalities (Lopes and others, 1995). Data on the chemistry of urban stormwater were used in this study to focus on the constituents that were most likely to be toxic to aquatic organisms.

Purpose and Scope

This report presents selected chemical and toxicological characteristics of stormwater and bed material from urban drainage basins, characteristics of streamflow and bed material from the Salt River, and characteristics of bed material from a drainage basin with undeveloped land use and the Agua Fria River. An evaluation also is presented of phases of stormwater and constituents in bed material that caused the toxic response.



Site	Station number	Station name
1	09512060	Salt River at Alma School Road, near Mesa
2	09512162	Indian Bend Wash at Curry Road, at Tempe
3	09512165	Salt River at Priest Drive, at Phoenix
4	09512184	Box culvert at 48th Street drain
5	09512200	Salt River tributary in South Mountain Park, at Phoenix
6	09512403	27th Avenue at Salt River
7	09513700	Agua Fria River tributary at Youngtown
8	09513885	43rd and Peoria

Figure 1. Study area and stormwater-monitoring stations, Maricopa County, Arizona.

Approach

Monitoring stations were installed in drainage basins with residential, commercial, light industrial, heavy industrial, and undeveloped land use (table 1). Criteria for the selection of drainage basins included (1) drainage basins with a predominant land use so that stormwater and bed material from different land uses could be

characterized, (2) an outfall at which a stage-discharge rating could be developed so that stream discharge could be computed, and (3) definite drainage-basin boundaries so that drainage-basin characteristics could be quantified. Drainage basins with mixed land use were sampled to characterize the composite effect of many land uses on the chemistry and toxicity of urban stormwater and bed material. Streamflow and bed material were

Table 1. Drainage-basin area, land use, and impervious area of drainage basins monitored for stormwater and bed-material characteristics, Maricopa County, Arizona

[ND, industrial land uses were not differentiated, and all industrial land use was assumed to be light. NA, insufficient data to calculate percentages]

Station number	Station name or local identifier	Area of drainage basin, in acres	Land use, in percent					Impervious area, in percent
			Residential	Light Industry	Heavy Industry	Commercial	Undeveloped	
Urban drainage basins								
09512162	Indian Bend Wash at Curry Road, at Tempe, Arizona	52,480	31	5.6	ND	2.4	61	16
09512184	Box culvert at 48th Street drain.....	39	0	85	0	8	7	80
09512200	Salt River tributary in South Mountain Park, at Phoenix, Arizona	1,120	0	0	0	0	99	1
09512403	27th Avenue at Salt River	45	6	0	94	0	0	15
09513885	43rd and Peoria.....	3.4	0	0	0	97	3	94
09513700	Agua Fria tributary at Youngtown, Arizona.....	81	90	0	0	10	0	33
333543112090501	Arizona Canal Diversion Channel.....	32,770	34	12	ND	5.9	53	24
Salt River								
09512165	Salt River at Priest Drive, at Phoenix, Arizona.....	846,336	NA	NA	NA	NA	NA	NA

sampled from the Salt River, and bed material was sampled from South Mountain Park to characterize the chemistry and toxicity of nonurban water and bed material and to determine whether urban activities could degrade water and bed-material quality. Stormwater was not sampled from South Mountain Park because of lack of flow; therefore, comparisons between urban stormwater and stormwater from an undeveloped drainage basin were not possible.

The concentration of constituents in sediment depends on the chemical composition of sediments and sediment-grain size; the largest concentrations are associated with fine-grained particles (Horowitz and Elrick, 1987). Samples of bed material were sieved, and particles less than 63 microns (silt and clay) were analyzed to characterize the grain size that contains most toxicants and to make comparisons between different sites. A previous study in these basins showed that suspended solids in urban stormwater are fine sand or less in size (Lopes and others, 1995).

The duration of stormflow typically is short and makes sampling difficult. Streamflow-gaging stations in urban drainage basins were instrumented with equipment that allowed remote monitoring of rainfall and stream discharge so that field crews could collect manual grab samples before flow stopped. Streamflow-gaging stations were instrumented with automatic-pumping samplers to collect samples for the duration of each storm and to reduce the personnel requirements for the study. The Salt River, Arizona Canal Diversion Channel (ACDC), and Indian Bend Wash were sampled using manual techniques, and stream discharge was monitored at or near the sampling locations.

Stormwater samples were collected between January 1993 and May 1994 from six drainage basins with residential, commercial, light industrial, heavy industrial, and mixed land use. During the study, there was insufficient flow to characterize stormwater from the drainage basin with undeveloped land use; therefore, only bed-material samples were collected from this drainage basin.

Streamflow samples were collected from the Salt River in December 1992 and January 1993. Bed-material samples were collected from the urban and undeveloped drainage basins and ephemeral channels in September 1992 and June 1993.

Acknowledgments

The Flood Control District of Maricopa County (FCDMC) granted access to gaging stations that were used to monitor stormwater chemistry. Mattie Cahill and Jess and Sons, an auto recycler, allowed access through their property at 27th Avenue.

DESCRIPTION OF THE STUDY AREA

The study area is about 1,000 mi² in a broad, flat basin in south-central Arizona, and much of the urbanization has occurred on land that was previously used for agriculture. The basin is between 1,000 and 1,300 ft above sea level and slopes downward from east to west. The highest peak in the surrounding mountains is about 2,700 ft above sea level in South Mountain Park.

The combined population of the Phoenix area is about 2,090,000 (Maricopa Association of Governments, 1993), which is about 90 percent of the total population of Maricopa County. Residential and open spaces are the most abundant land-use types and constitute about 62 and 18 percent of the Phoenix area, respectively (Maricopa Association of Governments, 1989). The remaining 20 percent includes other land uses such as commercial and industrial land use, parks, and schools.

Maricopa County is in the northern part of the Sonoran Desert climatic zone. The mean monthly maximum temperature is 105°F in the summer, and the mean monthly minimum temperature is 40°F in the winter (U.S. Department of Commerce, 1990). Mean annual rainfall at Sky Harbor International Airport is 7.11 in. Most of the annual rainfall occurs from two weather patterns that have the same mean storm rainfall (0.46 in.) and differ mostly in storm duration (Lopes and others, 1995). About 40 percent of the annual rainfall occurs between July and October from subtropical monsoons that originate from the Gulf of Mexico

and Gulf of California and typically are short duration, high-intensity thunderstorms. About 50 percent of the annual rainfall occurs between November and March from cold fronts that originate in the Gulf of Alaska and are typically long duration, low-intensity storms. The remaining 10 percent of the annual rainfall occurs between April and June and could be the result of either type of weather pattern.

Stormwater and bed material were sampled from a drainage basin with residential land use near the intersection of Oregon and Peoria Avenues in Youngtown (USGS streamflow-gaging station 09513700, Agua Fria River tributary at Youngtown, Arizona). This site is an open channel that is tributary to the Agua Fria River, and discharge was monitored by the USGS from 1961 to 1968. About 50 percent of the homes in the basin have desert landscaping and about 50 percent have irrigated lawns. Commercial areas in the northern part of the drainage basin consist of two small shopping malls with parking lots, a gas station, and an automobile-repair shop. Peoria Avenue was resurfaced with asphalt in January 1993. The street was flushed by several storms before samples were collected; however, the asphalt may have affected the chemistry and toxicity of all stormwater samples.

Stormwater and bed material were sampled from a drainage basin with commercial land use at the northwest corner of Peoria and 43rd Avenues in Phoenix (09513885, 43rd and Peoria). Runoff at this site flows through a weir and into the ACDC, which is a stormwater conveyance that discharges into Skunk Creek. About 50 percent of the pervious area is undeveloped and about 50 percent has desert landscaping with some irrigation. Commercial businesses include a restaurant and retail stores. Resurfacing of the parking lot in March 1993 may have affected the chemistry and toxicity of the stormwater samples collected on March 26, August 24, and October 6, 1993.

Stormwater and bed material were sampled from a drainage basin with light industrial land use in Tempe near the intersection of 48th Street and the 48th Street drain (09512184, Box Culvert at 48th Street Drain). Stormwater flows through a 120-inch-wide box culvert and discharges into the 48th Street drain, which is a stormwater conveyance that discharges to the Salt River. Most of the pervious

area has irrigated landscaping. Light industrial businesses include office parks, warehouses, small manufacturing shops, and heavy-equipment rental. Commercial land use includes a hotel and a restaurant.

Stormwater and bed material were sampled from a drainage basin with heavy industrial land use in Phoenix near the intersection of Broadway and 27th Avenue (09512403, 27th Avenue at Salt River). Stormwater flows through a 96-inch-diameter culvert and discharges into the Salt River. All businesses are automobile recycling and repair shops that operate on unpaved lots except for a precast concrete-products plant, a chemical-storage facility, and a mobile-home park.

Bed material was sampled from a drainage basin with undeveloped land use at South Mountain Park (09512200, Salt River tributary in South Mountain Park, at Phoenix, Arizona), where continuous stream-discharge records have been collected since 1961. In 1973, the drainage basin was designated as a mountain preserve and is used as an outdoor recreation area. The drainage basin is tributary to the Salt River; however, in 1979, a retention pond that reduces the amount of streamflow that reaches the river was constructed downstream from the streamflow-gaging station. During the study, there was insufficient flow at this site to collect stormwater samples.

Stormwater was sampled from the ACDC at 51st Avenue (333543112090501). Bed material was sampled from the ACDC where it intersects 51st, 59th, and 67th Avenues. ACDC is an urban stormwater conveyance that is concrete lined to 51st Avenue and grass lined from 51st Avenue to Skunk Creek. The grass-lined sections are irrigated and used for recreation.

Stormwater was sampled from Indian Bend Wash at the streamflow-gaging station, Indian Bend Wash at Curry Road, at Tempe, Arizona (09512162). Two bed-material samples were collected from Indian Bend Wash where it intersects Shea Boulevard and Curry Road. The drainage area at Curry Road is about 82 mi² and is tributary to the Salt River. About 39 percent of the drainage basin is developed; however, about 70 percent of the basin near the streamflow-gaging station is urbanized. Indian Bend Wash was sampled when streamflow consisted of runoff from the City of Scottsdale and the northern part of the

City of Phoenix. Runoff from undeveloped parts of the drainage basin contribute to streamflow only during large storms. Much of Indian Bend Wash is grass lined and used for recreation and includes several golf courses and manmade lakes.

Streamflow was sampled from the Salt River at the streamflow-gaging station, Salt River at Priest Drive, at Phoenix, Arizona (09512165). Bed material was sampled from the Salt River where it intersects Priest Drive, 24th Street, and 35th Avenue. The Salt River is ephemeral and typically flows when dams upstream from Maricopa County release water or during large storms in the Phoenix area. Streamflow samples were collected between December 1992 and February 1993 when water was released from dams on the Verde and Salt Rivers. Streamflow reached a maximum of 129,000 ft³/s on January 8, 1993. Most ephemeral channels in the Phoenix area were flooded, which probably scoured bed material and chemicals that had accumulated in the Salt River, ACDC, and Indian Bend Wash before samples were collected in June 1993.

Bed material was sampled from the Agua Fria River, which drains a mostly undeveloped area of about 2,000 mi² at the sampling sites. Samples were collected upstream, downstream, and at the mouth of the tributary that drains urban stormwater from Youngtown (09513700). Samples were collected in September 1992 and were not affected by floods.

DATA-COLLECTION METHODS

Runoff and precipitation data were collected at urban drainage basins between January 1993 and March 1994 using the following equipment:

Campbell Scientific Instruments, Inc.,
CR10 datalogger and SM192 storage
module;

Sierra-Misco Environment Ltd., model
2500 tipping-bucket rain gage;

Druck PDCR 940 pressure transducer;

Conoflow and pressure-regulator system;

Isco, Inc., Model 3700 automatic-pumping
sampler; and

Motorola MC310 cellular telephone.

The datalogger was programmed to record instrument readings, calculate stream discharge, and activate the automatic-pumping sampler when a specified volume of water had been discharged

from the drainage basin. The datalogger also initiated a telephone call to a hydrologist when precipitation or discharge was measured so that the hydrologist could make manual discharge measurements and collect grab samples during runoff. Data were recorded at 1-minute intervals when either rainfall or stream discharge was measured. Data were recorded once a day (at midnight) if dry conditions persisted.

Precipitation

Precipitation was measured in urban drainage basins and at South Mountain Park with a tipping-bucket rain gage that was calibrated to tip when 0.01 in. of rainfall was collected. A pulse was transmitted to the datalogger each time 0.01 in. of rainfall was collected. The number of pulses in each minute was recorded to measure rainfall intensity. Rainfall during each successive minute was recorded and summed to obtain accumulated rainfall.

Stream Discharge

Gage height was measured at urban drainage basins and South Mountain Park using a Conoflow and pressure-regulator system. The Conoflow and pressure regulator maintain a constant rate of nitrogen flowing through a tube that extends from the gaging station to an orifice at the bottom of the channel or culvert. Greater pressure is required to maintain a constant flow rate through the tube as stage increases. Pressure in the tube was measured with the pressure transducer, which was calibrated to within 0.02 ft and, except for 27th Avenue, was placed 3 to 5 ft underground to reduce effects of temperature on pressure-transducer measurements. At 27th Avenue, the transducer was placed in the culvert and insulated. Temperature of the pressure transducer varied by about 4°F/d, and temperature corrections were not necessary. Gage height at the ACDC was measured by the Flood Control District of Maricopa County using a pressure transducer. Gage height at Indian Bend Wash was measured by a float, and gage height at the Salt River at Priest Drive was measured using a wire-weight gage.

Stage-discharge ratings for all urban drainage basins were developed on the basis of channel geometry and slope by using the slope-conveyance method (Kennedy, 1984). A stage-discharge rating that was based on manual discharge measurements had been developed at South Mountain Park. Instantaneous discharge measurements were made when water samples were collected from the Salt River and the ACDC. Stream-discharge measurements were made at all streamflow-gaging stations using either a pygmy or Price AA meter and the 0.6-depth or 0.2- and 0.8-depth wading method or a calibrated bucket (Rantz and others, 1982). Stream-discharge measurements and associated gage heights compared well with developed stage-discharge ratings.

Instantaneous discharge rates at urban drainage basins and South Mountain Park were computed by programming the data logger with a log-normal regression equation that was fit to the stage-discharge rating of each site (Kolb, 1983). The log-normal equations compared well with all discharge ratings (correlation coefficients were 0.99 or greater). Stream-discharge volumes were computed by multiplying the mean of two consecutive discharge-rate measurements by 60 seconds to obtain the mean volume of stream discharge during that 1-minute interval. The mean volumes were summed to obtain the total volume of runoff.

Stormwater, Streamflow, and Bed-Material Samples

Stormwater samples were collected from urban drainage basins by automatic-pumping samplers and automatic-grab samplers to collect flow-weighted composite samples and samples of the initial runoff from a storm (first flush). The automatic-pumping sampler is a portable, nonrefrigerated unit calibrated to pump a specified volume of stormwater. Twenty-four 1-liter, teflon-lined, polyethylene bottles were used to hold discrete samples for chemical analyses. Samples were pumped when a specified volume of water discharged from the drainage basin. Discrete samples were later combined to obtain a flow-weighted composite sample. Samples for toxicity analyses were collected with another

automatic-pumping sampler that was equipped with 24 1-liter polyethylene bottles. The automatic-grab sampler is an 8-liter Nalgene carboy fitted with a float valve and placed in the ground so that the top of the carboy is level with the ground surface. Initial runoff from a storm flows into the carboy, and the float minimizes mixing of the first-flush water with the rest of the runoff. A steel grate, coated with an epoxy-based paint, was used to prevent the first-flush samplers from floating away and to keep traffic from damaging the samplers. Two or three automatic-grab samplers were installed at each site to collect enough water for chemical and toxicity analyses.

Streamflow samples were collected from the Salt River, ACDC, and Indian Bend Wash by using the equal-width-increment method and by collecting grab samples. Equal-width-increment samples were collected at equal distances perpendicular to the direction of streamflow. Samples were then composited to obtain a single sample that is representative of the stream at a discrete time.

Stormwater and streamflow samples were placed on ice and transported to the field office for processing. Samples for chemical analyses were poured into a teflon-lined, stainless-steel churn splitter to split the samples into the bottles required for each analysis. Samples for toxicity analyses were poured into 8-liter Nalgene carboys and chilled at 39°F. Samples for dissolved-constituent analyses were filtered using a 0.45-micron effective pore-size cellulose filter. Preservatives then were added to sample bottles as required. All components of the sampling equipment that came in contact with samples for chemical analyses were constructed of either glass, teflon, or stainless steel except for the distribution hose in the automatic-pumping sampler, which was silicon rubber. Samples for toxicity analyses came into contact with glass, teflon, stainless steel, polyethylene, and the Nalgene carboys. Equipment that was in contact with sample water was cleaned by washing with Liquinox followed by rinsing with large quantities of tap water, a rinse of ultrapure methanol, and a final rinse of deionized water.

Bed-material samples from urban drainage basins, South Mountain Park, and the Agua Fria River were collected at specific points. Bed-material samples were collected at equal increments

across each reach of the Salt River, ACDC, and Indian Bend Wash and were composited for chemical and toxicity analyses. Bed material was sieved, and particles that were less than 63 microns were collected for analyses. Samples for trace-metal, nutrient, inorganic plus organic carbon, inorganic carbon, and toxicity analyses were collected using plastic trowels and plastic sieves and screens. Samples for organochlorine and organophosphate pesticide analysis were collected using stainless-steel spoons and brass sieves and screens. Whole-sediment samples were collected using stainless-steel spoons for oil and grease and particle-size distribution analyses. All equipment that came in contact with sediment were washed with Liquinox followed by rinsing with large quantities of tap water and a final rinse of deionized water.

Quality-assurance procedures were followed throughout the study to identify potential problems in data collection caused by sampling methods or equipment contamination. These procedures included the use of duplicate and equipment-blank samples. Duplicate samples of stormwater and bed material were used to check the precision of laboratory analyses and the sample-splitting process. Analyses from duplicate samples indicated that laboratory analyses and the sample-splitting process were producing consistent results. Equipment-blank samples were collected by pumping and processing inorganic-free and organic-free water in the same manner as sample water and by rinsing the automatic-grab sampler and epoxy-painted grates with deionized water. Analyses from equipment blanks indicated that minimal cross contamination occurred between sampled storms and that the grates were not contaminating samples with trace metals.

The representativeness of samples collected with the automatic-pumping sampler was evaluated by collecting manual, depth-integrated samples simultaneously with automatically collected samples. Samples were analyzed for concentrations of nutrients, selected metals, and suspended solids. Results indicated that flow in shallow, swift channels is well mixed and can be accurately represented by samples collected at a single point.

Samples for chemical analyses were shipped to the USGS National Water Quality Laboratory for analysis. Acute toxicity, total hardness, pH, specific

conductance, total alkalinity, total chlorine, and total ammonia were analyzed by Aquatic Consulting and Testing, Inc., Tempe, Arizona. Dissolved and total alkalinity concentrations were measured in the Tempe office of the USGS.

METHODS OF MEASURING THE ACUTE TOXICITY OF WATER AND SEDIMENT

Acute toxicity tests that were performed on water samples were dilution series, static renewal tests, which were done using standard procedures (U.S. Environmental Protection Agency, 1990). Stormwater and streamflow samples were diluted with moderately hard reconstituted water (MHRW) to obtain mixtures that had 100, 50, 25, 12.5, and 6.25 percent (by volume) of stormwater or streamflow; mixtures were kept between 66° to 70°F during the test. MHRW was prepared from reagent grade salts and had a total hardness of 80 to 100 mg/L as CaCO₃ and pH of 7.4 to 7.8. Fathead minnows (*Pimephales promelas*; 5- to 10-days old) and the water flea *Ceriodaphnia dubia* (hereafter referred to as *C. dubia*; less than 24-hours old) were used as surrogate species and fed before testing. The mixtures were renewed every 24 hours to more accurately simulate a natural environment where the amount of toxicants in the water is not limited. Acute toxicity tests using fathead minnows lasted 96 hours and tests using *C. dubia* lasted 48 hours. The trimmed Spearman-Kärber, probit, or graphical methods (Rand and Petrocelli, 1985) were used to estimate the concentration of stormwater (in percent, by volume) that caused a certain mortality rate (for example, percentage of stormwater that killed 50 percent of the organisms, LC50). Water and bed-material samples were considered acutely toxic if 20 percent or more of the organisms died when exposed to 100 percent of the sample; a mortality rate of 20 percent is significantly different from the mortality rate measured in positive controls (U.S. Environmental Protection Agency, 1990).

The toxic strength of a constituent or sample can be expressed in toxic units (TU), which is the concentration of a constituent or sample divided by the LC50 value (Rand and Petrocelli, 1985). For

example, a stormwater sample with an LC50 value of 25 percent has 4 TU (100 percent stormwater divided by the LC50). TU values are used instead of LC50 values because TU values increase as toxicant concentrations increase, whereas LC50 values decrease as toxicant concentrations increase.

Fathead minnows and *C. dubia* were selected as surrogate species because: (1) no native species inhabit ephemeral streams in urbanized Maricopa County; (2) these are standard species used in toxicity testing; (3) substantial data exist on their mortality rates when exposed to a large number of toxicants; and (4) their mortality rates have been correlated with the mortality rates of other organisms (Mayer and Ellersieck, 1986). Mayer and Ellersieck (1986) compared mortality rates of species exposed to pesticides and determined that insects generally were the most sensitive species, followed by crustaceans, fish, and amphibians. Of the most commonly tested organisms, daphnids were more sensitive 58 percent of the time, rainbow trout, 35 percent; blue gill, 5 percent; and fathead minnows, 2 percent.

All aquatic toxicity tests were performed in duplicate to check precision of the test. Positive controls using only MHRW were performed with all samples to ensure that mortality was caused by exposure to stormwater rather than by poor health of the organisms. Negative controls, using copper sulfate, were performed monthly to check laboratory protocol and to ensure that the brood stock produced organisms that responded within acceptable limits to a toxicant with a known LC50 value. Analyses from duplicate samples, positive controls, and negative controls indicated that the laboratory was producing consistent results, that toxic constituents in samples caused the mortality, and that the mortality of organisms was predictable when exposed to various concentrations of a known toxicant.

Toxicity identification evaluations (TIE's) were performed on most stormwater samples that were toxic. TIE is a method of identifying the toxic components of a solution and consists of three procedures: (1) toxicity characterization (U.S. Environmental Protection Agency, 1991); (2) toxicity identification (U.S. Environmental Protection Agency, 1993a); and (3) toxicity confirmation (U.S. Environmental Protection Agency, 1993b). Toxicity characterization was the

only procedure performed for this study because toxicity-reduction strategies would probably focus on certain phases of stormwater rather than a specific constituent. Results of a previous study (Lopes and others, 1995) indicated that oil and grease, suspended solids, dissolved metals, and dissolved organic compounds were the most likely phases of stormwater that could be toxic. Volatile organic compounds were seldom detected, and the pH of most stormwater samples was typical of most rivers, indicating that these constituents were unlikely to cause toxicity in stormwater.

TIE's were performed by sequentially extracting oil and grease, suspended solids, dissolved metals, and dissolved organic compounds from toxic samples. After extracting potentially toxic phases, toxicity tests were performed on the solutions. A reduction in mortality indicated that the extracted phase caused a toxic response. Stormwater samples were split into five subsamples when a toxic response was measured during the definitive acute toxicity (original) test (fig. 2). Oil and grease were extracted from one of the subsamples by decanting the upper inch of water, passing it through activated carbon and glass wool, and returning the water to the container. Oil and grease were extracted, and then suspended solids were removed from the second subsample by filtering the subsample through a 0.45-micron polycarbonate filter. After extracting oil and grease and suspended solids from the third subsample, dissolved metals were chelated with various concentrations of ethylenediaminetetraacetic acid (EDTA), which reduces the toxicity of dissolved metals (U.S. Environmental Protection Agency, 1991). If more EDTA is added than is needed to chelate dissolved metals, the excess EDTA can be toxic. Oil and grease were extracted, and suspended solids were filtered from a fourth subsample using an 0.7-micron glass-fiber filter. Next, two aliquots of the fourth subsample were adjusted to either pH 3 or 9 with hydrochloric acid or sodium hydroxide and a third aliquot was untreated. The three aliquots were passed through a column of octadecyl to extract anionic, cationic, and neutral organic compounds, and then the pH of the aliquots was adjusted to the initial pH. The fifth subsample was the baseline and was untreated. The baseline was used to determine if sample degradation occurred between the time the original test and TIE

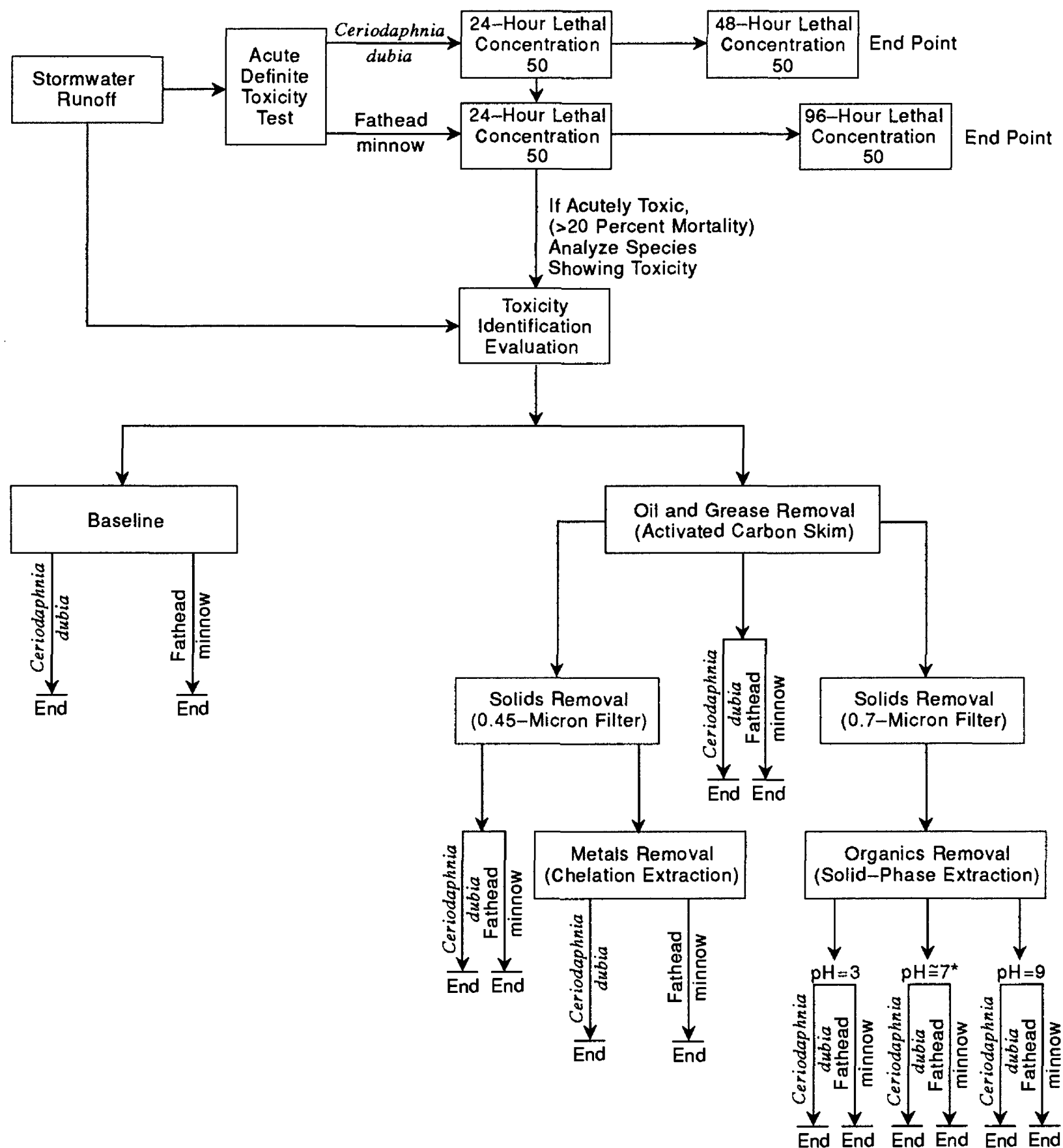
were performed and to measure reduction in toxicity due to each extraction. Acute toxicity tests were performed concurrently on the baseline, the extracted solutions, and the positive controls.

Quality assurance of each TIE included use of duplicates and positive controls. Duplicates were performed on the oil and grease and suspended-solids extractions and on the organics extraction using solutions after 25 and 150 mL had passed through the octadecyl column. The extraction procedure was performed on dilution water and was used as a positive control. Duplicates compared well, and significant mortality was not observed in the controls. This comparison and absence of mortality indicated that procedures for removing oil and grease, suspended solids, and organic compounds did not contribute to the toxicity of stormwater samples.

Acute sediment toxicity tests were performed on urban bed-material samples using guidelines of the American Society of Testing and Materials (1990). Sediment toxicity tests consisted of placing 100 g of sieved (less than 63 microns) bed material in each of five 1-liter glass beakers, adding 400 mL of hard reconstituted water (HRW; hardness of 160 to 180 mg/L as CaCO_3 ; pH 7.6 to 8.0), and letting the mixtures equilibrate for 24 hours. After equilibration, 20 *Hyaella azteca* (hereafter referred to as *H. azteca*; less than 16-days old, 2 to 3 mm in size) were added to each beaker and observed for 10 days. Organisms remained in the same mixture during the test and were fed TetraMin three times a week. *H. azteca* is an epibenthic amphipod that can live in a wide range of sediment grain size (Ingersoll and Nelson, 1990) and is a sensitive indicator of the presence of contaminants associated with sediments (Ingersoll and Nelson, 1990; Landrum and Scavia, 1983; Nebeker and others, 1984). The acute mortality rate, in percent, was calculated at the end of the test by counting the number of organisms that died.

All sediment toxicity tests were performed with five replicates to check the precision of the test and to ensure that equipment contamination did not cause the mortality. Positive controls consisted of silica sand and HRW, and had mortality rates of 2 and 8 percent. Sand was purchased from a hardware store and tested as a positive control, but was unsuitable because significant mortality rate (19 percent) was measured. Negative controls

TEST PROTOCOL FLOW DIAGRAM
(Modified Toxicity Identification Evaluation)



*Initial pH

Figure 2. Extraction sequence for toxicity identification.

consisted of copper sulfate, silica sand, and HRW and were used to ensure that organisms responded to a known toxicant. Negative controls done in September 1992 and June 1993 yielded the same LC50 for copper (70 µg/L). This value compares well with results of another study (Schubauer-Berigan and Dierkes, 1993), which calculated an LC50 for copper of 87 µg/L in very hard (300 mg/L as CaCO₃) reconstituted water. Analyses of replicates, positive controls, and negative controls indicated that the laboratory was producing consistent results, that mortality was caused by exposure to bed-material samples, and that the mortality of organisms was predictable when exposed to various concentrations of a known toxicant.

CHEMICAL CHARACTERISTICS OF URBAN STORMWATER, STREAMFLOW, AND BED MATERIAL

A total of 36 water samples were collected, including 14 first-flush samples, 14 flow-weighted composite samples, 4 equal-width-increment samples from urbanized tributaries, and 4 equal-width-increment samples from the Salt River. First-flush samples were collected at the drainage basins with a single land use during the first few minutes of flow. Equal-width-increment samples from the ACDC and Indian Bend Wash were collected near the peaks and tails of the hydrographs, and samples from the Salt River were collected at various discharge rates.

All water samples were analyzed for oil and grease, suspended solids, total chlorine, total ammonia, total hardness, dissolved trace metals, and acute toxicity. Organophosphate pesticides also were analyzed in flow-weighted composite and equal-width-increment samples. Organochlorine pesticides, acid/base/neutral organic compounds, and volatile organic compounds were not analyzed because concentrations of these constituents in flow-weighted composite and streamflow samples collected during an earlier study typically were less than detection limits (Lopes and others, 1995).

A total of 30 bed-material samples were collected including 13 samples from urban drainage basins, 11 samples from ephemeral channels, and 6 samples from South Mountain Park. All bed-

material samples were analyzed for oil and grease, organic and inorganic carbon, total recoverable trace metals and nutrients, organochlorine and organophosphate pesticides, and acute toxicity.

Lead, molybdenum, beryllium, and silver had the lowest concentrations of dissolved trace metals that were tested and were detected in one to four samples of stormwater and streamflow. Total chlorine and the pesticides—parathion, trithion, disyston, phorate, DEF, and fonofos—were detected in zero to four samples of stormwater and streamflow. Perthane, endosulfan, aldrin, endrin, heptachlor, ethion, malathion, methylparathion, parathion, trithion, polychlorinated naphthalene (PCN), methoxychlor, and mirex were detected in five or fewer bed-material samples. Chemical analyses indicated that these constituents were not significant in stormwater and bed material from the Phoenix area.

Urban Stormwater

Initial runoff from Youngtown, Peoria, and 48th Street was typically black in color, which could be from oil and grease, particulates from ground-up tires, or other sources. The strainer for the automatic-pumping sampler at Youngtown and Peoria was coated with oil and grease after most storms. Oil and grease were detected in about 80 percent of urban stormwater samples, and the median concentration was 4 mg/L for first-flush and for flow-weighted composite samples. Concentrations of oil and grease in samples from the ACDC and Indian Bend Wash ranged from less than 1 to 8 mg/L, and the maximum concentration measured in all urban stormwater samples was 10 mg/L.

Total ammonia was detected in about 80 percent of urban stormwater samples, and the median concentration measured in first-flush and flow-weighted composite samples was 1.3 and 0.6 mg/L, respectively. The concentration of total ammonia in samples from the ACDC and Indian Bend Wash ranged from less than 0.1 to 8.9 mg/L, and the maximum concentration measured in all stormwater samples was 11.0 mg/L from Peoria.

Median total hardness concentrations of first-flush and flow-weighted composite samples were 150 and 48 mg/L as CaCO₃, respectively, and

ranged from 36 to 500 mg/L and from 20 to 228 mg/L, respectively. Total hardness in samples from the ACDC and Indian Bend Wash ranged from 44 to 92 mg/L. About 50 percent of all stormwater samples were soft (less than 60 mg/L), 30 percent were moderately hard to hard (between 61 and 180 mg/L), and 20 percent were very hard (more than 180 mg/L). Soft stormwater typically was associated with flow-weighted composite samples. Hardness affects the toxicity of certain constituents (Rand and Petrocelli, 1985).

Median specific-conductance values of first-flush and flow-weighted composite samples were 296 and 142 $\mu\text{S}/\text{cm}$, respectively, and values ranged from 84 to 7,100 $\mu\text{S}/\text{cm}$ and from 44 to 414 $\mu\text{S}/\text{cm}$, respectively. Specific-conductance values in samples from Indian Bend Wash and the ACDC ranged from 138 to 251 $\mu\text{S}/\text{cm}$. In general, specific-conductance values were higher in first-flush

samples than flow-weighted composite samples, indicating that soluble constituents were washed from exposed surfaces during the initial part of a storm and diluted during the latter part of the storm (figs. 3 and 4). Specific conductance, however, increased during some storms at 48th Street and 27th Avenue. The increase could be due to an uneven distribution of rainfall in the drainage basin and runoff arriving at the gaging station at different times. Large concentrations of toxicants could be associated with high specific-conductance values, which did not always occur with the first flush.

Median concentrations of suspended solids were about 160 mg/L for first-flush and flow-weighted composite samples. Suspended-solids concentrations ranged from 45 to 1,190 mg/L in first-flush samples and from 40 to 544 mg/L in flow-weighted composite samples. Suspended-

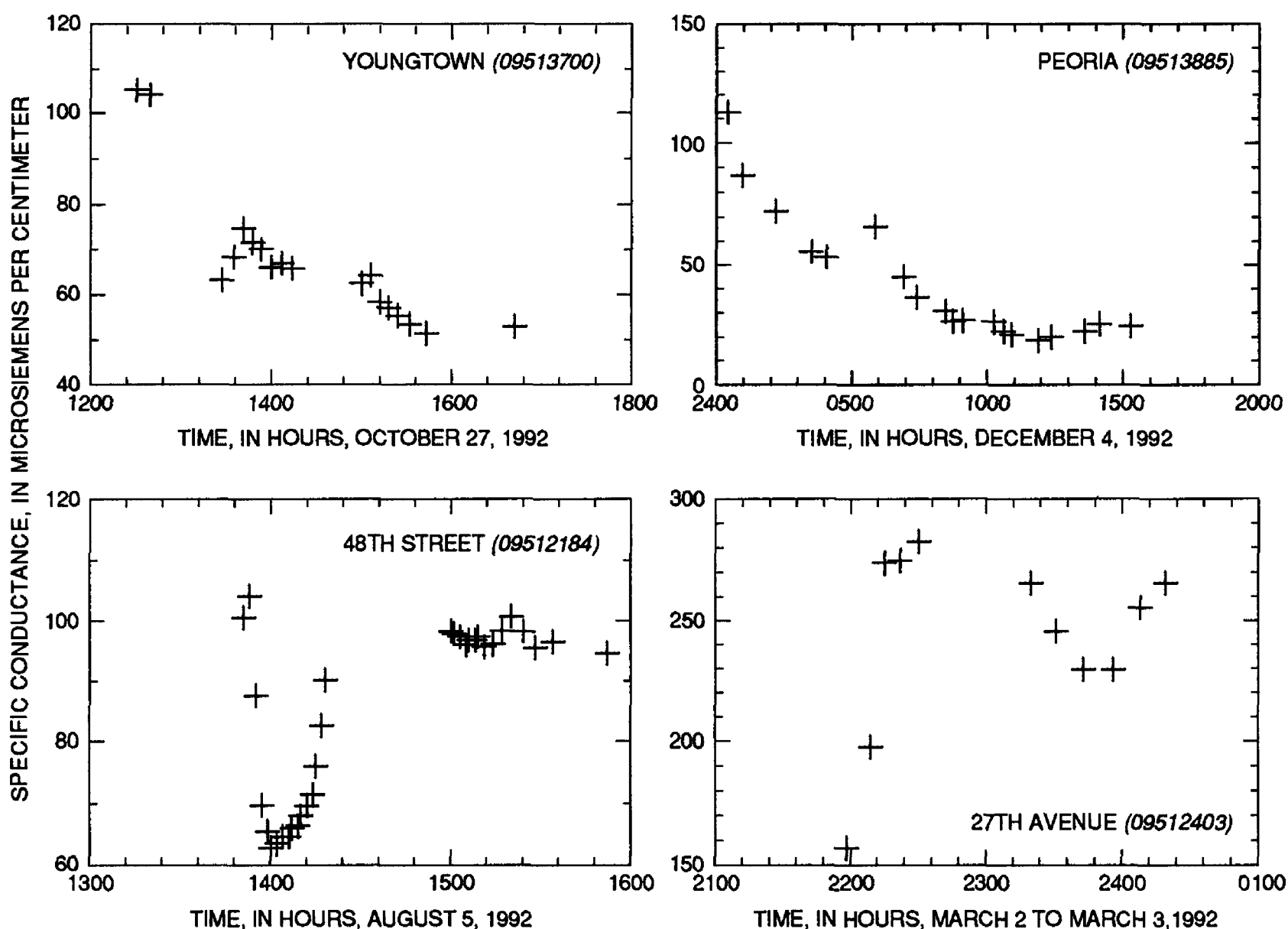


Figure 3. Specific conductance of stormwater as a function of time during a storm, Maricopa County, Arizona. Time refers to Mountain Standard Time and is given according to the 24-hour clock. U.S. Geological Survey stormwater-monitoring station number is in italics.

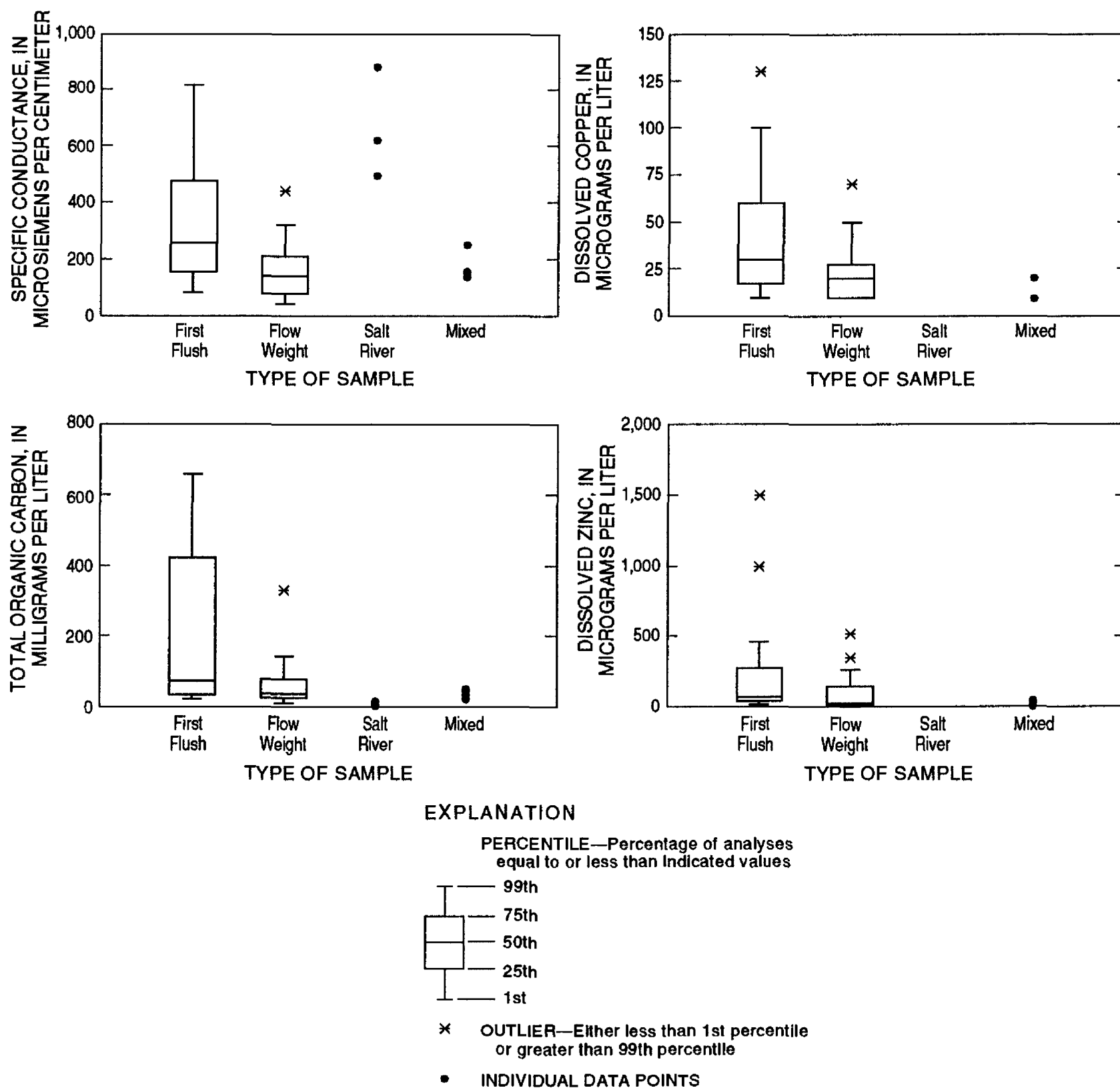


Figure 4. Ranges in specific-conductance values and constituent concentrations in stormwater and streamflow, Maricopa County, Arizona. The sample of the initial runoff (first flush) collected near 43rd Avenue and Peoria (USGS streamflow-gaging station 09813885), August 24, 1994, which had 5,000 milligrams per liter of total organic carbon and a specific conductance of 7,100 microsiemens per centimeter, was not included in the boxplots. Constituent concentrations measured in this outlier were too extreme to plot.

solids concentrations ranged from 30 to 394 mg/L in samples from the ACDC and Indian Bend Wash.

The median concentrations of total organic carbon in first-flush and flow-weighted composite samples were 80 and 36 mg/L, respectively, and are larger than concentrations found in bogs (Thurman, 1985). Concentrations of total organic carbon ranged from 23 to 49 mg/L in samples from the

ACDC and Indian Bend Wash. Total organic carbon in first-flush samples had a significant rank correlation with suspended-solids concentrations (0.85), which indicates that most organic carbon occurred as particulates. The rank correlation is the correlation of two variables using their relative ranking rather than absolute values, which gives outliers less weight in the correlation. Total organic

carbon in flow-weighted composite samples and samples from the ACDC and Indian Bend Wash did not correlate with suspended solids (-0.05). Samples commonly had a brown color after filtering with an 0.45-micron filter indicating that organic carbon was in the dissolved phase.

Concentrations of the organophosphate pesticides, diazinon and malathion, ranged from 0.01 to 1.1 $\mu\text{g/L}$ and were detected in 94 percent of urban stormwater samples. The largest concentrations of diazinon and malathion were measured in samples from the ACDC. Concentrations of chlorpyrifos ranged from 0.02 to 0.13 $\mu\text{g/L}$ and were detected in 66 percent of urban stormwater samples. Chlorpyrifos was the only other pesticide that was detected in samples from the ACDC and Indian Bend Wash. Methylparathion, dysiston, DEF, and phorate were detected in 7 to 42 percent of flow-weighted composite samples, and concentrations ranged from less than 0.01 to 0.11 $\mu\text{g/L}$. Organophosphate pesticides commonly are used in urban environments and are applied frequently because they readily degrade into nontoxic compounds (Fukato, 1987). The weak correlation of diazinon and malathion to suspended solids (0.25 and 0.20, respectively) and to total organic carbon (-0.18 and 0.09, respectively) indicated that these pesticides were associated with the dissolved phase.

Certain organic compounds are weak acids that appear to make urban stormwater slightly acidic. pH values of stormwater were typically between 6 and 7; whereas, most streams in Arizona have alkaline pH values (pH values are typically between 7.5 and 9.0; Smith and others, 1994). Total organic carbon significantly correlated with the pH of urban stormwater and streamflow samples (-0.58 ; fig. 5). The lowest pH value (4.9) was measured in a first-flush sample from Peoria and the highest values (between 7.0 and 7.8) were measured in samples from 27th Avenue and Indian Bend Wash. Peoria has 94 percent impervious area and little exposed soil to react with stormwater, whereas 27th Avenue and Indian Bend Wash have mostly pervious area. The pH of stormwater appears to be a function of the percentage of impervious area in a drainage basin.

pH measurements made on unfiltered samples for chemical analyses differed from pH measurements made on unfiltered samples for

toxicity testing by as much as 1.4 units. pH in unfiltered samples for chemical analyses were significantly biased to lower pH values, and the median difference was 0.4 units less than the pH in samples for toxicity testing (fig. 5). pH varied by about 0.1 to 0.2 units during the toxicity test; one sample changed by 0.7 units. The stable pH during the toxicity test indicated that stormwater and sediments reached equilibrium by the time tests started.

Concentrations of dissolved trace metals were typically larger in first-flush than in flow-weighted composite samples (fig. 4). These larger concentrations could be due to flushing of metal salts in the initial runoff, dilution of the flow-weighted composite samples, or because metals are more soluble in first-flush water. Statistical comparisons between first-flush samples and flow-weighted composite samples, however, indicated that first-flush samples had significantly larger concentrations only for dissolved barium, strontium, calcium, magnesium, and cadmium. Flow-weighted composite samples and samples from drainage basins with mixed land use had similar concentrations. An insufficient number of data, however, were collected to make statistical comparisons between stormwater samples from mixed land-use basins and samples from single land-use basins. Comparisons were made using a one-way analysis of variance (ANOVA) with Tukey's W multiple-comparison test (Statware, Inc., 1990).

Dissolved iron, manganese, barium, strontium, and zinc were detected in all stormwater samples. Median concentrations of dissolved iron and manganese were 80 and 66 $\mu\text{g/L}$, respectively, and concentrations ranged from 10 to 25,000 $\mu\text{g/L}$. Median concentrations of dissolved barium and strontium were 24 and 135 $\mu\text{g/L}$, respectively, and concentrations ranged from 6 to 770 $\mu\text{g/L}$. The median concentration of dissolved zinc was 42 $\mu\text{g/L}$, and concentrations ranged from 4 to 1,500 $\mu\text{g/L}$. Dissolved copper was detected in 75 percent of stormwater samples, had a median concentration of 20 $\mu\text{g/L}$, and concentrations ranged from 10 to 130 $\mu\text{g/L}$. Other dissolved trace metals were detected in about 50 percent or fewer of the stormwater samples.

Aerated streams with pH values between 6.5 and 8 typically have about 10 $\mu\text{g/L}$ of dissolved

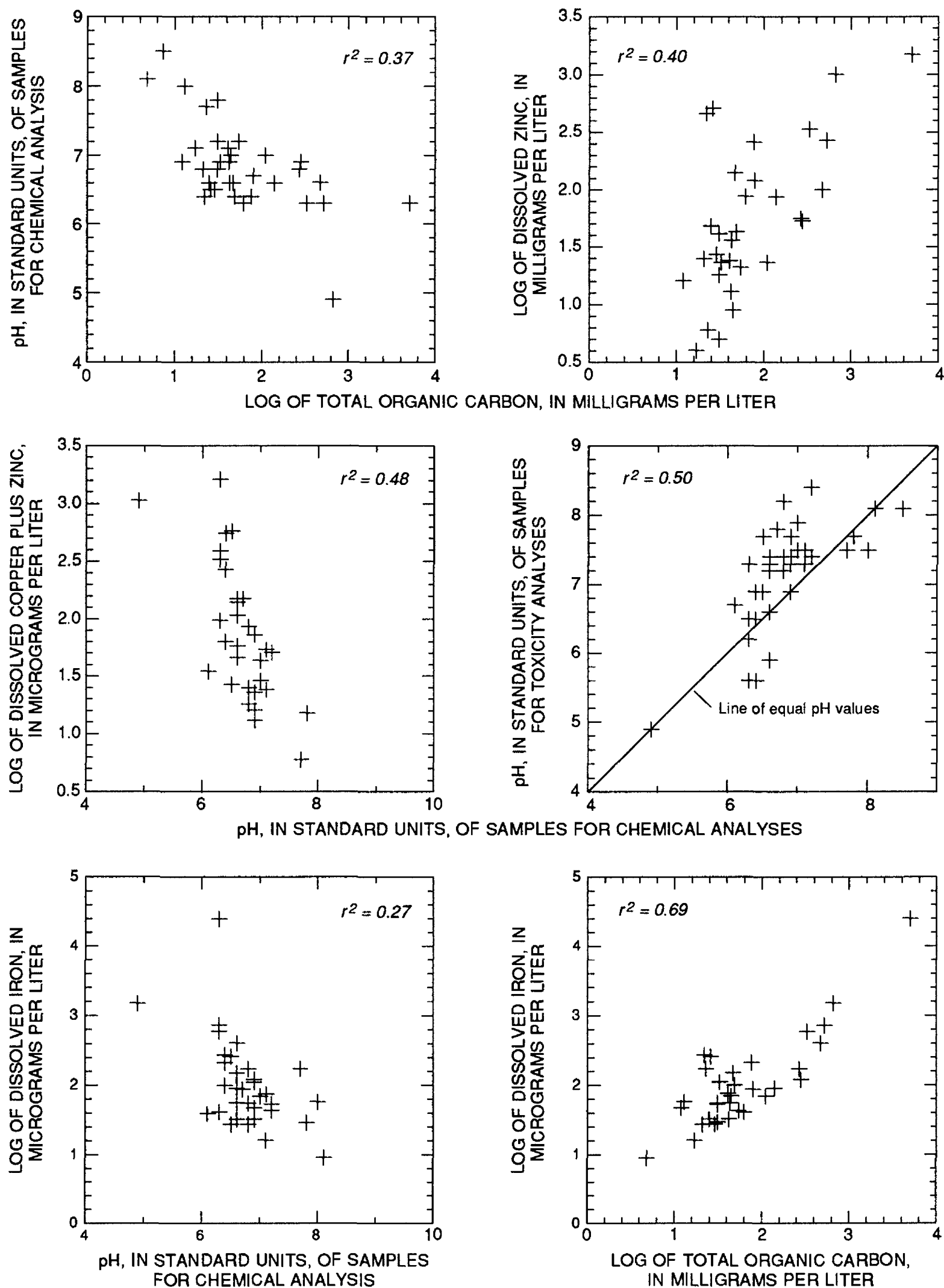


Figure 5. Concentration of selected constituents as a function of concentration of total organic carbon and as a function of pH, and pH measured in samples for toxicity analyses as a function of pH measured in samples for chemical analyses, Maricopa County, Arizona.

iron (Hem, 1985). The large concentrations of dissolved iron indicate that iron is either stabilized by organic complexes and (or) reducing conditions exist in stormwater. Oxidation of organic carbon by dissolved oxygen lowers the oxidation-reduction (redox) potential if concentrations of organic carbon are larger than dissolved oxygen concentrations. Iron and manganese solubility increase as the pH and redox potential decrease (Hem, 1985). For all samples, dissolved iron and manganese concentrations were positively correlated with total organic carbon (0.64 and 0.75, respectively), inversely correlated with dissolved oxygen (-0.93 and -0.72 , respectively), and inversely correlated with pH (-0.48 and -0.68 , respectively; fig. 5). The large concentrations of organic carbon appeared to lower the pH, and may form stable complexes with iron, lower the redox potential of stormwater, and control trace-metal solubility. Trace metals that adsorbed to iron and manganese oxyhydroxides would be released into solution when the oxyhydroxides dissolve. These releases could account for the strong correlation between iron, manganese, and dissolved trace-metal concentrations (correlations were between 0.4 and 0.9 for most trace metals). Iron and manganese oxyhydroxides have a large adsorption capacity and concentrate trace metals in sediments (Horowitz and Elrick, 1987).

The concentrations of dissolved trace metals measured in urban stormwater may be different from the concentrations that organisms were exposed to during the acute toxicity test. Samples for trace-metal analyses were filtered and preserved soon after collection; equilibrium between the solution and suspended solids probably was not established before samples were filtered. Samples for toxicity analyses were not filtered and stored at 39°F in sealed containers for 1 to 3 days. The lack of filtration would allow more time for solutions and suspended solids to equilibrate. Lowering the redox potential in the sealed containers would increase the solubility of certain trace metals. The redox potential probably changed again during the toxicity test when samples were aerated to maintain 40 percent or greater saturation of dissolved oxygen. The changes in pH and redox potential probably affected dissolved trace-metal concentrations, and the concentrations measured in

stormwater may not represent the concentrations that organisms were exposed to during the toxicity test.

Dissolved trace-metal concentrations, percentage of land use, and percentage of impervious area were not significantly correlated at a level of 5 percent, except for zinc. Correlations were calculated using concentrations of dissolved trace metals measured in flow-weighted composite samples collected from the urban drainage basins since 1991. Dissolved zinc positively correlated with the percentage of commercial land use (0.60), percentage of undeveloped land use (0.52), percentage of impervious area (0.75), and inversely correlated with percentage of industrial land use (-0.50) and percentage of residential land use (-0.42). Dissolved trace-metal concentrations and rainfall were inversely correlated (correlations ranged from -0.34 to -0.57) indicating that large storms dilute constituent concentrations.

Streamflow

Streamflow samples were collected during the flood that occurred in 1993 when dams upstream from Maricopa County released water into the Salt River. Stream discharge reached a maximum of $129,000\text{ ft}^3/\text{s}$ on January 8, 1993, at Salt River at Alma School Road near Mesa, Arizona (09512060), which is about 6 mi upstream from Salt River at Priest Drive (09512165). Three samples were collected at discharge rates of 10 (first-flush sample), 8,600, and $47,800\text{ ft}^3/\text{s}$ for constituent concentrations and acute toxicity. Another sample was collected at $25,500\text{ ft}^3/\text{s}$ after a landfill collapsed into the Salt River and was analyzed only for constituent concentrations.

Concentrations of oil and grease, organophosphate pesticides, and dissolved zinc, copper, cadmium, nickel, chromium, and cobalt were less than detection limits in samples from the Salt River, but were detected in 28 to 100 percent of urban stormwater samples. Total ammonia was detected only in the first-flush sample (0.36 mg/L). These data indicate that urban stormwater could degrade the quality of streamflow with oil and grease, pesticides, dissolved trace metals, and ammonia.

Total hardness of streamflow ranged from 152 to 216 mg/L as CaCO_3 , which was harder than most stormwater samples. Suspended-solids concentrations ranged from 9 to 318 mg/L and were significantly correlated with stream discharge (0.77). Specific conductance of streamflow steadily decreased from 880 to 299 $\mu\text{S}/\text{cm}$ during the floods (fig. 6) indicating that salts were flushed from the channel, diluted, or that runoff had a short

retention time in the reservoirs. Specific-conductance values of streamflow were larger than most values of urban stormwater indicating that streamflow is saltier than urban stormwater. The pH of streamflow was 8.9 in the first-flush sample and about 8 in other samples.

ANOVA results indicated that concentrations of dissolved strontium, magnesium, and lithium in streamflow samples were significantly larger than

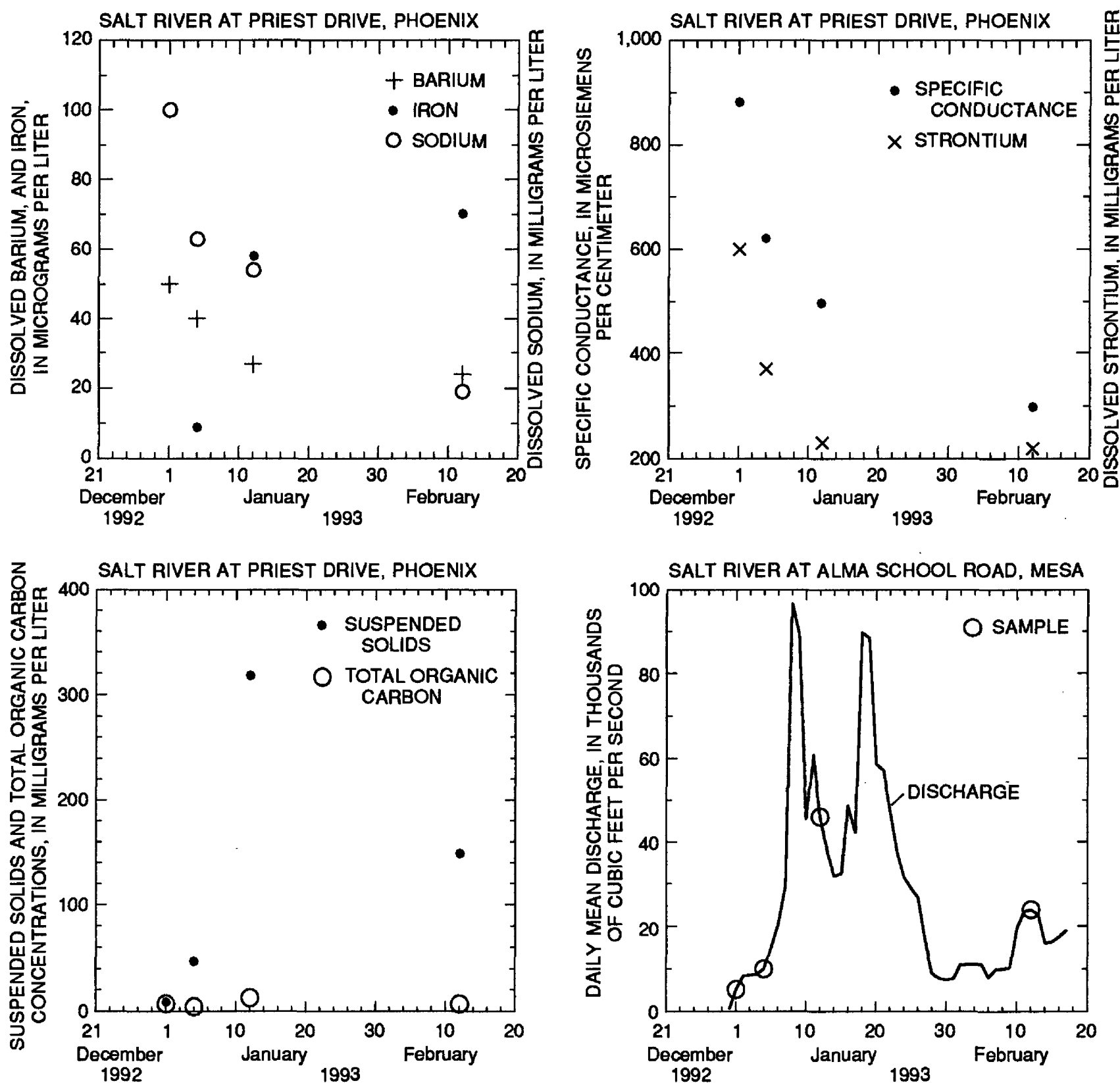


Figure 6. Concentrations of selected constituents and specific conductance in streamflow from the Salt River at Priest Drive, Phoenix, Arizona, as a function of time, and daily mean discharge of the Salt River at Alma School Road, near Mesa, Arizona, as a function of time.

in urban stormwater samples. Concentrations of these constituents steadily decreased with time during the flood and did not vary with discharge (fig. 6). Concentrations of dissolved iron and manganese steadily increased during the flood from less than 3 to 70 $\mu\text{g/L}$ and from 1 to 4 $\mu\text{g/L}$, respectively. Concentrations of dissolved molybdenum, vanadium, and silver were near the detection limits in the first-flush sample from the Salt River and were not detected in other samples.

The increase in iron and manganese concentrations at pH values of about 8 could be due to complexation with dissolved organic carbon, changes in the redox potential of streamflow, or an increase in the amount of colloids smaller than 0.45 μm during the flood. Dissolved oxygen concentration was near saturation, and total organic carbon concentrations ranged from 4.8 to 13 mg/L. These concentrations are typical of most rivers and lakes (Thurman, 1985). These data indicate that redox and complexation probably did not affect iron and manganese concentrations and that concentrations are probably due to colloids.

Bed Material

Urban activities appear to have increased the concentrations of certain constituents above natural concentrations. The natural concentration of constituents depends primarily on the geology of the source areas for bed material. Most urbanization has occurred on bed material deposited by the Salt, Verde, and Agua Fria Rivers, which originate from the White Mountains, Verde Valley, and Chino Valley, respectively. The geology of South Mountain Park is the primary factor affecting constituent concentrations in bed material from the drainage basin with undeveloped land use. Constituent concentrations in bed material from urban drainage basins should be similar to concentrations in bed material from ephemeral channels. Urban activities, however, release certain constituents that accumulate in bed material, which increases the concentration of these constituents above natural concentrations.

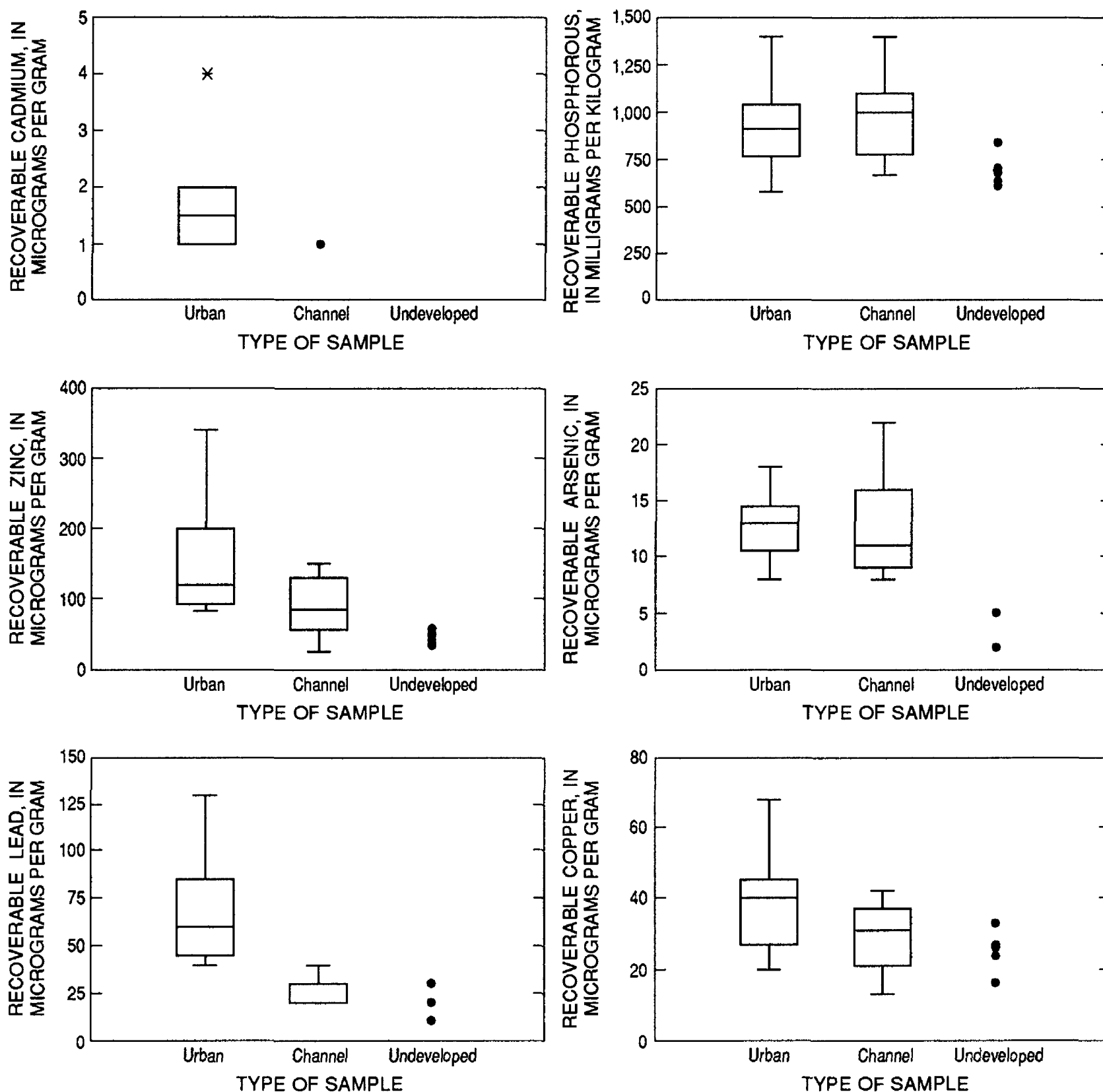
ANOVA results indicate that urban activities increased the recoverable concentrations of ammonia, lead, cadmium, and zinc in bed-material samples from urban drainage basins compared to

recoverable concentrations in samples from ephemeral channels (fig. 7). Ammonia could be from fertilizers, fecal matter, and other sources. Lead is probably from vehicles that use leaded gasoline. Cadmium and zinc could be from particulate metal in oil, brake pads, and other sources. Recoverable concentrations of other constituents, including pesticides, were not significantly different between bed-material samples from urban drainage basins and ephemeral streams.

ANOVA results also indicate that recoverable concentrations of manganese were larger and recoverable concentrations of arsenic were smaller in bed-material samples from South Mountain Park than samples from either urban drainage basins or ephemeral channels. Recoverable concentrations of copper were smaller in bed-material samples from South Mountain Park than in samples from urban drainage basins, and phosphorous concentrations were smaller in bed-material samples from South Mountain Park than in samples from ephemeral channels. Recoverable concentrations of other constituents, including pesticides, were not statistically different among bed-material samples from South Mountain Park, urban drainage basins, and ephemeral streams. Manganese is a common constituent in bed material, and the large recoverable concentrations probably are due to the geology of South Mountain Park.

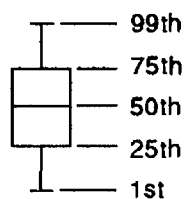
Organic carbon and oxyhydroxides of iron and manganese remove metals from solution because of their large adsorption capacities (Horowitz and Elrick, 1987; Drever, 1988). The percentage of organic carbon significantly correlated with recoverable concentrations of zinc and cadmium in bed material from urban drainage basins (table 2 and fig. 8) and indicates that most metals released by urban activities are associated with organic matter. Lead, however, was not significantly correlated with organic carbon, iron, or manganese. Lead is emitted in automobile exhaust as halide salts (Post and Buseck, 1984), which could explain the lack of correlation with adsorbing materials. Naturally occurring trace metals in bed material from urban drainage basins strongly correlated with iron and manganese and poorly correlated with organic carbon.

Trace metals in bed-material samples collected from ephemeral channels and South Mountain Park



EXPLANATION

PERCENTILE—Percentage of analyses equal to or less than indicated values



× OUTLIER—Either less than 1st percentile or greater than 99th percentile

• INDIVIDUAL DATA POINTS

Figure 7. Ranges in constituent concentrations in bed material from urban drainage basins, ephemeral channels, and South Mountain Park, which is an undeveloped drainage basin, Maricopa County, Arizona.

Table 2. Spearman rank correlations of trace metals and pesticides to recoverable iron and manganese and organic carbon in bed material, Maricopa County, Arizona

[Trace elements, in micrograms per gram are total recoverable; pesticides are given in micrograms per kilogram; ---, dashes indicate there were an insufficient number of detections to compute correlations; DDE, dichlorodiphenylethylene; DDT, dichlorodiphenyltrichloroethane]

Trace metal	Number of samples	Correlation to total recoverable Iron	Correlation to total recoverable manganese	Correlation to organic carbon, in percent	Pesticide	Number of samples	Correlation to total recoverable Iron	Correlation to total recoverable manganese	Correlation to organic carbon, in percent
Urban drainage basins									
Arsenic.....	13	¹ 0.79	0.36	-0.16	Chlordane	11	0.40	0.01	0.12
Cadmium	12	¹ -.61	-.28	¹ .60	DDE	12	.25	-.08	-.28
Chromium.....	13	¹ .67	.08	.14	DDT	12	.20	-.02	-.44
Copper	13	.46	¹ .62	.07	Dieldrin	11	.19	.26	.33
Lead	13	-.47	¹ -.70	-.06	Toxaphene	12	-.04	.12	-.22
Zinc.....	13	-.13	.11	¹ .80	Diazinon.....	12	¹ -.64	-.46	.47
Ephemeral streams									
Arsenic.....	11	.19	.20	.33	Chlordane	11	-.07	-.38	.43
Cadmium	3	-----	-----	-----	DDE	11	.03	-.38	.06
Chromium.....	11	¹ .81	.36	.52	DDT	11	-.37	-.43	-.43
Copper	11	.50	.54	.46	Dieldrin	9	.14	-.27	.15
Lead	11	.28	-.25	¹ .64	Toxaphene	10	-.52	-.54	-.52
Zinc.....	11	.33	.02	¹ .63	Diazinon	8	-.01	-.42	¹ .70
Undeveloped drainage basin									
Arsenic.....	6	.65	-.40	.13	Chlordane	4	.40	¹ 1.0	.40
Cadmium	0	-----	-----	-----	DDE	4	.60	.74	¹ 1.0
Chromium.....	6	-.13	.66	.65	DDT	2	-----	-----	-----
Copper	6	.23	¹ 1.0	.67	Dieldrin	3	.50	.50	¹ 1.0
Lead	6	.28	¹ .89	.62	Toxaphene	6	-.03	¹ .84	.54
Zinc.....	6	.48	¹ .93	.77	Diazinon	1	-----	-----	-----

¹Correlation was significant at a level of 5 percent.

significantly correlated with the percentage of organic carbon and (or) recoverable concentrations of iron and manganese (table 2). Streamflow may partially solubilize metal oxyhydroxides and release trace metals from adsorbing materials. The trace metals would adsorb onto the organic carbon and metal oxyhydroxides after flow ceases. The redistribution of metals between adsorbing materials could account for the difference in

correlations between naturally occurring constituents in bed material from urban drainage basins, ephemeral channels, and South Mountain Park.

Toxaphene, dichlorodiphenylethylene (DDE), dichlorodiphenyltrichloroethane (DDT), chlordane, and dieldrin were detected in 77 to 93 percent of all bed-material samples and had recoverable concentrations between 0.6 and 1,800 µg/kg.

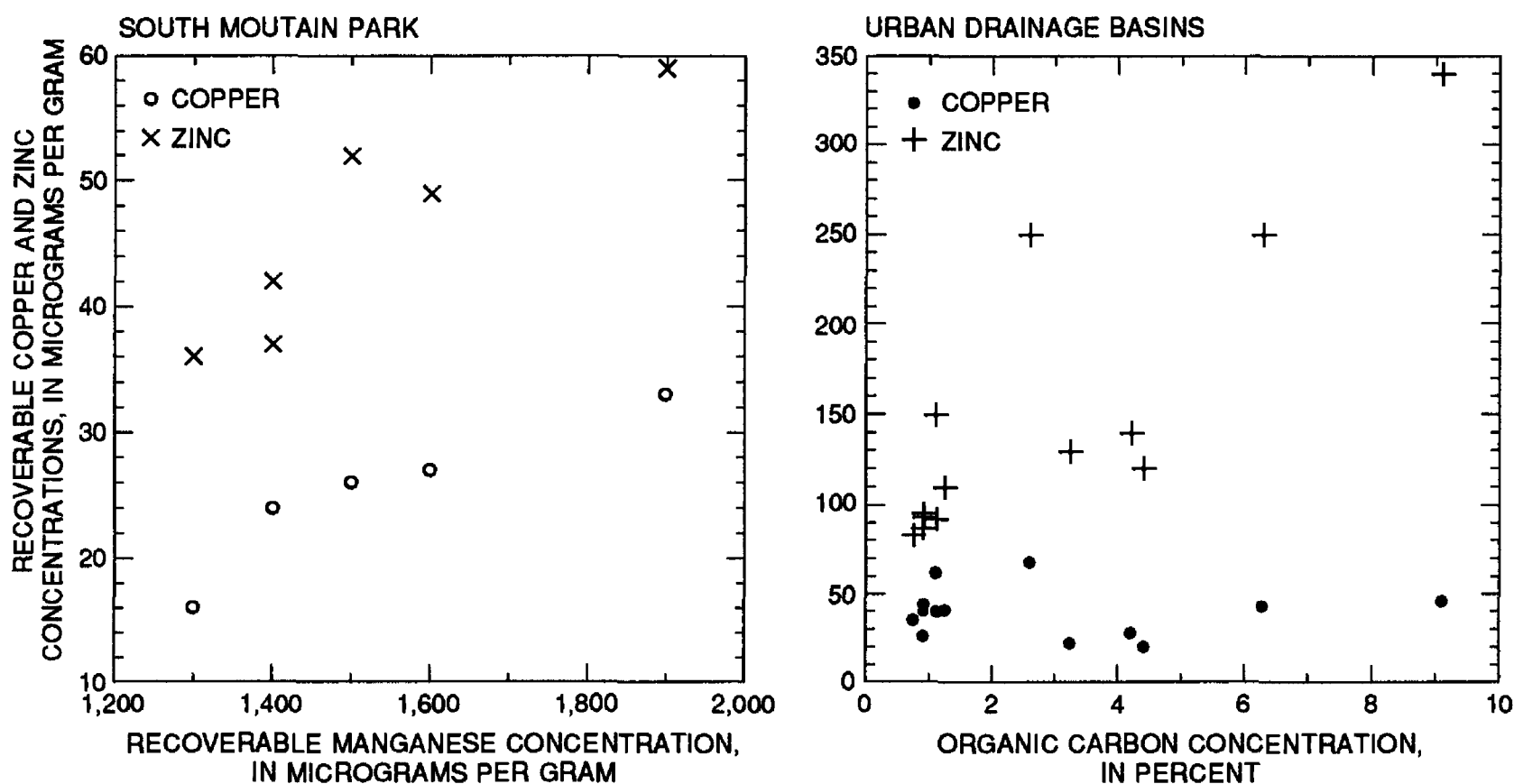


Figure 8. Constituent concentrations of manganese and organic carbon in bed material, Maricopa County, Arizona. Bed-material samples were sieved, and the fraction less than 63 microns was analyzed for constituent concentrations.

Toxaphene and DDT were heavily applied in the Phoenix area during the 1950's and 1960's when much of the area was used for agriculture (Hanks, 1988) and probably are residues from agricultural practices rather than urban activities. Overspraying of nearby fields, illegal dumping, or wind-blown particulates could account for the pesticides at South Mountain Park.

Diazinon was detected in 70 percent of all bed-material samples and had recoverable concentrations between 0.1 and 690 $\mu\text{g/kg}$. Other organophosphate pesticides were detected in less than 17 percent of all bed-material samples and, when detected, were typically in bed material from urban drainage basins.

Organochlorine pesticides were weakly correlated with the percentage of organic carbon and recoverable concentrations of iron and manganese (table 2) except for heptepoxy, which significantly correlated with iron (0.95) in urban bed material. Organochlorine pesticides also could be adsorbed onto clay minerals, but the mineralogy of bed-material samples was not determined. Diazinon was moderately to strongly correlated with the percentage of organic carbon.

The particle-size distribution of bed-material samples was variable and ranged from less than 1 to 75 percent silt and clay, which was the particle size that was analyzed for chemical concentrations and acute toxicity. Bed-material samples from urban drainage basins and ephemeral streams had similar median percentages of silt and clay-size particles. South Mountain, which has steep slopes and boulder terrain, had the coarsest bed material.

ACUTE TOXICITY OF URBAN STORMWATER, STREAMFLOW, AND BED MATERIAL

Acute toxicity was measured in all stormwater and bed-material samples and in three of four streamflow samples that were analyzed for constituent concentrations. TIE's, using either fathead minnows or *C. dubia*, were performed on 11 first-flush samples, 5 flow-weighted composite samples, and 3 samples from drainage basins with mixed land use.

Urban Stormwater

First-flush samples generally were more toxic than flow-weighted composite samples and urban stormwater was more harmful to fathead minnows than to *C. dubia*. About 71 percent of first-flush samples were toxic to fathead minnows, and 28 percent were toxic to *C. dubia* (table 3). About 36 percent of flow-weighted composite samples were toxic to fathead minnows, and 14 percent were toxic to *C. dubia*. Most of the toxic stormwater samples were from the drainage basins with commercial and residential land use. For toxic stormwater samples, however, the LC50 values of first-flush and flow-weighted composite samples were not significantly different. The enhanced mortality rate of fathead minnows by urban stormwater from most drainage basins indicated that the toxicants were more detrimental to fish and

could be present in stormwater throughout the Phoenix area.

Toxicants are not always in the initial runoff. The flow-weighted composite sample collected from 27th Avenue on March 25, 1994, was toxic to fathead minnows; however, the first-flush sample was nontoxic. The difference in toxicity between these samples could be due to the retention time of toxicants in the drainage basin, and specific-conductance values measured at this site indicated that the largest concentrations of constituents are not always associated with the first flush (fig. 3).

Two of four stormwater samples collected from drainage basins with mixed land use were toxic to fathead minnows, and one was toxic to *C. dubia*. An insufficient number of data, however, were collected to compare the toxicity of stormwater samples from drainage basins with mixed land use to first-flush or flow-weighted composite samples

Table 3. Summary of toxicity tests conducted on urban stormwater and streamflow, Maricopa County, Arizona

[LC20 is the percent of stormwater that causes 20-percent mortality; ----, dashes indicate there were an insufficient number of toxic samples to calculate descriptive statistics; descriptive statistics were calculated using the log-probability method (Helsel and Cohn, 1988)]

Type of sample	Species	Number of samples	Number of toxic samples ¹	Percent toxic samples	Median LC20	Mean LC20	Standard deviation of LC20
Stormwater from urban drainage basins with single land use Peoria, Youngtown, 48th Street, and 27th Avenue							
First flush	Fathead minnow	14	10	71	54	48	77
First flush	<i>Ceriodaphnia dubia</i>	14	4	28	100	79	65
Composite	Fathead minnow	14	5	36	76	70	82
Composite	<i>Ceriodaphnia dubia</i>	14	2	14	--	---	---
Stormwater from urban drainage basins with mixed land use Indian Bend Wash at Curry Road, at Tempe, Arizona, and Arizona Canal Diversion Channel							
Equal width	Fathead minnow	4	2	50	--	---	---
Equal width	<i>Ceriodaphnia dubia</i>	4	1	25	--	---	---
Streamflow from dams upstream from Phoenix, which is a nonurban source Streamflow in the Salt River at Priest Drive, at Phoenix, Arizona							
Equal width	Fathead minnow	3	0	0	--	---	---
Equal width	<i>Ceriodaphnia dubia</i>	3	0	0	--	---	---

¹Toxicity is defined as causing mortality in 100-percent stormwater of 20 percent or more.

and to determine if stormwater was more harmful to fathead minnows than to *C. dubia*.

The most toxic stormwater samples, indicated by TU values, were collected from Youngtown on March 26, 1993, and from Peoria on August 24, 1993. Toxicity probably was due to asphalt and resealant. The street at Youngtown was repaved with asphalt in late January 1993, and the parking lot at Peoria was resealed with oil on or about March 20, 1993. The accumulated rainfall between the resurfacing and sample collection was 1.52 in. at Youngtown and 1.03 in. at Peoria. The first-flush and flow-weighted composite samples collected from Youngtown on March 26 were toxic to *C. dubia* (TU values were 5.6 and 3.9, respectively); however, only the first-flush sample was toxic to fathead minnows (TU value was 1.5). Stormwater sampled at Youngtown after March 1993 was toxic only to fathead minnows and TU values ranged from 1.5 to 4.4. The first-flush sample collected from Peoria on August 24 was toxic to fathead minnows and *C. dubia* (TU values were 11.4 for both species); however, the flow-weighted composite sample was not toxic to either species. First-flush and flow-weighted composite samples collected from Peoria on October 6, 1993, were toxic only to fathead minnow (TU values were 3.8 and 1.6, respectively).

Stormwater at Youngtown and Peoria had a yellowish foam that was not present before the resurfacing. The foam was absent after several storms had occurred at each site, indicating that surfactants and possibly other constituents were leached from the asphalt and resealant. Surfactants are a complex group of ionic and nonionic chemicals that are most commonly used as cleaning agents, such as detergents and fabric softeners, and were identified as constituents that are toxic to *C. dubia* (Ankley and Burkhard, 1992). The TU values of samples from Youngtown indicated that toxicants leached from asphalt and resealant were harmful to *C. dubia* and that fathead minnows may be unaffected by these constituents. After about 3 in. of accumulated precipitation at Youngtown and Peoria, toxicants that were harmful to *C. dubia* appeared to be leached from the asphalt and resealant or concentrations were too small to have an acute effect. Sources other than asphalt and resealant could have caused the toxic response; however, these other sources would not account for

the foam and the difference in toxicity of stormwater to *C. dubia* after several storms occurred at each site.

Of the toxic stormwater samples, the least toxic samples were collected from Peoria on January 6, 1993 (before resealant was applied), 48th Street on March 26, 1993, and the ACDC on March 7, 1994. At least 20 percent of the organisms died; however, less than 50 percent died in 100 percent stormwater. LC50 and TU values could not be calculated for these samples and, although these samples were toxic, the data were not used in comparisons between toxicity and constituent concentrations.

Stormwater toxicity decreased in most samples between the time the original test and the TIE were conducted, which was about 2 to 4 days (compare baseline with original in fig. 9). The median mortality rate, in 100-percent stormwater, was 100 percent for the original test and 50 percent for the baseline test, indicating that about 50 percent of the toxicity was lost due to volatilization, adsorption, biodegradation, or other processes in the sample containers. The flow-weighted composite sample collected from 27th Avenue on February 7, 1994, and the sample collected from the ACDC on March 7, 1994, were nontoxic (less than 20 percent mortality) by the time the TIE's were performed, which invalidated the TIE results. The mortality rate of *C. dubia* did not change in stormwater samples collected soon after asphalt and resealant were applied at Youngtown (March 26, 1993) and at Peoria (August 24, 1993).

Confirming that specific constituents are causing toxicity can be done by adding TU values for each toxicant in a sample. If all toxicants are accounted for and are additive, the sum of TU values will equal the TU value of a stormwater sample. Confirmation also can be done by correlating and regressing the TU of several samples with constituent concentrations. Comparisons between TU values of stormwater and toxicants are questionable because: (1) effects from mixtures of toxicants are not always additive; (2) the LC50 of certain toxicants depends on many factors; and (3) concentrations of toxicants measured in stormwater samples may not represent concentrations that organisms were exposed to during the toxicity test. Certain mixtures of toxicants, such as copper and zinc (Rand and Petrocelli, 1985), are not additive because the

mixture can have a smaller (antagonistic) or larger (synergistic) effect than expected on the basis of exposure to individual toxicants. Reactions likely occurred that changed constituent concentrations during sample storage, and the concentrations that were measured in stormwater samples may not represent concentrations to which organisms were exposed. The LC50 of trace metals can change by an order of magnitude with a change of 1 pH unit (Schubauer-Berigan and Dierkes, 1993) and also is dependent on other factors such as hardness, salinity, and speciation (Rand and Petrocelli, 1985; Goyer and Mehlman, 1977). A detailed evaluation of which constituents could be toxic to fathead minnows is not possible because of the changes in sample chemistry that occur during storage and the many factors that control constituent toxicity. A general evaluation, using rank correlations and linear relations between TU values of stormwater and constituent concentrations, was done to identify toxic constituents. Constituents that were not analyzed also could be contributing to the toxicity of stormwater and bed material.

Organic compounds and (or) toxicants that are pH dependent appeared to be the primary toxicants in most stormwater samples. Extraction of neutral and cationic organic compounds caused the largest reduction in mortality rates of either species. Metal hydroxides, however, could have precipitated when samples were adjusted to pH 9 and could have been filtered by the octadecyl column. Comparison of mortality rates between extractions at pH 3, pH 9, and the initial pH indicated which group of organic compounds caused the toxic response. An equal reduction in mortality rates at the three pH levels indicates that toxicant extraction is unaffected by pH, and neutral organic compounds probably caused the toxic response. A reduction in mortality rates at pH 3 or pH 9 without a reduction in mortality rates at the initial pH indicates that the toxicants are anionic or cationic organic compounds, respectively.

Of the organic constituents that were analyzed, TU values for fathead minnows were significantly correlated with total organic carbon (0.67), were not significantly correlated with oil and grease (0.20), and were inversely correlated with the sum of organophosphate pesticides concentrations (-0.89). A positive correlation between TU values and constituent concentrations indicates that the most

toxic stormwater was associated with the largest constituent concentration, which is the expected trend if the constituent is toxic. A negative correlation indicated that the mortality of fathead minnows was not due to organophosphate pesticides. The acute toxicity of organophosphate pesticides to fathead minnows was found to be small in waters of varying hardness (20 and 400 mg/L as CaCO₃; Henderson and others, 1960). Mortality of fathead minnows probably was due to organic constituents that were not analyzed.

Few stormwater samples were toxic to *C. dubia*, so correlations between TU values and constituent concentrations were not significant. The LC50 of diazinon to *C. dubia* is 0.35 µg/L (Norberg-King and others, 1989). TU values for diazinon in stormwater samples ranged from 0.03 to 1.6 and exceeded 1 TU in six samples. Samples with TU values greater than 1 would indicate that at least 50 percent of the organisms should have died in these samples. Of the samples with diazinon exceeding 1 TU, however, only one was toxic to *C. dubia*. Stormwater samples may not have been toxic because organophosphate pesticides readily degrade and (or) pesticides were not biologically available to *C. dubia*.

Trends in the mortality rate when increasing amounts of EDTA were added to the subsamples indicated that dissolved metals contributed to the toxicity of certain stormwater samples. Toxicity from dissolved metals was indicated when small amounts of EDTA were added to the samples and the mortality rate decreased; the mortality rate increased when larger amounts of EDTA were added to samples. This trend of decreasing then increasing mortality rates occurs when EDTA chelates with dissolved metals and reduces metal toxicity; then EDTA causes a toxic response when more EDTA is added than is needed to chelate the dissolved metals. Dissolved metals appeared to contribute to stormwater toxicity in samples collected from Youngtown, 27th Avenue, Peoria, and the ACDC (fig. 9).

Concentrations of dissolved zinc weakly correlated with TU values for fathead minnows. Dissolved zinc and copper had a roughly linear relation between log-transformed values (table 4 and fig. 10), and were greater in some stormwater samples than the LC50 values of zinc and copper to fathead minnows in very hard water

A

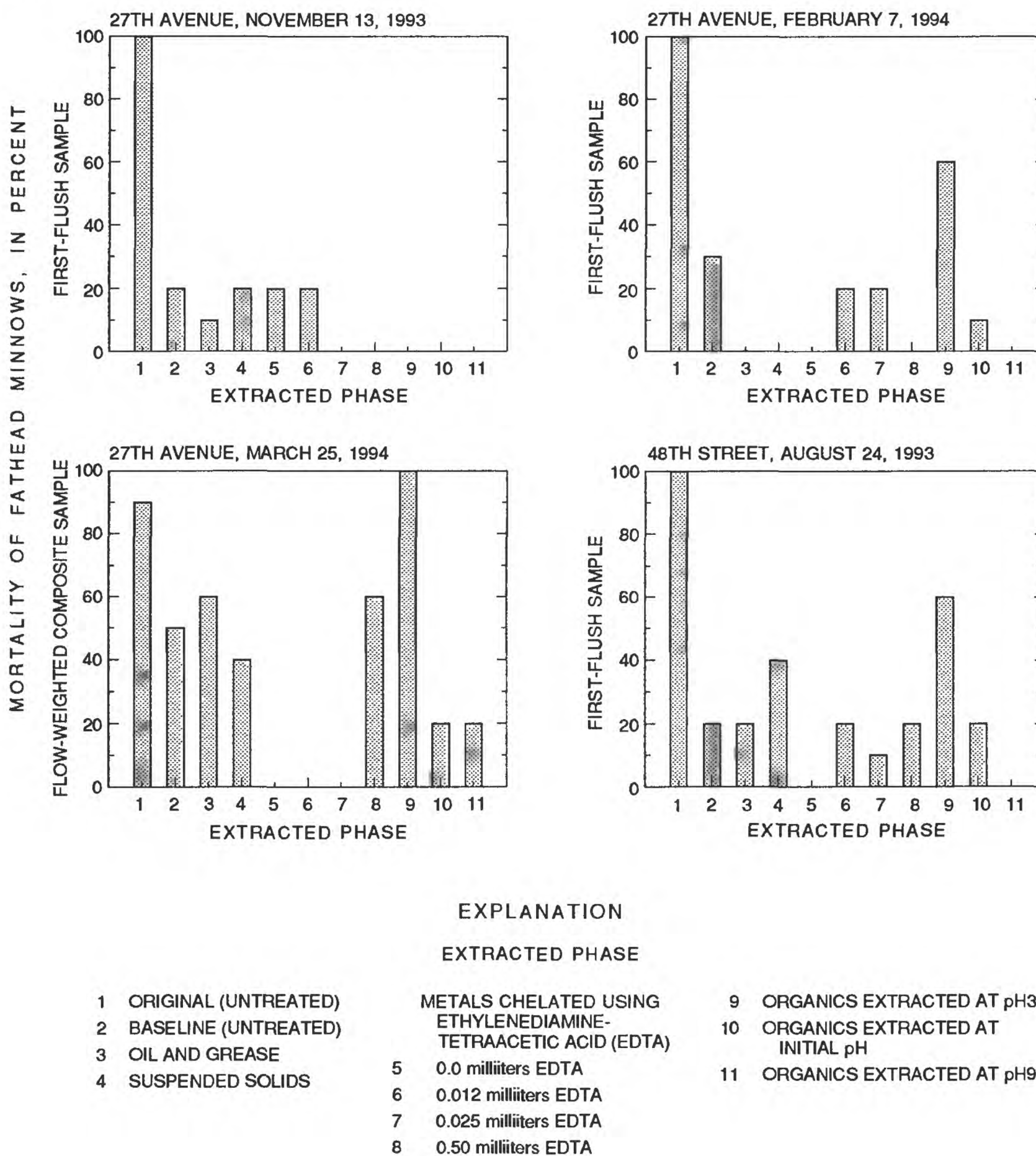


Figure 9. Mortality rates of fathead minnows and *Ceriodaphnia dubia* as a function of phases extracted from stormwater samples, Maricopa County, Arizona, March 26, 1993, through March 25, 1994. A, 27th Avenue and 48th Street. B, Youngtown. C, Youngtown and Arizona Canal Diversion Channel. D, Peoria.

B

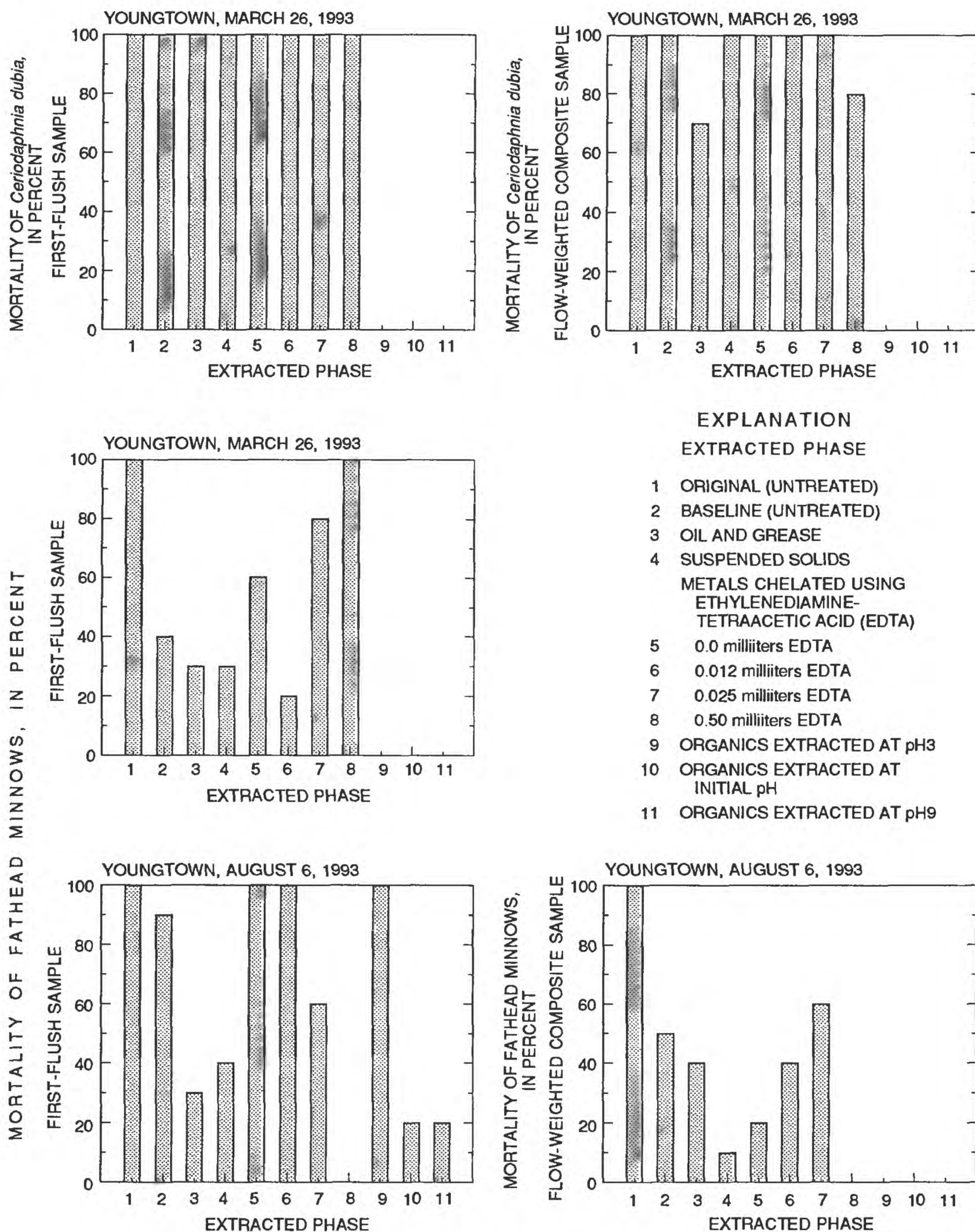


Figure 9. Continued.

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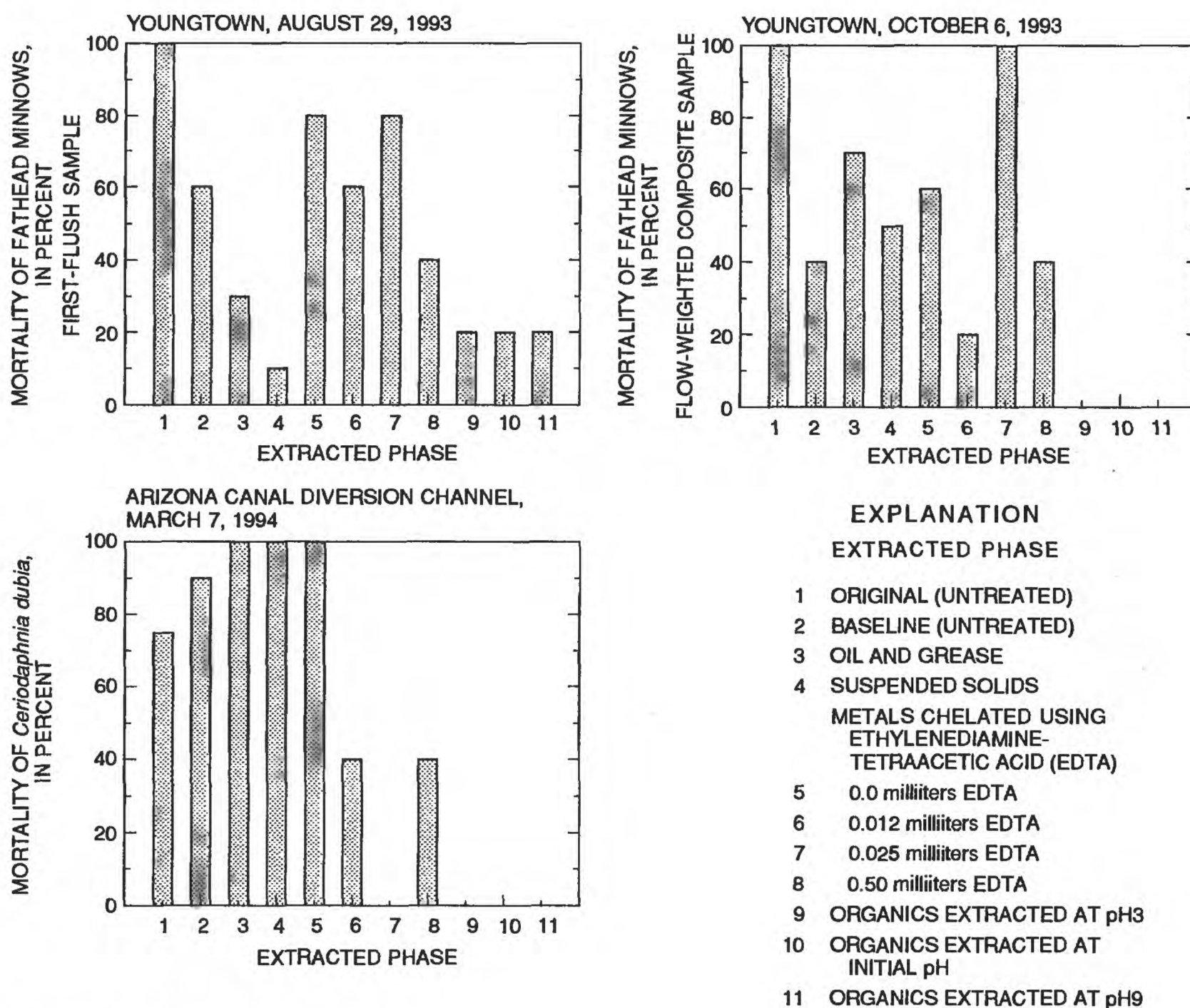


Figure 9. Continued.

(Schubauer-Berigan and Dierkes, 1993). The toxicity of trace metals generally increases as hardness decreases (Rand and Petrocelli, 1985). About 50 percent of the stormwater samples were soft; the softness indicated that concentrations of dissolved zinc and copper in stormwater could be greater than 1 TU. Other trace-metal concentrations were weakly correlated with TU values for fathead minnows.

Extraction of oil and grease and suspended solids usually did not reduce stormwater toxicity, but these constituents could have contributed to the toxicity of two samples from Youngtown and one

sample from Peoria (fig. 9). Concentrations of oil and grease and suspended solids were not significantly correlated with TU values for either fathead minnows or *C. dubia*.

Streamflow

Streamflow samples collected from the Salt River were not toxic to either fathead minnows or *C. dubia*, and most toxic constituents that were analyzed were not detected in streamflow. These data indicate that the Salt River could support an

D

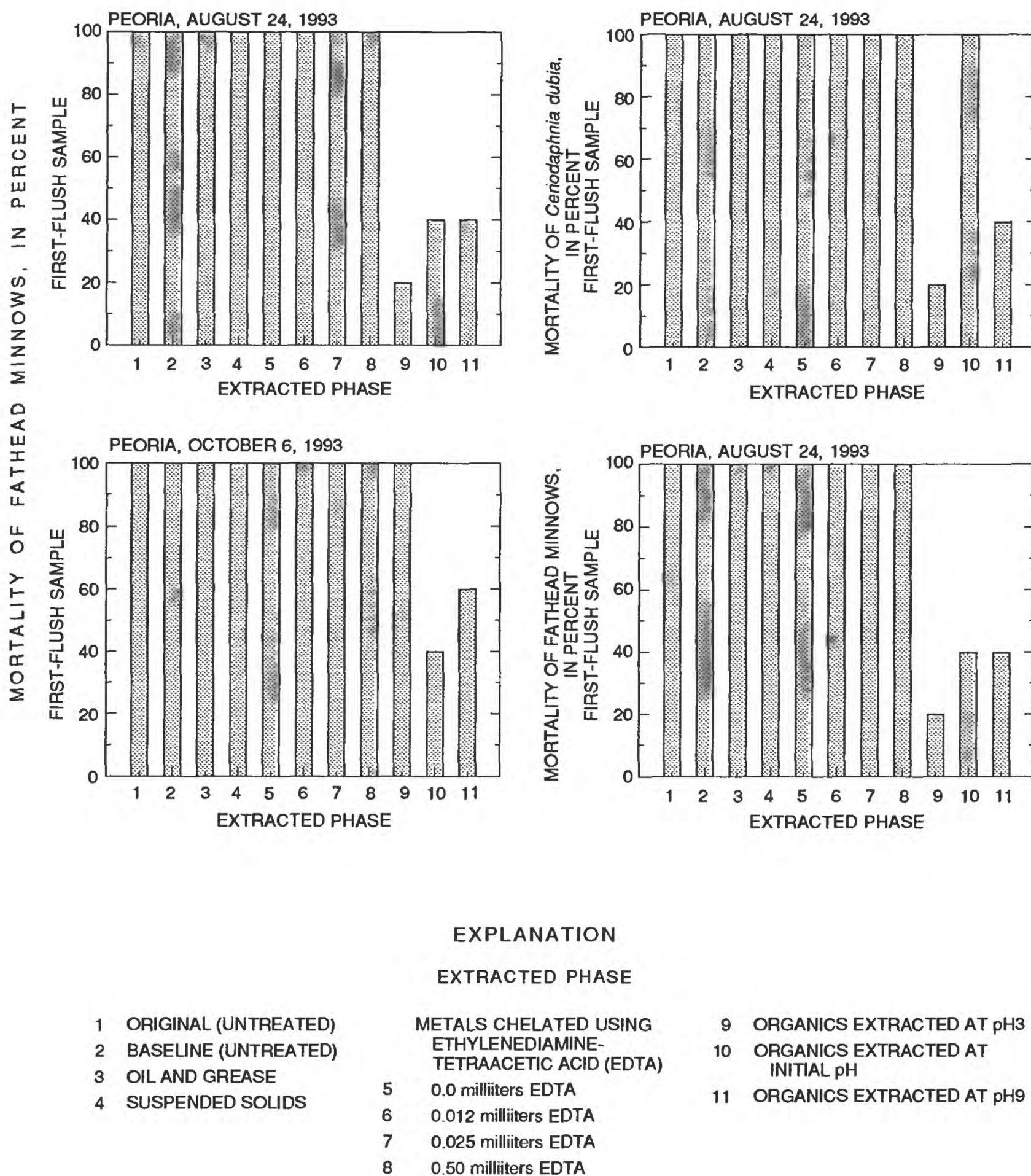


Figure 9. Continued.

Table 4. Spearman rank correlations of dissolved trace metals and organophosphate pesticides to toxic units of stormwater for fathead minnows, Maricopa County, Arizona

[The toxic unit of stormwater to fathead minnows is equal to 100 divided by the LC50. The LC50 is the concentration of stormwater, in percent, by volume, that killed 50 percent of the organisms. Significance is the probability that the variables are independent]

Trace metal and organophosphate pesticide	Number of samples	Correlation of toxic unit of stormwater for fathead minnows	Significance
Copper, dissolved.....	13	0.15	0.62
Zinc, dissolved.....	15	.39	.14
Sum of dissolved zinc and copper.....	15	.44	.10
Iron, dissolved.....	15	.45	.09
Manganese, dissolved.....	15	.40	.14
Vanadium, dissolved.....	11	.26	.44
Cadmium, dissolved.....	7	.41	.36
Cobalt, dissolved.....	7	.63	.13
Total organic carbon.....	14	.67	.01
Diazinon, total.....	4	-.50	.50
Sum of total organophosphate pesticides, total.....	4	-.89	.11

aquatic ecosystem if the river were perennial or intermittent and that the quality of streamflow could be degraded by toxic constituents in urban stormwater. Constituents, however, could be diluted to nontoxic concentrations by the time stormwater flows into the streams and would be diluted by streamflow if the Salt River were perennial.

Bed Material

The most toxic bed-material samples, indicated by the mean and median mortality rate of *H. azteca* (table 5), were collected from South Mountain Park. ANOVA results indicated that the mortality rates of *H. azteca* in bed-material samples from South Mountain Park were significantly higher than in samples from ephemeral channels. Significant differences in mortality rates were not indicated between samples from South Mountain Park and urban drainage basins and between urban drainage basins and ephemeral channels. Of the bed-material samples collected, 83 percent from South Mountain Park were toxic (greater than 20 percent mortality), 85 percent from urban drainage basins were toxic, and 45 percent from ephemeral streams were toxic.

The high mortality rate of *H. azteca* in bed-material samples from South Mountain Park appeared to be due to trace metals. Several mining

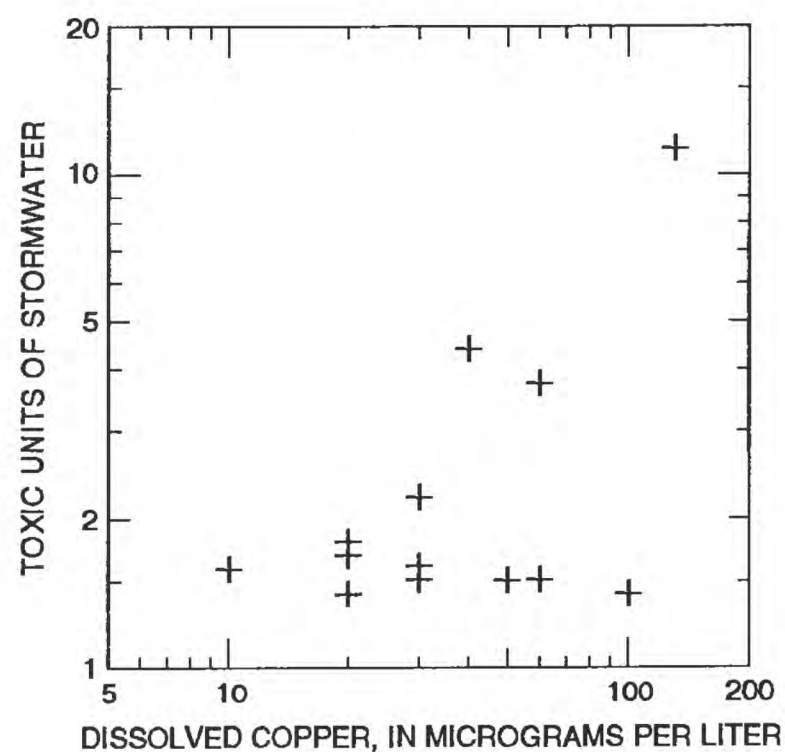
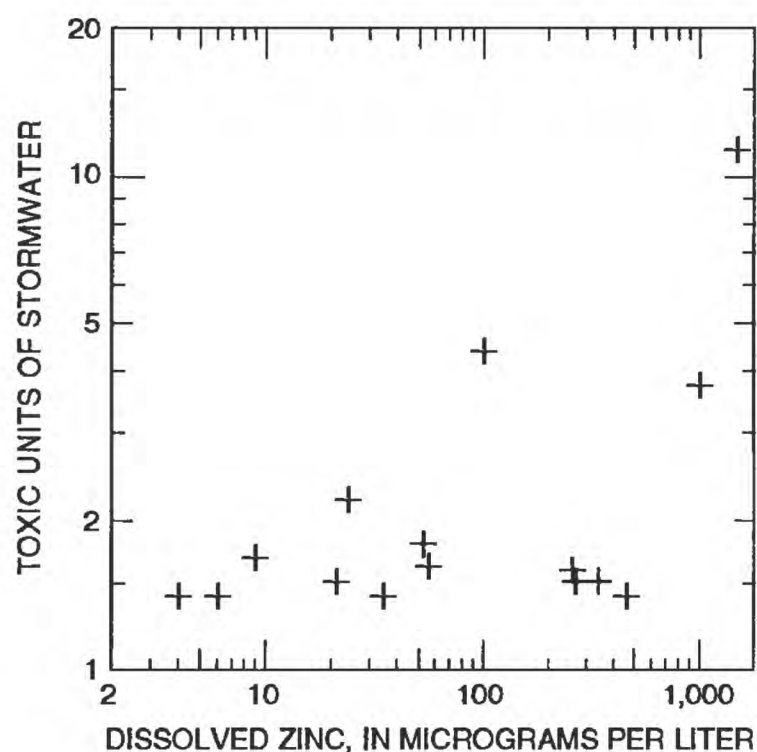


Figure 10. Toxic units of stormwater to fathead minnows as a function of dissolved trace-metal concentrations, Maricopa County, Arizona.

Table 5. Summary of acute toxicity tests of sediment, Maricopa County, Arizona.

Number of samples	Number of toxic samples ¹	Percent toxic samples	Median percent mortality	Mean percent mortality	Standard deviation of percent mortality
Bed material from urban drainage basins with single land use Peoria, Youngtown, 48th Street, and 27th Avenue					
13	11	85	61	57	33
Bed material from ephemeral channels Indian Bend Wash, Arizona Canal Diversion Channel, Agua Fria River, and Salt River at Priest Drive					
11	5	45	16	28	33
Bed material from a drainage basin with undeveloped land use South Mountain Park					
6	5	83	81	76	30

¹Toxicity is defined as causing a mortality rate of 20 percent or more.

prospects are in the drainage basin, and erosion of mine spoils could affect bed-material chemistry. Mortality rates ranged from 76 to 100 percent, except for one sample (17 percent). Samples from the Salt River had 95-, 86-, and 10-percent mortality and were collected after floods scoured urban debris that had accumulated in the channel; the toxicity of these samples probably was caused by naturally occurring toxicants.

Mortality rates of *H. azteca* in bed-material samples from South Mountain Park strongly correlated with recoverable concentrations of zinc (0.71; table 6, fig. 11) and moderately correlated with recoverable concentrations of copper (0.46). Few samples, however, were collected from South Mountain Park; therefore, the correlations were not significant at a level of 5 percent. The high pH values and hard water used in the sediment toxicity test were similar to conditions that increase the toxicity of zinc, cadmium, and nickel to *H. azteca*

and decrease the toxicity of copper and lead (Schubauer-Berigan and Dierkes, 1993). Organochlorine pesticides were detected in most bed-material samples from South Mountain Park, urban drainage basins, and ephemeral channels. Mortality rates of *H. azteca*, however, poorly correlated with pesticide concentrations.

The dissolved concentrations of zinc and copper that organisms were exposed to depends on the solubility of zinc in hard water with a pH of about 8, and on the adsorption properties of the bed material. Recoverable concentrations of zinc and copper strongly correlated with concentrations of manganese in bed material from South Mountain Park (0.93 and 1.0, respectively). The median recoverable concentration of zinc in bed material from South Mountain Park was 45 µg/g, and the median recoverable concentration of zinc in a beaker with 100 grams of bed material and 400 milliliters of hard water was 11,250 µg/L. About 2.6 percent of the recoverable zinc would have to be in solution for organisms to be exposed to the reported LC50 of 290 µg/L for *H. azteca* in very hard water (Schubauer-Berigan and Dierkes, 1993).

Bed-material samples collected from areas where urban stormwater accumulates appeared to be more toxic than samples collected from areas where stormwater does not accumulate. The mortality rate for *H. azteca* was 3 percent in a bed-material sample collected from an undeveloped lot in the drainage basin with light industrial land use; the mortality rates were 77- to 100-percent in samples collected from an urban stormwater conveyance and a dry puddle that accumulates runoff from a freeway. The mortality rate for *H. azteca* was 28 percent in a bed-material sample collected from a dirt lot in the drainage basin with commercial land use and 56 percent in a bed-material sample collected from a dry puddle that accumulates runoff from the parking lot. The difference in mortality rates between these samples indicated that accumulation of constituents in stormwater increased the toxicity of bed material.

Bed-material samples from Youngtown were collected at the streamflow-gaging station in September 1992 and June 1993. Samples from the Agua Fria River were collected at the tributary that drains Youngtown and upstream and downstream from the tributary in September 1993. The mortality

Table 6. Spearman rank correlations of selected recoverable constituents in bed material to mortality rates of *Hyalella azteca*, Maricopa County, Arizona

[mg/kg, milligrams per kilogram; (µg/g), micrograms per gram; (µg/kg), micrograms per kilogram; ----, dashes indicate that there were too few data to compute correlations; DDE, dichlorodiphenylethylene; DDT, dichlorodiphenyltrichloroethane]

Constituents	Mortality rate of <i>Hyalella azteca</i> , in percent			Constituents	Mortality rate of <i>Hyalella azteca</i> , in percent		
	Urban drainage basin	Ephemeral channels	Undeveloped basin		Urban drainage basin	Ephemeral channels	Undeveloped basin
Ammonia (mg/kg).....	0.04	0.37	−0.32	Zinc (µg/g).....	¹ 0.70	−0.30	0.71
Arsenic (µg/g).....	−.52	¹ .60	.39	Chlordane (µg/kg).....	−.19	−.28	.40
Cadmium (µg/g).....	¹ .81	-----	-----	DDE (µg/kg).....	−.28	−.42	.60
Chromium (µg/g).....	−.35	−.31	.13	DDT (µg/kg).....	−.48	.00	-----
Cobalt (µg/g).....	¹ −.72	−.22	-----	Diazinon (µg/kg).....	.51	−.04	-----
Copper (µg/g).....	.29	.26	.46	Dieldrin (µg/kg).....	−.13	−.52	.50
Lead (µg/g).....	.45	−.46	.59	Toxaphene (µg/kg).....	−.16	−.26	.08
Mercury (µg/g).....	.02	.23	.31				

¹Significant at a level of 5 percent

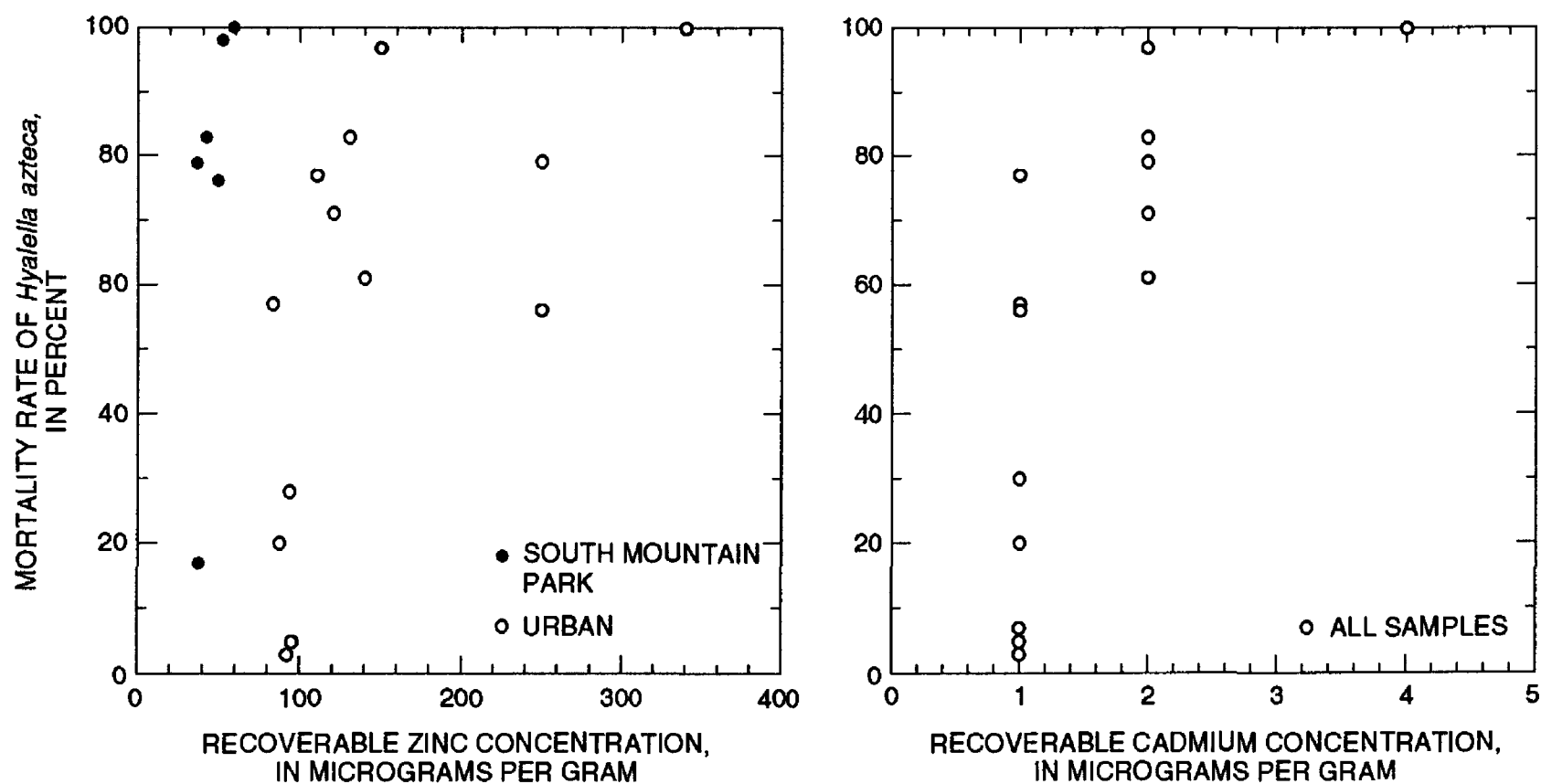


Figure 11. Mortality rates of *Hyalella azteca* as a function of recoverable constituent concentrations in bed material, Maricopa County, Arizona. Bed-material samples were sieved, and the fraction less than 63 microns was analyzed for constituent concentrations and acute toxicity.

rate of *H. azteca* was 61 to 83 percent in bed-material samples from Youngtown and was 16 to 29 percent in samples from the Agua Fria River.

Comparisons between mortality rates for *H. azteca* in bed-material samples from Youngtown and in samples from the Agua Fria River also indicated that urban stormwater affects the toxicity of bed material. The difference in mortality rates indicated that toxicants from urban activities accumulate in bed material and that toxicants are either degraded or diluted by bed material from nonurban sources.

Mortality rates in bed-material samples from urban drainage basins significantly correlated with recoverable concentrations of cadmium (0.81) and zinc (0.70), which were significantly larger in urban bed-material samples than in samples from ephemeral channels. The median recoverable concentrations of cadmium and zinc in urban bed material were 1.5 and 120 $\mu\text{g/g}$, respectively, and recoverable concentrations were moderately to strongly correlated with the percentage of organic carbon. The median recoverable concentration of cadmium and zinc in each beaker of the toxicity test was 375 and 30,000 $\mu\text{g/L}$, respectively. Estimates indicate that about 1 percent of the recoverable cadmium and zinc would have to desorb from the organic carbon for *H. azteca* to be exposed to the LC50 of 5 $\mu\text{g/L}$ for cadmium and 290 $\mu\text{g/L}$ for zinc in very hard water (Schubauer-Berigan and Dierkes, 1993).

The recoverable concentration of zinc that appeared to cause a certain mortality rate was about three times larger in urban bed-material samples than in samples from South Mountain Park (fig. 11); recoverable cadmium was not detected at South Mountain Park. Larger recoverable concentrations of zinc in urban bed material would be needed to cause the same effect as bed material from South Mountain Park. These concentrations are caused by the different adsorption properties and solubility of zinc, organic carbon, and manganese in hard water. Zinc was associated with organic carbon in urban bed-material samples and with manganese in samples from South Mountain Park. Sulfide precipitation, iron oxides, manganese oxides, and organic carbon are effective at controlling dissolved trace-metal concentrations and, in general, iron oxyhydroxides are the most important adsorbing material, followed by organic

carbon and manganese oxyhydroxides (Horowitz and Elrick, 1987; Jenne, 1976; Jenne and Luoma, 1977). The material that toxic constituents are adsorbed onto appears to control the concentration in solution to which organisms are exposed, and thus the bioavailability of the constituent.

Toxicants in bed material from urban drainage basins could be diluted or degraded to nontoxic concentrations in ephemeral channels. The mortality rates of *H. azteca* in bed-material samples from the ACDC, Indian Bend Wash, and Agua Fria River were less than 30 percent, although most of the streamflow in these channels is urban stormwater. Low mortality rates could be caused by dilution of bed material from urban sources with bed material from nonurban sources, biodegradation, leaching, or other processes. Metals accumulated in urban bed material could be leached, and organic constituents could be degraded in the ACDC and Indian Bend Wash, which are grass lined and irrigated. ANOVA results indicated that bed-material samples collected from grass-lined channels were significantly less toxic than samples collected from bare soil. A population of bacteria could be supported in these channels and degrade organic constituents, and irrigation could leach trace metals into lower soil horizons.

Recurrent sampling at the drainage basins with residential and heavy industrial land use indicated that the toxicity of bed material can change significantly. *H. azteca* in a bed-material sample collected adjacent to a catch basin at 27th Avenue in September 1992 had a 97-percent mortality rate, and a sample collected at the same catch basin in June 1993 had a 5-percent mortality rate. The mortality rates for duplicate samples collected at Youngtown in September 1992 compared well and had 71- and 83-percent mortality, indicating that the sampling procedure and toxicity analysis produced consistent results. *H. azteca* in a sample collected at Youngtown in June 1993 had a 61-percent mortality rate, which did not appear to be significantly different from those of the duplicate samples.

SUMMARY

The chemistry and toxicity of urban stormwater, streamflow, and bed material were characterized to determine if urban stormwater

could degrade the quality of streams in Maricopa County. Toxic phases of stormwater (oil and grease, suspended solids, dissolved metals, and dissolved organics) were identified to aid water-quality managers identify and minimize the sources of toxicants. Samples of stormwater were collected from drainage basins with residential, commercial, industrial, and mixed land use; streamflow was sampled from the Salt River; and bed material was sampled from urban drainage basins, South Mountain Park (an undeveloped drainage basin), and ephemeral channels that receive urban stormwater.

Selected constituents and acute toxicity were analyzed in water and bed-material samples. Acute aquatic toxicity tests were done using *C. dubia* and fathead minnows and acute sediment toxicity tests were done using *H. azteca*. TIE's, using either *C. dubia* or fathead minnows, were done on toxic stormwater samples. Statistical analyses were used to determine the effect of urbanization on the quality of water and bed material and to identify toxic constituents.

Concentrations of oil and grease, organophosphate pesticides, and dissolved zinc, copper, cadmium, nickel, chromium, and cobalt were less than detection limits in four streamflow samples collected from the Salt River, but were detected in 28 to 100 percent of 32 urban stormwater samples. Total ammonia was detected in one streamflow sample and 80 percent of stormwater samples. These data indicated that urban stormwater could degrade the quality of streamflow with oil and grease, pesticides, dissolved trace metals, and ammonia.

Recoverable concentrations of ammonia, lead, cadmium, and zinc in bed-material samples from urban drainage basins were significantly larger than recoverable concentrations in samples from ephemeral channels. Ammonia could be from fertilizers, fecal matter, and other sources. Lead is probably from vehicles that use leaded gasoline. Cadmium and zinc could be from particulate metal in oil, brake pads, and other sources. Recoverable concentrations of other constituents, including pesticides, were not significantly different between bed-material samples from urban drainage basins and ephemeral streams.

The percentage of organic carbon was significantly correlated with recoverable concen-

trations of zinc and cadmium in bed material from urban drainage basins; this correlation indicates that most metals released by urban activities are associated with organic matter. Lead, however, which is emitted in automobile exhaust as halide salts, could explain the lack of correlation with adsorbing materials. Recoverable concentrations of other trace metals in bed-material samples from urban drainage basins and trace metals in samples from ephemeral channels and South Mountain Park strongly correlated with recoverable concentrations of iron and manganese.

First-flush samples appeared to be more toxic than flow-weighted composite samples, and urban stormwater was more harmful to fathead minnows than to *C. dubia*. Streamflow samples were not toxic to either species, indicating that urban stormwater could degrade the quality of the Salt River. In a comparison of species mortality rates, daphnids were typically more sensitive than fish to toxicants. The enhanced mortality rate of fathead minnows to urban stormwater from most drainage basins indicated that the toxicants were more detrimental to fish and could be present in stormwater throughout the Phoenix area.

The most toxic stormwater samples were collected from the drainage basins with residential and commercial land use, and the toxicity probably was from asphalt and resealant. The street and parking lot at these basins were resurfaced before sampling, and the toxicity of stormwater to test species changed after storms rinsed the asphalt and resealant. Stormwater sampled from these drainage basins soon after resurfacing was toxic to *C. dubia*. Stormwater sampled after several storms had rinsed the street and parking lot was toxic only to fathead minnows. Stormwater at these drainage basins had a yellowish foam that was not present before the resurfacing. The absence of foam after several storms had occurred at each site indicated that surfactants and (or) other constituents that leached from the asphalt and resealant were toxic to *C. dubia*.

About 50 percent of the toxicity of stormwater samples was lost between the time the original toxicity test and TIE's were performed (about 2 to 4 days). Reduction in stormwater toxicity was caused by volatilization, adsorption, biodegradation, or other processes in the sample containers. The toxicity of stormwater appeared to

be caused mostly by organic constituents; dissolved metals also appeared to contribute to stormwater toxicity. Statistical analyses indicated that toxicity was not caused by organophosphate pesticides.

The most toxic bed-material samples were collected from South Mountain Park, and the mortality rates were significantly higher than in samples from ephemeral channels. The mortality rate of *H. azteca* was not significantly different between bed-material samples from undeveloped and urban drainage basins and between urban drainage basins and ephemeral channels. Within urban drainage basins, bed-material samples collected from areas where stormwater accumulates appeared to be more toxic than samples collected from areas unaffected by stormwater.

Mortality rates in bed-material samples from South Mountain strongly correlated with recoverable concentrations of zinc and moderately correlated with recoverable concentrations of copper. The high mortality rate of *H. azteca* samples from South Mountain probably was caused by naturally occurring trace metals. Mortality rates in bed-material samples from urban drainage basins significantly correlated with recoverable concentrations of cadmium and zinc, which were significantly larger in urban bed-material samples than in samples from ephemeral channels. Organochlorine pesticides were detected in most bed-material samples from South Mountain Park, urban drainage basins, and ephemeral channels. Mortality rates, however, were poorly correlated with pesticide concentrations.

The bioavailability of trace metals in bed material could be controlled by the adsorption properties and solubility of trace metals, organic carbon, and manganese. Zinc and cadmium were associated with organic carbon in urban bed-material samples, and zinc was associated with manganese in samples from South Mountain Park. The material that toxic constituents are adsorbed onto appears to control the concentration in solution to which organisms are exposed, and thus the bioavailability of the constituent.

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TABLES 7-11

Table 7. Selected water-quality data for stormwater and streamflow samples, Maricopa County, Arizona[$\mu\text{S}/\text{cm}$; microsiemens per centimeter at 25°C ; $\mu\text{g}/\text{L}$, micrograms per liter; mg/L , milligrams per liter]

Station number (See table 1 and figure 1 for station names)	Date	pH, water, whole, field (standard units)	pH, water, whole, lab (standard units)	Oxygen, dissolved (mg/L)	Specific conductance, lab ($\mu\text{S}/\text{cm}$)	Barium, dissolved ($\mu\text{g}/\text{L}$ as Ba)	Cobalt, dissolved ($\mu\text{g}/\text{L}$ as Co)	Iron, dissolved ($\mu\text{g}/\text{L}$ as Fe)	Lead, dissolved ($\mu\text{g}/\text{L}$ as Pb)	Manganese, dissolved ($\mu\text{g}/\text{L}$ as Mn)	Molybdenum, dissolved ($\mu\text{g}/\text{L}$ as Mo)
09512162	02-08-94	7.6	7.7	----	155	16	<3	170	<10	10	<10
09512165	12-31-92	8.9	8.5	----	880	50	<3	<3	<10	1	10
	01-04-93	7.9	8.1	10.2	620	40	<3	9	<10	2	<10
	01-12-93	8.2	8.0	8.1	496	27	<3	58	<10	2	<10
	02-11-93	8.0	8.1	8.7	299	24	<3	70	<10	4	<10
09512184	02-28-93	----	6.6	----	84	15	<3	32	<10	22	<10
	02-28-93	6.9	6.9	8.4	62	12	<3	47	<10	15	<10
	03-26-93	----	6.7	----	176	25	<3	86	<10	80	<10
	03-26-93	6.8	6.6	3.8	158	29	<3	90	<10	59	<10
	08-24-93	----	6.4	----	248	40	5	270	10	200	<10
	08-25-93	6.9	6.5	----	321	42	5	260	<10	210	<10
09512403	11-13-93	----	7.2	----	232	34	<3	43	<10	21	<10
	11-13-93	8.4	7.8	----	151	16	<3	29	<10	13	<10
	02-07-94	----	7.1	----	259	34	<3	75	<10	32	<10
	02-07-94	7.4	7.1	----	230	28	<3	16	<10	22	<10
	03-25-94	----	7.2	----	452	70	<3	53	<10	100	<10
	03-25-94	8.2	7.0	----	208	22	<3	70	<10	18	<10
09513700	03-26-93	----	6.8	----	414	140	5	170	<10	280	<10
	03-26-93	6.7	7.0	7.0	192	43	<3	69	<10	94	<10
	08-06-93	----	6.3	----	714	260	10	730	<10	1,300	<10
	08-06-93	6.7	6.3	4.6	439	140	6	600	<10	1,100	<10
	08-29-93	----	6.9	----	333	80	<3	120	<10	290	<10
	08-29-93	7.3	6.9	----	79	12	<3	32	<10	55	<10
	10-06-93	----	6.6	----	558	170	7	400	<10	660	<10
	10-06-93	7.6	6.8	----	117	20	<3	55	<10	21	<10
09513885	01-06-93	----	6.1	----	99	12	<3	39	<10	72	<10
	01-06-93	6.8	6.8	8.6	44	6	<3	27	<10	28	<10
	02-08-93	----	6.3	----	87	15	<3	41	<10	90	<10
	02-08-93	6.3	6.5	8.2	49	7	<3	27	<10	47	<10
	08-24-93	----	6.3	----	7,100	24	20	25,000	<50	630	<40
	08-24-93	6.5	6.6	----	121	10	<3	150	<10	140	<10
	10-06-93	----	4.9	----	819	83	10	1,500	<10	900	<10
	10-06-93	7.0	6.4	----	133	16	<3	210	<10	160	<10
333543112090501	02-07-94	7.1	6.9	----	155	18	<3	110	<10	18	<10
	03-07-94	6.8	6.6	----	138	18	<3	56	<10	21	<10
	05-25-94	7.4	6.4	----	251	35	4	100	10	23	10

Table 7. Selected water-quality data for stormwater and streamflow samples, Maricopa County, Arizona—Continued

Station number (See table 1 and figure 1 for station names)	Date	Strontium, dissolved ($\mu\text{g/L}$ as Sr)	Vanadium, dissolved ($\mu\text{g/L}$ as V)	Beryllium, dissolved ($\mu\text{g/L}$ as Be)	Copper, dissolved ($\mu\text{g/L}$ as Cu)	Calcium, dissolved (mg/L as Ca)	Magnesium, dissolved (mg/L as Mg)	Lithium, dissolved ($\mu\text{g/L}$ as Li)	Silica, dissolved (mg/L as SiO_2)	Zinc, dissolved ($\mu\text{g/L}$ as Zn)	Cadmium, dissolved ($\mu\text{g/L}$ as Cd)
09512162	02-08-94	110	<6	<0.5	<10	8.5	3.5	10	4.4	6	<1.0
09512165	02-31-92	600	7	<0.5	<10	43	20	61	7.1	<3	<1.0
	01-04-93	370	<6	<0.5	<10	37	12	42	16	<3	<1.0
	01-12-93	230	<6	<0.5	<10	29	8.3	35	15	<3	<1.0
	02-11-93	220	<6	<0.5	<10	27	8.6	13	15	<3	<1.0
09512184	02-28-93	40	<6	<0.5	10	8.3	0.78	<4	1.2	48	<1.0
	02-28-93	36	<6	<0.5	<10	6.9	0.62	<4	1.5	16	<1.0
	03-26-93	84	8	<0.5	30	17	2.0	6	1.5	120	<1.0
	03-26-93	73	<6	<0.5	20	14	1.5	6	1.9	87	<1.0
	08-24-93	160	16	<0.5	100	27	4.3	14	3.2	460	5.0
	08-25-93	180	15	<0.5	70	30	4.8	23	3.3	510	1.0
09512403	11-13-93	180	6	<0.5	30	19	2.7	7	5.2	21	<1.0
	11-13-93	99	<6	<0.5	10	13	1.6	7	3.1	5	2.0
	02-07-94	190	<6	<0.5	30	21	3.7	9	3.9	24	<1.0
	02-07-94	150	<6	0.6	20	19	2.7	7	3.6	4	1.0
	03-25-94	430	<6	<0.5	10	49	8.0	22	13	41	<1.0
	03-25-94	130	9	<0.5	20	16	2.2	8	3.9	9	<1.0
09513700	03-26-93	530	20	<0.5	30	71	6.2	8	2.9	56	2.0
	03-26-93	190	12	<0.5	20	25	2.4	4	3.5	23	<1.0
	08-06-93	770	51	0.8	60	130	11	12	4.7	270	<1.0
	08-06-93	420	44	<0.5	50	77	7.3	8	5.3	340	<1.0
	08-29-93	370	27	<0.5	20	64	5.8	14	9.6	53	5.0
	08-29-93	50	11	<0.5	<10	10	0.94	<4	2.9	13	<1.0
	10-06-93	540	25	<0.5	40	95	8.4	11	6.7	100	2.0
	10-06-93	92	9	<0.5	<10	12	1.2	<4	1.7	18	<1.0
09513885	01-06-93	70	<6	<0.5	<10	9.3	1.4	<4	2.4	35	<1.0
	01-06-93	28	<6	<0.5	<10	4.9	0.53	<4	1.8	25	<1.0
	02-08-93	71	<6	<0.5	10	11	1.5	<4	2.2	87	<1.0
	02-08-93	31	<6	<0.5	<10	4.9	0.66	<4	1.1	27	3.0
	08-24-93	290	46	<2	130	90	32	56	170	1,500	15
	08-24-93	63	9	0.6	10	11	1.4	<4	1.1	140	<1.0
	10-06-93	370	21	<0.5	60	67	9.9	11	2.3	1,000	3.0
	10-06-93	83	10	<0.5	10	13	2.0	<4	1.4	260	<1.0
333543112090501	02-07-94	140	<6	<0.5	<10	13	2.3	<4	3.0	23	<1.0
	03-07-94	110	<6	<0.5	10	12	1.5	<4	2.1	36	<1.0
	05-25-94	240	7	<0.5	20	24	4.3	9	4.8	43	<1.0

Table 7. Selected water-quality data for stormwater and streamflow samples, Maricopa County, Arizona—Continued

Station number (See table 1 and figure 1 for station names)	Date	Sodium, dissolved (mg/L as Na)	Nickel, dissolved (µg/L as Ni)	Chromium, dissolved (µg/L as Cr)	Silver, dissolved (µg/L as Ag)	Oil and grease, total recov., gravimetric (mg/L)	Residue, total at 105°C, suspended (mg/L)	Carbon, organic, total (mg/L as C)	Diazinon, total (µg/L)	Ethion, total (µg/L)
09512162	02-08-94	12	<10	<5	<1.0	---	394	23	0.12	<0.01
09512165	12-31-92	100	<10	<5	2.0	<1	9	7.3	-----	-----
	01-04-93	63	<10	<5	<1.0	<1	47	4.8	<0.01	<0.01
	01-12-93	54	<10	<5	<1.0	<1	318	13	<0.01	<0.01
	02-11-93	19	<10	<5	<1.0	<1	149	6.7	<0.01	<0.01
09512184	02-28-93	2.9	<10	<5	2.0	<1	66	25	-----	-----
	02-28-93	1.9	<10	<5	3.0	<1	50	12	0.07	<0.01
	03-26-93	8.2	<10	<5	<1.0	4	340	80	-----	-----
	03-26-93	5.7	<10	<5	<1.0	4	175	140	0.01	<0.01
	08-24-93	13	20	<5	<1.0	3	58	22	-----	-----
	08-25-93	18	20	<5	<1.0	3	124	26	0.08	<0.01
09512403	11-13-93	12	<10	<5	<1.0	5	128	54	-----	-----
	11-13-93	9.3	<10	<5	<1.0	4	216	31	0.20	<0.01
	02-07-94	13	<10	<5	<1.0	4	246	41	-----	-----
	02-07-94	14	<10	<5	<1.0	6	168	17	0.48	<0.01
	03-25-94	27	<10	<5	<1.0	5	124	31	-----	-----
	03-25-94	10	<10	<5	<1.0	4	524	44	<0.20	<0.01
09513700	03-26-93	7.2	30	6	<1.0	5	271	270	-----	-----
	03-26-93	6.4	10	7	<1.0	3	277	110	0.18	<0.01
	08-06-93	9.8	120	20	<1.0	5	572	520	-----	-----
	08-06-93	7.9	70	9	<1.0	4	212	330	0.09	<0.01
	08-29-93	6.3	40	6	<1.0	4	152	280	-----	-----
	08-29-93	1.6	10	<5	<1.0	<1	140	42	0.14	<0.01
	10-06-93	8.0	60	9	<1.0	10	156	470	-----	-----
	10-06-93	2.1	<10	<5	<1.0	2	544	31	0.36	<0.01
09513885	01-06-93	6.2	<10	<5	<1.0	3	45	-----	-----	-----
	01-06-93	1.5	<10	<5	<1.0	5	240	21	0.38	<0.01
	02-08-93	4.2	<10	<5	<1.0	4	96	62	-----	-----
	02-08-93	1.7	<10	<5	<1.0	5	144	29	0.15	<0.01
	08-24-93	1,600	<50	40	<4.0	---	1,190	5,000	-----	-----
	08-24-93	5.7	<10	<5	<1.0	2	42	47	0.06	<0.01
	10-06-93	62	40	5	<1.0	4	348	660	-----	-----
	10-06-93	8.4	<10	<5	<1.0	1	40	76	0.12	<0.01
333543112090501	02-07-94	7.7	<10	<5	<1.0	8	172	33	0.58	<0.01
	03-07-94	6.2	<10	<5	3.0	5	176	43	0.54	<0.01
	05-25-94	17	<10	<5	2.0	<1	30	49	0.43	<0.01

Table 7. Selected water-quality data for stormwater and streamflow samples, Maricopa County, Arizona—Continued

Station number (See table 1 and figure 1 for station names)	Date	Mala- thion, total (µg/L)	Methy- para- thion, total (µg/L)	Para- thion, total (µg/L)	Total tri- thion (µg/L)	Di- syaton, total (µg/L)	Phorate, total (µg/L)	Chlor- pyrifos, total recov. (µg/L)	DEF, total (µg/L)	Fonofos (dy- fonate), water, whole, total recov. (µg/L)
09512162	02-08-94	0.37	<0.01	<0.01	<0.01	<0.01	<0.01	0.13	<0.01	<0.01
09512165	12-31-92	-----	-----	-----	-----	-----	-----	-----	-----	-----
	01-04-93	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01
	01-12-93	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01
	02-11-93	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01
09512184	02-28-93	-----	-----	-----	-----	-----	-----	-----	-----	-----
	02-28-93	0.09	<0.01	<0.01	<0.01	<0.01	<0.01	<0.05	<0.01	<0.05
	03-26-93	-----	-----	-----	-----	-----	-----	-----	-----	-----
	03-26-93	0.13	0.03	<0.01	<0.01	0.01	<0.01	<0.01	<0.01	<0.01
	08-24-93	-----	-----	-----	-----	-----	-----	-----	-----	-----
	08-25-93	0.04	0.05	<0.01	<0.01	<0.01	<0.01	0.01	<0.01	<0.01
09512403	11-13-93	-----	-----	-----	-----	-----	-----	-----	-----	-----
	11-13-93	0.03	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	0.11	<0.01
	02-07-94	-----	-----	-----	-----	-----	-----	-----	-----	-----
	02-07-94	0.08	<0.01	<0.01	<0.01	<0.01	0.02	0.01	0.05	-----
	03-25-94	-----	-----	-----	-----	-----	-----	-----	-----	-----
	03-25-94	<0.10	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01
09513700	03-26-93	-----	-----	-----	-----	-----	-----	-----	-----	-----
	03-26-93	1.0	<0.01	<0.01	<0.01	<0.01	<0.01	<0.03	<0.01	<0.01
	08-06-93	-----	-----	-----	-----	-----	-----	-----	-----	-----
	08-06-93	0.08	0.02	<0.01	<0.01	<0.01	<0.01	0.01	<0.01	<0.01
	08-29-93	-----	-----	-----	-----	-----	-----	-----	-----	-----
	08-29-93	0.05	0.01	<0.01	<0.01	<0.01	<0.01	0.02	<0.01	<0.01
	10-06-93	-----	-----	-----	-----	-----	-----	-----	-----	-----
	10-06-93	0.04	0.01	<0.01	<0.01	<0.01	<0.01	0.03	0.11	<0.01
09513885	01-06-93	-----	-----	-----	-----	-----	-----	-----	-----	-----
	01-06-93	0.20	<0.01	<0.01	<0.01	<0.01	<0.01	0.02	<0.01	<0.01
	02-08-93	-----	-----	-----	-----	-----	-----	-----	-----	-----
	02-08-93	0.09	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01
	08-24-93	-----	-----	-----	-----	-----	-----	-----	-----	-----
	08-24-93	0.06	0.01	<0.01	<0.01	<0.01	<0.01	0.03	<0.01	<0.01
	10-06-93	-----	-----	-----	-----	-----	-----	-----	-----	-----
	10-06-93	0.04	<0.01	<0.01	<0.01	<0.01	<0.01	0.01	0.03	<0.01
333543112090501	02-07-94	1.1	<0.01	<0.01	<0.01	<0.01	<0.01	0.05	<0.01	<0.01
	03-07-94	0.83	<0.01	<0.01	<0.01	<0.01	<0.01	0.06	<0.01	<0.01
	05-25-94	0.30	<0.01	<0.01	<0.01	<0.01	<0.01	0.02	<0.01	<0.01

Table 8. Selected chemical data for bed-material samples, Maricopa County, Arizona

[$\mu\text{g/g}$, micrograms per gram; mg/kg , milligrams per kilogram; $\mu\text{g/kg}$, micrograms per kilogram; g/kg , gram per kilogram; mm , millimeter; DDD, dichlorodiphenyldichloroethane; DDE, dichlorodiphenylethylene; DDT, dichlorodiphenyltrichloroethane; PCB, polychlorinated biphenyl; PCN, polychlorinated naphthalene]

Station number (See table 1 and figure 1 for sta- tion names)	Date	Moisture content, dry wt. (percent of total)	Iron, recov. ($\mu\text{g/g}$ as Fe)	Cadmium, recov. ($\mu\text{g/g}$ as Cd)	Chro- mium, recov. ($\mu\text{g/g}$ as Cr)	Cobalt, recov. ($\mu\text{g/g}$ as Co)	Copper, recov. ($\mu\text{g/g}$ as Cu)	Lead, recov. ($\mu\text{g/g}$ as Pb)	Mercury, recov. ($\mu\text{g/g}$ as Hg)	Manga- nese, recov. ($\mu\text{g/g}$ as Mn)
09512090	06-09-93	4	12,000	<1	20	170	30	30	0.04	390
09512162	06-09-93	3	14,000	<1	20	20	20	20	<0.01	540
09512165	06-14-93	3	15,000	<1	20	20	40	20	0.07	800
09512190	06-09-93	2	12,000	<1	20	10	40	30	0.08	400
09512200	09-04-92	2	7,500	<1	10	<5	30	30	0.03	1,500
	09-24-92	--	-----	---	---	---	---	---	-----	-----
09512405	06-09-93	2	13,000	<1	20	20	40	20	0.14	810
09513700	09-09-92	2	3,700	2	20	<5	20	70	0.02	190
	09-25-92	--	-----	---	---	---	---	---	-----	-----
	06-03-93	4	11,000	2	30	10	30	70	0.03	290
332007112054301	09-04-92	3	6,300	<1	10	10	20	10	0.03	1,400
	09-24-92	--	-----	---	---	---	---	---	-----	-----
332025112050801	09-04-92	2	6,700	<1	10	<5	20	20	0.02	1,400
	09-24-92	--	-----	---	---	---	---	---	-----	-----
332109112041401	09-08-92	2	6,200	<1	10	<5	30	30	0.03	1,600
	09-24-92	--	-----	---	---	---	---	---	-----	-----
332116112040501	09-08-92	2	6,900	<1	8	<5	20	10	0.02	1,300
	09-24-92	--	-----	---	---	---	---	---	-----	-----
332119112041001	09-08-92	3	8,600	<1	10	10	30	30	0.04	1,900
	09-24-92	--	-----	---	---	---	---	---	-----	-----
332405111583801	06-14-93	2	3,800	4	7	10	50	50	0.04	810
332414111583701	06-02-93	2	12,000	2	20	10	70	90	0.05	360
332425112062401	09-09-92	2	5,300	1	9	10	30	50	0.02	370
	09-25-92	--	-----	---	---	---	---	---	-----	-----
332425112064001	09-09-92	1	5,600	1	9	10	40	120	0.02	320
	09-25-92	--	-----	---	---	---	---	---	-----	-----
332425112065501	09-08-92	1	5,500	2	10	10	60	130	0.05	370
	09-25-92	--	-----	---	---	---	---	---	-----	-----
	06-08-93	2	15,000	1	20	20	40	60	0.04	440
332509111584001	06-02-93	2	12,000	1	10	10	40	40	0.02	450
332513111582601	06-02-93	3	13,000	1	20	20	40	40	0.04	530
333435112181401	09-25-92	4	8,700	<1	10	10	20	20	0.02	370
333438112181401	09-25-92	3	9,100	<1	9	<5	10	20	0.02	280
333439112181701	09-25-92	3	12,000	<1	10	10	30	20	0.05	550
333458112090401	06-03-93	3	25,000	<1	40	20	40	40	0.04	780
333459112090401	06-03-93	3	17,000	1	30	20	40	50	0.08	640
333543112090501	06-03-93	0.3	16,000	1	30	20	40	40	0.05	500
333627112110601	06-03-93	6	13,000	1	20	20	30	30	0.04	460
333704112120501	06-03-93	4	13,000	1	20	20	40	30	0.04	530

Table 8. Selected chemical data for bed-material samples, Maricopa County, Arizona—Continued

Station number (See table 1 and figure 1 for station names)	Date	Zinc, recov. (µg/g as Zn)	Arsenic, total (µg/g as As)	Nitrogen, NO ₂ +NO ₃ , total (mg/kg as N)	Phos- phorus, total (mg/kg as P)	Nitrogen, NH ₄ , total (mg/kg as N)	Nitrogen, NH ₄ +org., total (mg/kg as N)	Perthane, total (µg/kg)	Endo- sulfan, total (µg/kg)	Aldrin, total (µg/kg)
09512090	06-09-93	150	9	46	690	11	910	<1.00	<0.3	<0.1
09512162	06-09-93	60	9	15	960	11	460	<1.00	<0.1	<0.1
09512165	06-14-93	60	16	18	1,000	58	2,000	<1.00	<0.4	<0.1
09512190	06-09-93	140	10	35	670	14	710	<1.00	<0.2	<0.1
09512200	09-04-92	50	5	31	610	8.5	1,300	<10.0	<0.2	<0.2
	09-24-92	----	---	-----	-----	-----	-----	-----	-----	-----
09512405	06-09-93	90	22	25	1,400	54	720	<1.00	<0.2	<0.1
09513700	09-09-92	130	13	5.0	670	16	1,300	<1.00	<0.1	<0.1
	09-25-92	----	---	-----	-----	-----	-----	-----	-----	-----
	06-03-93	140	11	43	580	17	1,200	<3.00	<0.3	<0.3
332007112054301	09-04-92	40	5	64	700	16	1,200	<1.00	<0.1	<0.1
	09-24-92	----	---	-----	-----	-----	-----	-----	-----	-----
332025112050801	09-04-92	40	5	19	640	7.4	780	<1.00	<0.1	<0.1
	09-24-92	----	---	-----	-----	-----	-----	-----	-----	-----
332109112041401	09-08-92	50	2	60	690	11	1,100	<1.00	<0.1	<0.1
	09-24-92	----	---	-----	-----	-----	-----	-----	-----	-----
332116112040501	09-08-92	40	5	16	680	5.5	490	<1.00	<0.1	<0.1
	09-24-92	----	---	-----	-----	-----	-----	-----	-----	-----
332119112041001	09-08-92	60	5	69	840	11	1,300	<1.00	<0.1	<0.1
	09-24-92	----	---	-----	-----	-----	-----	-----	-----	-----
332405111583801	06-14-93	340	8	170	890	130	8,100	<1.00	<0.1	<0.1
332414111583701	06-02-93	250	15	140	1,100	140	1,700	<1.00	<0.1	<0.1
332425112062401	09-09-92	90	13	43	940	12	590	<10.0	<0.2	<0.2
	09-25-92	----	---	-----	-----	-----	-----	-----	-----	-----
332425112064001	09-09-92	80	10	54	910	16	470	<10.0	<0.2	<0.2
	09-25-92	----	---	-----	-----	-----	-----	-----	-----	-----
332425112065501	09-08-92	150	11	13	930	18	580	<10.0	<0.2	<0.2
	09-25-92	----	---	-----	-----	-----	-----	-----	-----	-----
	06-08-93	100	16	310	860	43	570	<1.00	<0.1	<0.3
332509111584001	06-02-93	110	13	22	880	200	960	<1.00	<0.1	<0.1
332513111582601	06-02-93	90	14	28	980	200	860	<1.00	<0.1	<1.0
333435112181401	09-25-92	40	12	19	1,100	4.3	490	<1.00	<0.1	0.1
333438112181401	09-25-92	30	11	2.0	790	3.0	400	<1.00	<0.1	<0.1
333439112181701	09-25-92	60	8	28	1,000	7.9	850	<1.00	<0.1	<0.1
333458112090401	06-03-93	90	14	33	1,400	17	840	<20.0	<2.0	<2.0
333459112090401	06-03-93	250	18	120	1,300	120	2,000	<3.00	<0.3	<0.3
333543112090501	06-03-93	130	10	18	1,100	16	840	<1.00	<0.1	<1.0
333627112110601	06-03-93	100	18	27	980	11	860	<1.00	<0.1	<1.0
333704112120501	06-03-93	120	14	30	1,300	22	1,100	<1.00	<0.1	<0.7

Table 8. Selected chemical data for bed-material samples, Maricopa County, Arizona—Continued

Station number (See table 1 and figure 1 for station namea)	Date	Chlor- dane, total (µg/kg)	DDD, total (µg/kg)	DDE, total (µg/kg)	DDT, total (µg/kg)	Dieldrin, total (µg/kg)	Endrin, total (µg/kg)	Hepta- chlor, total (µg/kg)	Hepta- chlor epoxide, total (µg/kg)	Lindane, total (µg/kg)
09512090	06-09-93	110	3.9	24	1.4	5.2	<0.1	<0.2	0.3	<0.1
09512162	06-09-93	1.0	0.4	20	1.6	<0.1	<0.1	<0.1	<0.1	<0.1
09512165	06-14-93	3.0	0.3	7.6	0.7	0.8	<0.1	<0.1	0.1	<0.2
09512190	06-09-93	53	2.2	8.6	6.5	7.6	<0.3	<0.2	0.2	<0.1
09512200	09-04-92	8.0	<0.2	51	4.9	0.7	<4.0	<0.2	<0.2	<0.2
	09-24-92	-----	-----	-----	-----	-----	-----	-----	-----	-----
09512405	06-09-93	16	2.4	13	9.5	3.9	<0.1	<0.1	<0.1	<0.1
09513700	09-09-92	<100	<0.1	150	<0.1	<0.1	<0.1	<0.1	<0.1	<0.1
	09-25-92	-----	-----	-----	-----	-----	-----	-----	-----	-----
	06-03-93	35	2.3	32	3.2	3.9	<0.3	<0.3	<0.3	<0.3
332007112054301	09-04-92	<1.0	<0.1	16	<0.1	<0.1	<0.1	<0.1	<0.1	<0.1
	09-24-92	-----	-----	-----	-----	-----	-----	-----	-----	-----
332025112050801	09-04-92	2.0	<0.1	12	0.7	0.1	<0.7	<0.1	<0.1	<0.1
	09-24-92	-----	-----	-----	-----	-----	-----	-----	-----	-----
332109112041401	09-08-92	12	<2.0	<0.1	<1.0	<0.1	<0.1	<0.1	<0.1	<0.1
	09-24-92	-----	-----	-----	-----	-----	-----	-----	-----	-----
332116112040501	09-08-92	<2.0	<0.1	<0.1	<0.1	<0.1	<2.0	<0.1	<0.1	<0.1
	09-24-92	-----	-----	-----	-----	-----	-----	-----	-----	-----
332119112041001	09-08-92	16	<2.0	30	<0.1	0.2	<0.1	<0.1	<0.1	<0.1
	09-24-92	-----	-----	-----	-----	-----	-----	-----	-----	-----
332405111583801	06-14-93	9.0	0.7	7.3	0.6	6.3	<0.1	<0.1	<0.1	<0.1
332414111583701	06-02-93	37	2.0	29	3.4	2.5	<0.1	<0.2	0.2	0.3
332425112062401	09-09-92	13	<0.2	91	11	0.3	<4.0	<0.2	<0.2	<0.2
	09-25-92	-----	-----	-----	-----	-----	-----	-----	-----	-----
332425112064001	09-09-92	7.0	<0.5	39	5.0	0.8	<2.0	<0.2	<0.2	<0.2
	09-25-92	-----	-----	-----	-----	-----	-----	-----	-----	-----
332425112065501	09-08-92	22	<0.2	97	26	<0.2	<5.0	<0.2	<0.2	<0.2
	09-25-92	-----	-----	-----	-----	-----	-----	-----	-----	-----
	06-08-93	10	8.2	130	40	3.9	<0.1	<0.1	<0.1	0.5
332509111584001	06-02-93	15	1.2	30	4.2	1.5	<0.1	<0.1	0.1	0.1
332513111582601	06-02-93	280	8.6	30	9.5	150	<0.1	<0.1	0.8	0.1
333435112181401	09-25-92	4.0	<0.1	34	4.8	0.7	2.4	<0.1	<0.1	<0.1
333438112181401	09-25-92	9.0	<0.1	13	17	<0.1	3.3	<0.1	<0.1	<0.1
333439112181701	09-25-92	3.0	<0.1	11	1.2	0.2	1.0	<0.1	<0.1	<0.1
333458112090401	06-03-93	130	80	620	300	70	<60	<0.4	1.5	<0.3
333459112090401	06-03-93	11	0.5	140	9.0	2.1	<0.3	<0.3	<0.3	<0.3
333543112090501	06-03-93	32	2.1	35	3.8	4.8	<0.1	<0.2	0.2	0.2
333627112110601	06-03-93	26	2.2	20	3.5	3.7	<0.1	<0.1	<0.1	0.2
333704112120501	06-03-93	13	0.6	24	2.2	2.0	<0.1	<0.1	<0.1	0.1

Table 8. Selected chemical data for bed-material samples, Maricopa County, Arizona—Continued

Station number (See table 1 and figura 1 for atation namea)	Date	Toxa- phene, total (µg/kg)	Diazinon, total (µg/kg)	Ethlon, total (µg/kg)	Malathion, total (µg/kg)	Methyl parathion, total (µg/kg)	Parathion, total (µg/kg)	Trithion, total (µg/kg)	PCB, total (µg/kg)
09512090	06-09-93	40	7.6	<0.1	0.4	<0.1	<0.1	<0.1	5
09512162	06-09-93	<10	0.2	<0.1	<0.1	<0.1	<0.1	<0.1	<1
09512165	06-14-93	20	0.4	<0.1	<0.1	<0.1	<0.1	<0.1	13
09512190	06-09-93	20	0.2	<0.1	0.4	<0.1	<0.1	<0.1	20
09512200	09-04-92	140	<0.1	<0.1	<0.1	<0.1	<0.1	<0.1	<10
	09-24-92	-----	-----	-----	-----	-----	-----	-----	-----
09512405	06-09-93	30	0.2	<0.1	<0.1	<0.1	<0.1	<0.1	17
09513700	09-09-92	830	9.4	<2.0	5.8	<2.0	<2.0	<2.0	<1
	09-25-92	-----	-----	-----	-----	-----	-----	-----	-----
	06-03-93	80	0.6	<0.1	<0.1	<0.1	<0.1	<0.1	11
332007112054301	09-04-92	130	<0.1	<0.1	<0.1	<0.1	<0.1	<0.1	<1
	09-24-92	-----	-----	-----	-----	-----	-----	-----	-----
332025112050801	09-04-92	30	<0.1	<0.1	<0.1	<0.1	<0.1	<0.1	6
	09-24-92	-----	-----	-----	-----	-----	-----	-----	-----
332109112041401	09-08-92	170	<0.1	<0.1	<0.1	<0.1	<0.1	<0.1	<1
	09-24-92	-----	-----	-----	-----	-----	-----	-----	-----
332116112040501	09-08-92	90	0.1	<0.1	<0.1	<0.1	<0.1	<0.1	<1
	09-24-92	-----	-----	-----	-----	-----	-----	-----	-----
332119112041001	09-08-92	160	<0.1	<0.1	<0.1	<0.1	<0.1	<0.1	<1
	09-24-92	-----	-----	-----	-----	-----	-----	-----	-----
332405111583801	06-14-93	110	3.2	<0.1	<0.1	<0.1	<0.1	<0.1	12
332414111583701	06-02-93	80	3.5	<0.1	0.4	0.2	<0.1	<0.1	19
332425112062401	09-09-92	390	0.8	<0.1	<0.3	1.0	<0.1	<0.1	120
	09-25-92	-----	-----	-----	-----	-----	-----	-----	-----
332425112064001	09-09-92	80	0.2	<0.1	<0.1	0.4	<0.1	<0.1	<1
	09-25-92	-----	-----	-----	-----	-----	-----	-----	-----
332425112065501	09-08-92	380	0.3	<0.1	0.3	0.8	<0.1	<0.1	240
	09-25-92	-----	-----	-----	-----	-----	-----	-----	-----
	06-08-93	240	0.4	<0.1	<0.1	<0.1	<0.1	<0.1	30
332509111584001	06-02-93	50	<0.1	<0.1	<0.1	<0.1	<0.1	<0.1	13
332513111582601	06-02-93	110	0.2	<0.1	<0.1	<0.1	<0.1	<0.1	65
333435112181401	09-25-92	160	0.4	<0.1	<0.1	<0.1	<0.1	<0.1	<1
333438112181401	09-25-92	280	<0.1	<0.1	<0.1	<0.1	<0.1	<0.1	<1
333439112181701	09-25-92	70	<0.1	<0.1	<0.1	<0.1	<0.1	<0.1	<1
333458112090401	06-03-93	1,800	0.3	<0.1	<0.1	<0.1	0.1	<0.1	16
333459112090401	06-03-93	200	0.3	<0.1	<0.1	<0.1	<0.1	<0.1	4
333543112090501	06-03-93	50	3.7	<0.1	<0.1	<0.1	<0.1	<0.1	8
333627112110601	06-03-93	40	<0.1	<0.1	<0.1	<0.1	<0.1	<0.1	6
333704112120501	06-03-93	40	5.0	<0.1	<0.1	<0.1	<0.1	<0.1	4

Table 8. Selected chemical data for bed-material samples, Maricopa County, Arizona—Continued

Station number (See table 1 and figure 1 for station names)	Date	PCN, total (µg/kg)	Methoxychlor, total (µg/kg)	Mirex, total (µg/kg)	Carbon, inorganic, total (g/kg as C)	Carbon, inorganic+organic, total (g/kg as C)	Oil and grease, total, gravimetric (mg/kg)	Bed material, sieve diameter, percent finer than 0.062 mm	Bed material, sieve diameter, percent finer than 0.125 mm
09512090	06-09-93	<1.0	0.5	<0.1	4.5	18	300	18	32
09512162	06-09-93	<1.0	<0.1	<0.1	4.9	10	<100	47	71
09512165	06-14-93	<1.0	<0.1	<0.1	4.6	20	<100	61	77
09512190	06-09-93	<1.0	<1.0	<0.1	5.3	15	<100	24	70
09512200	09-04-92	<1.0	<1.0	<0.2	2.5	16	-----	0	1
	09-24-92	-----	-----	-----	-----	-----	<1,000	---	---
09512405	06-09-93	<1.0	<1.0	<0.1	7.2	14	<100	75	83
09513700	09-09-92	<1.0	<0.1	<0.1	9.6	42	-----	31	68
	09-25-92	-----	-----	-----	-----	-----	2,200	---	---
	06-03-93	<3.0	<0.3	<0.3	10	52	920	37	60
332007112054301	09-04-92	<1.0	-----	<0.1	8.5	19	-----	0	1
	09-24-92	-----	-----	-----	-----	-----	<1,000	---	---
332025112050801	09-04-92	<1.0	<0.1	<0.1	2.3	8.4	-----	2	3
	09-24-92	-----	-----	-----	-----	-----	1,100	---	---
332109112041401	09-08-92	<1.0	0.8	<0.1	2.3	12	-----	2	4
	09-24-92	-----	-----	-----	-----	-----	<1,000	---	---
332116112040501	09-08-92	<1.0	<0.1	<0.1	3.1	6.7	-----	2	5
	09-24-92	-----	-----	-----	-----	-----	1,200	---	---
332119112041001	09-08-92	<1.0	0.3	<0.1	3.8	15	-----	60	66
	09-24-92	-----	-----	-----	-----	-----	<1,000	---	---
332405111583801	06-14-93	<1.0	<0.1	<0.1	49	140	110	4	6
332414111583701	06-02-93	<1.0	<0.1	<0.1	11	37	230	25	42
332425112062401	09-09-92	<1.0	<1.0	<0.2	11	20	-----	36	59
	09-25-92	-----	-----	-----	-----	-----	<1,000	---	---
332425112064001	09-09-92	<1.0	<1.0	<0.2	9.5	17	-----	21	42
	09-25-92	-----	-----	-----	-----	-----	<1,000	---	---
332425112065501	09-08-92	<1.0	<1.0	<0.2	10	21	-----	17	31
	09-25-92	-----	-----	-----	-----	-----	<1,000	---	---
	06-08-93	<1.0	<1.0	<0.2	9.9	19	200	18	32
332509111584001	06-02-93	<1.0	<0.1	<0.1	7.5	20	<100	23	39
332513111582601	06-02-93	<1.0	<0.1	<0.1	8.8	20	<100	16	26
333435112181401	09-25-92	<1.0	<0.1	<0.1	5.5	11	<1,000	6	15
333438112181401	09-25-92	<1.0	<0.1	<0.1	5.9	11	<1,000	38	56
333439112181701	09-25-92	<1.0	<0.1	<0.1	2.5	10	<1,000	59	80
333458112090401	06-03-93	<1.0	<2.0	<0.1	0.8	10	<100	27	41
333459112090401	06-03-93	<3.0	<1.0	<0.3	2.2	65	1,200	12	17
333543112090501	06-03-93	<1.0	<1.0	<0.3	5.4	18	<100	22	31
333627112110601	06-03-93	<1.0	<1.0	<0.1	5.7	25	<100	10	20
333704112120501	06-03-93	<1.0	<0.1	<0.1	5.3	22	160	11	16

Table 8. Selected chemical data for bed-material samples, Maricopa County, Arizona—Continued

Station number (See table 1 and figure 1 for station name)	Date	Bed material, sieve diameter, percent finer than 0.250 mm	Bed material, sieve diameter, percent finer than 0.500 mm	Bed material, sieve diameter, percent finer than 1.00 mm	Bed material, sieve diameter, percent finer than 2.00 mm	Bed material, sieve diameter, percent finer than 4.00 mm	Bed material, sieve diameter, percent finer than 8.00 mm	Bed material, sieve diameter, percent finer than 16.0 mm	Bed material, sieve diameter, percent finer than 32.0 mm
09512090	06-09-93	67	97	99	100	100	---	---	---
09512162	06-09-93	85	93	97	98	99	100	---	---
09512165	06-14-93	92	99	100	100	100	---	---	---
09512190	06-09-93	96	98	99	100	100	100	---	---
09512200	09-04-92	4	12	26	52	91	99	100	---
	09-24-92	---	---	---	---	---	---	---	---
09512405	06-09-93	95	99	100	100	100	100	---	---
09513700	09-09-92	81	87	93	97	99	100	---	---
	09-25-92	---	---	---	---	---	---	---	---
	06-03-93	73	81	90	96	98	99	100	---
332007112054301	09-04-92	1	3	21	47	71	93	100	---
	09-24-92	---	---	---	---	---	---	---	---
332025112050801	09-04-92	8	24	47	65	77	85	95	100
	09-24-92	---	---	---	---	---	---	---	---
332109112041401	09-08-92	9	25	50	73	90	97	100	---
	09-24-92	---	---	---	---	---	---	---	---
332116112040501	09-08-92	10	16	24	43	79	97	100	---
	09-24-92	---	---	---	---	---	---	---	---
332119112041001	09-08-92	77	93	100	100	---	---	---	---
	09-24-92	---	---	---	---	---	---	---	---
332405111583801	06-14-93	8	10	19	49	75	85	94	100
332414111583701	06-02-93	60	78	91	97	98	99	100	---
332425112062401	09-09-92	76	86	91	93	94	96	100	---
	09-25-92	---	---	---	---	---	---	---	---
332425112064001	09-09-92	60	68	74	77	80	88	100	---
	09-25-92	---	---	---	---	---	---	---	---
332425112065501	09-08-92	47	64	82	91	94	96	100	---
	09-25-92	---	---	---	---	---	---	---	---
	06-08-93	44	50	54	57	60	67	85	100
332509111584001	06-02-93	53	66	78	86	91	95	97	100
332513111582601	06-02-93	36	47	62	73	79	86	94	100
333435112181401	09-25-92	38	66	86	95	97	99	100	---
333438112181401	09-25-92	67	79	90	96	99	100	---	---
333439112181701	09-25-92	93	98	99	100	---	---	---	---
333458112090401	06-03-93	51	58	71	81	88	96	100	---
333459112090401	06-03-93	25	33	41	50	68	92	100	---
333543112090501	06-03-93	41	53	68	78	85	92	100	---
333627112110601	06-03-93	31	45	60	73	82	89	96	100
333704112120501	06-03-93	23	33	49	65	77	83	87	100

Table 9. Results of acute toxicity analyses of urban stormwater and streamflow, Maricopa County, Arizona

[LC50 and LC20, concentration of stormwater (in percent, by volume) that killed 50 or 20 percent of the organisms, respectively; LOEC, lowest observed effect concentration; NOEC, no observed effect concentration; >, greater than]

Date	Land use	Sample type	Species	LC50	LC20	LOEC	NOEC
Indian Bend Wash at Curry Road, at Tempe, Arizona (09512162)							
02-08-94	Stream channel	Streamflow	Fathead minnow	70.7	58	100	50
02-08-94	Stream channel	Streamflow	<i>Ceriodaphnia dubia</i>	>100	>100	>100	100
Salt River at Priest Drive, at Phoenix, Arizona (09512165)							
12-31-92	Stream channel	Streamflow	Fathead minnow	>100	>100	>100	100
12-31-92	Stream channel	Streamflow	<i>Ceriodaphnia dubia</i>	>100	>100	>100	100
01-04-93	Stream channel	Streamflow	Fathead minnow	>100	>100	>100	100
01-04-93	Stream channel	Streamflow	<i>Ceriodaphnia dubia</i>	>100	>100	>100	100
01-12-93	Stream channel	Streamflow	Fathead minnow	>100	>100	>100	100
01-12-93	Stream channel	Streamflow	<i>Ceriodaphnia dubia</i>	>100	>100	>100	100
Box culvert at 48th Street drain (09512184)							
02-28-93	Light industry	First flush	Fathead minnow	>100	>100	>100	100
02-28-93	Light industry	First flush	<i>Ceriodaphnia dubia</i>	>100	>100	>100	100
02-28-93	Light industry	Flow weight	Fathead minnow	>100	>100	>100	100
02-28-93	Light industry	Flow weight	<i>Ceriodaphnia dubia</i>	>100	>100	>100	100
03-26-93	Light industry	First flush	Fathead minnow	>100	>100	>100	100
03-26-93	Light industry	First flush	<i>Ceriodaphnia dubia</i>	>100	100	100	50
03-26-93	Light industry	Flow weight	Fathead minnow	>100	>100	>100	100
03-26-93	Light industry	Flow weight	<i>Ceriodaphnia dubia</i>	>100	100	6.25	6.25
08-24-93	Light industry	First flush	Fathead minnow	70.7	58	100	50
08-24-93	Light industry	First flush	<i>Ceriodaphnia dubia</i>	>100	>100	>100	100
08-24-93	Light industry	Flow weight	Fathead minnow	>100	>100	>100	100
08-24-93	Light industry	Flow weight	<i>Ceriodaphnia dubia</i>	>100	>100	>100	100
27th Avenue at Salt River (09512403)							
11-13-93	Heavy industry	First flush	Fathead minnow	66	55	100	50
11-13-93	Heavy industry	First flush	<i>Ceriodaphnia dubia</i>	>100	>100	>100	100
11-13-93	Heavy industry	Flow weight	Fathead minnow	>100	>100	>100	100
11-13-93	Heavy industry	Flow weight	<i>Ceriodaphnia dubia</i>	>100	>100	>100	100
02-07-94	Heavy industry	First flush	Fathead minnow	45.2	26	50	25
02-07-94	Heavy industry	First flush	<i>Ceriodaphnia dubia</i>	>100	>100	>100	100
02-07-94	Heavy industry	Flow weight	Fathead minnow	70.7	58	100	50
02-07-94	Heavy industry	Flow weight	<i>Ceriodaphnia dubia</i>	>100	>100	>100	100
03-25-94	Heavy industry	First flush	Fathead minnow	>100	>100	6.25	<6.25
03-25-94	Heavy industry	First flush	<i>Ceriodaphnia dubia</i>	>100	>100	>100	100
03-25-94	Heavy industry	Flow weight	Fathead minnow	59	36	100	50
03-25-94	Heavy industry	Flow weight	<i>Ceriodaphnia dubia</i>	>100	>100	>100	100
Agua Fria River tributary at Yonngtown, Arizona (09513700)							
03-26-93	Residential	First flush	Fathead minnow	61.6	55	25	12.5

Table 9. Results of acute toxicity analyses of urban stormwater and streamflow, Maricopa County, Arizona

Date	Land use	Sample type	Species	LC50	LC20	LOEC	NOEC
Agua Fria River tributary at Youngtown, Arizona (09513700)—Continued							
03-26-93	Residential	First flush	<i>Ceriodaphnia dubia</i>	17.7	14.7	25	12.5
03-26-93	Residential	Flow weight	Fathead minnow	>100	>100	>100	100
03-26-93	Residential	Flow weight	<i>Ceriodaphnia dubia</i>	25.7	10	12.5	6.25
08-06-93	Residential	First flush	Fathead minnow	66	55	100	50
08-06-93	Residential	First flush	<i>Ceriodaphnia dubia</i>	>100	>100	>100	100
08-06-93	Residential	Flow weight	Fathead minnow	66	55	100	50
08-06-93	Residential	Flow weight	<i>Ceriodaphnia dubia</i>	>100	>100	>100	100
08-29-93	Residential	First flush	Fathead minnow	55.5	35	50	25
08-29-93	Residential	First flush	<i>Ceriodaphnia dubia</i>	>100	>100	>100	100
08-29-93	Residential	Flow weight	Fathead minnow	>100	>100	>100	100
08-29-93	Residential	Flow weight	<i>Ceriodaphnia dubia</i>	>100	>100	>100	100
10-06-93	Residential	First flush	Fathead minnow	22.7	13.8	25	12.5
10-06-93	Residential	First flush	<i>Ceriodaphnia dubia</i>	>100	>100	>100	100
10-06-93	Residential	Flow weight	Fathead minnow	>100	>100	>100	100
10-06-93	Residential	Flow weight	<i>Ceriodaphnia dubia</i>	>100	>100	>100	100
43rd and Peoria (09513885)							
01-06-93	Commercial	First flush	Fathead minnow	70.7	58	100	50
01-06-93	Commercial	First flush	<i>Ceriodaphnia dubia</i>	70.7	58	100	50
01-06-93	Commercial	Flow weight	Fathead minnow	>100	66	100	50
01-06-93	Commercial	Flow weight	<i>Ceriodaphnia dubia</i>	>100	>100	>100	100
02-08-93	Commercial	First flush	Fathead minnow	>100	>100	>100	100
02-08-93	Commercial	First flush	<i>Ceriodaphnia dubia</i>	>100	>100	>100	100
02-08-93	Commercial	Flow weight	Fathead minnow	>100	>100	>100	100
02-08-93	Commercial	Flow weight	<i>Ceriodaphnia dubia</i>	>100	>100	>100	100
08-24-93	Commercial	First flush	Fathead minnow	8.8	7.5	6.25	<6.25
08-24-93	Commercial	First flush	<i>Ceriodaphnia dubia</i>	8.8	7.5	6.25	<6.25
08-24-93	Commercial	Flow weight	Fathead minnow	>100	>100	>100	100
08-24-93	Commercial	Flow weight	<i>Ceriodaphnia dubia</i>	>100	>100	>100	100
10-06-93	Commercial	First flush	Fathead minnow	26.6	18	25	12.5
10-06-93	Commercial	First flush	<i>Ceriodaphnia dubia</i>	>100	>100	>100	100
10-06-93	Commercial	Flow weight	Fathead minnow	62.5	31.4	25	12.5
10-06-93	Commercial	Flow weight	<i>Ceriodaphnia dubia</i>	>100	>100	>100	100
Arizona Canal Diversion Channel (333543112090501)							
02-07-94	Drainage channel	Streamflow	Fathead minnow	>100	>100	>100	100
02-07-94	Drainage channel	Streamflow	<i>Ceriodaphnia dubia</i>	>100	>100	>100	100
03-07-94	Drainage channel	Streamflow	Fathead minnow	>100	85	100	50
03-07-94	Drainage channel	Streamflow	<i>Ceriodaphnia dubia</i>	>100	73	25	12.5
05-25-94	Drainage channel	Streamflow	Fathead minnow	>100	>100	>100	100
05-25-94	Drainage channel	Streamflow	<i>Ceriodaphnia dubia</i>	>100	>100	>100	100

Table 10. Results of toxicity identification evaluations of urban stormwater and streamflow, Maricopa County, Arizona

[EDTA, ethylenediaminetetraacetic acid; mL, milliliter; Mx, mixed land use; LI, light industry; HI, heavy industry; R, residential; C, commercial; EWI, equal-width increment; FF, first flush; FW, flow weight; ----, dashes indicate no data]

Mortality of species, in percent														
Dste	Land use	Sam- ple type	Species	Original	Base- line	Oil and grease skim	Solids filtration	EDTA (0.0 mL)	EDTA (0.012 mL)	EDTA (0.025 mL)	EDTA (0.050 mL)	Anionic organics (pH 3)	Neutral organics (initial pH)	Cationic organics (pH 9)
Indian Bend Wash at Curry Road, at Tempe, Arizona (09512162)														
02-08-94	Mx	EWI	Fathead minnow	100	10	0	0	0	0	0	0	0	0	20
Box culvert at 48th Street drain (09512184)														
08-24-93	LI	FF	Fathead minnow	100	20	20	40	0	20	10	20	60	20	0
27th Avenue at Salt River (09512403)														
11-13-93	HI	FF	Fathead minnow	100	20	10	20	20	20	0	0	0	0	0
02-07-94	HI	FF	Fathead minnow	100	30	0	0	0	20	20	0	60	10	0
02-07-94	HI	FW	Fathead minnow	100	0	30	10	0	0	0	0	80	20	0
03-25-94	HI	FW	Fathead minnow	90	50	60	40	0	0	0	60	100	20	20
Agua Fria River tributary at Youngtown, Arizona (09513700)														
03-26-93	R	FF	Fathead minnow	100	40	30	30	60	20	80	100	0	0	0
03-26-93	R	FF	<i>Ceriodaphnia dubia</i>	100	100	100	100	100	100	100	100	0	0	0
03-26-93	R	FW	<i>Ceriodaphnia dubia</i>	100	100	70	100	100	100	100	100	0	0	0
08-06-93	R	FF	Fathead minnow	100	90	30	40	100	100	60	---	100	20	20
08-06-93	R	FW	Fathead minnow	100	50	40	10	20	40	60	---	0	0	0
08-29-93	R	FF	Fathead minnow	100	60	30	10	80	60	80	40	20	20	0
10-06-93	R	FF	Fathead minnow	100	40	70	50	60	20	100	40	0	0	0
43rd and Peoria (09513885)														
08-24-93	C	FF	Fathead minnow	100	100	100	100	100	100	100	100	20	40	40
08-24-93	C	FF	<i>Ceriodaphnia dubia</i>	100	100	100	100	100	100	100	100	20	100	40
10-06-93	C	FF	Fathead minnow	100	100	100	100	100	100	100	100	100	40	60
10-06-93	C	FW	Fathead minnow	80	80	60	70	100	80	20	40	100	100	20
Arizona Canal Drainage Channel (333543112090501)														
03-07-94	Mx	EWI	Fathead minnow	25	10	10	0	0	0	20	60	0	0	0
03-07-94	Mx	EWI	<i>Ceriodaphnia dubia</i>	75	90	100	100	100	40	0	40	0	0	0

Table 11. Results of acute toxicity analyses of sediment, Maricopa County, Arizona

[ACDC, Arizona Canal Diversion Channel]

Station number	Site description	Date	Land use	Soil cover	Mortality rate of <i>Hyaella azteca</i> , in percent ¹
332425112065501	27th Street, 1st catch basin.....	09-08-92	Heavy industry	Bare soil	97
332425112065501	27th Street, 1st catch basin.....	06-08-93	Heavy industry	Bare soil	5
332425112064001	27th Street, 2nd catch basin	09-09-92	Heavy industry	Bare soil	57
332425112062401	27th Street, 5th catch basin	09-09-92	Heavy industry	Bare soil	20
09513700	Agua Fria River tributary at Youngtown, Arizona.....	09-09-92	Residential	Bare soil	83
09513700	Agua Fria River tributary at Youngtown, Arizona ²	09-09-92	Residential	Bare soil	71
09513700	Agua Fria River tributary at Youngtown, Arizona	06-03-93	Residential	Bare soil	61
332513111582601	48th Street, undeveloped lot.....	06-02-93	Light industry	Bare soil	3
332509111584001	48th Street, near freeway	06-02-93	Light industry	Bare soil	77
332414111583701	48th Street, near restaurant.....	06-02-93	Light industry	Bare soil	79
332405111583801	48th Street, near gaging station.....	06-14-93	Light industry	Bare soil	100
333459112090401	Peoria, parking lot.....	06-03-93	Commercial	Bare soil	56
333458112090401	Peoria, dirt lot.....	06-03-93	Commercial	Bare soil	28
09512162	Indian Bend Wash at Curry Road, at Tempe, Arizona.....	06-09-93	Channel ³	Grass	0
09512090	Indian Bend Wash at Shea.....	06-09-93	Channel	Grass	5
333543112090501	ACDC at 51st Avenue.....	06-03-93	Channel	Grass	3
333627112110601	ACDC at 59th Avenue	06-03-93	Channel	Grass	7
333704112120501	ACDC at 67th Avenue	06-03-93	Channel	Grass	30
332007112054301	South Mountain Park, Prospects	09-04-92	Undeveloped	Bare soil	17
332025112050801	South Mountain Park, above track.....	09-04-92	Undeveloped	Bare soil	83
09512200	Salt River tributary in South Mountain Park, at Phoenix, Arizona	09-04-92	Undeveloped	Bare soil	98
332119112041001	South Mountain Park, retention pond #1	09-08-92	Undeveloped	Bare soil	100
332116112040501	South Mountain Park, retention pond #2	09-08-92	Undeveloped	Bare soil	79
332109112041401	South Mountain Park, retention pond #3	09-08-92	Undeveloped	Bare soil	76
333439112181701	Agua Fria above Youngtown	09-25-92	Channel	Bare soil	26
333438112181401	Agua Fria at Youngtown	09-25-92	Channel	Bare soil	16
333435112181401	Agua Fria below Youngtown	09-25-92	Channel	Bare soil	29
09512165	Salt River at Priest Drive, at Phoenix, Arizona	06-14-93	Channel	Bare soil	95
09512190	Salt River at 24th Street	06-09-93	Channel	Bare soil	10
09512405	Salt River at 35th Avenue	06-09-93	Channel	Bare soil	86

¹*Hyaella azteca* was exposed to 100 grams of bed material, sieved to less than 63 micron, and 400 milliliters of hard reconstituted water.²Duplicate sample.³Channel samples were taken from ephemeral channels.