

# **Hydrologic and Water-Quality Data at Selected Sites in the Upper Animas River Watershed, Southwestern Colorado, 1997-99**

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# Hydrologic and Water-Quality Data at Selected Sites in the Upper Animas River Watershed, Southwestern Colorado, 1997-99

By M. Alisa Mast, Jonathan B. Evans, Kenneth J. Leib, and Winfield G. Wright

U.S. GEOLOGICAL SURVEY

Open-File Report 00-53

Denver, Colorado 2000

# U.S. DEPARTMENT OF THE INTERIOR BRUCE BABBITT, Secretary

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U.S. GEOLOGICAL SURVEY Charles G. Groat, Director

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# **CONTENTS**



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# **PLATE**

[Plate is in pocket attached to inside back cover]

1. Map of upper Animas River watershed and locations of water-quality sampling sites at selected springs and streams in southwestern Colorado

# **FIGURES**



# **TABLES**



[CD-ROM containing tables 8-28 and abbreviations is in pocket attached to inside front cover]

Hydrologic and water-quality data tables and abbreviations  $8-19$ . Daily mean streamflow for:





## **CONVERSION FACTORS AND ABBREVIATIONS**



Temperature in degrees Fahrenheit (°F) may be converted to degrees Celsius (°C) using the following equation:

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

Additional abbreviations used in this report:

micrometer  $(\mu m)$ microsiemens per centimeter at 25 degrees Celsius ( $\mu$ S/cm) milligram per liter (mg/L) microgram per liter  $(\mu g/L)$ milliliter (mL) millimeter (mm)

# **Hydrologic and Water-Quality Data at Selected Sites in the Upper Animas River Watershed, Southwestern Colorado, 1997-99**

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# **Abstract**

The water quality and aquatic resources of the upper Animas River watershed are affected by more than 1,500 abandoned mines from historical metal-mining activities in the late 1800's and early 1900's. In 1997, the U.S. Geological Survey implemented the Abandoned Mine Lands Initiative to provide scientific information to Federal land-management agencies responsible for remediation of abandoned mine sites on publicly owned land. This report presents hydrologic and water-quality data collected as part of the Abandoned Mine Lands Initiative during water years 1997-99 in the upper Animas River watershed, Colorado. Water-quality samples were collected from four streamflow-gaging stations, nine sites along the main stem and two major tributaries, 55 abandoned mine sites, and 194 selected stream and spring sites. Data include field measurements of streamflow, specific conductance, pH, water temperature, and dissolved oxygen and laboratory analyses of major inorganic and selected traceelement constituents. Daily mean discharge values are tabulated for the four streamflowgaging stations. A CD-ROM containing hydrologic and water-quality data is in the pocket of this report.

# **INTRODUCTION**

The upper Animas River watershed (UARW) lies in the southwestern part of the Colorado mineral belt and was heavily mined in the late 1800's and early 1900's for gold, silver, zinc, and lead. Numerous

prospects, draining mine adits, waste-rock piles, and mill tailings left behind by historical mining activities have severely affected the water quality and aquatic resources of the Animas River and its tributaries. Recently, there has been much interest by local stakeholders, and by State and Federal agencies, in remediating abandoned mine sites in the watershed with the goal of restoring aquatic ecosystems along some reaches of the Animas River. Effective and costefficient remediation of abandoned mine lands requires knowledge of the degree of contamination of the environment as well as an understanding of the processes by which abandoned mine lands disturb the natural ecosystem. In 1997, the U.S. Geological Survey (USGS) implemented the Abandoned Mine Lands Initiative (AMLI) to provide this type of scientific information to the Federal land-management agencies responsible for remediation of abandoned mine sites in the UARW (Nimick and von Guerard, 1998). This is part of a larger strategy by the U.S. Department of the Interior and the U.S. Department of Agriculture to coordinate activities between Federal and State agencies and local stakeholders for the cleanup of Federal lands affected by abandoned mine lands. One of the main components of the AMLI is to describe the geological, hydrological, and biological characteristics of the watershed. This information will be used to improve understanding of the processes that control the occurrence and transport of contaminants and to identify mining sites that have the most adverse effect on the water quality and aquatic resources of the Animas River.

### **Purpose and Scope**

The purpose of this report is to present hydrologic and water-quality data and methods of data

collection at selected sampling sites in the UARW in southwestern Colorado (fig. 1). Data were collected during water years 1997-99 as part of the AMLI. Bimonthly to weekly water-quality samples were collected at four streamflow-gaging stations and nine sites along the main-stem upper Animas River and two major tributaries to monitor seasonal variations in stream chemistry and solute fluxes in the watershed. The four streamflow-gaging stations are the Animas River below Silverton (station 09359020; A72), Animas River at Silverton (station 09358000; A68), Cement Creek at Silverton (station 09358550; C48), and Mineral Creek at Silverton (station 09359010; M34). One hundred and thirty-eight water-quality samples were collected at 55 selected mine sites during the study period to characterize spatial and seasonal variability in mine-water chemistry and discharge. Water-quality samples at 194 selected streams and springs were sampled throughout the watershed during summer base-flow conditions to help distinguish natural from mining-related sources of metals to surface water. Data presented in this report include field measurements of streamflow, specific conductance, pH, water temperature, and dissolved oxygen and laboratory analyses of concentrations of major inorganic and selected trace-element constituents. Daily mean discharge values for streamflow also are presented for the four streamflow-gaging stations. Supplemental data are on the CD-ROM in the pocket of the report cover.

#### **Acknowledgments**

Many aspects of this study would not have been possible without the support of the Upper Animas River Stakeholders Group; the U.S. Department of the Interior, Bureau of Land Management and Bureau of Reclamation; and the U.S. Department of Agriculture Forest Service. The authors are grateful for assistance and guidance from USGS employees Greg Alexander, James Bennett, Mark Gress, Dave Grey, Paul von Guerard, David Roth, Jenny A. Taylor, and Philip Verplanck.

# **DESCRIPTION OF STUDY AREA**

The upper Animas River watershed is in the San Juan Mountains in southwestern Colorado. The river drains  $146 \text{ mi}^2$  of rugged mountainous terrain ranging in elevation from 9,200 ft near the town of Silverton, Colorado, to 13,894 ft at the summit of Vermilion Peak (fig. 1). The upper Animas River is a southwardflowing tributary of the San Juan River and has a channel length of about 15.5 mi above the streamflowgaging station at Animas River below Silverton (A72) and an average stream gradient of about 184 ft/mi. The main channel of the river is perennial, and mean monthly discharge ranges from about 63 ft $^{3}/s$  in February to  $1,196$  ft<sup>3</sup>/s in June. Snowmelt is the major hydrologic event of the year, and more than 60 percent of the annual streamflow occurs in May, June, and July. Average annual runoff for the watershed was 28 inches from 1992 to 1998 (Crowfoot and others, 1999). The Animas River has two major tributaries, Cement Creek and Mineral Creek, which together account for about 50 percent of the annual discharge from the watershed. Climate of the area is characterized by long, cold winters and short (3-4 months), cool summers. Average monthly air temperature at Silverton, Colorado, ranges from 16.0°F in January to 55.3°F in July (URL http://www.wrcc.sage.dri.edu, accessed March 2000). Precipitation averages 45 inches annually, of which about 70 percent accumulates in a seasonal snowpack between November and April (Wirt and others, 1999). Most of the remaining precipitation falls during monsoonal thundershowers in late summer and early fall. The watershed lies in the Southern Rocky Mountain Forest ecoregion, and much of the area is above treeline, which is at an average elevation of 11,600 ft. The lower slopes are covered by a dense forest of Engelmann spruce and subalpine fir (Bailey and others, 1994).

The UARW lies within the San Juan volcanic field of Tertiary age, which includes the San Juan, Uncompahgre, and Silverton calderas that formed about 28 million years ago (Lipman and others, 1976; Bove and others, 1999). Most of the bedrock in the study area consists of thick sequences of volcanic rocks that are composed of lava flows, ash-flow tuffs, tuff breccias, conglomerates, and mudflow breccias (Lipman and others, 1973; Ringrose, 1982). The lavas and tuffs generally are of intermediate composition and are composed of plagioclase, biotite, augite, quartz, and hornblende. Nearly all the volcanic rocks in the study area were subject to some degree of hydrothermal alteration after deposition (Burbank and Luedke, 1969). Hydrothermal activity related to the Silverton caldera caused regional propylitization of the volcanic rocks, which altered the rocks to chlorite and



**Figure 1.** Upper Animas River watershed and locations of four USGS streamflow-gaging stations and nine water-quality sampling sites along<br>the main stern and major tributaries. **Figure 1.** Upper Animas River watershed and locations of four USGS streamflow-gaging stations and nine water-quality sampling sites along the main stem and major tributaries.

epidote with calcite and pyrite often disseminated throughout the groundmass. The propylitized rocks were subsequently mineralized and hydrothermally altered during three later episodes of magmatism (11-23 million years ago), which were the source of most ore deposition in the area (Casadevall and Ohmoto, 1977). Most economic ore deposits occur in polymetallic vein systems that follow structures formed during subsidence and resurgence of the San Juan, Uncompahgre, and Silverton calderas (Burbank and Luedke, 1969; Ringrose, 1982). The area was first prospected for precious metals in 1860, and most mining activities occurred in the late 1800's and early 1900's. Production was primarily from vein type deposits that yielded gold, silver, zinc, and lead. Over 1,500 prospect pits, abandoned mines, and dismantled mill sites are now scattered throughout the watershed as a result of past mining activities.

# **METHODS OF STUDY**

This section includes a description of the methods used to collect and analyze water-quality samples collected in the UARW as part of the AMLI. Methods of operation for streamflow-gaging stations

are standardized within the USGS (Rantz and others, 1982) and will not be included in this report. Daily mean streamflow values at the four streamflow-gaging stations for 1997-99 are presented in the "Hydrologic and Water-Quality Data" section of this report and also are in Crowfoot and others (1998), Crowfoot and others (1999), and Crowfoot and others (U.S. Geological Survey, written commun., 2000).

# **Water-Quality Sampling**

Water-quality samples were collected seasonally (weekly to bimonthly) at the four streamflow-gaging stations and at nine upstream sites along the main stem Animas River and major tributaries. The sites are listed in table 1 and locations shown in figure 1. Seasonal (monthly to bimonthly) samples also were collected at seven mine sites: the Avalanche Mine, Bandora Mine, Bonner Mine, Elk Tunnel, Evelyn Mine, Forest Queen Mine, and Mighty Monarch Mine. The mine sites are listed in table 2 and locations shown in figure 2. Field measurements at the sampling sites included stream discharge, specific conductance, pH, water temperature, and dissolved oxygen. Waterquality measurements were made using methods

**Table 1.** Streamflow-gaging stations and water-quality sampling sites on the main stem and major tributaries in the upper Animas River watershed

[Site locations are shown in figure 1; identification number is USGS downstream order number; elevation in feet; watershed area in square miles; --, not reported]



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**Table 2.** Water-quality sampling sites at selected mines in the upper Animas River watershed



[Site locations are shown in figure 2; elevation in feet; site MS24 eliminated from this report]

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[Site locations are shown in figure 2; elevation in feet; site MS24 eliminated from this report] Site number MS39 MS40 MS41 MS42 MS43 MS44 MS45 MS46 **MS47** MS48 MS49 MS50 MS51 MS52 MS53 MS54 MS55 MS56 Site name Mine in upper Minnie Gulch Mine in upper North Fork Cement Creek Mine above Brooklyn #6 Mine near Horseshoe Bend Mine near Hwy 550 south of Middle Fork Mineral Creek Mine northwest of Burro Bridge Mine near McMillan Peak Mine west of Burro Bridge Mine near Eureka Gulch Mine near Burro Bridge Old Hundred Mine Queen Anne Mine Silver Crown Mine Silver Ledge Mine Silver Wing Mine Treasure Mountain Mine Vermilion Mine Yukon Mine Latitude 37°50'17" 37°53'50" 37°51'53" 37°52'26" 37°50'25" 37°51'20" 37°52'46" 37°51'02" 37°53'07" 37°51'01" 37°49'24" 37°54'52" 37°52'18" 37°52'36" 37°54'12" 37°54'49" 37°56'10" 37°50'58" Longitude 107°31'33" 107°37'37" 107°42'35" 107°43'58" 107°43'31" 107°43'27" 107°41'43" 107°43'45" 107°35'28" 107°43'30" 107°35'04" 107°37'47" 107°44'35" 107°38'38" 107°33'19" 107°34'11" 107°36'01" 107°40'32" Elevation 12,000 12,650 11,890 10,420 10,080 10,440 12,450 10,200 10,680 10,040 9,980 12,200 11,800 10,970 10,440 11,640 12,400 9,985

Table 2. Water-quality sampling sites at selected mines in the upper Animas River watershed—Continued





similar to those described by Wilde and Radtke (1998). Field meters were calibrated in the morning on the day of the sample collection. The pH meters were calibrated with pH-2 and pH-4 buffers or pH-4 and pH-7 buffers to bracket the range of pH values measured in the field. Specific-conductance (SC) meters were calibrated with two standards that bracketed the range of values expected at the sampling sites. Dissolved oxygen (DO) meters were calibrated to the saturated DO of water according to the water temperature and barometric pressure at each sampling site. The DO meters also were checked against a solution with a DO concentration of 0 mg/L. Instantaneous discharge at the time of sampling was determined from the stage reading and stage-discharge-rating table at the four streamflow-gaging stations. At the nine upstream sites and seven mine sites, instantaneous discharge was measured using a current meter according to methods described by Rantz and others (1982). Standard USGS sampling techniques as described in Ward and Hair (1990) and Shelton (1994) were used to collect water-quality samples. Waterquality samples at the stream sites were collected using depth-integrating samplers utilizing the equalwidth-increments (EWI) method (Edwards and Glysson, 1988) then transferred into precleaned 2-L or 3-L polyethylene bottles. Mine-site samples were collected in precleaned polyethylene bottles by using the dip method.

Filtering and processing of water samples was done in a field laboratory in Silverton, Colorado, within 5 hours of sample collection. Standard USGS equipment (Horowitz and others, 1994) was used. Samples requiring filtered water were passed through a 0.45-urn cellulose filter housed in a 47-mm polycarbonate filtering unit using a peristaltic pump. The filter housing and pump tubing were soaked in dilute nitric acid and rinsed six times with deionized water between samples. Occasionally samples containing substantial amounts of sediment or colloids, were filtered using a 0.45-um capsule filter instead of the plate filter. Filtered and unfiltered aliquots for dissolved and total major inorganic and trace-element constituents were preserved with concentrated ultrapure nitric acid (1 mL per 250 mL of sample). A filtered unpreserved aliquot was collected for sulfate analysis, and an unfiltered unpreserved aliquot was collected for the alkalinity determination. For selected samples, a filtered aliquot was preserved with 6 molar hydrochloric acid for ferrous iron analysis. Collection and processing procedures similar to those described above were used for the remaining mine sites listed in table 2 and the selected spring and stream sites listed in table 3 and shown in plate 1, except that samples were filtered and processed at the sampling site rather than in the laboratory.

# **Analytical Techniques**

Analytical techniques used for water-quality analyses and the corresponding analytical detection limits are listed in table 4. Alkalinity was determined by incremental titration on unfiltered samples with pH greater than 4.5. The analysis was conducted at the sampling site or within 5 hours of sample collection at a USGS field laboratory in Silverton, Colorado. Major inorganic and selected trace-element constituents were analyzed on filtered and unfiltered (raw) samples by inductively coupled plasma-atomic emission spectroscopy (ICP-AES) and potassium was analyzed by atomic absorption spectroscopy (AA) at a USGS research laboratory in Boulder, Colorado. The unfiltered samples were digested in the laboratory with concentrated hydrochloride acid at 60 C for 8 hours according to the method of Hoffman and others (1996). The digested samples were filtered through 0.45-um polycarbonate filters into acid-washed polyethylene bottles prior to ICP-AES and AA analysis. Sulfate was determined by a turbidimetric method using a Hach SulfoVer reagent and ferrous iron  $(Fe^{+2})$ was determined by a colorimetric method using a Hach 1, 10-phenanthroline reagent. The sulfate and  $Fe<sup>+2</sup>$  determinations were measured on filtered samples using a Hach DR2000 spectrophotometer with a 25-mL sample cell. The quality of all laboratory analyses was assessed through the analysis of laboratory blanks, sample duplicates, USGS standard reference water samples (Long and Farrar, 1995), and calculated ion balances.

# **QUALITY ASSURANCE**

Quality-assurance procedures used during the study period included daily calibration of waterquality meters, cleaning of sampling equipment and rinsing of sample bottles, and collection of qualitycontrol samples (field equipment blanks and replicate samples). Results of 15 field equipment blanks and



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[UA, upper Animas River above Silverton; CC, Cement Creek; MC, Mineral Creek; site locations are shown on plate 1]

[UA, upper Animas River above Silverton; CC, Cement Creek; MC, Mineral Creek; site locations are shown on plate 1]



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[UA, upper Animas River above Silverton; CC, Cement Creek; MC, Mineral Creek; site locations are shown on plate 1]



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[UA, upper Animas River above Silverton; CC, Cement Creek; MC, Mineral Creek; site locations are shown on plate 1]



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#### **Table 4.** Analytical techniques and corresponding detection limits used for water-quality analyses

[ICP-AES, inductively coupled plasma-atomic emission spectroscopy; AA, atomic absorption spectroscopy; mg/L, milligrams per liter; ug/L, micrograms per liter]



18 replicate samples collected during the study period are presented in tables 5–7 and are discussed below.

# **Field Equipment Blanks**

A field equipment blank is a sample prepared using blank (deionized) water that is passed through all the sampling and processing equipment (Spahr and Boulger, 1997). This type of sample is used to check for potential contamination of the environmental samples during sampling collection, processing, handling, and analysis. Analytical results for the 15 field equipment blanks collected during the study period are listed in table 5. The measured concentrations for most constituents were at or below the detection limits of the analytical methods. Several samples did have detectable concentrations of molybdenum, although most concentrations were less than twice the analytical detection limit.

# **Concurrent and Sequential Replicates**

Concurrent and sequential replicate samples are collected to estimate variability in the environmental data. In the concurrent replicate, one sample is collected concurrently with the environmental sample by a second collection team or by using a second set of collection equipment. In the sequential replicate, a replicate is collected as close in time as possible to the environmental sample. Each sample is processed through all the normal steps of a regular water-quality sample (Spahr and Boulger, 1997). For each step of sample processing, the environmental sample is processed before the replicate sample. The replicate samples were processed using a clean filter and equipment.

Ten sequential and eight concurrent replicates were collected during the study period. Comparison of analytical results between environmental samples and sequential replicates and between environmental samples and concurrent replicates for major inorganic and trace-element constituents are listed in tables 6 and 7. The sequential replicates were collected using the same sampling team and the same collection procedures and analytical methods discussed previously in this report. The concurrent replicates were collected as part of another USGS data-collection program, and samples were analyzed at the USGS National Water Quality Laboratory in Lakewood, Colorado (Crowfoot and others, 1998; Crowfoot and others, 1999; Crowfoot and others, written commun., 2000). The majority of the differences between the environmental and the sequential replicates agree

Table 5. Chemical analyses of field equipment blanks collected during the study period **Table 5.** Chemical analyses of field equipment blanks collected during the study period

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[concentrations in micrograms per liter except calcium, magnesium, sodium, potassium, sulfate, and silica in milligrams per liter; diss., dissolved; --, no data; <, less than] [concentrations in micrograms per liter except calcium, magnesium, sodium, potassium, sulfate, and silica in milligrams per liter; diss., dissolved; , no data; <, less than]



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 $\mathbf{t}$  $\sim1$  Table 6. Comparison of chemical analyses of environmental samples and sequential replicates collected during the study period **Table 6.** Comparison of chemical analyses of environmental samples and sequential replicates collected during the study period

[diss., dissolved; Env., environmental sample; Rep., replicate sample; R, river; Cr, creek; mg/L, milligrams per liter; µg/L, micrograms per liter] [diss., dissolved; Env., environmental sample; Rep., replicate sample; R, river; Cr, creek; mg/L, milligrams per liter; |J.g/L, micrograms per liter]



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[diss., dissolved; Env., environmental sample; Rep., replicate sample; R, river; Cr, creek; mg/L, milligrams per liter; µg/L, micrograms per liter] [diss., dissolved; Env., environmental sample; Rep., replicate sample; R, river; Cr, creek; mg/L, milligrams per liter; (ig/L, micrograms per liter]

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Table 6. Comparison of chemical analyses of environmental samples and sequential replicates collected during the study period--Continued **Table 6.** Comparison of chemical analyses of environmental samples and sequential replicates collected during the study period-Continued

[diss., dissolved; Env., env [diss., dissolved; Env., environmental sample; Rep., replicate sample; R, river; Cr, creek; mg/L, milligrams per liter; µg/L, micrograms per liter] sample; Rep., replicate sample; R, river; Cr, creek; mg/L, milligrams per liter; µg/L, micrograms per liter]



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Table 7. Comparison of chemical analyses of environmental samples and concurrent replicates collected at the Animas River below Silverton during the<br>study period **Table 7.** Comparison of chemical analyses of environmental samples and concurrent replicates collected at the Animas River below Silverton during the study period



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within  $\pm 10$  percent. Many of the trace-element concentrations were less than the analytical detection limit.

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