



Hydrologic Benchmark Network Stations in the Eastern U.S. 1963-95 (USGS Circular 1173-A)

Abstract and Map Index	List of all HBN Stations	Introduction to Circular	Analytical Methods
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Scape Ore Swamp near Bishopville, South Carolina (Station 02135300)

This report details one of the approximately 50 stations in the Hydrologic Benchmark Network (HBN) described in the four-volume U.S. Geological Survey Circular 1173. The suggested citation for the information on this page is:

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All of the tables and figures are numbered as they appear in each circular. Use the navigation bar above to view the abstract, introduction and methods for the entire circular, as well as a map and list of all of the HBN sites. Use the table of contents below to view the information on this particular station.

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Site Characteristics and Land Use

The Scape Ore Swamp HBN Basin is in the Congaree Sand Hills region of the Coastal Plain physiographic province of South Carolina ([Figure 24. Map showing study area in the Scape Ore Swamp Basin and photograph of a typical tributary stream](#)). The 249-km² basin drains an area characterized by small hills with long, gentle slopes and

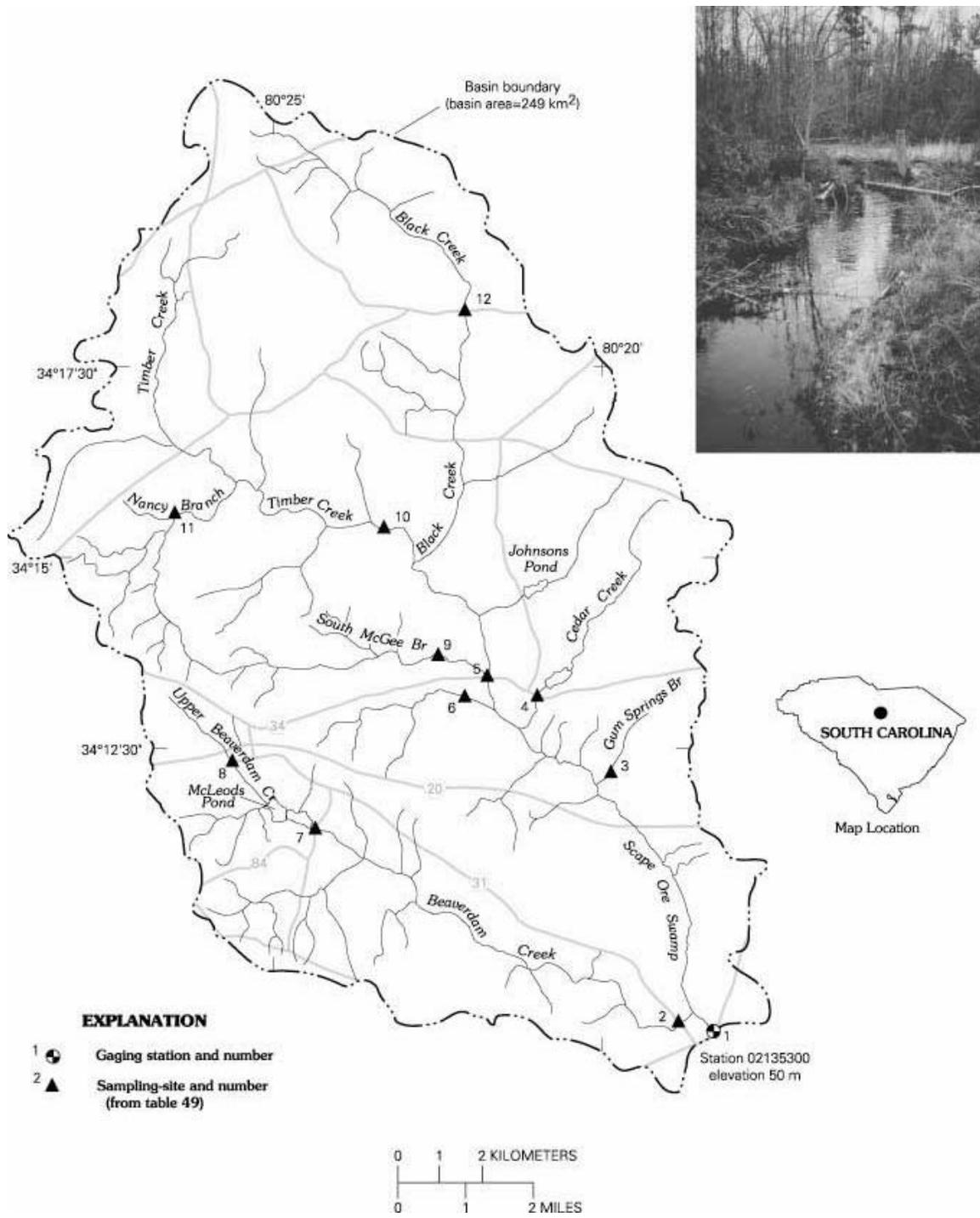


Figure 24. Map showing study area in the Scape Ore Swamp Basin and photograph of a typical tributary stream

broad summits that ranges in elevation from 50 to 120 m. The USGS gaging station is 8 km southwest of the town of Bishopville, S.C., at latitude 34°09'02" and longitude 80°18'18". Scape Ore Swamp is a southeast-flowing tributary of the Pee Dee River with a channel length of about 28 km upstream from the gage and an average stream gradient of 1.4 m/km. The main channel is perennial, and mean monthly discharge varies over a relatively narrow range from a minimum of 1.66 m³/s in July to 4.87 m³/s in March. Average annual runoff from the basin was 38 cm from 1968 through 1995 (U.S. Geological Survey, Water Resources Data, South Carolina). Climate of the area is relatively temperate with warm, humid summers and mild winters. Average daily summer and winter temperatures are 25.5°C and 5.5°C, respectively (Mitchell, 1990). Winter ground frost is common, but snowfall is rare. Precipitation averages 115 cm annually and is fairly evenly distributed throughout the year (Mitchell, 1990).

The basin lies in the Outer Coastal Plain Mixed Forest ecoregion, which is characterized by temperate evergreen forests (Bailey and others, 1994). Forests on hillslopes and upland areas are dominated by southern pine and upland hardwood communities; the dominant species are loblolly pine, longleaf pine, slash pine, shortleaf pine, oak, and hickory. The flood plains are covered by lowland hardwood forests of water tupelo, swamp tupelo, water oak, and bald cypress. Most soils in the basin are classified as Ultisols and mapped in the Alpin and Blanton soil series (Mitchell, 1990). These soils are formed in sandy marine sediments and are generally found on broad ridgetops and side slopes of sand hills. A typical profile has a sandy surface layer, 10 to 30 cm thick, overlying a sandy to loamy subsoil down to a depth of 200 cm. Because of the high sand content, these soils are highly permeable and acidic (pH 4.5 to 6.0) and have a low nutrient-holding capacity. Poorly drained soils in relatively narrow strips along the flood plains are mapped in the Johnson Series (Mitchell, 1990). A typical Johnson soil has a black loam surface layer about 75 cm thick overlying a grayish sandy loam sublayer. The dominant minerals in all soils are quartz and kaolinite (Mitchell, 1990).

The basin is underlain by Upper Cretaceous sediments of the Tuscaloosa Formation (Cooke, 1936). The Tuscaloosa Formation consists of fluvial and marine sediments assumed to be derived from weathered crystalline rocks of the Piedmont. Cooke (1936) described the sediments as light-colored arkosic sands composed of quartz, feldspar, and muscovite, interlayered with lenses of light-colored clay. Argillaceous sands of the Eocene Black Mingo Formation underlie the basin near the gage but are covered by sandy Pleistocene terrace deposits.

The Scape Ore Swamp Basin drains parts of Lee and Kersaw Counties. Land ownership in the basin is 96 percent private and 4 percent Federal, State, and county (Mike Newman, Natural Resources Conservation Service, written commun., 1996). More than 200 km of primary and secondary roads provide access to most areas of the basin. Because of swampy conditions along most of the stream channels, roads tend to follow the sand hills summits rather than parallel the drainages. A 13-km segment of Interstate 20 crosses the center of the basin in the east-west direction. Land use in the basin is 60 percent forest and 40 percent crop and pasture land. Farmland is more commonly situated

on the hillslopes and broad summits of the sand hills, whereas forested areas are concentrated along the drainages. Most farms in the basin are relatively small (average size 70 ha) and generally produce soybeans, corn, peanuts, and cotton; pasture lands are covered by a mixture of Coastal Bermuda, Common Bermuda, annuals, and native grasses (Mike Newman, written commun., 1996). About 30 small impoundments located in the basin are primarily used for irrigation and recreational fishing. Residents in the basin use ground water as their primary domestic water supply. A major resource problem in the basin is soil loss from highly erodible cropland, which results in damage to streams, roads, and property (Lynn Newton, Natural Resources Conservation Service, written commun., 1996). The U.S. Department of Agriculture instituted the Conservation Reserve Program (CRP) in 1985 as a means of reducing soil erosion by paying farmers to plant permanent vegetation types, such as grasses and pines. About 20 percent of agricultural land in the basin is currently managed under the CRP (Wayde Ross, Natural Resources Conservation Service, written commun., 1996).

Historical Water-Quality Data and Time-Series Trends

The data set analyzed for this report includes 212 water-quality samples that were collected from August 1970 through September 1995. Sampling frequency was monthly from 1971 through 1982 and bimonthly from 1983 through 1995. Although not documented, water-quality samples in the early part of the record probably were analyzed at USGS laboratory in Raleigh, N.C., which was operated until 1973 (Durum, 1978). After establishment of the USGS Central Laboratory System, samples were analyzed at the Atlanta laboratory from 1973 through 1985 and at the NWQL in Arvada, Colo., from 1986 through 1995. Daily discharge records for Scape Ore Swamp (station 02135300) are available beginning in July 1978.

Calculated ion balances for 205 samples with complete major ion analyses are shown in [Figures 25a](#) and [25b](#). *Temporal variation of discharge, field pH, major ion concentrations, and ion balance at Scape Ore Swamp, South Carolina*. Ion balances ranged from -45 to +31 percent and more than 65 percent of samples had values within the ± 10 percent range, indicating that the analytical measurements were generally of good quality. The average charge balance for all samples was 1.1 percent, indicating measured cations and anions were close to balanced. Time-series plots of the major dissolved constituents were inspected for evidence of method-related effects (fig. 25). Several patterns are evident at the Scape Ore station. For example, elevated sulfate concentrations were reported for most samples during the late 1980's. This pattern coincides with the use of a turbidimetric titration for sulfate analyses at the NWQL between March 1986 and December 1989 (Fishman and others, 1994). In 1989, the NWQL determined that sulfate concentrations can be overestimated by this technique and changed the method to ion chromatography in 1990 (Office of Water Quality Technical Memorandum No. 90.04, Turbidimetric Sulfate Method, issued December 21, 1989, at URL <http://water.usgs.gov/public/admin/memo/>). Uncharacteristically high calcium concentrations also were evident during the period 1974 through 1976. Interestingly, a similar pattern of high calcium concentrations also was observed at the Wild River

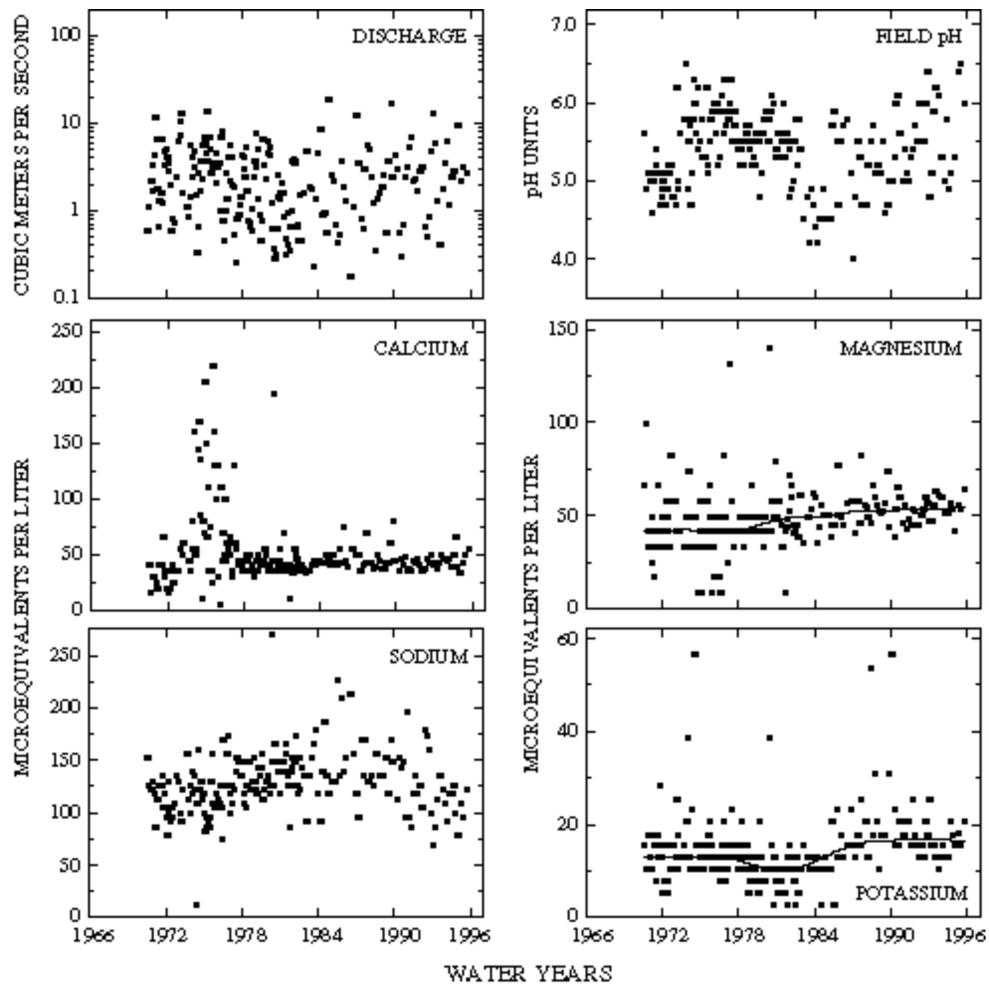


Figure 25a. Temporal variation of discharge, field pH, major ion concentrations, and ion balance at Scape Ore Swamp, South Carolina

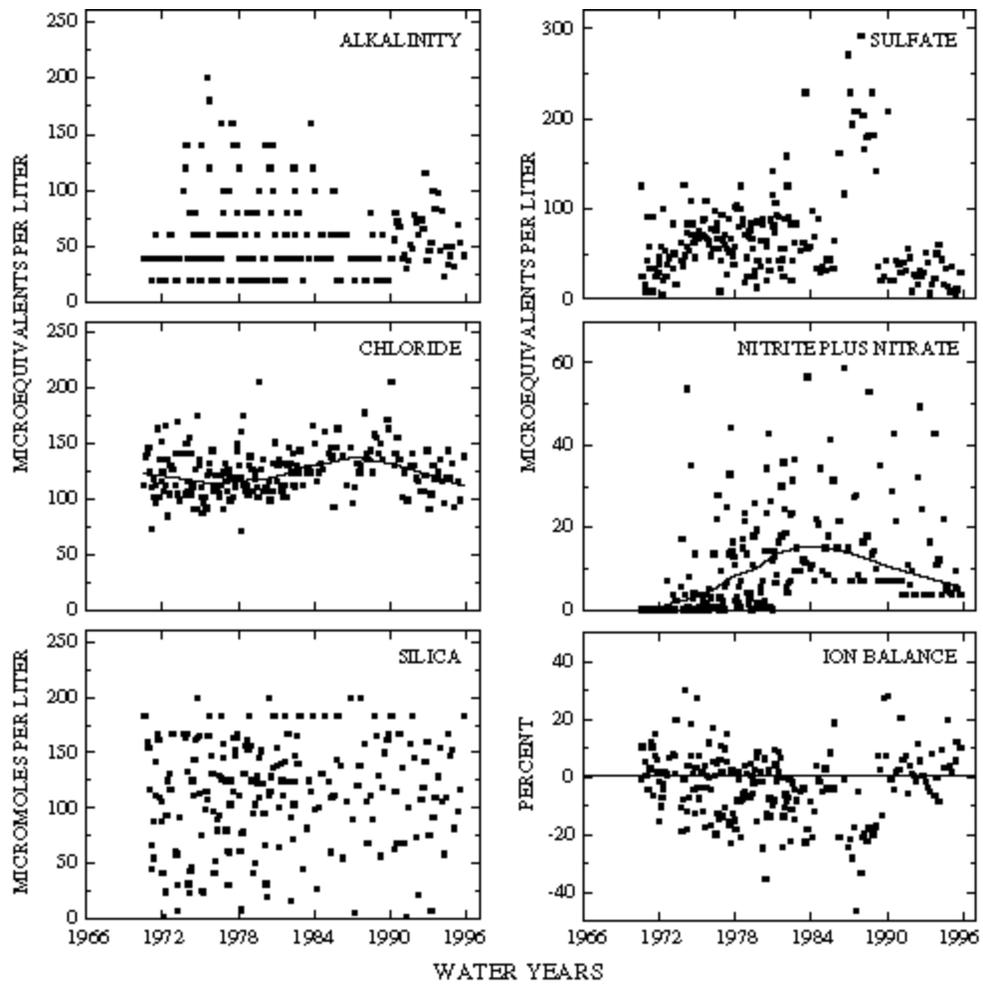


Figure 25b. Temporal variation of discharge, field pH, major ion concentrations, and ion balance at Scape Ore Swamp, South Carolina - Continued

station in Maine during the same period of record. Because samples from these two sites were analyzed at different laboratories during this period, it seems likely that high calcium concentrations may be related to collection or processing procedures common to both stations. The only documented procedural change during this period was the introduction of a polypropylene churn splitter for compositing and splitting samples that was required for use at all HBN stations beginning in water year 1977 (Office of Water Quality Technical Memorandum No. 76.24–T, Sample Splitter for Water-Sediment Samples, issued August 16, 1976, at URL <http://water.usgs.gov/public/admin/memo/>). Although problems with previous sample splitters were not documented by the Office of Water Quality (OWQ), the fact that the anomalous pattern disappeared around the beginning of 1977 indicates that previous splitters or compositing devices may have been a potential source of contamination. Temporal patterns in pH also reveal potential bias in the field determinations. Between 1983 and 1985, field pH values were 0.5 to 1 pH unit smaller than values typically reported for this station. A possible cause of this shift was the gel-filled pH probe used by field personnel in the USGS office in South Carolina during this period of record (John Barton, USGS, oral commun., 1996). Some instrument-electrode systems are known to give erroneous readings when measuring pH in low-conductivity waters, and the electrode commonly is the critical component (Office of Water Quality Technical Memorandum No. 81.08, Electrodes for pH Measurement in Low-Conductivity Waters, issued February 10, 1981, at URL <http://water.usgs.gov/public/admin/memo/>).

Median concentrations and ranges of major constituents in stream water collected at the HBN gaging station and VWM concentrations in wet-only deposition measured at the Santee National Wildlife Refuge NADP station about 60 km south of the basin are presented in table 46. Precipitation chemistry at the NADP station was dilute and acidic with a VWM pH of 4.6 for 10 years of record. The dominant cations were hydrogen, which contributed 45 percent of the cation charge, and sodium, which contributed 18 percent. Sulfate accounted for 54 percent of the total anions, and nitrate and chloride each accounted for 23 percent. These results suggest that precipitation at the NADP station probably is a mixture of both anthropogenic emissions of sulfate and nitrogen compounds and salt derived from marine aerosols.

Table 46. Minimum, first quartile, median, third quartile, and maximum values of physical properties and major ion concentrations measured in water-quality samples from Scape Ore Swamp, South Carolina, August 1970 through September 1995, and volume-weighted mean concentrations in wet precipitation collected at Santee National Wildlife Refuge, South Carolina

[Concentrations in units of microequivalents per liter, discharge in cubic meters per second, specific conductance in microsiemens per centimeter at 25 degrees Celsius, pH in standard units, and silica in micromoles per liter; n, number of stream samples; VWM, volume-weighted mean; inst., instantaneous; spec. cond., specific conductance; <, less than; --, not reported]

Parameter	Stream Water						Precipitation VMA ^a
	Minimum	First quartile	Median	Third quartile	Maximum	n	
Discharge, inst.	0.18	0.85	2.0	4.0	19	212	--
Spec. cond., field	20	25	26	29	43	209	--
pH, field	4.0	5.1	5.4	5.7	6.5	211	4.6 ^b
Calcium	5.0	36	43	50	220	211	4.5
Magnesium	<8.2	41	46	55	140	211	3.0
Sodium	13	110	130	150	270	211	10
Potassium	<2.6	10	13	17	56	211	2.3
Ammonium	<.7	1.4	2.1	3.6	13	96	5.6
Alkalinity, laboratory	<20	37	40	71	200	211	--
Sulfate	4.2	33	56	83	290	210	29
Chloride	69	110	120	140	200	210	12
Nitrite plus nitrate	<.7	4.3	8.6	18	59	181	12 ^c
Silica	1.7	80	120	160	220	211	--

^aValues are volume-weighted mean concentrations for 1984—94.

^bLaboratory pH.

^cNitrate only.

Stream water in the Scape Ore Swamp is dilute and weakly buffered; total ionic constituents ranged from 210 to 860 meq/L, and alkalinities were generally below 70 meq/L. The major cation in stream water was sodium, and the dominant anion was chloride. The low concentrations of the major weathering products, particularly alkalinity, are attributed to the slow weathering rates of the quartz-rich soils and sediments in the basin. The median chloride concentration in stream water was 120 meq/L, which was about 10 times larger than the VWM concentration of chloride in precipitation. Based on the difference between annual precipitation and runoff, evapotranspiration can account for no more than a threefold increase in the chloride concentration of precipitation, indicating that a substantial amount of stream-water chloride is derived from sources other than precipitation. Considering the proximity of this site to the coast, dry deposition of marine aerosols probably is the dominant source of chloride (and sodium) in stream water. Chloride also may be derived from manmade sources, such as runoff from agricultural areas or discharge from domestic wastewater systems. The median concentration of sulfate in stream water was twice the VWM concentration in precipitation, which was somewhat less than expected considering the effects of evapotranspiration and additional inputs of sulfate in dry deposition. This difference indicates that a substantial portion of atmospherically deposited sulfate is retained in the basin. Possible mechanisms of sulfate retention include adsorption on basin soils or sulfate reduction in wetland areas along the stream channel. The basin also is apparently an important sink of atmospherically deposited nitrogen based on the smaller concentrations of both nitrate and ammonium in stream water compared to precipitation. The low stream-water nitrate concentrations also indicate that agricultural areas in the basin do not significantly affect stream-water nutrient concentrations at the gage. Because weathering supplies only minor amounts of base cations and alkalinity to surface waters in the basin, basin retention of sulfate and nitrate probably is the dominant process buffering surface waters from acidic deposition (Elwood and others, 1991).

The solute composition of stream water was further evaluated by analyzing correlations between solutes and stream discharge (table 47). Discharge was poorly correlated with most dissolved constituents with the exception of weak inverse correlations with pH ($\rho = -0.588$), sodium ($\rho = -0.0462$), and alkalinity ($\rho = -0.536$). Solute concentrations in stream water also were poorly correlated with each other with the exception of a weak positive correlation between calcium and magnesium ($\rho = 0.551$). The lack of strong correlations among these constituents probably is related both to hydrologic and geochemical factors. The highly permeable soils and sediments on the hillslopes and upland surfaces allow precipitation to infiltrate quickly into the ground-water system. In turn, streamflow is supplied by ground-water discharge, which maintains a relatively constant flow rate and stream composition throughout the year. Additionally, the quartz-rich soils and sediments contain few weatherable minerals and, as a consequence, produce soil and ground waters that are not significantly different in composition from that of the incoming precipitation.

Table 47. Spearman rank correlation coefficients (rho values) showing the relation among discharge, pH, and major ion concentrations, Scape Ore Swamp, South Carolina, 1980 through 1995

[Q, discharge; Ca, calcium; Mg, magnesium; Na, sodium; K, potassium; Alk, alkalinity; SO₄, sulfate; Cl, chloride; Si, silica]

	Q	pH	Ca	Mg	Na	K	Alk	SO ₄	Cl
pH	-0.588								
Ca	.149	-0.235							
Mg	.047	-.083	0.551						
Na	-.462	.165	.294	0.294					
K	.311	-.193	.350	.409	-0.052				
Alk	-.536	.504	-.089	.043	.159	-0.157			
SO ₄	.243	-.357	.102	.204	.199	.044	-0.321		
Cl	.114	-.28	.198	.390	.264	.259	-.203	0.171	
Si	-.251	-.005	.156	.291	.301	.071	.014	-.003	0.426

Results of the seasonal Kendall test for trends in discharge and major dissolved constituents from 1970 through 1995 are shown in table 48. Statistically significant upward trends were observed for unadjusted magnesium, potassium, and chloride concentrations at the 0.01 confidence level. The flow-adjusted concentrations of magnesium were not tested for trends because the flow model was not statistically significant at a = 0.1. The unadjusted and flow-adjusted trends were similar for potassium and chloride, indicating that the trends were not driven by variations in stream discharge. The LOWESS curve in figure 25 shows a pattern of increasing chloride concentrations between 1972 and 1986 and decreasing concentrations after 1986. A similar pattern also was observed for sodium, although a trend was not detected during the entire period of record. Land-use changes may be a likely mechanism for producing the similar patterns in chloride and sodium because both ions are commonly indicators of manmade effects. The most substantial land-use change in the basin was the conversion of about 20 percent of cropland to permanent vegetation as a result of the Conservation Reserve Program (CRP). Interestingly, chloride and sodium concentrations began to decrease shortly after the CRP was instituted in 1985. In contrast to chloride, potassium concentrations declined through the early part of the record, then increased somewhat abruptly in 1986. Most of the increase in magnesium concentrations occurred in the early 1980's. It is unlikely that the sudden increase in potassium is associated with the CRP. A decrease in agricultural land use is more likely to cause a decline in stream-water potassium concentrations,

particularly since potassium is a common component of fertilizer. It is perhaps more likely that the sudden increase in potassium was caused by a analytical-related factor. The increase in potassium, for example, occurred at the same time that the analytical determinations were switched from the Central Laboratory in Atlanta to the NWQL in Arvada, Colo. A method change also may provide a plausible explanation for the upward trend in magnesium. For example, the analytical method for the determination of several constituents, including magnesium, was changed from AA spectroscopy to ICP spectroscopy in 1983 (Office of Water Quality Technical Memorandum No. 82.18, National Water- Quality Networks, issued September 28, 1982, at URL <http://water.usgs.gov/public/admin/memo/>), which correlates with the increase in magnesium concentrations in the early 1980's.

Table 48. Results of the seasonal Kendall test for trends in discharge and unadjusted and flow-adjusted pH and major ion concentrations, Scape Ore Swamp, South Carolina, August 1970 through September 1995

[Trends in units of microequivalents per liter per year, except discharge in cubic meters per second per year, pH in standard units per year, and silica in micromoles per liter per year; <, less than; --, not calculated]

Parameter	Unadjusted		Flow adjusted	
	Trend	p-value	Trend	p-value
Discharge	-0.01	0.554	--	--
pH, field	<.01	.849	<-0.01	0.906
Calcium	.1	.214	(^a)	--
Magnesium	.7	.000	(^a)	--
Sodium	<.1	.914	-.2	.509
Potassium	.1	.008	.2	.007
Alkalinity, laboratory	<.1	.318	.2	.485
Sulfate	-.7	.132	-.9	.093
Chloride	.8	.002	.9	.001
Nitrite plus nitrate	<.1 ^b	.601	--	--
Silica	.6	.103	.5	.093

^a Flow model not significant at $\alpha = 0.10$.

^b Trend calculated for 1973—95, using a trend test for censored data.

Synoptic Water-Quality Data

Results of the surface-water synoptic sampling of January 10 and 11, 1991, are presented in table 49; locations of sampling sites are shown in figure 24. Discharge at the gage was 2.27 m³/s compared to the median daily discharge of 3.68 m³/s for January (Lawrence, 1987), indicating that the basin was sampled during base-flow conditions for that time of year. Solute concentrations measured at the gage (site 1) during the synoptic sampling were generally below the first-quartile concentrations reported for the gage during the entire period of record (table 46). The tributary streams were extremely dilute, and sodium and chloride were the dominant ions. Measured alkalinities were below the detection limit at 9 of the 12 sites, and pH ranged from 4.92 to 6.54, indicating that most surface waters were slightly acidic. The source of most stream-water acidity probably is organic acids, although some may be derived from strong acids in precipitation. Although DOC was not measured in this study, all samples had positive ion balances (range 3.8 to 14 percent), inferring that organic anions were an important component of stream water during the sampling period.

In general, stream chemistry did not vary markedly among the sampling sites. For example, specific conductance ranged from 18 to 34 µS/cm, and most dissolved constituents did not vary by more than a factor of three. This result primarily reflects the uniform composition and slow weathering rate of soils and sediments in the basin. Some subtle variations in stream chemistry are evident, however, which may reflect the influence of factors other than geology. For example, samples that were collected in the Beaverdam Creek subbasin (sites 2, 7, and 8) had elevated concentrations of calcium, alkalinity, and nitrate. These solutes are common components of fertilizer and sewage, and their elevated concentrations may be indicative of human disturbance in this subbasin. Substantial differences also were observed in Beaverdam Creek upstream from (site 8) and downstream from (site 7) McLeods Pond. For example, alkalinity concentrations almost tripled downstream from the pond, whereas nitrate concentrations decreased by a factor of 10. This change in stream chemistry indicates that the impoundment may have a strong influence on stream-water chemistry in this tributary. Chloride, which is commonly used as an indicator of human activities, was remarkably uniform over the basin and ranged from 110 to 140 meq/L. In contrast, sodium concentrations varied by more than a factor of two. Interestingly, streams in the northern one-half of the basin (sites 4, 5, 9, 10, 11, and 12) had an average sodium-to-chloride ratio of 0.6, whereas sites in the southern one-half of the basin (sites 2, 3, 6, 7, and 8) had an average ratio of 1.0. Sulfate also differed slightly between tributaries that drain the upper and lower parts of the basin. For example, sulfate averaged 40 meq/L in the northern tributaries compared to 20 meq/L in the southern tributaries. These differences in stream chemistry may be related, in part, to land-use patterns in the basin. Subbasins in the northern one-half of the basin are primarily forested, whereas those in the southern one-half have a greater percentage of agricultural land.

Table 49. Physical properties and major ion concentrations from surface-water sampling sites in the Scape Ore Swamp Basin, South Carolina, January 9—10, 1991

[Site locations shown in fig. 24; Q, discharge in cubic meters per second; SC, specific conductance in microsiemens per centimeter at 25 degrees Celsius; pH in standard units; Ca, calcium; Mg, magnesium; Na, sodium; K, potassium; Alk, alkalinity; SO₄, sulfate; Cl, chloride; NO₃, nitrate; concentrations in microequivalents; Si, silica in micromoles per liter; <, less than, --, not measured]

Site	Station number	Q	SC	pH	Ca	Mg	Na	K	Alk	SO ₄	Cl	NO ₃	Si	Criteria ^a	Remarks
1	02135300	2.27	26	4.98	32	40	96	15	<10	31	130	6.4	63	--	--
2	340910080184900	.62	27	5.88	35	42	140	19	39	17	140	19	87	MT, LU	Significant development
3	341230080195100	.042	22	5.45	19	30	120	13	<10	21	120	2.9	120	BG	Black Mingo Shale
4	341321080210300	.093	18	5.81	22	34	78	9.7	<10	23	110	2.9	90	MT, LU	Reservoir in subbasin
5	341335080214900	--	25	4.90	33	45	83	15	<10	38	130	2.9	87	MT, LU	Upstream from most development
6	341322080221300	--	23	5.28	21	30	110	14	<10	20	120	1.4	130	LU	I—20 crosses subbasin
7	341141080241800	.062	29	6.54	60	55	130	28	110	15	130	2.9	38	LU	Downstream from large impoundment
8	341232080253400	.006	34	5.89	60	67	160	13	43	25	140	30	92	LU	Upstream from large impoundment
9	341354080223200	.15	25	5.14	31	40	96	18	<10	38	140	5.0	130	LU	No I—20 influence
10	341528080231900	.71	26	4.92	40	47	87	18	<10	42	130	2.1	110	MT, LU	Forested, little development
11	341539080263100	.13	27	5.13	45	48	74	26	<10	58	120	4.3	100	MT, LU	Forested, little development
12	341812080221000	.27	25	4.99	36	54	65	21	<10	40	120	7.9	67	LU	Primarily forested

^aCriteria used in selection of sampling sites: BG = bedrock geology, MT = major tributary, LU = land use.

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- Lawrence, C.L., 1987, Streamflow characteristics at hydrologic bench-mark stations: U.S. Geological Survey Circular 941, 123 p.
- Mitchell, Cleveland J., Jr., 1990, Soil survey of Kershaw County area, South Carolina: U.S. Department of Agriculture Soil Conservation Service, 165 p.
- U.S. Geological Survey, published annually, Water resources data, South Carolina, Columbia: U.S. Geological Survey Water Data Report.

Appendix A. List of Map Resources

a. U.S. Geological Survey topographic maps:

- Bishopville West, South Carolina (1: 24,000), gaging station on this quadrangle
- Cassatt, South Carolina (1: 24,000)
- Lucknow, South Carolina (1: 24,000)
- Spring Hill, South Carolina (1: 24,000)
- Camden, South Carolina (1: 100,000)

b. Geologic maps:

- Cooke, Wythe B., 1936, Geology of the Coastal Plain of South Carolina: U.S. Geological Survey Bulletin 867, 196 p., scale 1: 500,000.

c. Soil surveys:

- Mitchell, Cleveland J., Jr., 1990, Soil survey of Kershaw County area, South Carolina: U.S. Department of Agriculture Soil Conservation Service, 165 p., 65 maps.
- U.S. Department of Agriculture, 1963, Soil survey of Lee County, South Carolina: U.S. Department of Agriculture Soil Conservation Service.

d. Other maps:

- U.S. Department of Agriculture, 1991, Important farmlands, Lee County, South Carolina: Columbia, S.C., U.S. Department of Agriculture Soil Conservation Service in cooperation with South Carolina Land Resources Conservation, scale 1: 100,000.

Appendix B - NWIS Site-Identification Numbers

Table B-1. NWIS site identification numbers and site names for water-quality sampling sites.

Site	Identification Number	Site Name
1	02135300	SCAPE ORE SWAMP NEAR BISHOPVILLE, SC
2	340910080184900	BEAVERDAM CREEK AT MOUTH
3	341230080195100	GUM SPRINGS BRANCH
4	341321080210300	CEDAR CREEK
5	341335080214900	SCAPE ORE SWAMP ABOVE I-20
6	341322080221300	I-20 BRANCH
7	341141080241800	BEAVERDAM CREEK AT MCLEODS POND
8	341232080253400	UPPER BEAVERDAM CREEK
9	341354080223200	SOUTH MCGEE BRANCH
10	341528080231900	TIMBER CREEK
11	341539080263100	NANCY BROOK
12	341812080221000	BLACK CREEK AT COUNTY LINE

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