# CHEMICAL CONSTITUENTS IN SNOW FROM SIERRA NEVADA, CALIFORNIA AND WASATCH MOUNTAINS, UTAH

By

S. M. Rogers and J. H. Feth 1/

#### INTRODUCTION

A study of inorganic chemical constituents determinable in snow has been made by the Water Resources Division, U. S. Geological Survey. Ninety-eight snow samples have been analyzed at the Geological Survey's Pacific Coast field center, Menlo Park, California, as part of a larger investigation of the geochemistry of selected segments of the hydrologic cycle. Attention has been centered on evaluation of chloride concentration, but where the volume of the sample allowed, a more detailed study of the chemistry of the snow has been made. Results of the analyses to March 1, 1959, are shown in table 1.

Chemical analyses were made by S. M. Rogers, C. E. Roberson, J. P. Schuch, and H. C. Whitehead, chemists, Quality of Water Branch.

#### SAMPLING PROCEDURES

In early stages of the work the samples for chemical analysis were taken by forcing snow into a 1-liter serum bottle where it was permitted to melt. Later, the samples were collected in chemically clean polyethylene buckets, covered with a plastic-sheet protector, and permitted to melt before being decanted into glass for storage until analysis. The samples from the Wasatch Mountains, Utah, were collected by personnel of the U. S. Soil Conservation Service during snow-course measurements, the snow cores being transferred directly from the coring tube to widemouth bottles which were then tightly covered with plastic-lined screw-on lids.

#### ANALYTICAL PROCEDURES

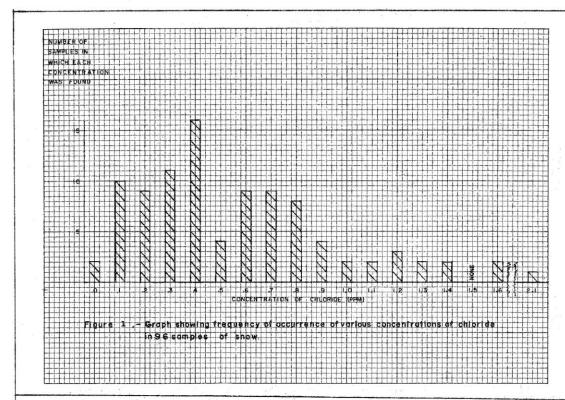
Water resulting from melting of the snow samples was analyzed by prescribed methods— used in laboratories of the Quality of Water Branch. Chloride was determined in later stages of the study, however, by a colorimetric method (Bergman and Sanik, 1957) in which the color is developed by displacement of the thiocyanate ion from mercuric thiocyanate by the chloride ion in the presence of the ferric ion. The color is stable and proportional to the original chloride—ion concentration,

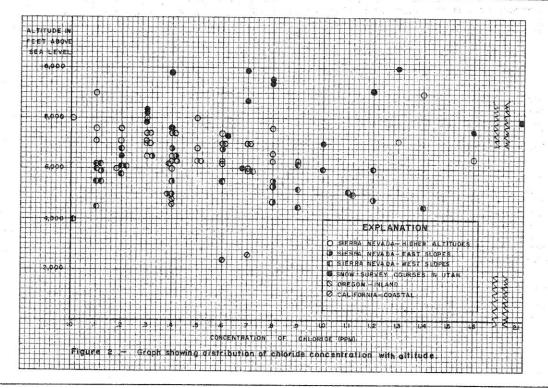
## INFERENCES DERIVED FROM THE DATA

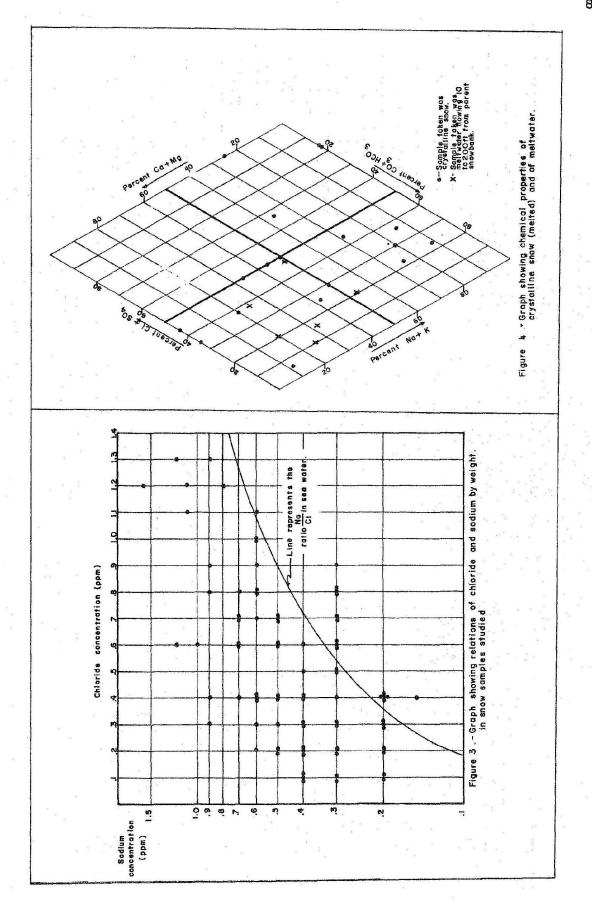
Figure 1 shows the distribution of chloride concentrations found in 96 samples of snow. The concentration ranges from below the threshold of detection to 2.1 ppm (parts per million). The mean value calculated was 0.6 ppm. Two additional analyses (table 1, Nos. 671, 960) show chloride concentrations of 3.7 and 5.0 ppm. The sample containing 3.7 ppm of chloride was collected adjacent to a road in Oregon giving access to a lodge, and may have been contaminated. The sample having 5.0 ppm of chloride was from the Provo River drainage basin in Utah. The origin of the high chloride value is uncertain.

The general range of chloride values found, and unpublished analytical data in the files of the U. S. Geological Survey at Menlo Park representing the chemical quality of lakes and streams in the Sierra Nevada, suggest that a mean value for the chloride concentration of snowfall in those mountains is about 0.5 ppm. Data from the Wasatch Mountains, Utah, are less complete. All samples analyzed were taken in the first week of January 1959 and may not represent the average quality of snows in the area. The tentative conclusions drawn from the samples analyzed, however, is that the range of chloride content of snow is slightly higher in Utah than along the Sierra Nevada, perhaps because moderate amounts of sodium chloride dust are blown aloft from the Great Salt Lake Desert and become incorporated in the snow crystals as they form.

<sup>1/</sup> Chemist and Geologist, respectively, U. S. Geological Survey, Menlo Park, California, 2/. U. S. Geol. Survey Water-Supply Paper 1454, manuscript in preparation.







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717	4.2 mi. west of Peddlers Hill, Galif.	0.	z.	-2	E.	0	E	1.8	.2	0	-0f	0.	.10	00.	2.0	3	5.6	9
716	4 mi. east of Camp Connell, Calif., Galif. 88	0.	۶.	.1	г,	0.	3	1.6	۲.	0.	00.	9.	0.	.03	2.0	77	5.6	9
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721	Shore of Mirror Lake, Yosemite National Park, Calif.	т.	0.	6:	ω,	1.9	9	4.	6.	r!	01.	0.	0.	70.	1.0	01	6.0	
742	Spooner Grade, 9.5 mi. east of Junct. U.S. 50 and U.S. 395	0.	-2	6.	7.	.1	η	9.	٠.		.01	,	.1	00.	1.0	7	5.7	-
71.7	Donner Grade, bottom U.S. 40	7.	0.	٠.	٠.	0	7	£.	7.	.1	01.	. 0	0.	00.	1.0	7	5.9	. 6
149	1 mi. west of Soda Springs, Calif., U.S. 40	4.	o.	.3	۴.	1.0	7	14	ω.	0'	80.	Q.	0.	.02	1.0	9	8.3	
752	Echo Summit, Calif., U.S. 50	1	1	4.	3		8	.2	٤٠	,		,		1	7.	η	0.9	
753	Echo Lake (Needle Point), Calif.	1	1	•2	۲,	1	1	.3	2,	,	,	ı	,		1.5	:3	5.8	8
						1000												

a/ Dash indicates determination not made.

 $\underline{b}/$  Hardness as  ${\rm CaCD}_3$  .  $\underline{c}/$  Specific conductance (micromios at  $25^{\circ}{\rm C})$ 

Table 1.- Mineral constituents and related physical measurements of snow samples after meathing - Constitued.

[Analytical results in warts ner million except as indicated.]

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758	Echo Summit Lodge, Calif.	,	,	6.	η.	-1:	5	-77	80.			,		,	1,0	7	6.5
759	Summit of Highway, Mt. Ross, Nev.		,	.1	.2	,	2	.2	۲.	,	1				1.0	3	5.7
260	2.7 mi. west of Mt. Ros	,	1	cu.	₹	1	CI.	ŗ	.1	ı	1	1	1	1	4.	3	5.7
762	Norden Lake, Calif., U.S. 40	,	,	-₹.	۲.	,	2	-5	ν.	,	1	,	,	,	1.0	ή	0.9
763	Sugar Howl Skd area, Calif., U.S. MO	,		2.	1.	,	2	۲.	e.	١,	٠,	,	,	,	эċ.	2	5.9
764	Squaw Valley, Calif., just south of ski lodge	١.		1,4	۲.	,	6	۲.	1.6	,	۲	,	,		1.0	30	6.3
765	Rainbow Tavern, Calif., U.S. 40	1	1	r.	7.	1	3.		.7	,		,	ı	1	1.5	80	5.9
166	Mt. Rose Highway Summit, Nev.					1	#	-	1.4	١.		,	,	1		11	5.9
767	Echo Lake, Calif.		1	2.	۲.		7.	9.	-1	,	,		,	,	1.0	ъ	5.3
927	\$ mi. east of Kyburz, Calif., U.S. 50	٥.	5.	1.1	7.	0	н	٠-	1,1	o.	٥.	0.	o.	• 05	2,0	Ħ	5.5
928	h m. west of Twin Bridges, Calif., U.S. 50	0.	14	9	-7:	0	3		7	o.	0.	0.	0.	.05	2.0	7	5.6
930	Echo Surmit, Dalif., U.S. 50	7.	0.	1.2	ŗċ	0.	0	2.8	ė.	.05	1.	o.	٠.	,00,	1.0		5.7
932	Spooner Snumit, Nev., U.S. 50	o.	4	9.	.5	٥.	8	1.6	۲.	%	0,	٥.	۲.	6.	1.0	-2	5.6
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953	Above Hetch Hetchy Reservoir			۲.					9.								
956	Duchesne River area, Indian Canyon, Utah			1,1					1.2							771	
156	Weber River area, Smith and Moorehouse reservoir, Utah			۶.					4.			-				6	
958	Duchesne River area, Trial Lake, Utah			.v.					4.							4	
959	Wasatch Mts., main range, Parley's Canyon Summit, Utah			6.					1.6						-	2,1	
960	Provo River area, Dardels-Strawberry Summit, Utah	-		4.					5.0					-		27	
825	Weber River area, Beaver Creek R.S., Utah			4.				-	9.							1.2	
61.6	Lower Bear River area, Garden City Summit, Utan	Ę		e.					2,1								
980	Virgin River area, Midway Valley, Utah			e,			- 3		89								-
100	963 Fremont River area, Farnsworth Lake, Utah			9.					.7							1.8	

984 Great Salt Lake ares, Lower Farmington Canyon, Utah	9•		1.0	18	
985 Recalante Aiver area, Widtsoe-Escalante Summit, Utah	9*		<b>బ</b>		
986 Duchesne River area, Julius Park, Utah	1.2		1.3		
983 Provo River area, Timpanogas Divide, Utah	ন্		.3		
993 Ogden River area, Lower Ben Lomond, Utah	2.		.7		
994 Ogden River area, Snow Basin, Utah	zů		2.		
1012 Price River area, Gooseberry Reservoir, Utah	7.	عدرين وراسية والمراجعة	.7		
1013 3.1 mi. west of Kybura, Calif., U.S. 50	7.		F.		
1014 0.6 mi. east of Twin Bridges, Galif., U.S. 50	τ.		Ţ.		
1015 Echo Summit, Galif., U.S. 50	e C		£.		
1016 Spooner Burnit, Nev., U.S. 50	7.				
1017 Geiger Mountain Summit, Nov., Nov. 17	m,		9.		
1018 Squew Valley, Calif., at Ski Lift	u,		7.		
JOJ9 Sugar Bowl Ski area, Calif., U.S. 40	.2		.3		
1020 Rainbow Tavern, Calif., U.S. 40	.2	دوم سندها والزياسة سرعان ويوزياسيدان سيدان مدينة ويوانيان والمستدان والمستدان والمتعاربة والمتعاربة	7.		
1022 Price Hiver area, Dry Valley Divide, Utah	9.	rini e in de la companya de la comp	£.		
1023 Price River area, Mud Creek, No. 2, Utah	χį		.3		
1025 Spooner Summit, Nev., U.S. 50	6.		1.3		
1026 10 mi. south of Gardnerville, Nev., U.S. 395	L*		æ.		
1027 3/4 mi. north of Silver City, Wev.	-2		<b>†</b>		78.0
1028 3/1 mi. north of Silver City, Nev.	2.		7.		
1029 0.1 mi. north of Virginia City, Nev.	<b>α</b> .		村•		
1030 1.9 mi. north of Virginia City, Nev.	F.		્ય•	AND THE REAL PROPERTY OF THE P	
1031 1 mi. north of Geiger Mountain Summit, Nev.	-2		.3		
1035 Saratoga, Calif., Junct. Hwy 9 and 5 (Skyline)	ē.				
1036 Santa Onus, Calif., Wunct. Jamison Creek Road and Empire Grado	ε,		9.		

An attempt (fig. 2) to relate chloride content to altitude yielded inconclusive results. The samples from the east slopes of the Sierra Nevada, including those in closely adjacent parts of Nevada, show a general tendency toward increasing chloride concentration with decreasing altitude. The west-slope samples do not show any apparent trend. The scattering of data points for Utah shows an opposing trend to that of the east slope of the Sierra Nevada, but the restricted number and collection period of those samples limit the reliability of the apparent trend.

Chloride was determined (table 1) in all 98 samples. Thereafter the extent of the analytical work done was determined principally by the volume of sample remaining. Reasonably detailed analyses were made of about 60 samples.

Results of the analyses show that many of the snows contained a variety of dissolved matter detectable by available methods. Specific conductance, measured in micromhos at 2500, ranged from 3 to 44. Only a few snow samples exceeded 10 micromhos, and 24 of 63 determinations were in the range of 3 to 5 micromhos. This group of 24 approaches common distilled water in chemical purity.

The bicarbonate and sulfate commonly occurred in concentrations of less than 5 ppm. A few tenths of a part per million of fluoride was found in some samples, and nitrate was found in 15 of the 46 samples analyzed for that constituent. Boron and, a little less often, bromide and iodide are reported in concentrations of a few hundredths of a part per million from a large number of the snows tested.

The concentration of sodium is plotted in relation to chloride concentrations in figure 3, where the scatter of points shows that characteristically the reported sodium content of the snow is higher in relation to chloride than it is in sea water. The evidence available suggests that, during transport of airborne oceanic salts, the sodium and chloride do not occur in molecular proportions. This phenomenon has been noticed by earlier investigators (Emanuelsson, Eriksson, and Egner, 1954, p. 265; Rossby, 1955). Results of the present study tend to confirm the occurrence, although they do not help to explain the mechanism.

Chemical changes occur in nature during melting of the snow and during even the very early stages of runoff, according to data shown in figure 4. The diagram is after Piper (1945). Each point shown represents a sample of snow (dots) or of snowmelt runoff (X's) taken not more than 200 feet from the parent snowbank. The position of each mark on the diagram is in effect a summary statement of the principal chemical characteristics of the water represented. Thus samples plotting in the right-hand quarter of the diagram contain more than 50 percent chloride plus sulfate (total anions = 100 percent) and more than 50 percent sodium plus potassium (total cations = 100 percent), all expressed as chemical equivalents. They are commonly sodium chloride waters. Samples plotting in the lower part of the diagram are characterized by high sodium and bicarbonate content. The left-hand quarter of the diagram illustrates waters in which calcium and magnesium together make up most of the cations, and carbonate and bicarbonate make up most of the anions. In snow waters the "carbonate and bicarbonate" probably is wholly bicarbonate, as none of the pH values of these waters exceeds 8.2, the threshold value for the occurrence of carbonate.

Figure 4 is significant in showing that the chemical character of snows studied varies rather widely, although, like most natural waters, no snow sample analyzed falls in the top (calcium-magnesium-chloride-sulfate) part of the diagram. But of the 6 melt waters sampled near their parent snowbanks, 4 plot near the center of the calcium bicarbonate quarter of the diagram, and two are closely adjacent to the dividing line. The tendency shown probably reflects an actual relationship prevailing in nature. During melting, it is probable that carbon dioxide (CO2) is dissolved in the water droplets as they form, in quantities sufficient to overbalance the chloride present. In the earliest stages of runoff—almost immediately upon contact with soil or weathered rock—the water acquires enough calcium and magnesium to cause those ions to become dominant on the cation side of the chemical diagram (fig. 4).

#### CONCLUSIONS

The chloride content of 98 samples of snow, principally from the Sierra Nevada and from the Wasatch Mountains, has been determined. Some 60 melted snow samples were analyzed for other constituents as well. Preliminary results suggest that chloride ranges in concentration from below the limit of detection to 5 ppm but is seldom found in concentrations greater than 2 ppm. Snow

samples from Utah tend to contain slightly more chloride than those from the Sierra Nevada, but samples are not numerous enough to provide an assured relationship. The data suggest that the chloride content of snow decreases with altitude on the east slope of the Sierra Nevada, but other areas do not show recognizable trends.

Study of a few samples of melt water in relation to their nearby parent snowbanks suggests that the process of melting and the first few tends of feet of surface flow cause measurable chemical changes, calcium and bicarbonate ions percentages showing increases over percentages of the same ions in the snow. When concentrations are as low as found in these analyses, however, the percentage changes may not be as noteworthy as they appear. Analytical error and sampling procedures can account for part of the values reported.

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