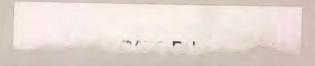
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CHANGES IN QUALITY OF GROUND WATER IN THE LINCOLN AREA, MONTANA, 1974-79 by Robert B. Leonard, Wayne A. Wood, and Arnold J. Boettcher

Open-File Report 80-1108

Prepared in cooperation with the

Lewis and Clark County Commissioners

Helena, Montana November 1980

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METRIC CONVERSION TABLE

The following factors can be used to convert inch-pound units to the International System (SI) of metric units.

Multiply inch-pound unit	By	To obtain SI unit
acre	0.4047	square hectometer
foot (ft)	0.3048	meter
mile	1.609	kilometer
acre-foot	1233	cubic meter
ton (short)	0.9072	megagram

temperature, degrees Celsius (°C) = 0.556 (°F-32)

CHANGES IN QUALITY OF GROUND WATER IN

THE LINCOLN AREA, MONTANA, 1974-79

By

Robert B. Leonard, Wayne A. Wood, and Arnold J. Boettcher

ABSTRACT

Water samples from representative shallow wells in the vicinity of Lincoln, Montana, were collected during March and June 1979 for analysis to evaluate changes in quality since 1974-75 and possible contamination by domestic or agricultural waste. All samples were suitable for drinking water with respect to the constituents tested. Tests for fecal coliform bacteria were negative. The concentrations of dissolved solids, composed mainly of calcium, magnesium, bicarbonate, and sulfate, were less than 300 milligrams per liter. Concentrations of sodium, potassium, and chloride were 3.5 milligrams per liter or less.

From March to June, the maximum concentration of nitrite plus nitrate as nitrogen decreased from 0.57 to 0.29 milligram per liter and the mean concentrations decreased from 0.16 to 0.11 milligram per liter, probably as a result of dilution of ground water by snowmelt. The concentrations in samples from the same wells generally were less during 1979 than during 1974-75. Isolation of sewage effluent in shallow zones by hardpan, consumption of nutrients by abundant phreatophytes, filtration of bacteria by the soil or aquifer, and dilution by underflow and recharge evidently combine to maintain the quality of the well waters within acceptable limits.

INTRODUCTION

Domestic and public ground-water supplies in the vicinity of Lincoln, Mont. (fig. 1), are obtained chiefly from wells completed in permeable alluvium in which the water table is normally at a depth of less than 8 feet below the land surface (Coffin and Wilke, 1971). Most wells are less than 40 feet deep (Wilke, 1976, sheet 3) and are located near the buildings they serve.

Septic tanks are the principal means of domestic sanitary waste disposal. The drainfields discharge into the alluvium only a few feet above the water table; therefore, contamination of adjacent water supplies with domestic waste effluent is a possibility.

Analyses of water samples collected from selected wells during 1974-75 (Wilke, 1976) revealed that concentrations of common chemical constituents were significantly less than the maximum concentrations recommended by the U.S. Environmental Protection Agency (1977) for drinking water. Whether the analyses

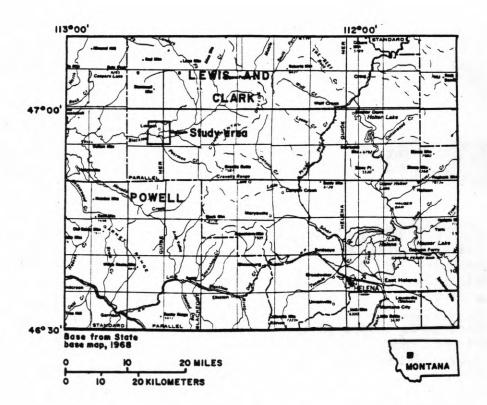


Figure 1.--Location of study area.

represented natural ground water or ground water slightly affected by septic tank waste is unknown.

The presence of sodium, chloride, potassium, and the nutrients nitrogen and phosphorus in concentrations exceeding normal background concentrations in natural ground water would constitute evidence for contamination with human or agricultural waste. Disproportionate increases of the concentrations of those constituents with time and with respect to other constituents, and the presence of fecal coliform bacteria, would substantiate the evidence.

The purpose of this report is to document the changes in water quality from 1974-75 to 1979 in representative shallow wells and to evaluate the extent of contamination, if any, by domestic or agricultural waste. Water samples were collected during 1979 at many of the same sites sampled during 1974-75.

The writers gratefully acknowledge the cooperation of landowners in the Lincoln area who permitted access for the collection of water samples and measurement of water levels. The investigation was funded in part by the Lewis and Clark County Commissioners.

WELL-NUMBERING SYSTEM

In this report, locations are numbered according to geographic position within the rectangular grid system used by the U.S. Bureau of Land Management (fig. 2). The location number consists of 13 characters. The first three characters specify the township and its position north (N) of the Montana Base Line. The next three characters specify the range and its position west (W) of the Montana Principal Meridian. The next two characters are the section number. The next three characters designate the quarter section (160-acre tract), quarter-quarter section (40-acre tract), and quarter-quarter-quarter section (10-acre tract) in which the well is located. The subdivisions of the section are designated A, B, C, and D in a counterclockwise direction, beginning in the northeast quadrant. The last two characters form a sequential number within a 10-acre tract, assigned in order of well inventory. For example, as shown on figure 2, well 14N09W24BACO1 was the first well inventoried in the SW1/4 NE1/4 NW1/4 sec. 24, T. 14 N., R. 9 W.

METHODS OF INVESTIGATION

Water samples were collected from 34 existing systems served by wells (fig. 3) during March 19-21 and June 27-28, 1979, to obtain data representing low and high water levels. The water levels measured during June (3.40 to 8.42 feet below land surface) were higher than in March in corresponding wells; however, landowners reported even higher levels in May due to early runoff from snowmelt and to lack of normal rainfall in June.

The samples were collected from taps or faucets after stabilization of specific conductance and water temperature indicated that flow was representative of the aquifer. Water temperature, pH, and specific conductance were measured, and samples were filtered and otherwise prepared onsite for laboratory analysis. Chemical analyses were made by the U.S. Geological Survey's laboratory in Denver, Colo., according to methods described by Skougstad, Fishman, Friedman, Erdmann, and Duncan (1979). Samples collected at 10 sites were analyzed for fecal coliform bacteria by membrane-filter techniques. (Greeson and others, 1977).

RESULTS

Results of onsite measurements of specific conductance, pH, and temperature and laboratory analyses for sulfate, nitrogen, and phosphorus are included in table 1. Results of more complete analyses including common ions, nutrients, and iron are given in table 2. Results for samples collected from the same sources during 1974 and 1975 (Wilke, 1976) are included for comparison. Fecal coliform bacteria, indicators of contamination by human or other warm-blooded animal waste, were absent from all samples collected at 10 sites (fig. 3) during March and again during June 1979.

The concentrations of dissolved solids, consisting mainly of calcium, magnesium, bicarbonate, and sulfate, were less than 300 mg/L (milligrams per liter).

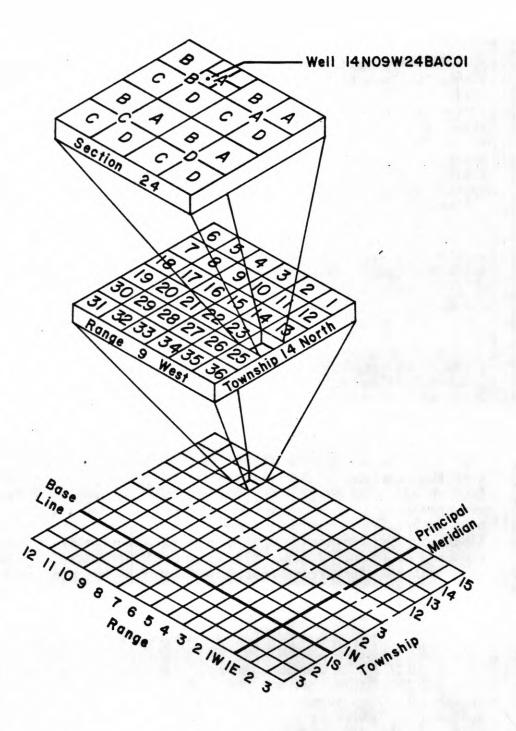


Figure 2.--Well-numbering system.

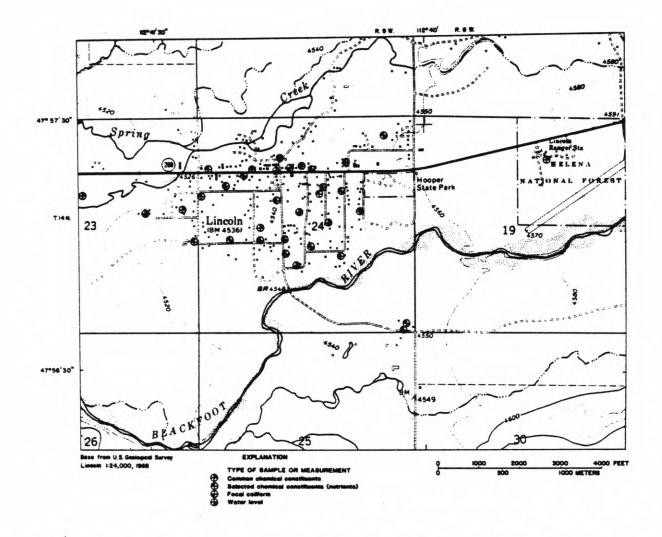


Figure 3.--Locations of wells and types of analyses for water samples collected during March and June 1979.

Changes from sampling to sampling appear to be insignificant and without readily discernable trends. Maximum concentrations of sodium were 3.5 mg/L; potassium, 0.9 mg/L; and chloride, 2.9 mg/L (table 2).

Results of analyses for phosphorus and nitrogen, common constituents of sewage and fertilizers, are the most abundant and amenable to statistical comparison. However, phosphorus is readily precipitated by the calcium and magnesium in soil water; therefore, excessive concentrations would not be expected even in samples from contaminated wells. The maximum concentration of phosphorus was 0.06 mg/L (table 1); most variations were within the range of laboratory precision. Conversely, nitrogenous wastes normally are rapidly oxidized by soil bacteria to soluble nitrite and nitrate, which are readily transported by ground water.

To facilitate comparison, concentrations of soluble nitrogen in samples collected during March 1979 were plotted versus concentrations at the same sites during May or June 1975 (fig. 4). If the concentration between the two sampling periods was unchanged, the regression would define a line inclined at 45° through the origin with a slope equal to 1.0. Data points above the line indicate increases in concentration and those below the line indicate decreases. The maximum concentrations of 0.75 mg/L for 1975 and 0.57 mg/L for 1979 in samples from well 14N09W24BAD02 (table 1), were anomalously large for the area, but were significantly less than the maximum of 10 mg/L recommended by the U.S. Environmental Protection Agency (1976) for drinking water. The background concentration of nitrogen in pristine ground water in the area is unknown, but probably does not exceed 0.2 mg/L. The corresponding mean concentrations of 0.23 mg/L for 1975 and 0.16 mg/L for 1979, as well as the slope of the regression line (0.47), indicate a general decrease in concentration from the spring 1975 to the spring 1979. Concentrations decreased slightly in 14 of the 25 sampled sources.

The plot for June 1979 versus March 1979 (fig. 5) also shows an apparent general decrease in concentration with time. For the March and June surveys, the maximum concentrations were 0.57 mg/L during March and 0.29 mg/L during June in samples from 14N09W24BAD02 (table 1) and the mean values were 0.16 mg/L for March and 0.11 mg/L for June. Concentrations decreased in 23 of the 28 samples. The relatively small coefficient of determination (0.40) between the March and June 1979 regressions (fig. 5) compared with that (0.59) between the May or June 1975 and March 1979 regression (fig. 4), as well as the decrease in concentration, probably reflects the effect of non-uniform dilution by snowmelt of the ground water sampled during June 1979.

The changes in concentration of nitrogen with time were substantiated using statistical comparison tests (Paul Woods, U.S. Geological Survey, written commun., July 1980). However, the small apparent decreases in the uniformly small concentrations of nitrogen and other ions could represent differences in laboratory or onsite techniques. In any event, the results clearly indicate no significant deterioration in the quality of the sampled waters with respect to the constituents for which the analyses were made. All samples were suitable for drinking water with respect to the constituents tested.

A resistant "hardpan," reportedly penetrated near the water table during installation of driven wells, may locally isolate the upper part of the aquifer from the shallow soil water. If the aquifer is not isolated, the absence of fecal coliform bacteria indicates that the soil or the aquifer is an effective' filter. Abundant phreatophytes probably use a significant part of the wastederived nutrients in the shallow ground water. Finally, recharge to the aquifer and lateral movement of fresh water within it are evidently adequate to dilute significantly any waste that enters near the supplies tested. These factors combine to diminish and obscure the effects of percolation of sewage effluent into the aquifer.

The results of this study do not necessarily indicate that the water from all wells in the Lincoln area is suitable for domestic use. Even for the limited number of wells tested, increased pumpage might accelerate percolation of deleterious waste into the aquifer.

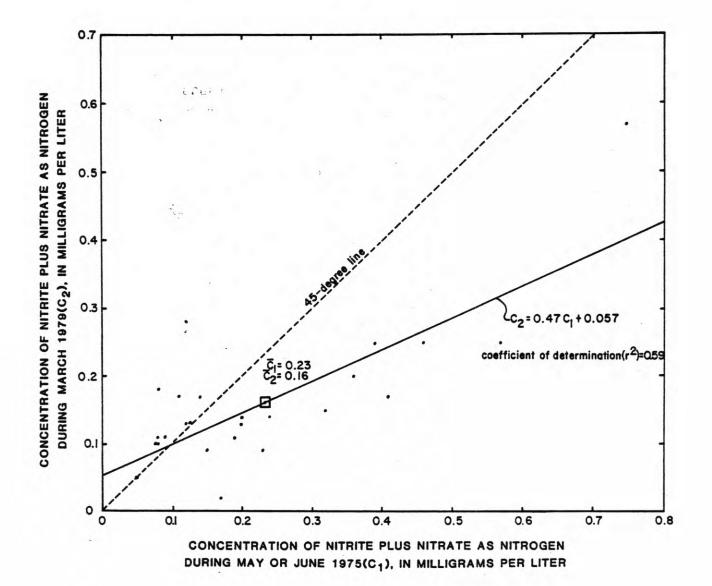


Figure 4.--Relation of concentrations of nitrite plus nitrate in ground water during March 1979 to concentrations during May or June 1975 from the same wells.

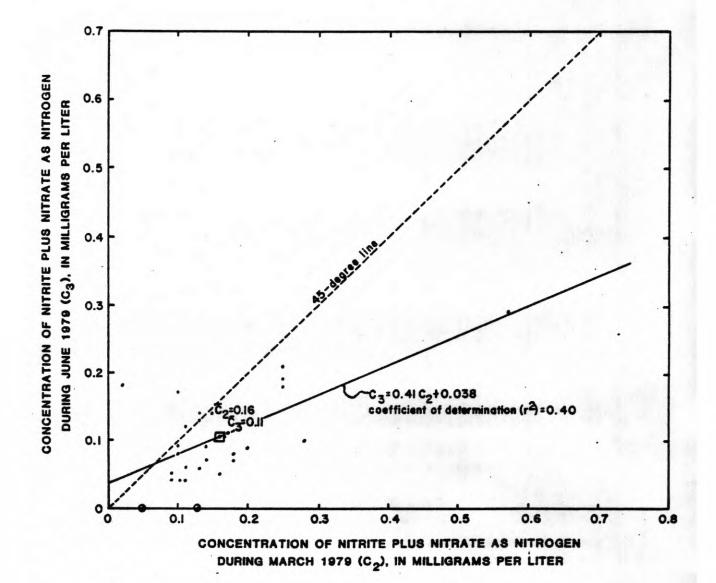


Figure 5.--Relation of concentrations of nitrite plus nitrate in ground water during June 1979 to concentrations during March 1979 from the same wells.

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- Greeson, P. E., Ehlke, T. A., Irwin, G. A., Lium, B. W., and Slack, K. V., 1977, Methods for collection and analysis of aquatic biological and microbiological samples: U.S. Geological Survey Techniques of Water-Resources Investigations, book 5, chapter A4, 332 p.
- Skougstad, M. W., Fishman, M. J., Friedman, L. C., Erdmann, D. E., and Duncan, S. S., 1979, Methods for determination of inorganic substances in water and fluvial sediments: U. S. Geological Survey Techniques of Water-Resources Investigations, book 5, chapter A1, 626 p.
- U.S. Environmental Protection Agency, 1976, National interim primary drinking water regulations: U.S. Environmental Protection Agency, Office of Water Supply Publication, EPA-570/9-76-003, 159 p.

_____1977, National secondary drinking water regulations (proposed): Federal Register, v. 42, no. 62, Mar. 31, Part I, p. 17143-17147.

Wilke, K. R., 1976, Ground-water levels and chemical quality of ground water in Lincoln, Montana: U.S. Geological Survey Open-File Map 76-333.

Table 1.--Results of analyses for miscellaneous chemical constituents in ground water

[Except as indicated otherwise, constituents are dissolved and values are reported in milligrams per liter. Analysis by: MDHES, Montana Department of Health and Environmental Sciences; USGS, U.S. Geological Survey. Water level below land surface and water temperature: R, reported. Micromhos, micromhos per centimeter at 25 degrees Celsius; ^oC, degrees Celsius]

			3. 3.		Water level below land	Specific conduc- tance
Well number	Latitude	Longitude	Date	Time	surface (feet)	(micro- mhos)
14N09W23DAA01	465702	1124121	10-09-74 05-28-75 03-19-79		4.5	335
14N09W24AA B01	465728	1124022	10-09-74 05-28-75 03-20-79 06-28-79	1500 1115	5.14	380 336
14N09W24ABC02	465720	1124039	10-09-74 05-28-75 03-19-79 06-27-79	 1000 1010	4.89 3.40	 300 330
14N09W24ABD02	465721	1124028	10-09-74 05-28-75 03-19-79 06-27-79	 1520 0950	=	 360 331
14N09W24ACA03	465713	1124030	10-09-74 05-28-75 03-19-79 06-28-79	1455 1605	 7.13 5.24	295 309
14N09W24ACB03	465717	1124037	10-09-74 05-28-75 03-20-79 06-27-79	 1315 1145	Ξ	 335 319
14N09W24ACB04	465713	1124037	10-09-74 05-28-75 03-21-79 06-28-79	 0945 1655	Ξ	 305 314
14N09W24BAC01	465720	1124100	10-09-74 05-28-75 03-19-79 06-28-79	 1030 1215	7.3	310 326
14N09W24BAC03	465723	1124052	10-09-74 05-28-75 03-20-79 06-27-79	1055 1100	Ξ	345 320
14N09W24BAD02	465720	1124044	10-09-74 05-27-75 03-19-79 06-28-79	 1255 1155		375 343

Onsite pH (units)	Water tem- pera- ture (°C)	Sul- fate (SO ₄)	N1trogen NO ₂ + NO ₃ as N	Ammo- nia, total as N	Phos- phorus as P	Analysis by
 7.7	 10.0	 8.6	0.14 .41 .17	<0.01	0.004	MDHES MDHES USGS
 7.7	 8.0 7.0	 6.9 11	.10 .12 .13 .14	<.01 	.001 .01 .00	MDHES MDHES USGS USGS
 7.5	 7.0 7.0	 10 9.6	.24 .57 .25 .18	<.01 	.001 .00 .00	MDHES MDHES USGS USGS
 8.0	9R 7.0 7.0	 8.2 10	.15 .46 .25 .21	<.01 	.002 .00 .00	MDHES MDHES USGS USGS
 7.5	 12.0 7.5	 8.4 12	.13 .19 .11 .06	.01 	.001 .01 .00	MDHES MDHES USGS USGS
 7.6	8.0 7.0	 9.2 11	.12 .20 .14 .07	<.01 	.005 .01 .00	MDHES MDHES USGS USGS
 7.4	 7.0 7.0	8.2 12	.11 .08 .18 .07	<.01	.002 .01 .06	MDHES MDHES USGS USGS
 7.8	 9.0 7.0	 8.9 11	.21 .12 .13 .13	< .01 	.002 .00 .00	MDHES MDHES USGS USGS
7.6	 9.0 8.0	8.5 10	.20 .12 .28 .10	< .01	.002 .01 .00	MDHES MDHES USGS USGS
 7.5	7.0	 9.1 12	•53 •75 •57 •29	< .01	.003 .00 .00	MDHES MDHES USGS USGS

1	Location				Water level below land	Specific conduc- tance
Well number	Latitude	Longitude	Date	Time	surface (feet)	(micro- mhos)
14N09W24BBC01	465720	1124116	10-09-74 05-28-75 06-28-79	1240	6.9 	368
14N09W24BCA03	465715	1124110	10-09-74 05-28-75 03-21-79	1300	Ξ	350
14N09W24BCA04	465717	1124104	10-09-74 05-28-75 03-21-79 06-28-79	 1325 1225	 5.82 5.19	 300 333
14N09W24BCB01	465712	1124119	10-09-74 05-28-75 03-19-79 06-28-79	 1120 1345	7R 	320 335
14N09W24BDC01	465712	1124055	10-09-74 05-28-75 03-19-79 06-27-79	1330 1110	4R 	320 317
14N09W24CAA02	465703	1124049	10-09-74 05-28-75 03-21-79 06-28-79	 1440 1420	 8.81 7.10	340 328
14N09W24CAB04	465702	1124058	10-09-74 05-28-75 03-19-79 06-27-79	 1230 1210	 7.05 5.58	355 339
14N09W24CAD04			10-09-74 05-28-75 03-19-79 06-28-79	1255 1435	9.95 8.42	325 326
14N09W24CBA01	465703	1124108	10-09-74 05-28-75 03-21-79 06-28-79	1505 1800	8R 4.34	330 307
14N09W24DBB03	465700	1124039	10-09-74 05-28-75 03-19-79 06-28-79	 1415 1550	 6.44	 345 191
14N09W24DBD01	465759	1124029	10-09-74 05-28-75 03-19-79 06-27-79	1435 1335	7.5 	345 270

Table 1.--Results of analyses for miscellaneous chemical constituentsground water--Continued

-

Onsite pH (units)	Water tem- pera- ture (°C)	Sul- fate (SO ₄)	Nitrogen NO ₂ + NO ₃ as N	Ammo- nia, total as N	Phos- phorus as P	Analysis by
 7.5	 7.5	 11	0.09 .05 .05	<0.01	.002 .00	MDHES MDHES USGS
=	 7.0	 7.9	.14 .14 .17	< .01	.001 .01	MDHES MDHES USGS
 7.6	 12.0 7.0	 7.3 12	.19 .17 .02 .18	< .01	.001 .00 .00	MDHES MDHES USGS USGS
 7.7	 10.0 10.0	 8.5 12	.12 .09 .11 .12	< .01	.002 .00 .01	MDHES MDHES USGS USGS
 7.5	12.0 6.5	 9.2 10	.11 .24 .14 .09	< .01	.001 .00 .00	MDHES MDHES USGS USGS
 7.5	 9.0 8.5	 7.5 12	.08 .36 .20 .09	< .01	.002 .01 .01	MDHES MDHES USGS USGS
 7.6	 8.5 6.5	 9.4 11	.16 .32 .15 .14	< .01	.001 .00 .00	MDHES MDHES USGS USGS
 7.6	 7.5 6.0	 8.8 12	.12 .15 .09 .05	< .01	.002 .00 .01	MDHES MDHES USGS USGS
 7.6	 7.5 7.0	 7.9 12	.11 .08 .10 .09	< .01	.002 .01 .00	MDHES MDHES USGS USGS
 7.5	 6.0 6.5	 9.8 12	.09 .23 .09 .04	< .01	.001 .00 .00	MDHES MDHES USGS USGS
 7.8	 6.0 6.5	 10 9.6	.07 .05 .05 .00	< .01	.001 .00 .00	MDHES MDHES USGS USGS

Table 2 .-- Results of analyses for common chemical constituents in ground water

	Location			1	Water level below land	Specific conduc- tance		Water tem- pera-	Hard- ness
Well number	Latitude	Longitude	Date	Time	surface (feet)	micro- mhos	pH (units)	(°C)	as CaCO3
14N08W19ABD01	465722	1123916	0974		11.20				
			03-20-79	1530		345	7.81	5.0	160
			06-28-79	1045		286	8.1	9.0	150
14N09W23ADC03	465709	1124137	0675			336			180
			03-21-79	1040	4.04	335	7.81	6.0	170
			06-27-79	1355	3.68	329	7.7	8.0	180
14N09W24ACC03	465707	1124033	0974						
			0575						
			03-20-79	1355	5.25	315	7.61	8.0	150
			06-28-79	1720	4.01	318	7.6	6.5	160
14N09W24ACD02	465709	1124023	0974		5.50				
			0675		3.20	321			160
			03-20-79	1420	6.51	325	7.81	11.0	
			06-28-79	1630		305	7.6	9.5	160
14N09W24BAC04	465720	1124052	0974						
			0575						
			03-20-79	1035		335	7.81	9.0	170
			06-27-79	1045		321	7.7	8.0	170
14N09W24BAD03	465720	1124047	0675			335			180
			03-20-79	1005		325	7.71	9.0	160
			06-27-79	1025		337	7.6		
14N09W24BDD01	465709	1124044	0974		5.10				
			0575		3.40				
			03-20-79	1200		345	7.61	7.0	
			06-28-79	1945	4.37	300	7.5	7.0	160
14N09W24CAB02	465705	1124058	0974		5.90				
			0675			351			180
			06-28-79	1815		347	7.5	6.5	170
14N09W24CAD05	465657	1124045	03-20-79	1245	9.03	350	7.91	7.0	180
			06-28-79	1530	7.12	303	7.6	6.5	150

[Except as indicated otherwise, pH data are onsite values, constituents are dissolved, and values are reported in milligrams per liter. Micromhos, micromhos per centimeter at 25 degrees Celsius; $^{\circ}C$, degree Celsius; $\mu g/L$, micrograms per liter]

¹Laboratory determination.

Date	Hard- ness, noncar- bonate, as CaCO ₃	Calcium (Ca)	Magne- sium- (Mg)	Sodium (Na)	Percent	Sodium adsorp- tion ratio (SAR)	Sodium + po- tas- sium (as Na)	Potas- sium (K)
0974	·							
03-20-79	50	41	14	2.3	3	0.1		0.9
06-28-79	10	37	14	2.2	3	.1	3	.9
0675		46	15	3.1	·	.1		.6
03-21-79	0	43	15	2.0	3	.1		.7
06-27-79	7	46	15	2.0	2	.1	3	.8
0974			· 					
0575								
03-20-79	1	39	13	1.8	3	.1		.7
06-28-79	0	40	14	2.1	3	.1	3	.8
0974								
0675		43	15	2.6		.1		.6
03-20-79								
06-28-79	8	40	14	2.0	3	.1	3	.9
0974								
0575								
03-20-79	0	43	14	2.2	3	.1		.9
06-27-79	10	45	14	2.1	3	.1	3	.8
0675		49	14	3.5		.1		•7
03-20-79	20	41	14	2.2	3	.1		.8
06-27-79								
0974								
0575								
03-20-79								
06-28-79	4 .	41	15	2.2	3 .	1	3	.8
0974		-					-	
0675		53	12	3.1		.1		.(
06-28-79	4	45	15	2.5	3	.1	3	• !
03-20-79	28	48	14	2.2	3	.1	3	.1
06-28-79	8	38	13	2.0	3	.1	3	

Date	Bicar- bonate (HCO ₃)	Car- bonate (CO ₃)	Alka- linity, total as CaCO ₃	Sulfate (SO ₄)	Chlo- ride (Cl)	Fluo- ride (F)	Silica (S10 ₂)	Dis- solved solids residu at 180°C
0974								
03-20-79			110	13	1.3	0.1	11	154
06-28-79			140	10	.7	.1	12	162
0675	210	0	180	8.9	1.0	.1		
03-21-79			170	8.1	8	.1	12	183
06-27-79			170	10	.8	.1	9.9	183
0974								
0575								
03-20-79			150	9.3	.7	.1	10	167
06-28-79			160	11	.6	.1	11	174
0974								
0675	200	0	160	9.0	1.8	.1		
03-20-79			160					
06-28-79			150	10	.8	.1	12	170
0974								
0575								
03-20-79			170	9.1	1.1	.1	11	174
06-27-79			160	11	1.1	.1	9.7	181
0674	220	0	180	7.8	2.0	.1		
03-20-79			170	11	2.9	.1	10	182
06-27-79			170					
0974								
0575							、	
03-20-79			160	9.6	.9	.1	9.8	172
06-28-79			160	11	.6	.1	11	177
0675					1.0	.1		
03-20-79	220	0	180	6.1				
06-28-79			170	11	1.4	.1	11	190
03-20-79			150	10	.8	.1	9.7	170
06-28-79	-	-	140	9.4	.5	.1	10	165

Date	Dis- solved solids, sum of consti- tuents	Dis- solved solids (tons per acre- foot)	Nitrogen, NO ₂ + NO ₃ as N	Ammo- nia, total as N	Phos- pho- rus as P	Iron (Fe) (µg/L)	Analysis by
0974	-	-	0.11				MDHES
03-20-79	150	0.21	.16		0.01		USGS
06-28-79	161	.22	.05				USGS
0675	288	·	.08	<0.01	.001		MDHES
03-21-79	184	.25	.11		.01	20	USGS
06-27-79	187	.25	.04			10	USGS
0974			.09				MDHES
0575			.20		.001		MDHES
03-20-79	165	.23	.13		.00	20	USGS
06-28-79	176	.24	.06				USGS
0974							MDHES
0675	273		.08	<.01	.002		MDHES
03-20-79			.10	-	.00		USGS
06-28-79	171	.23	.17				USGS
0974		_	.16				MDHES
0575			.11		.002		MDHES
03-20-79	184	.24	.17		.01	20	USGS
06-27-79	181	.25	.11			10	USGS
0674	296	·	•39	.01	.005		MDHES
03-20-79	167	.25	.25		.01	10	USGS
06-27-79	_	-	.19				USGS
0974							MDHES
0575							MDHES
03-20-79		.23	.18		.01		USGS
06-28-79	178	.24	.08				USGS
0675	294		•39	<.01	.003		MDHES
03-20-79		-					USGS
06-28-79	190	.26	.17				USGS
03-20-79	176	.23	.13		.01	10	USGS
06-28-79	158	.22	.00				USGS

