WATER QUALITY OF SELECTED LAKES IN EASTERN SHERIDAN COUNTY MONTANA

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INTRODUCTION

Dozens of lakes are located in eastern Sheridan County
Montana (Plate 1). Although a few individual water analysis of area
lakes have been conducted, background water quality has never been
analyzed or compiled for a majority of these lakes. Realizing the
importance of maintaining water quality in the county, area
residents proposed conducting a baseline sampling program to assess
the existing lake water quality and to provide background data in
the event of future impacts.

These lakes are very diverse in size, depth, water quality, and uses. The size of the lakes ranges from small unnamed ponds covering only a few acres to large lakes such as Medicine Lake covering several square miles. While most of the lakes are shallow ephemeral water bodies that are dry during extended droughts, many are deeper perennial water bodies. The deepest lake in the region is Brush Lake which is over 60 feet deep (Donovan, in press). Although very little previous work has been conducted comparing the water quality of lakes in this region, observations based on vegetation, presence or absence of deposits of sodium sulfate salts, and traditional uses all point towards a wide range of lake water quality. Uses of the lakes include stock watering, recreation, wildlife habitat, and a potential for extraction of mineral resources (glauber salts) in some of the lake basins. Many of the lakes are important to the water balance of the region both as focused recharge sources and as buffers influencing ground-water levels.

PHYSIOGRAPHY AND CLIMATE

The study area covers the eastern one-third of Sheridan County (Figure 1). This region consists of a slightly elevated generally hummocky land surface in the north and west bounded by relatively flat broad swales in the south and east. The swales are named the Clear Lake and Medicine Lake swales and appear to be associated with relict pre-Quaternary river systems. located in both the hummocky upland areas and the relatively flat broad swales. Most of the larger lakes are located in the broad swales. Ephemeral streams such as Lake Creek occupy parts of the larger swales. Surface drainage is very poorly developed in much of the rest of the area. Typically, the individual closed-lake basins form catchments collecting runoff from the immediate surrounding area. Only under extremely high water conditions are many of these lakes physically connected by surface water. Most of the lakes have formed in topographic depressions such as kettles and meltwater channels that developed following the retreat of the last late-Wisconsinan glaciers(10,000-12,000 years B.P.).

Eastern Sheridan County has a semiarid continental climate, characterized by cold dry winters, moderately hot and dry summers, and cool dry falls. The average precipitation in Westby, near the center of the study area is about 14.2 inches per year, with about 65% of the precipitation falling from May through August (Table 1). Potential evaporation is significantly higher than precipitation. Measured evaporation from a class A pan near Froid, MT averaged 52 inches annually from 1984 to 1988. Evaporation estimated from

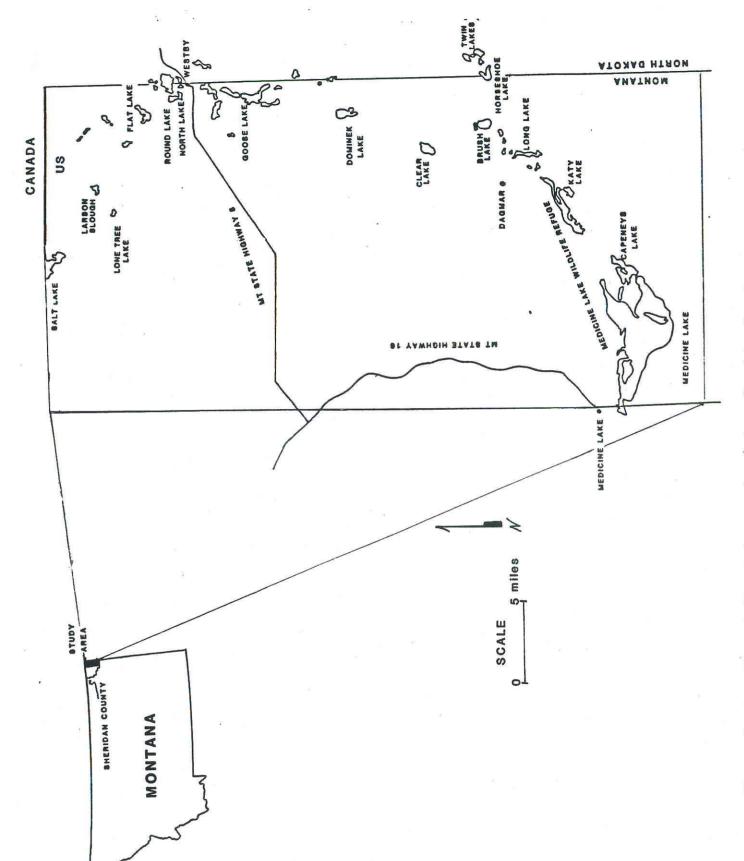


Figure 1. Location of the study area in eastern Sheridan County.

Table 1. Measured total monthly precipitation in the Westby area. Data collected by the National Oceanic and Atmospheric Administration

														16					
TOTAL	15.05	12.33	14.15	9.55	6.88	10.57	17.79	12.26	69.6	10.13	15.11		11.84		12, 14			14.19	
DEC	.55	.29	.82	.30	.22	. 29	90.	.0	2	.10	16		.34		3	1		.43	
NON	.26	.81	0	.29	0	.30	77	.11	.43	.24	. 28		.28		20	ì		.34	
OCT	1.06	.81	.82	.01	.42	.93	1.41	.18	.27	1.65	.03		92.		09)		69.	
SEP	1.58	.78	1.85	2.00	1.25	1.25	4.97	1.04	1.54	.42	.16		1.67		1 53			1.42	
AUG	72.7	2.04	1.93	.88	.29	2.35	67.	.61	.95	.72	2.13		1.50		1.55			2.07	
JUL	1.14	1.12	1.31	2.27	.07	1.02	3.22	5.17	1.32	.50	5.21		1.71		2 03			1.92	
NOC	2.63	3.97	1.14	1.58	3.31	.93	2.56	1.20*	2.33	1.89	3.01		2.15	33 85	2 23	3		2.89	
MAY	.39	1.47	3.07	1.16	.59	1.29	3.12	2.50*	1.07	1.30	2.23		1.60		1 65	3		1.93	
APR	1.30	.50	.82	0	0	1.15	52.	*10.	777	1.38	.57		.63		29	3		1.15	
MAR	.43	.38	.85	.73	.07	.78	.17	1,16*	07.	-39	1.23		.54		04	2		77.	
FEB	.31	%	94.	.07	.03	.22	.37	.12*	.20	.21	.01		.21		10	<u>.</u>		.45	
JAN	99.	.10	1.08	.26	.63	90.	52:	.15*	101	1.33	.10		.45		67	74.		97.	
YEAR	1980	1981	1982	1983	1984	1985*	1986*	1987	1988	1989	1990	10 Year	Average	1960-69	11 Year	1980-90	20 Year	Average 1970-1989	

Westby data not available in 1985, 1986, and Jan-Jun of 1987, data from the station in nearby Plentywood was used for calculations.

daily water level fluctuations in an aquifer hydraulically connected to Medicine Lake indicate similar evaporation rates (Donovan, 1988).

HYDROGEOLOGIC SETTING

Bedrock in the study area consists of interbedded sand, silt, clay, and lignite of the Paleocene Fort Union Formation. In this part of Sheridan County, bedrock is generally covered by younger sediments. Because of these intervening deposits, the interaction between lake water and ground water in the Fort Union Formation or deeper formations appears to be insignificant.

All of the lakes in this area are underlain by late-Wisconsinan glacial deposits. These deposits include outwash sand and gravel, lacustrine silt and clay, and till consisting of a mixture of sand, silt, clay, and boulders. Ground water in the glacial deposits is in direct connection with water in the lakes.

Post-glacial lake deposits typically overlie the glacial deposits within the lake basins. These consist of marls, tufas, evaporitic salts, clays, and organic debris. Very little information is currently available on the thickness and variability of these lake deposits. Current research by Joe Donovan (Penn State University) has focused on investigating the occurrence and origin of some of the Holocene lake deposits in Brush Lake and White Lake. These lake deposits form zones of water transfer between the water in the lakes and ground water in the surrounding sediments. The water chemistry of the individual lakes is dependent on geochemical reactions which include: dissolution of

minerals exposed to lake water, precipitation of minerals from lake water, and similar reactions involving ground water moving in and out of the lakes. The concentrations of dissolved solids in these lakes depends chiefly on the flux of water moving out of an individual lake. As the flux of water through a lake is restricted, the concentration of dissolved solids is increased due to evaporation.

PURPOSE

Protection of the regions water resources was the primary driving force behind this investigation. The initial objective was to obtain representative water samples from a cross-section of lakes in this part of the state. Based on the water quality data, a lake classification scheme was developed. In addition, an attempt was made at determining what impacts, if any, have occurred to area lakes as a result of past land use practices.

Land uses that may have potentially impacted area water resources include agriculture related activities and oil field related activities. Farming could impact lakes if fertilizer, pesticides, and animal wastes are flushed into the lakes by runoff or ground water seepage. Expected impacts would include increased nutrients causing noxious organisms to flourish resulting in obnoxious odors, poisoning of wildlife, and shoreline degradation depending on the extent of contamination.

Oil-field activities that could impact lake water quality all relate to the extraction and production of hydrocarbons. Potential contaminants include additives used in the drilling process, saltwater in the drilling muds or produced along with the

hydrocarbons, acid used in developing producing zones, and waste hydrocarbons. Previous ground water studies (Reiten, 1991, Payne and Reiten, 1991, and Reiten and Tischmak, in press) have determined produced brines as a major source of industrial contamination in the Montana portion of the Williston Basin. Impacts have included soil sterilization and aquifer water quality degradation. Several wells and stock ponds that previously had produced potable water have been degraded and are currently unusable. Environmentally unsound brine storage and disposal practices used during the 1960's and 1970's were the major causes of water quality degradation.

METHODS

Water samples were collected from about 50 lakes in eastern Sheridan County. A few water samples were also collected from borderline lakes in Williams and Divide Counties, North Dakota. Both named and unnamed lakes were sampled during this survey. Informal names were attached to the unnamed lakes to simplify interpretation and discussion. All of the lakes surveyed are identified on Plate 1. The location of the sampling sites are identified by Township, Range, Section, and Tract in the accompanying tables.

About 90 field water samples were collected and complete water analyses were run on 30 water samples. Twenty-nine of the analyses were conducted at the MBMG water lab in Butte and the remaining analysis was conducted at the water lab at Penn State University. Standard analytical procedures were used in analyzing all of the water samples.

All of the samples were tested for specific conductance and temperature at the sampling site. The conductivity meters used were calibrated using known standards to develop correction curves. Other parameters including dissolved oxygen, pH, and chloride concentration were measured for selected lake samples. Calibrated field instruments were used to measure dissolved oxygen and pH. Ouantab chloride titrators were used to measure concentration. At sample sites tested for only field parameters, water was collected by simply dipping an adequate volume from near the shoreline. At sample sites with complete lab analyses run, water was collected using the sampling system described below. A lake water sampling system was constructed using a tire inner tube mounted with a plywood platform in the center of the tube. tubing was inserted through the platform extending 15 to 25 centimeters below the platform. A screen was wrapped around the inlet portion of the tygon tubing to prevent clogging of the About 50-feet of tygon tubing was attached to the orifice. sampling platform enabling the water sample to be collected away from shore.

The sampling platform was thrown or floated out from shoreline and attached to a peristaltic pump for sample collection. This sampling system ensured collecting water from relatively similar depths in all lakes. Field parameters were measured immediately. Laboratory samples collected included raw unfiltered lake water and water filtered through 0.45 micron filters and preserved by refrigeration. Samples were acidified upon submittal to the MBMG lab in Butte. The Penn State sample was acidified following collection and also refrigerated.

RESULTS

The results of measurements of field water quality parameters are summarized in Table 2. The water samples were initially classified on the basis of field specific conductance (SC). The SC is a measure of the electrical conductivity of the water sample which is generally directly proportional to the dissolved solids concentration. Field values of SC were used to allow the comparison of a larger number of samples than if the comparison were restricted to samples analyzed in the lab. Five ranges of SC values were set up as categories for classifying the lakes. The five categories and associated SC ranges are listed below:

GROUP I 0 to 5000 umhos/cm

GROUP II 5000 to 10000 umhos/cm

GROUP III 10000 to 25000 umhos/cm

GROUP IV 25000 to 50000 umhos/cm

GROUP V more than 50000 umhos/cm

Plate 1 shows the distribution of these 5 lake water categories in the study area.

The range of field pH for all five water groups was from 8.60 to 10.00 based on samples collected for laboratory analysis (Figure 2). Water in Group I, III, IV, and V lakes displayed a wide range of pH values within each group. Water in Group II lakes had relatively uniform pH values ranging from 8.90 to 9.30. Field chloride concentrations in all lakes ranged from below detection limits to more than 9,300 mg/L.

TABLE 2. SHERIDAN COUNTY LAKES BASELINE FIELD WATER QUALITY SAMPLED IN AUGUST/SEPTEMBER 1990

SAMPLE	LABGS	LABGS	LABGS	FIELD	LABGS	LABGS	FIELD	FIELD	FIELD	LAB	FIELD	FIELD	FIELD	FIELD	FIELD	FIELD	LAB	FIELD	FIELD
*CONTAMINATION INDEX	0.074	0.062	0.028		0.055	0.108	0.021			0.016							0.009		
FIELD *C CL MG/L	7276	3592	1016		2266		109			7.4							59		
DO AS PERCENT OF SATURATION	108	101	165		132	86			10	26							110		
DO TEMP C	21.2	29.0	30.0		26.8	29.0				22.0							24.0		
DISSOLVED OXYGEN (DO) MG/L	00.6	7.60	12.50		10.50	7.40				8.60						2	05.6		
4 0 0	9.33	9.61	62.6		9.63	9.65	9.07			9.25							8.94		
SPECIFIC CONDUCTANCE umhos/cm	00086	58450	36580	39420	41210	02967	5310	4420	4180	4580	4760	4870	4320	929	0695	5100	6300	4210	0295
LOCATION	15910308cbcc	15910308DBDA	15910308DDAB	15910309CCCA	15910317AADD	15910317ABDB	16310334CBBB	31N56E05CDBC	31N56E06BBAD	31N56E09BBDD	31N56E13CCDC	31N57E04CCBB	31N57E05DDDB	32N56E31CCBB	32N56E33CBBB	32N57E13CDAA	32N57E13DBDB	32N57E13DBDB	32N57E31CAAD
SAMPLE	08/23/90	08/23/90	08/23/90	08/23/90	08/23/90	08/23/90	08/56/90	08/52/90	08/55/90	08/52/90	08/58/90	08/58/90	08/55/90	08/55/90	08/52/90	08/22/90	08/54/90	08/22/90	08/52/90
SAMPLE	V	¥	٧	8	۷	¥		۵	ບ	ш	ii.	x	G	8	¥	4	89	89	٦,
LAKE NAME	091060 NODAK 1	091065 NORTH TWIN	091061 MIDDLE TWIN	091064 SOUTH TWIN	091063 SOUTH TWIN	091062 VAN VOAST	MILKY	091034 MEDICINE	091033 MEDICINE	091027 MEDICINE	091035 MEDICINE	091037 GAFFNEY	091036 MEDICINE	091032 MEDICINE	091031 MEDICINE	NO. 12	NO. 12	NO. 12	091038 MEDICINE
SITE	091060	091065	091061	091064	091063	091062	091075 MILKY	091034	091033	091027	091035	091037	091036	091032	091031	091039 NO.	091003 NO.	091030 NO.	091038

SAMPLE	LAB	FIELD	FIELD	LAB	FIELD	FIELD	FIELD	LAB	FIELD	FIELD	FIELD	FIELD	LAB	FIELD	FIELD	LAB	FIELD	FIELD	FIELD	FIELD
*CONTAMINATION INDEX	0.024			0.016				0.005					0.036			0.024				
FIELD ' CL MG/L	128			78									823			242				
DO AS PERCENT OF SATURATION	87							126					134			89				
DO TEMP C	22.1			22.0				25.0					25.5			27.0				
DISSOLVED OXYGEN (DO) MG/L	7.70			05.6				11.10					11.00			5.50				
HQ O	9.30			9.15				9.92					10.01			9.72				
SPECIFIC CONDUCTANCE umhos/cm	5340	2440	5360	5200	5370	4750	2260	2230	2300	2190	2280	5610	22990	5910	24440	31190	20000	9300	5545	5190
LOCATION	32N57E32BBAB	32N58E02CCC	32N58E03AACB	32N58E03ACAD	32N58E03DACC	32N58E04CDCD	32N58E04DAAA	32N58E04DAAD	32N58E04DAAD	32N58E04DABA	32N58E04DACB	32N58E08DCCC	32N58E09AAAB	32N58E09BABA	32N58E10BBBD	32N58E17CDDB	32N58E17CDDB	32N58E18AADB	32N58E18ACAB	32N58E18ACCB
SAMPLE	08/29/90	08/22/90	08/22/90	08/25/90	08/22/90	08/22/90	08/22/90	08/54/90	08/22/90	08/22/90	08/22/90	08/22/90	08/54/90	08/22/90	08/22/90	08/54/90	08/22/90	08/22/90	08/22/90	08/22/90
SAMPLE	1	U	A	٥	8	9	8	¥	A	U	۵	u.	89	Ξ	A	¥	A	ш	۵ .	U
LAKE	DEEP	LONG	LONG	LONG	TONG	091044 NO. 12	FRED	FRED	FRED	FRED	FRED	091043 NO. 12	091005 BERGER POND	091045 NO. 12	091046 BERGER POND	KATY	KATY	NO. 12	091041 NO. 12	091040 NO. 12
SITE	091026 DEEP	091049 LONG	091047 LONG	091006 LONG	091048 LONG	091044	091051 FRED	091008 FRED	091090 FRED	091052 FRED	091053 FRED	091043	091005	091045	091046	091004 KATY	091088 KATY	091042 NO.	091041	091040

LAKE	SAMPLE	SAMPLE	LOCATION	SPECIFIC	HG.	DISSOLVED	DO	DO AS		*CONTAMINATION	SAMPLE	щ Ņ
	POINT	DATE		CONDUCTANCE umhos/cm		OXYGEN (DO) MG/L	C	PERCENT OF SATURATION	CL MG/L	INDEX	TYPE	Ä
	υ	08/23/90	33N58E03ADAA	12020							FIELD	9
	8	08/23/90	33N58E05AABC	11900	9.36	09.6	24.0	113	128	0.011	LAB	
		08/56/90	33N58E22ABAB	15890	60.6	7.10	16.3	22		0.017	LAB	
		08/60/80	33N58E22DBDA01	01 6144	9.20	00.6	22.2	102		0.012	LABGS	SS
	A	08/30/90	33N58E24DDDA	83400	96.6	4.00	26.0	53		0.092	LAB	
	A	08/23/90	33N58E24DDDA	72750							FIELD	9
		08/31/90	33N58E26BADD	14500	9.04	11.00	18.0	116		0.018	LAB	
	A	08/54/90	33N58E27AABB	27800	9.23	06.9	22.0	87		0.013	LAB	
		06/20/60	33N58E27ACBC	16420	9.54	11.00	20.0	126		0.016	LAB	
	ပ	08/22/90	33N58E27CBAA	0996							FIELD	۵.
	A	08/54/90	33N58E27CBCC	9350	9.54	5.50	24.2	92	101	0.011	LAB	
	A	08/22/90	33N58E27CBCC	0296							FIELD	9
	8	08/22/90	33N58E27CBDB	8990	3						FIELD	Q
	8	08/22/90	33N58E28CDBA	15990							FIELD	٩
	A	08/54/90	33N58E28DCBC	15120	9.72	12.60	24.0	148		600.0	LAB	
	۷	08/22/90	33N58E28DCBC	16190							FIELD	Δ.
	A	08/25/90	33N58E29DACC	3070	9.03	7.00	25.0	84		600.0	LAB	
	8	08/52/90	33N58E29DCDD	3260					39		FIELD	0
091010 MALLARD POND	v	08/25/90	33N58E33BCBC	2590	9.48	9.12	22.0	103		0.010	LAB	
	۷	08/22/90	33N58E33BCDA	1475							FIELD	Q.

SITE	LAKE	SAMPLE	SAMPLE	LOCATION	SPECIFIC CONDUCTANCE umhos/cm	퓹	DISSOLVED OXYGEN (DO) MG/L	DO TEMP C	DO AS PERCENT OF SATURATION	FIELD CL MG/L	*CONTAMINATION INDEX	SAMPLE
091055 MAI	091055 MALLARD POND	8	08/25/90	33N58E33BCDC	2610							FIELD
091050 BLACKWATER	ACKWATER	V	08/25/90	33N58E35CCAD	88700							FIELD
091066 CLEAR	EAR	V	08/23/90	34N58E33BCCB	8690							FIELD
091071 S. GOOSE	GOOSE	u.		35N58E01BAAA	69345					3290	0.047	FIELD
091069 S. GOOSE	GOOSE	۷	08/56/90	35N58E11DAAD	21000					899	0.032	FIELD
091016 S. GOOSE	GOOSE	8	08/27/90	35N58E12DADA	28790	6.45	8.40	19.8	92	006	0.031	LAB
091092 S. GOOSE	GOOSE	8	08/56/90	35N58E12DADA	28740	9.45	8.40	19.8	65	006	0.031	FIELD
091013 BEEHIVE	EHIVE	4	08/56/90	35N58E13AADC	11150	60.6	07.9	18.5	89	546	0.022	LAB
091068 ISLAND	LAND	V	08/56/90	35N58E36AAAA	3590							FIELD
091024 MUDCRACK	CRACK		08/52/90	36N57E34BCBA	27690	9.38	8.60	21.8	102	0727	0.171	LAB
091018 GAULKE	JLKE	V	08/26/90	36N58E01DAAD	7170	9.15	10.20	19.3	110	29	0.008	LAB
091074 GAULKE	JLKE	8	08/56/90	36N58E01DDAC	7210							FIELD
091076 ROUND	ONC	Ø	08/56/90	36N58E01DDDD	28390	9.52	El .			1210	0.043	FIELD
091087 MCELROY	ELROY		08/58/90	36N58E09DDBA	358							FIELD
091078 NORTH	чтн г		08/27/90	36N58E12CAAD	8540	9.16	::::::::::::::::::::::::::::::::::::::			308	0.036	FIELD
091077 ROUND	OND	8	08/56/90	36N58E12DBCA	38790	9.59				1521	0.039	FIELD
091015 F00TBALL	JTBALL		08/56/90	36N58E13AADD	25860	9.65	20.00	20.0	524	2621	0.101	LAB
091073 NE WESTBY	WESTBY		08/56/90	36N58E13AADD	30120	9.72	6			3638	0.121	FIELD
091017 S. WESTBY	WESTBY		08/27/90	36N58E24DADD	89900	9.88	10.00	27.0	127	2071	0.023	LAB
091014 N. GOOSE	GOOSE	O	08/56/90	36N58E25DADC	70800	9.31	10.00	21.0	115	3638	0.051	LAB

SITE ID	LAKE	SAMPLE	SAMPLE DATE	LOCATION	SPECIFIC CONDUCTANCE umhos/cm	DH DH 0	DISSOLVED OXYGEN (DO) MG/L	DO TEMP C	DO AS PERCENT OF SATURATION	FIELD * CL MG/L	*CONTAMINATION INDEX	SAMPLE
091025 NW GOOSE	SOOSE		08/53/90	36N58E26CBAD	83300	9.61	4.30	23.2	55	9626	0.113	LAB
091072 N. GOOSE (GW)	300SE (GW)	ш	08/27/90	36N58E36CDCC	18960					595	0.031	FIELD
091070 N. GOOSE	SOOSE	۵	08/27/90	36N58E36CDDC	20940	9.31	71			3290	0.046	FIELD
091022 SALT	ĭ		08/28/90	37N56E01BCAC	29560	9.25	7.70	24.3	92	1521	0.026	LAB
091085 NO FREEZE	REEZE	,	08/28/90	37N57E01DCAC	81270	19				2516	0.031	FIELD
091084 N.WIDGEON SLGH	DGEON SLGH		08/28/90	37N57E07DADC	1510							FIELD
091019 LARSON SLOUGH	NO SLOUGH		08/27/90	37N57E15DCBC	1150	8.68	10.50	22.5	120	52	0.045	LAB
091023 LONE TREE	TREE		08/28/90	37N57E20DCAC	31680	8.96	11.60	23.8	137	1521	0.048	LAB
091083 WEED POND	POND		08/27/90	37N58E07DCCD	8130	8.42				1055	0.130	FIELD
091082 MIRROR	tor		08/27/90	37N58E18AADC	42310	9.11				2621	0.062	FIELD
091086 CHEVRON	RON		08/28/90	37N58E18CCBB	7230					1751	0.242	FIELD
091081 SCUM	_		08/27/90	37N58E20ADDC	42890	9.58	10			3638	0.085	FIELD
091020 RUSTY	Ł		08/28/90	37N58E30CCBB	27910	9.16	00.6	23.2	105	745	0.027	LAB
091080 FLAT	(dE_		08/27/90	37N58E33ADCC	7380	9.16				009	0.081	FIELD
091079 CURVE	Œ		08/27/90	37N58E34CCDD	5820	90.6				545	0.093	FIELD

* The contamination index refers to the ratio of chloride concentration to field SC. Field chloride concentrations are used in this ratio when available; when unavailable lab chloride concentrations are used.

100000 WITH FIELD SPECIFIC CONDUCTANCE SPECIFIC CONDUCTANCE (umhos/cm) 1 10000 10.00 8.40+ 1000 -09.6 9.40-9.00-8.80-8.60-9.20-10.20-9.80 10.40 -FIELD PH

COMPARISON OF FIELD PH

Comparison of the field pH values with log10 SC in water from selected Sheridan County lakes. Grouping of lakes based on SC are depicted by Roman numerals I through ${
m V.}$ Figure 2.

The results of the laboratory analyses of the lake water samples are summarized in Table 3. A comparison of the relationship between \log_{10} SC and \log_{10} calculated dissolved solids is relatively linear for lake water in Groups I and II. This relationship is shown in Figure 3. At these concentrations (0 - 10,000 umhos/cm) the field SC tends to be consistently higher than dissolved solids. The relationship changes to a nearly 1:1 ratio of field SC to dissolved solids for Groups III and IV (10,000 - 50,000 umhos/cm). The relationship breaks down for high salinity Group V waters (>50,000 umhos/cm) and field SC is significantly lower than dissolved solids.

LAKE WATER QUALITY

The five different categories of lakes based on SC makes a good starting point for a more detailed look at the types of dissolved constituents found in the lake water. Plots of major cations and anions can also be developed to produce graphical images that aid in identifying the dominant ions in a particular water sample. One such type of plot is the semi-logarithmic Schoellor diagram (Freeze and Cherry, 1979). Figure 4 depicts five Schoellor plots indicating the mean concentrations of major cations and anions by Group in the lakes sampled.

The shape of the curves maps the relative proportions of the dominant cations and anions within each category of lake water. In general, the shape of the curves are very similar for all 5 groups. The upward shift from Group I to Group V corresponds to the overall increase in dissolved minerals.

The concentration of a particular ion can be evaluated by noting the position it plots on the Y-axis of the Schoellor plot.

TABLE 3. WATER QUALITY OF SELECTED LAKES, SHERIDAN COUNTY, MONTANA (negative values are below detection limits)

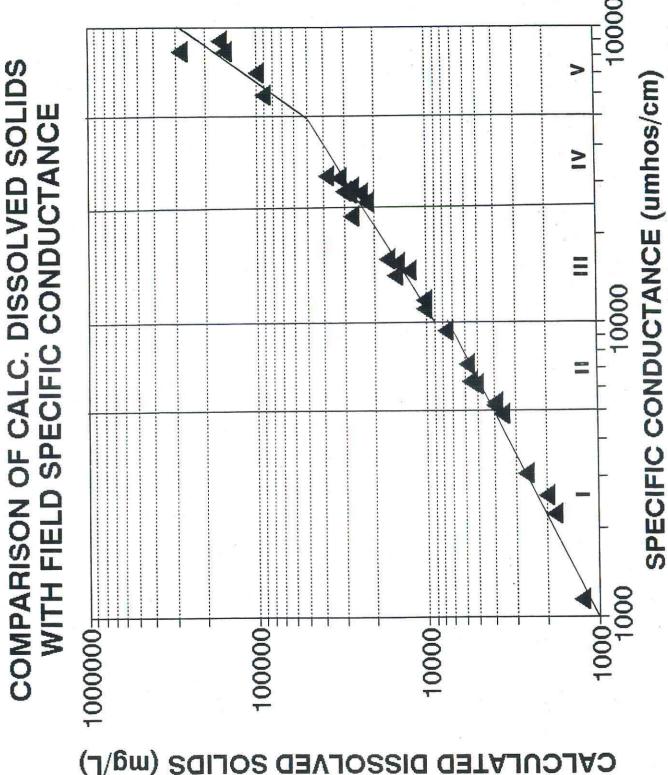
FLOU- RIDE (mg/L)	0.3	0.1	0.2	0.2	0.1	7.	0.2	0.0	0.0	1.0	0.1	0.1	0.1	7.0	2.8	10.5	0.1	0.3	7.0	0.2	0.1		4.	0.0	0.0	0.0	0.0	0.2	0.0	1.0
FI NITRATE R (mg/L) (m	0.32	1.08	0.17	0.25	0.07	0.10	0.10	0.07	0.07	0.10	0.07	0.07	1.19	0.07	90.0	0.20	0.82	0.38	-0.07	1.47	0.07	0.10	0.20	0.10	0.20	0.10	0.10	0.07	0.20	0.10
SULFATE NI (mg/L) (14800.0											
CHLORIDE (mg/L)																			1070.0						•					
CARBON- ATE (mg/L)	192.0	134.0	197.0	254.0	305.0	4030.0	3310.0	624.0	304.0	880.0	648.0	1420.0	137.0	122.0	0.49	54800.0	240.0	0.967	816.0	388.0	173.0	372.0	10400.0	3670.0	108.0	47.2	81.6	28.8	216.0	461.0
BICAR- BONATE (mg/L)	1240.0	1200.0	1464.0	1870.0	700.0	0.0696	7420.0	2420.0	1810.0	3760.0	2920.0	3580.0	1200.0	632.0	1599.0	71000.0	1601.0	3790.0	2740.0	2650.0	1120.0	979.0	15900.0	9300.0	309.0	212.0	502.0	349.0	2290.0	1880.0
SILICA (mg/L)	3.85	16.20	5.64	12.20	15.20	7.00	1.33	5.48	2.38	-0.10	2.52	4.45	6.84	13.00	6.40	22.00	0.00	12.50	3.27	3.36	3.19	6.89	15.00	-0.10	0.30	1.98	8.41	24.20	4.31	8.35
MANGANESE (mg/L)	0.012	-0.002	0.020	0.011	-0.002	-0.002	0.008	900.0	0.012	0.002	0.013	0.007	0.009	0.007	0.015	0.012	-0.002	0.010	0.060	0.018	0.005	0.012	0.008	0.003	0.015	0.016	0.033	0.053	0.010	0.011
IRON M (mg/L)	0.144	0.024	0.197	0.131	0.013	-0.004	0.027	0.037	0.122	0.049	0.028	-0.004	0.076	0.016	0.040	762.0	0.038	0.178	0.118	960.0	0.073	0.180	0.126	0.036	0.046	0.310	0.893	0.091	0.092	090.0
POTASSIUM (mg/L)	2 65	9.96	9.69	88.7	23.1	350.0	554.0	272.0	246.0	360.0	194.0	320.0	89.5	33.5	93.6	750.0	369.0	358.0	224.0	181.0	35.3	130.0	520.0	1050.0	704.0	322.0	459.0	35.0	714.0	163.0
SODIUM (mg/L)	1050.0	1030.0	1280.0	1190.0	430.0	0.0479	11240.0	2990.0	3780.0	7360.0	2220.0	4110.0	610.0	471.0	1379.0	108000.0	3410.0	0.0994	8840.0	3100.0	1800.0	7170.0	56000.0	32700.0	4,7000.0	5780.0	24400.0	219.0	8740.0	7950.0
MAGNESIUM (mg/L)	141.00	517.00	89.30	161.00	151,00	139.00	111,00	322.00	706.00	411.00	335.00	194.00	191.00	131.00	270.90	29.00	00.446	621.00	274.00	272.00	118.00	302.00	30.90	512.00	1600.00	1940.00	3000,00	59.70	2270.00	1280.00
CALCIUM (mg/L)	16.60	18.80	16.60	9.52	6.16	3.05	1.92	7.00	8.12	3.03	3.93	2.88	12.40	12.30	2.00	1.63	24.60	6.26	8.83	10.20	13.60	24.60	70 7	7.13	376.00	299.00	346.00	09.46	27.70	19.50
LAB	9000326	9000346	9000337	9000324	9000347	9000338	9000336	9000323	9000335	9000331	9000334	9000340	9000342	9000325	90P0107	9000365	9000367	9000366	9000327	9000344	9000328	9000333	9000343	9000330	9000345	9000332	9000348	9000339	90001329	9000341
DATE	08/56/80	08/54/90	08/53/90	08/25/90	08/54/90	08/54/90	08/54/90	08/23/90	08/56/90	08/24/90	08/54/90	08/54/90	08/25/90	08/25/90	08/06/80	08/30/90	08/31/90	09/03/90	08/27/90	08/26/90	08/26/90	08/26/90	08/27/90	08/56/90	08/26/90	08/29/90	08/28/90	08/27/90	08/28/90	08/28/90
SITE	91027	91003	91026	91006	91008	91005	91004	91001	91012	91002	91007	91009	91011	91010	91094	91021	91029	91028	91016	91013	91018	91015	91017	91014	91025	91024	01022	01010	01073	91020
LOCATION	31N56E09BBDD01	32N57E13DBDB01	32N57E32BBAB01		32N58E04DAAD01	32N58E09AAB01	32N58E17CDDB01	33N58E05AABC01	1000							33N59E24DDDA01	33N59F26BADD01	920				36N58E13AADD01								
LAKE NAME	MEDICINE	NUMBER 12	DEEP LAKE	SOUTH LONG	FRED	BERGER POND	KATY	CLEAR	BEER BOTTLE	HORSEFLY	BETTY	TERESA	NHOP	MALLARD POND	BRUSH	HORSESHOE	FAST LIHITE	UEST UNITE	S. GOOSE	BFFHIVE	GAULKE	FOOTBALL	S. WESTRY	N. GOOSE	NU GOOSE	MID CPACK	CAL T	SALI APCON CLONCE	LAKSON SLOOGN	RUSTY

TABLE 3. WATER QUALITY OF SELECTED LAKES, SHERIDAN COUNTY, MONTANA (continued) (negative values are below detection limits)

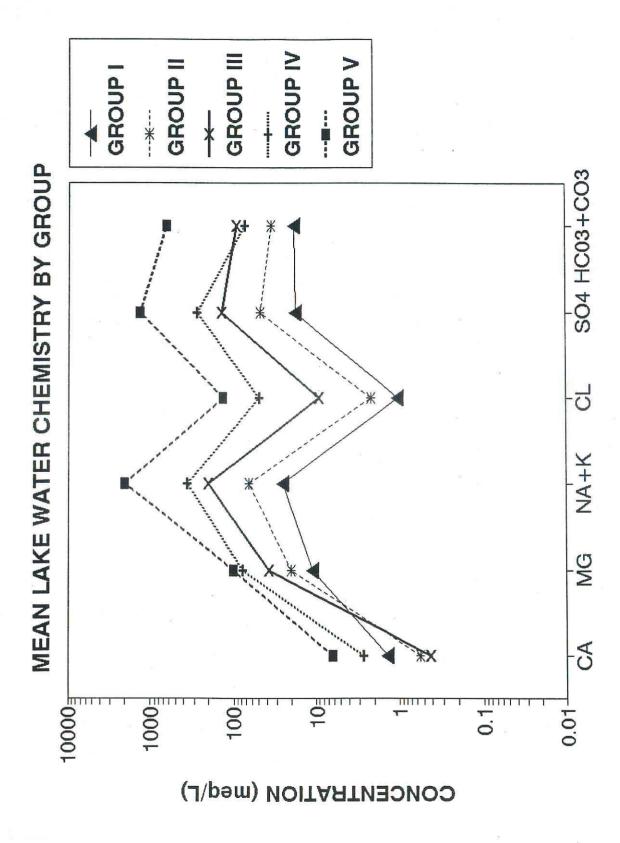
ORTHO-	PHOSPHATE	(mg/L)				0.00				0.00					- 3	0.00	0.00	22.50		-0.10						(#)						3	3
ō	ZIRCONIUM PHO	(mg/L)	-0.006	-0.006	-0.006	-0.006	-0.006	-0.006	0.007	900.0	0.007	0.013	0.009	-0.006	-0.006	-0.006	0.000	-0.114	-0.006	-0.006	-0.006	0.007	-0.006	-0.006	-0.006	-0.006	-0.006	-0.006	-0.00	-0.006	-0.006	-0.006	
	ZINC ZI	_	-0.006																		2					Ą.							
	VANAD I UM	(mg/L)	0.017	0.013	900.0	0.015	-0.004	-0.004	-0.004	0.012	0.009	0.005	0.008	-0.004	-0.004	0.015		-0.004	0.004	-0.004	0.008	-0.004	0.008	0.005	-0.004	-0.004	0.009	0.011	-0.004	-0.005	-0.004	-0.004	
SODIUM	ADSORPT I ON	RATE	18.32	9.61	27.54	19.76	7.41	176.03	227.64	35.61	30.41	07.77	25.92	63.01	9.29	8.59	17.87	4230.14	23.62	39.99	113.44	39.87	34.36	86.39	2079.91	308.67	235.78	26.92	92.38	4.34	39.20	77.47	
	LAB A	풉	67.6	8.87	9.35	9.41	10.02	9.74	9.78	9.74	9.05	67.5	9.34	62.6	20.6	9.52	9.11	10.01	8.94	9.23	62.6	9.11	6.43	09.6	9.95	9.38	00.6	8.72	8.86	8.89	9.41	90.6	
	FIELD	Hd	9.22	8.94	9.30	9.12	6.92	10.01	9.72	9.36	60.6	9.23	9.54	9.72	9.03	87.6	9.20	96.6	70.6	9.54	9.45	60.6	9.15	6.65	9.88	9.31	9.61	9.38	9.25	8.68	8.96	9.16	
LAB	CONDUCTIVITY	(mg/L)	4727.0	6525.0	5520.0	5396.0	2282.0	27769.0	32748.0	11374 0	15949.0	24490.0	9350.0	74957.0	3572.0	2684.0	5070.0	98083.0	14846.0	17952.0	27374.0	12579.0	7291.0	17952.0	60143.0	57987.0	61732.0	28872.0	50185.0	1746.0	33661.0	28791.0	
	FIELD	TEMPERATURE	21.4	25.3	21.5	22.0	25.0	25.5	26.5	23.3	16.7	19.0	21.0	24.7	25.2	21.9	22.2	25.8	18.6	20.4	19.9	18.4	19.6	20.0	23.0	20.8	22.5	22.4	23.2	22.5	24.8	22.9	
FIELD	CONDUCTIVITY	(mg/L)	4850	9300	5340	5200	2230	22990	31190	11900	15890	27800	9350	15120	3070	2590	6144	83400	14500	16420	28790	11150	7170	25860	89900	70800	83300	27690	29560	1150	31680	27910	
SUM OF	SOLIDS	(mg/L)	4225	6155	4710	2565	2157	31997	36163	11525	15762	25939	9259	14416	3223	2305	2944	318026	16157	18926	28777	11652	6383	23055	170341	104349	153897	27882	91528	1451	39682	31327	
CALCULATED DTSSOLVED	SOLIDS	(mg/L)	3596	5546	3967	3998	1802	27080	32399	10297	14843	24031	7777	12600	2614	1984	5131	282001	15345	17003	27386	10308	5815	22558	162274	99630	153741	27775	91273	1274	38520	30373	
		LAB NUMBER	9000326	9000346	9000337	9000324	9000347	9000338	9000336	9000323	9000335	9000331	9000334	9000340	9000342	9000325	9000107	9000365	9000367	9000366	9000327	9000344	9000328	9000333	9000343	9000330	9000345	9000332	8720000	9000339	9000329	9000341	
		LAKE NAME	MEDICINE	NUMBER 12	DEEP LAKE	SOUTH LONG	FRED	BERGER POND	KATY	CLEAR	BEER BOTTLE	HORSEFLY	BETTY	TERESA	NHOP	MALLARD POND	RRIISH	HORSESHOE	EAST WHITE	WEST WHITE	S. GOOSE	BEEHIVE	GAULKE	FOOTBALL	S. WESTBY	N. GOOSE	NIL GOOSE	MID CRACK	CALT	I ADSON STOUGH	I ONF TRFF	RUSTY	
																		10)														

TABLE 3. WATER QUALITY OF SELECTED LAKES, SHERIDAN COUNTY, MONTANA (continued) (negative values are below detection limits)

LAB ALU		-		CADIMUM	CHROMIUM	COPPER	_	MOL YBDENUM		BROMIDE	NICKEL	LEAD S	SELENIUM	STRONTIUM	TITANIUM
(mg/L) (mg/L)	?	(mg/L)	(mg/L)	(mg/L)	(mg/L)	(mg/L)	(mg/L)	(mg/L)	(mg/L)	(mg/L)	(mg/L)	(mg/L)	(mg/L)	(mg/L)	(mg/L)
	104	0.035		-0.005	-0.005	0.017	0.149	-0.04	0.05	09.0	-0.02	-0.05	-0.001	0.44	-0.004
-0.004		0.050	1.47	-0.005	-0.005	-0.004	0.580	+0.0-	0.03	0.29	0.02	0.05	-0.001	0.35	-0.004
-0.004		.046		-0.005	-0.005	0.013	0.140	-0.04	0.04	1.70	-0.02	-0.05	-0.008	0.37	-0.004
		.022		-0.005	900"0	600.0	0,370	*0.0-	0.02	1.60	-0.02	-0.05	0.007	0.16	-0.004
-0.004		.004		-0.005	-0.005	-0.004	0.175	*0.0-	0.01	-0,10	-0.02	-0.05	-0.001	0.02	-0.004
-0.004		045		-0.005	-0.005	0.004	1.260	-0.04	-0.01	16.30	-0.02	-0.05	-0.001	0.03	-0.004
		164		-0.005	-0.005	0.017	1.700	+0.0-	0.01	19.70	-0.02	-0.05	0.001	0.08	-0.004
-0.004		034		-0.005	0.007	-0.004	1.380	-0.04	-0.01	1.70	-0.02	-0.05	-0.001	0.01	-0.004
-0.004		990		900.0	-0.005	-0.004	2.110	-0.04	0.01	2.90	0.02	0.13	-0.001	0.16	-0.004
-0.004		015		-0.005	-0.005	-0.004	2.370	+0.0-	0.01	09.9	-0.02	0.08	-0.001	0.14	-0.004
-0.004		033		-0.005	-0.005	0.005	0.940	*0.0-	0.01	1.10	-0.02	-0.05	-0.001	0.01	-0.004
-0.004	2.0	940		-0.005	-0.005	0.024	1.120	-0.04	0.01	1.15	-0.02	0.02	-0.001	0.03	-0.004
0.042)23		-0.005	-0.005	-0.004	0.346	-0.04	0.03	-0.10	-0.02	-0.05	-0.001	0.18	-0.004
		Ξ		-0.005	-0.005	-0.004	0,201	-0.04	0.01	-0.10	-0.05	-0.05	0.008	0.08	-0.004
000.0		8		000.0	000.0	000.0	0.200	0.00	0.00	1.23	0.00	0.00	0.000	0.20	0000
-0.004		8		-0.005	-0.005	-0.004	1.280	0.07	0.08	170.00	-0.02	-0.05	0.001	0.20	0.052
-0.004		-		0.011	-0.005	900.0	2.180	-0.04	0.03	4.30	-0.02	0.13	-0.001	0.21	-0.004
-0.004		5		900.0	-0.005	0.005	2.070	-0.04	0.02	3.60	0.02	0.05	-0.001	0.15	-0.004
-0.004		5		-0.005	-0.005	0.029	1,210	-0.04	0.01	2.90	-0.02	90.0	-0.001	0.03	-0.004
-0.004		23		0.005	900.0	0.036	0.720	-0.04	0.02	0.70	-0.02	0.05	-0.001	0.04	-0.004
*************************************		13		-0.005	-0.005	-0.004	0.425	-0.04	0.01	1.10	-0.05	-0.05	900.0	0.18	-0.004
-0.004		120		-0.005	-0.005	-0.004	0.635	-0.04	0.05	52,60	-0.02	0.08	-0.001	0.34	-0.004
0.040		225		-0.005	-0.005	0.106	1,250	-0.04	-0.01	76.00	-0.02	-0.05	0.001	0.04	-0.004
-0.004		147		-0.005	-0.005	-0.004	5.190	-0.04	0.01	48.70	-0.05	-0.05	-0.001	-0.09	-0.004
0.067		020		0.007	-0.005	0.050	8.320	-0.04	0.04	35.10	-0.05	-0.05	0.001	6.31	0.016
		017		0.010	-0.005	0.032	1.650	0.20	0.05	6.08	0.02	0.36	0.002	4.70	0.017
0.032		0.039		0.026	-0.005	0.024	3.190	-0.04	0.02	13.40	0.07	0.34	0.002	7.89	0.025
0.026		0.014		-0.005	-0.005	-0.004	0.050	-0.04	0.05	0.10	-0.05	-0.05	-0.001	0,40	-0.005
700-0-	_	.088		0.015	-0.005	0.115	3,360	-0.04	0.03	18.00	0.05	0.33	-0.009	1.26	-0.004
0.04 -0.004 0.	_	3.164	6.01	-0.005	-0.005	-0.004	1,500	-0.04	0.02	8.90	-0.02	0.18	0.001	0.14	-0.004



Sheridan County lakes showing the change in this relationship as lake water becomes more Grouping of lakes based on SC are depicted by Roman numerals I through V. Comparison of the log calculated dissolved solids with log10 SC in water from selected mineralized. Figure 3.



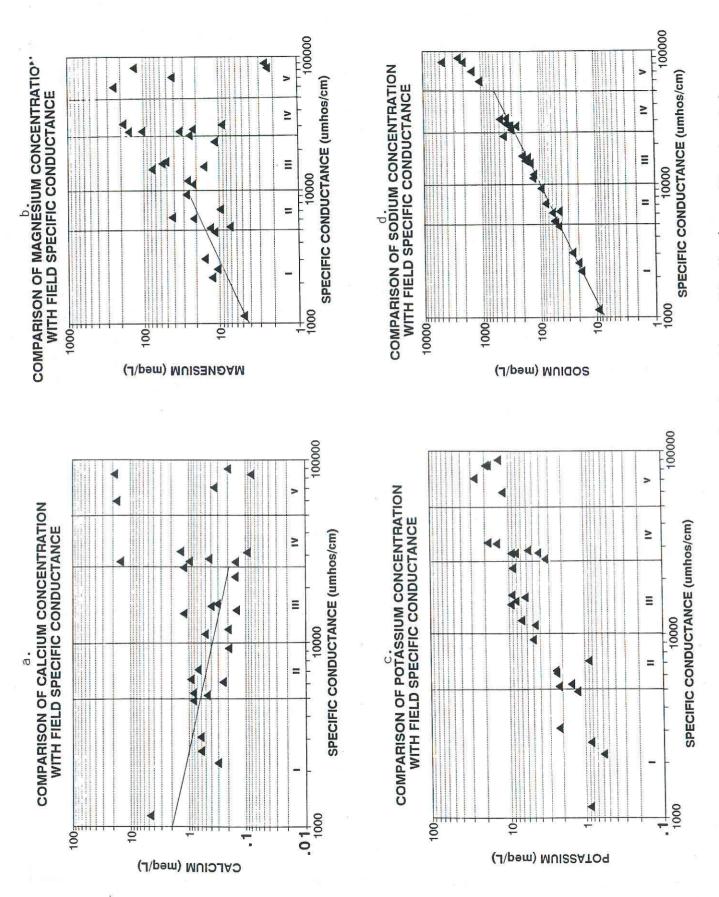
Semilogarithmic plots indicating the mean concentrations of major cations and anions by Group in the lakes sampled. Figure 4.

The relationship between different ions can be compared by observing the slope of a line connecting the mean concentrations for each ion. Sodium and sulfate are the dominant cation and anion in each of the five different water groups. Bicarbonate plus carbonate ions are proportionally only slightly less concentrated than sulfate ions in Group I, Group II, and Group III lake waters, but are in relatively low concentrations in Group IV and Group V lake waters. The concentrations of magnesium are intermediate and in similar proportions in all five water groups. Calcium and chloride ions are all in relatively low concentrations in all five water groups. The ranking of cation concentrations is NA > MG > CA and for anion concentrations is SO4 > HCO3 + CO3 > CL for all five water groups.

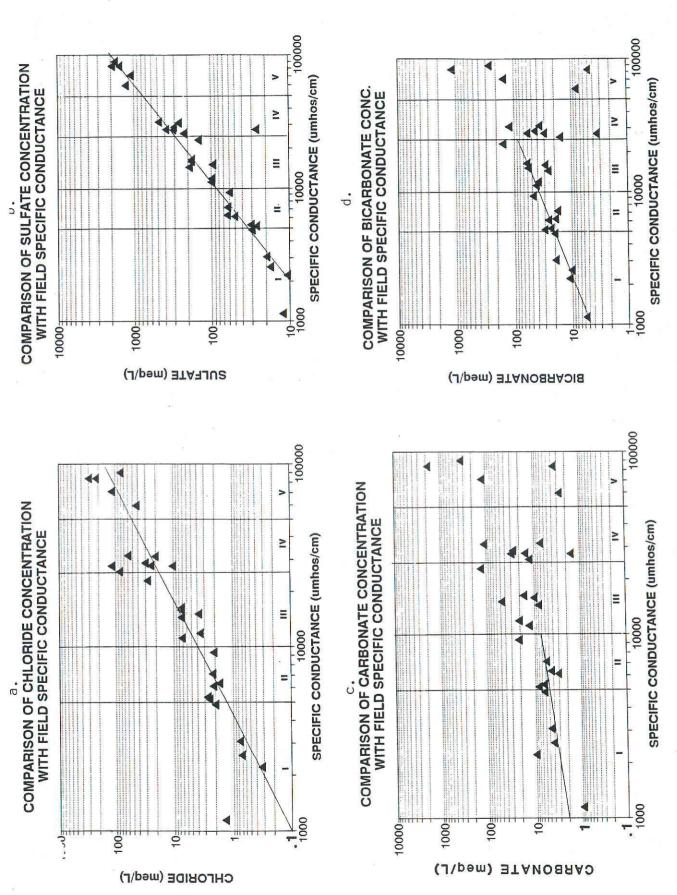
A series of plots showing the relationship of the \log_{10} concentrations of the major cations and anions with the \log_{10} SC are displayed in Figures 5 and 6. With SC plotted along the x-axis the distribution of the ionic concentration by the associated lake Group is apparent.

Calcium concentrations vary most in both the freshest (Group I) and most mineralized (Group IV and V) lakes (Figure 5a). A best fit line projected through Group I, Group II, and Group III data points indicate a decreasing trend in dissolved calcium up to an SC of 25,000 umhos/cm. The trend breaks down in highly concentrated water of Group IV and V lakes. This trend probably relates to the deposition of marl on the lake bottoms as the waters become supersaturated with calcium and bicarbonate ions.

The variability of magnesium concentrations in lake water increases as the water becomes more mineralized (Figure 5b).



Grouping of lakes based on SC are depicted by Comparison of the \log_{10} concentrations of major cations with the \log_{10} SC; a) calcium, b) magnesium, c) potassium, and d) sodium. Roman numerals | through V. 5 Figure



Grouping of lakes based on SC are depicted Comparison of the log10 concentrations of major anions with the log10 SC; a) chloride, b) sulfate, c) carbonate, and d) bicarbonate. by Roman numerals I through ${\sf V}.$ Figure 6.

A slight increasing trend of magnesium concentration with SC shown by the best fit line can be observed in Group I and Group II waters. The increased variability of magnesium concentration within Group III, Group IV, and Group V water masks any further trend. There is less of tendency for magnesium depletion due to carbonate precipitation than there was for calcium.

The log₁₀ relationship between potassium concentration and SC is nearly linear, and it is similar to the relationship between sodium and SC (Figure 5c). However, the concentration of potassium is about one order of magnitude less than sodium in the freshest waters and about 2 orders of magnitude less than sodium in the most mineralized waters. Similar processes for potassium and for sodium result in the observed relationships including availability of potassium in the soils, solubility of potassium, and increased concentration of potassium caused by the arid climate.

The log₁₀ relationship between sodium concentration and SC is approximately linear, with sodium steadily increasing as the water becomes more mineralized (Figure 5d). The steady increase in sodium reflects the accumulation of sodium salts in the soils of the region and the high solubilities of the sodium salts. Both abundance of sodium in the glacial deposits and the aridity of the climate influence the accumulation of sodium in the areas soils and eventually in the lakes.

The log₁₀ relationship between chloride concentration and SC is approximately linear with chloride steadily increasing as the water becomes more mineralized (Figure 6a). The variability of the data points is largest in Group IV lake water. Chloride salts tend to be a minor component of near surface sediments and soils

in this region. Data points plotting significantly above the best fit trend line may indicate water contaminated by chloride-rich brines. The general trend of increasing chloride concentration as the lakes become more mineralized probably is the result of evaporation concentrating the very soluble chloride salts in lakes with limited flow.

The log₁₀ relationship between sulfate concentration and SC is approximately linear with sulfate steadily increasing as the water becomes more mineralized (Figure 6b). The steady increase in sulfate mirrors the sodium increases shown in Figure 5c. Sulfate is derived from the oxidation of pyrite and other sulfide minerals in the near surface sediments. It commonly accumulates as very soluble sodium sulfate salts resulting in the similarity between increases in sodium and sulfate concentrations as the lake water becomes more mineralized.

The log₁₀ relationship between carbonate concentration and SC is again approximately linear but the linear trend becomes less evident above conductivities of 10,000 umhos/cm (Figure 6c). Similar to bicarbonate, the carbonate ions are depleted in the lake water as a result of marl deposition.

The log₁₀ relationship between bicarbonate concentration and SC is approximately linear up to concentrations of 25,000 umhos/cm (Group I through Group III water) (Figure 6d). Below an SC of 25,000 umhos/cm the bicarbonate concentration increases as the water becomes more mineralized. Above an SC 25,000 umhos/cm the variability of bicarbonate concentration masks any linear trend. As lakes become supersaturated with bicarbonate ions, calcium carbonate marls are deposited depleting both of these ions from the

dissolved minerals in the lake water.

Dissolved trace constituents in the sampled lakes generally fall into 3 categories based on trends of their concentrations. found in very elements are majority of these concentrations, usually below detection limits. Constituents fitting into this category include silver, cadmium, chromium, molybdenum, nickel, selenium, titanium, zinc, and zirconium. Other constituents including aluminum, barium, copper, lead, strontium, and vanadium were found in a relatively wide range of concentrations (Figure 7). Comparisons of the log10 concentrations of these constituents with \log_{10} SC do not show any well developed trends. The final group of trace constituents includes arsenic, boron, bromide, and lithium (Figure 8). Comparing the log10 SC shows a linear trend of increasing concentration with increasing mineralization of lake water.

WATER QUALITY CHANGES DURING THE 1980'S

A lake water quality survey was conducted in September 1984 as part of hydrogeologic investigations in northeastern Montana (Donovan, 1988). The survey consisted of SC measurements from lakes in the Medicine Lake swale and the Clear Lake swale between Westby and Medicine Lake. Many of these same lakes were again sampled in August 1990, resulting in the water quality comparisons listed in Table 4. A total of 22 of the originally surveyed lakes were re-sampled. Where several field samples were collected in different parts of the larger lakes, the mean SC values and standard deviations were compiled for the comparison. Smaller lakes typically only had one measurement. Based on SC, lake water quality deteriorated in 20 of the 22 lakes surveyed.

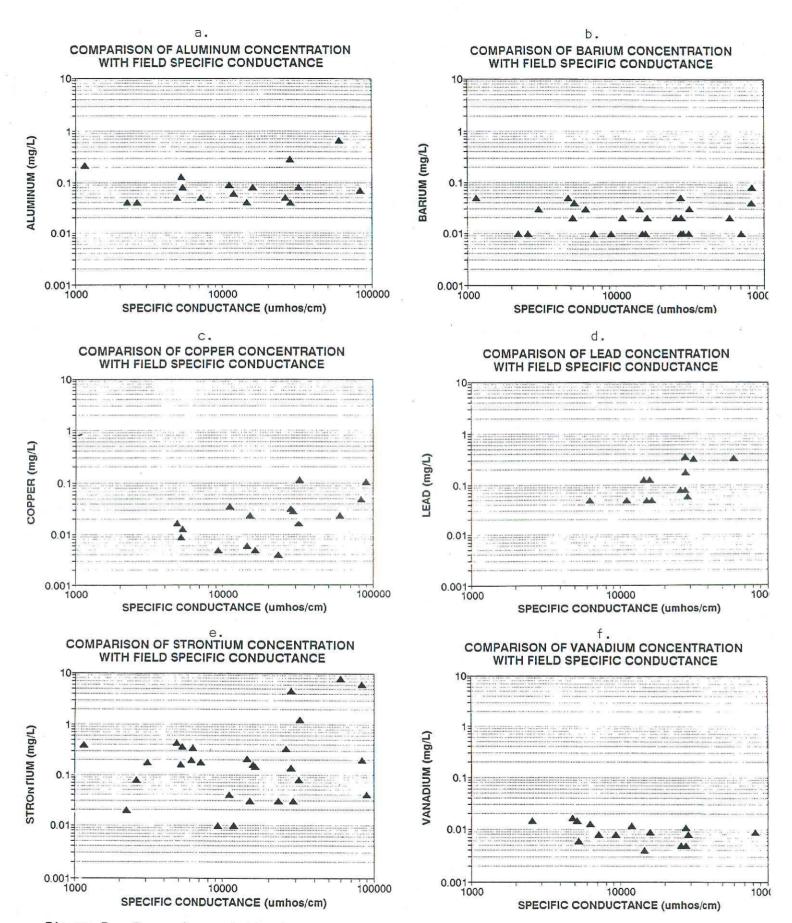


Figure 7. Comparison of the \log_{10} concentrations of a) aluminum, b) barium, c) copper, d) lead, e) strontium, and f) vanadium with \log_{10} SC.

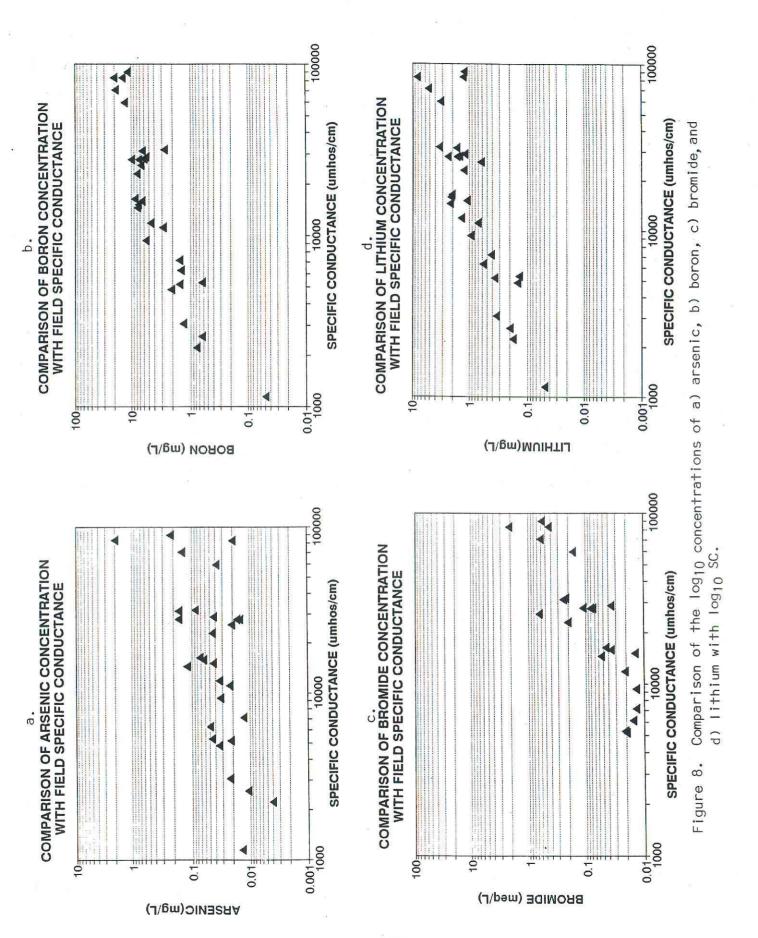


TABLE 4. LAKE WATER QUALITY COMPARISON 1984-1990

	120		10110101		september 1704 survey	ĺ
			South Westby Lake T. 36 N., R. 58 E., Sec. 24 DADD	12 13	а — A	
Sample Point	SC (nmhos)	Similar 1984 Sample Point		Sample Point	(solun) 3S*	
⋖	89,900	2		- 0 E	56,092 52,650 59,032	
					55,920 3,194	
		Change in SC 1984 Percent change	ange in SC 1984 - 1990 = +33,980 umhos Percent change in SC = 61% increase		2	1
		1. 36	Northwest Goose Lake S N., R. 58 E., Sec. 26 CBAD			
Sample Point	SC (numhos)	Similar 1984 Sample Point	N con	Sample Point	Sc (umhos)	
V	83,300			Dry		
		1, 3	Mudcrack Lake 36 N., R. 58 E., Sec. 34 B		^	
Sample Point	SC (umhos)	Similar 1984 Sample Point		Sample Point	SC (umhos)	
A	27,690			τ	56,200	
		Change in Percent	Change in SC 1984 - 1990 = -28,510 umhos Percent change in SC = 51% decrease		¥	1
		T. 35 N.,	Goose Lake 5 N., R. 58 E., Sec., 1, 11, 12, 13 T. 36 N., R.58 E., Sec. 25, 36			
Sample Point	(soumn) as	Similar 1984 Sample Point		Sample Point	SC (umhos)	
A South	21,000	2		1 North	31,450	
B South	28,790	2,3			29,670	
B South C North	70,800	, -		4 South	27,890	
D North F South	70,940	5 7			28,190	

TABLE 4. LAKE WATER QUALITY COMPARISON 1984-1990 (continued)

	ראכפרות	for the control of th
Combined $\overline{X} = 48,270 \text{ sd} = 24,370$ North Lake $\overline{X} = 70,870 \text{ sd} = 100$ South Lake $X = 36,970 \text{ sd} = 21,890$	Goose Lake (Continued) 70	Combined $\bar{X} = 27,000$ sd = 4,510 North Lake $\underline{X} = 29,080$ sd = 3,360 South Lake $X = 26,140$ sd = 4,970
	Combined lake change in SC 1984 - 1990 = +21,270 umhos Combined lake percent change in SC = 78% increase North lake change in SC 1984 - 1990 = +41,790 umhos	,
	South take change in SC 1984 - 1990 = +10,830 umhos South take percent change in SC = 41% increase	
	Beehive Lake T. 35 N., R. 58 E., Sec. 13 AADC	
Sample Point SC (umhos)	Similar 1984 Sample Point	Sample Point SC (umhos)
A 11,150	2	1 4,790 2 4,980
		$\bar{X} = 4,885$ sd = 134
	Change in SC 1984 - 1990 = +6,265 umhos Percent change in SC = 128% increase	
	Clear Lake T. 34 N., R. 58 E., Sec. 32, 33 T. 33 N., R. 58 E., Sec. 4, 5	
Sample Point SC (umhos)	Similar 1984 Sample Point	Sample Point SC (umhos)
A 8,690 B 11,900 C 12,020	1 2 3 3 3	1 5,890 2 6,090 3 6,390
$\frac{x}{x} = 10,870$ sd = 1,890	Change in SC 1984 - 1990 = +4,750 umhos Percent change in SC = 78% increase	$\frac{X}{X} = 6,120$ $sd = 252$

TABLE 4. LAKE WATER QUALITY COMPARISON 1984-1990 (continued)

August 1990 aur vey	Irvey				for the court of t
			Beer Bottle Lake T. 35 N., R 58 E., Sec. 22 AB	e e	
Sample Point	SC (umhos)	Similar 1984 Sample Point	2	Sample Point	SC (nmhos)
A	15,890	-			10,940
	8		Change in SC 1984 - 1990 = +4855 umhos Percent change in SC = 44% increase	ıı ıı	11,035
			Brush Lake T. 35 N., R. 58 E., Sec. 22		
Sample Point	SC (umhos)	Similar 1984 Sample Point		Sample Point	SC (umhos)
V	6,140	5		F ⊗ 8	5,810 5,660 5,360
			*	I II I	5,610
		191	Change in SC 1984 - 1990 = $+530$ umhos Percent change in SC = 9% increase		
			Horseshoe Lake T. 33 N., R. 58 E., Sec. 2 24 DDDA		
Sample Point	SC (umhos)	Similar 1984 Sample Point	ı	Sample Point	SC (umhos)
A (8/23/90) A (8/30/90)	72,750			F 2	63,216
2 = ps	7,530	o	Change in SC 1984 - 1990 = +14,720 umhos Percent change in SC = 23% increase	и и I× ps	63,360 200
			East White Lake T. 33 N., 58 E., 26 BADD		
Sample Point	SC (umhos)	Similar 1984 Sample Point	1	Sample Point	SC (umhos)
۷	14,500	-		2 2	10,960 10,960
		3	Change in SC 1984 - 1990 = $+3,540$ umhos Percent change in SC = 32% increase		

TABLE 4. LAKE WATER QUALITY COMPARISON 1984-1990 (continued)

August 1990 Survey	rvey	Lake Name Location	September 1984 Survey
		Horsefly Lake T. 33 N., R. 58 E., Sec. 27 AABB	8
Sample Point	SC (umhos)	Similar 1984 Sample Point	Sample Point SC (umhos)
A	27,800		1 34,390
		Change in SC 1984 - 1990 = -6.590 umhos Percent change in SC = 19% decrease	
		West White Lake T. 33 N., R. 58 E., Sec. 27 ACBC	
Sample Point	SC (umhos)	Similar 1984 Sample Point	Sample Point SC (umhos)
V	16,420		1 7,150 2 6,940
			$\frac{X}{X} = 7,045$ $sd = 148$
		Change in SC 1984 - 1990 = +9,375 umhos Percent change in SC = 133% increase	
		Betty Lake T. 33 N., R. 58 E., Sec. 27 CBCC	
Sample Point	SC (umhos)	Similar 1984 Sample Point	Sample Point SC (umhos)
A (8/22/90) A (8/24/90) B C	9,670 9,350 8,990 9,660		1 4,600
	9,420 320		
		Change in SC 1984 - 1990 = +4,820 umhos Percent change in SC = 105% increase	
:-6			

TABLE 4. LAKE WATER QUALITY COMPARISON 1984-1990 (continued)

August 1990 Survey		Lake Name Location	September 1984 Survey
		Teresa Lake T. 33 N., R. 58 E., Sec. 28 C	
Sample Point SC (umhos)	hos) Similar 1984 Sample Point		Sample Point SC (umhos)
A(8/22/90) 16,190 A(8/24/90) 15,120 B 15,990	2 - 1 - 2		1 7,510 2 6,980
$\frac{x}{x} = 15,770$ sd = 569			$\frac{X}{X} = 7,250$ $sd = 375$
	Ch.	Change in SC 1984 - 1990 = $+8,520$ umhos Percent change in SC = 118% increase	
		John Lake T. 33 N., 58 E., Sec. 29 D	
Sample Point SC (umhos)	hos) Similar 1984 Sample Point		Sample Point SC (umhos)
A 3,070 B 3,260	- 2		1 2,555 2 2,767
$\frac{x}{x} = \frac{3}{165}$			$\bar{x} = 2,660$ sd = 150
	5	Change in SC 1984 - 1990 = +505 umhos Percent change in SC = 19% increase	
*		Mallard Pond T. 33 N., R. 58 E., Sec. 33 B	
Sample Point SC (umhos)	hos) Similar 1984 Sample Point		Sample Point SC (umhos)
A 1,475 B 2,610 C 2,590			Not Sampled <2,000 (Est)
$\overline{X} = 2,220$ $sd = 650$			
		Black Water Lake T. 35 N., R. 58 E., Sec. 33 C	V
Sample Point SC (umhos)	hos) Similar 1984 Sample Point		Sample Point SC (umhos)
88 700			Not sampled DRY

TABLE 4. LAKE WATER QUALITY COMPARISON 1984-1990 (continued)

Sample Point SC (umhos) Similar 1 A 5,360 B 5,370 C 5,440		×
SC (umhos) 5,360 5,370 5,440	Long Lake T. 33 N., R. 58 E., Sec. 32 T. 32 N., R. 58 E., Sec. 2, 3	
	Similar 1984 Sample Point	Sample Point SC (umhos)
	3 2 2	1 4,070 2 4,390 3 4,240
$\frac{x}{sd} = \frac{5,340}{101}$		$\frac{x}{X} = 4,230$ $sd = 160$
	Change in SC 1984 - 1990 = +1,110 umhos Percent change in SC = 26% increase	
	Fred Lake T. 32 N., R. 58 E., Sec. 4 DA	
Sample Point SC (umhos) Similar 1	Similar 1984 Sample Point	Sample Point SC (umhos)
A(8/22/90) 2,300 A(8/24/90) 2,230 B 2,260 C 2,190 D 2,280	1 2	1 1,790 2 1,770
$\bar{x} = 2,250$ sd = 43		$\frac{x}{x} = 1,780$ $sd = 14$
	Change in SC 1984 - 1990 = +470 umhos Percent change in SC = 26% increase	
	Berger Pond T, 32 N., R. 58 E., Sec. 9 A and 10 B	
Sample Point SC (umhos) Similar 1	Similar 1984 Sample Point	Sample Point SC (umhos)
A 24,440 B 22,990	3	1 1,725 2 18,080 3 17,900
$\bar{X} = 23,720$ sd = 1,025		$\frac{X}{X} = 17,740$ $sd = 437$
	Change in SC 1984 - 1990 = $+5,980$ umhos Percent change in SC = 34% increase	

August 1990 Survey		Lake Name Location	September 1984 Survey
		No. 12 Lake T. 32 N., R. 57 E., Sec. 4, 13 T. 32 N., R. 58 E., Sec. 8, 9, 18	
Sample Point SC (umhos)			Sample Point SC (umhos)
A 5,100 B (8/22/90) 4,210 C 5,190 C 5,190 E 6,300 F 6,300 F 7,50 H 5,910	- 285		1 2,230 2 3,180 3 2,390 4 3,014 5 3,540
$\bar{x} = 5,440$ sd = 700			$\frac{X}{X} = 2,870$ $sd = 549$
		Change in SC 1984 - 1990 = $+2.570$ umhos Percent change in SC = 90% increase	
		Katy Lake T. 32 N., R. 58 E., Sec. 17	
Sample Point SC (umhos)	os) Similar 1984 Sample Point	int	Sample Point SC (umhos)
A (8/22/90) 20,000 A (8/24/90) 31,190	::1 		1 12,810 2 12,620
$\bar{x} = 25,600$ sd = 7,910			$\frac{X}{x} = 12,720$ sd = 134
		Change in SC 1984 - 1990 = +12,880 umhos Percent change in SC = 101% increase	
		Deep Lake T. 32 N., R. 57 E., Sec. 32 B	
Sample Point SC (umhos)	nos) Similar 1984 Sample Point	int	Sample Point SC (umhos)
1 5,340	-		
			x = 4,540
		Change in SC 1984 - 1990 = +800 umhos Percent change in SC = 18% increase	3
			The variety of the control of the co

TABLE 4. LAKE WATER QUALITY COMPARISON 1984-1990 (continued)

September 1984 Survey	Sample Point SC (umhos)	1 2,990 2 3,100 3 3,232 4 3,075	$\overline{X} = 3,100$ $sd = 100$			Sample Point SC (umhos)	1 2,290 2 2,354 3 1,782 4 2,227 5 3,053 6 1,678 7 1,637 9 1,590 9 1,590
Lake Name Location	Gaffney Lake T. 31 N., R. 57 E., Sec. 3, 4 T. 32 N., R. 57 E., Sec. 32, 33			Change in SC 1984 - 1990 = +1,770 umhos Percent change in SC = 57% increase	Medicine Lake T. 31 N., R. 56, 57 E. T. 32 N., R. 56, 57 E.		version)
	Similar 1984 Sample Point	4			= n	Similar 1984 Sample Point	2 1(Big Muddy Diversion) 1 9 8 4
rvey	SC (umhos)	4,870			×	SC (umhos)	4,690 650 4,180 4,420 4,580 4,760 4,320 4,670
August 1990 Survey	Sample Point	±	×			Sample Point	₹ ⊞ ∪ □ Ш ⊩ ∪ ¬

 $\bar{X} = 4,030$ sd = 1,380

= 2,060

sd ×1

Change in SC 1984 - 1990 = +1,970 umhos Percent change in SC = 96% increase

^{*} SC refers to the specific conductance of a lake water sample

^{**} $\overline{\chi}$ refers to the mean (average) SC of water samples from a specific lake

^{***} sd refers to the Standard Deviation (dispersion about the mean) of the accumulated SC data

The percent of water quality change ranged from +19 percent to +96 percent in Group I lakes; from +9 percent to +105 percent in Group II lakes; from +32 percent to +128 percent in Group III lakes; and from +23 percent to +61 percent in Group V lakes. The only lakes showing water quality improvement were two Group IV lakes where SC decreased 19 percent and 51 percent respectively while two of the Group IV lakes had SC increases of 23 percent and 61 percent respectively. In summary , no clear trends of water quality changes are apparent based on lake water quality category, lake size, or geographic position of the lakes.

likely cause of the overall water The most deterioration is the result of the prevailing drought conditions during the 1980's. Reduced precipitation produced less direct runoff into the lakes and less recharge into the ground water system feeding into the lakes. The drought was accompanied by high evaporation rates further concentrating dissolved constituents The decrease in lake volume was not into the lake water. accompanied by a uniform decrease in the mass of dissolved minerals. While decreased rainfall and runoff would also decrease the mass of dissolved constituents brought into the lakes, most of the dissolved minerals (largely sodium sulfate salts) are highly soluble and do not precipitate until highly concentrated brines have formed. As a result, the SC and calculated dissolved solids have in general, increased during the 1980's. Measurements at Medicine Lake by the U.S. Fish and Wildlife Service indicate a change in the lake water level from 1.28 feet below desired management level to 3.99 feet below desired management level between September 1984 and August 1990, confirming the impacts of the drought.

The two lakes showing water quality improvements were Mudcrack Lake and Horsefly Lake. The cause of the decrease in salinity is unclear. Local variabilities in where the samples were collected may have caused the anomalous freshening of the lake waters. Mudcrack Lake appears to have been impacted by oil-field brines and perhaps some of the sodium chloride salts have been removed from the basin since the first sample was collected. Possible methods of salt removal include flushing and dilution by rainfall and runoff, or by the wind blowing salt crystals out of the basin when the lake was dry.

EVIDENCE OF CONTAMINATION

While the intent of compiling background water quality samples was to identify baseline conditions in the region, existing uses may have already impacted lake water quality. Two likely activities that may have potentially impacted the water resources are farming and oil field development.

Evidence of farming impacts to water quality were not apparent based on the results of these water analyses. High levels of nitrates and phosphates would be the most obvious impacts if large quantities of animal wastes or fertilizers had been flushed into these lakes. Relatively low concentrations of nitrates and phosphates were identified in all of the lakes sampled. The high concentrations of dissolved oxygen in many of the lakes are probably the result of high algal activity and photosynthesis. It is possible that nutrients may be increasing the algal production in some of the lakes. But these lakes are probably naturally

entraphic oligitPophic, and impacts from agriculture are probably insignificant.

There is stronger evidence for impacts to water quality from oil field wastes. Large volumes of highly concentrated sodium chloride brines have been pumped to the land surface since the early 1960's. The brines are associated with oil producing zones and are produced along with the oil. Shallow ground water contamination by sodium chloride salts has been identified in the vicinity of many of these lakes (Reiten and Tischmak, in press). Occasionally chloride levels increase along with dissolved solids and SC due to evaporation in surface-water bodies. To distinguish between naturally-elevated chloride concentrations and those elevated by impacts of oil field brines, a contamination index (CI) was designed. For field applications, this index is defined as the ratio of field chloride concentration to field SC. Previous work in eastern Sheridan County developed an empirical lower limit indicating brine impacts at CI = 0.035. Based on this limit and the data compiled in Table 2 twenty-five lakes appear to have been impacted by brines. Although valid over wide ranges of water quality, the index tends to be less effective for predicting contamination in highly mineralized waters. In highly mineralized waters, SC loses its linear relationship with calculated dissolved solids and typically under predicts the concentration of dissolved minerals. Under these conditions the contamination index will improperly indicate contamination.

A more accurate index can be developed when complete water analyses are run on the lake water samples. Comparing the percent millequivalents of chloride of the total anions from different

lakes can also indicate the relative magnitude of brine impacts. The empirical limit of percent millequivalents chloride set as an indicator of brine contamination is 5.5%. Using this limit, brine impacted lakes and associated percentage of chloride ions by lake are Mudcrack Lake- 27%, Football Lake - 25.2%, Northwest Goose Lake - 13.2%, Lone Tree Lake - 10.1%, North Goose Lake - 7.8%, South Goose Lake - 7.4%, Larson Slough - 6.9%, and Berger Pond - 5.9%.

Impacts to Mudcrack Lake, Northwest Goose Lake, North Goose Lake, and South Goose Lake all appear to be related to sources of contamination in the Goose Lake oilfield that were identified in a previous study (Reiten and Tischmak, in press). Environmentally unsound methods of brine disposal that were commonly used at these sites prior to 1975 are the main causes of the existing contamination. The use of evaporation pits, trenching of reserve pits, and pipeline leaks all have contributed to the brine contamination. It is unlikely that the large extent and high level of surface-water and ground-water contamination existing in the Goose Lake Field could be derived from these relatively small volume sources of brine. Much of the existing contamination appears to confirm landowner reports that large volumes of brine were disposed of by simply allowing the brine to flow onto the ground surface. Similar brine disposal methods probably account for the contamination of other impacted lakes in the region.

CONCLUSIONS

Lake water in eastern Sheridan County covers a range from fresh to highly mineralized. The lakes were classified on the basis of specific conductance (SC) into 5 groups; Group I 0-5,000

umhos/cm, Group II 5,000-10,000 umhos/cm, Group III 10,000-25,000 umhos/cm, Group IV 25,000 - 50,000 umhos/cm and Group V >50,000 umhos/cm. In spite of this diversity in water quality, the water in these lakes is relatively uniform in proportions of major cations and anions. Sodium and sulfate are the dominant ions in the regions lake water. This dominance is probably the result of the availability of these constituents in near surface geologic sediments and high solubilities of sodium sulfate salts. The aridity of this region also accounts for concentrating these salts in the soil and lake water. The ranking of cation concentrations is NA > MG > CA and for anion concentrations is $SO_4 > HCO_3 + CO_3 > CL$ for all five water groups.

Lake water quality in general declined from September 1984 to August 1990 based on conductivity surveys conducted on 22 lakes between Westby and Medicine Lake. No clear trends of water quality changes were apparent based on lake water quality category, lake size, or geographic position of the lakes. Drought conditions during the 1980's appears to have decreased the volume of water in the lakes while the mass of dissolved solids remained relatively stable causing the increase in SC.

The water quality information does not indicate any negative impacts to lake water from agricultural activities. In contrast, the levels of chloride concentrations in several of the lakes are probably the result of improper disposal of oil field brines. The use of evaporation pits, trenching of reserve pits, pipeline leaks and other environmentally unsound methods of brine disposal are the main causes of the existing contamination. While many of these problems are actually related to site specific releases, the

extent, diversity and large number of sites create a nonpointsource effect of the brine contamination.

Considerable evidence exists implicating oilfield brines as the cause of high chloride concentrations in several of the lakes sampled. However, the levels of degradation have not impacted the lakes to the extent that shallow aquifers have been impacted. The high concentrations of other soluble salts tends to limit the uses of many of the lakes regardless of the observed influx of chloride salts. The relatively slow velocities of ground-water flow may account for concentrating the brine contamination in the shallow aquifers. The main slug of brines may not have yet been discharged into the lakes. Periodic resampling of several of these lakes would be a reasonable means to monitor the extent and magnitude of degradation caused by oil field wastes.

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