

**EXPLANATION**

- Dissolved-constituent concentrations**
- Circles represent wells completed in the deep alluvium.
- less than or equal to 360 mg/L between 360 and 500 mg/L
  - greater than or equal to 500 mg/L
- Squares represent wells completed in bedrock (Belt Supergroup).
- less than or equal to 360 mg/L between 360 and 500 mg/L
  - greater than or equal to 500 mg/L
- Township boundary  
Section boundary  
Major road  
Secondary road  
Stream  
Federal/State lands  
Other lands

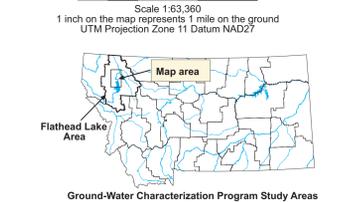


Table 1. Summary of water-quality data for 87 wells sampled during this study in the deep aquifer.

Common constituents	Min	Mean	Max	SMCL	MCL	Number exceeding
Dissolved constituents-sum (mg/L)*	160	387	766	--	--	--
Total dissolved solids (mg/L)**	102	249	531	500	--	1
Sodium (mg/L)	1.3	14.5	169.0	250	--	0
Potassium (mg/L)	<0.1	1.3	4.4	--	--	--
Calcium (mg/L)	7.0	48.9	92.1	--	--	--
Magnesium (mg/L)	7.3	21.8	55.0	--	--	--
Chloride (mg/L)	<0.5	4.2	25.0	--	--	0
Bicarbonate (mg/L)	114.9	271.7	521.2	--	--	--
Sulfate (mg/L)	<2.5	8.1	45.0	--	--	0
Nitrate (mg/L as N)	<0.25	1.1	8.2	--	10	0
Fluoride (mg/L)	<1.0	0.1	2.7	--	4	0
Iron (mg/L)	<0.003	0.2	4.1	0.3	--	15
Manganese (mg/L)	<0.002	0.08	0.97	0.05	--	24

Selected trace elements	Min	Mean	Max	SMCL	MCL	Number exceeding
Arsenic (ug/L)	<1.0	0.7	19.5	--	50	0
Barium (ug/L)	<10.0	241.9	4590	--	1000	1
Chromium (ug/L)	<2.0	4.6	16.6	--	100	0
Nickel (ug/L)	<2.0	2.7	8.8	--	100	0

Other constituents	Min	Median	Max	SMCL	MCL	Number exceeding
Sodium adsorption ratio (SAR)	0	0.3	7	--	--	--
Hardness as CaCO <sub>3</sub>	56	207	401	--	--	--
pH	6.7	7.7	8.9	6.5-8.5	--	1
Radon (pCi/L) 34 analyses	210	835	8360	--	300***	--

--, Not standard available or not applicable.  
\* Dissolved constituents is the sum of major cations (Na, Ca, K, Mg, Mn, Fe) and anions (HCO<sub>3</sub>, CO<sub>3</sub>, SO<sub>4</sub>, Cl, SiO<sub>3</sub>, NO<sub>3</sub>, F) in mg/L.  
\*\* Total dissolved solids reported as equivalent weight of evaporation residue.  
\*\*\* Proposed standard.  
MCL - U.S. EPA maximum contaminant level for public water supplies.  
SMCL - Secondary contaminant level for public water supplies.

**Dissolved-Constituent Map of the Deep Aquifer, Kalispell Valley, Flathead County, Montana**

by John I. LaFave

**INTRODUCTION**

The Kalispell valley (upper Flathead River valley) occupies part of an intermontane basin that is bounded by the Salish Mountains to the west, the Whitefish Range to the north and the Swan Range to the east. Flathead Lake marks the southern boundary. The valley covers roughly 325 square miles and is home to about 25,000 people. The mountains surrounding the valley are formed mostly by metamorphosed sedimentary rocks of the Precambrian Supergroup (Belt and include metabasites, gneisses and quartzites (Johns, 1970). Normal faults on the east and west sides of the valley have down-dropped the valley floor relative to the surrounding mountains, and the valley has been filled with a mix of alluvium and glacial till (Smith, 2000). During the Pleistocene Epoch (the last 100,000 years), the valley floor was covered by a valley-fill material composed of a mixture of alluvium, sand, and gravel deposited by glacial meltwater streams and later covered by glacial till (partly by direct glacial deposit directly by glaciers, and by glaciofluvial deposits (till) and alluvium in glacial lakes).

Although some surface water and shallow ground water are used within the Kalispell valley, ground water obtained from the deep aquifer is the most important source of water for municipal, domestic and agricultural needs. For this map, the deep aquifer is defined as sand and gravel deposits generally found at depths greater than 100 feet below the land surface, and is composed of backwash of the Flathead River (LaFave, 2000). The purpose of this map is to show the spatial distribution of dissolved constituents in the ground water of the deep aquifer and to identify the major sources.

Aquifers are saturated geologic materials that yield sufficient water to supply wells and springs. Non-aquifer materials (also known as confining beds) also may be saturated, but have low permeability and do not produce usable amounts of water to wells or springs. Confined aquifers (1965) first studied the deep aquifer in the Kalispell valley, called the "deep aquifer" (Smith, 1965). Summary descriptions of the deep aquifer are presented in Smith and Johns (1996), and Smith and Frisk (1999).

Two factors most likely account for the dissolved-constituent distribution. 1) The aquifer in the west part of the valley is composed of alluvium with interbedded sand and gravel deposits, and is generally composed of sand and gravel deposits. The coarsest sand and gravel deposits are located in the center of the valley and are composed of sand and gravel deposits. The coarsest sand and gravel deposits are located in the center of the valley and are composed of sand and gravel deposits. The coarsest sand and gravel deposits are located in the center of the valley and are composed of sand and gravel deposits.

potentiometric surface, to be in hydraulic communication with the deep sand and gravel deposits (ground water) in the deep aquifer is generally away from the mountains towards the center of the valley and from the southwest towards Flathead Lake (LaFave, 2000).

**SAMPLING METHODS**

Sixty-four sites and 87 domestic, stock, municipal, and recovery wells were analyzed for major and trace elements between April 1994 and September 1994. Field measurements of water specific conductance, pH, and temperature also were obtained at the sampled wells. To ensure acquisition of a representative sample, each well was pumped to sample collection until the field parameters stabilized and at least three samples were collected. Analyses were performed by the Montana Bureau of Mines and Geology (MBMG) Analytical Laboratory. The laboratory data were supplemented by dissolved-constituent concentrations estimated from field measurements of specific conductance at an additional 18 wells.

**GROUND-WATER QUALITY**

The overall water chemistry in the deep aquifer is similar to that of the surface water. The major ions are calcium, magnesium and bicarbonate, sodium, sulfate, and chloride. The water is hard (Figure 2). There is a discernible difference in chemical composition of water specific to each well. To ensure acquisition of a representative sample, each well was pumped to sample collection until the field parameters stabilized and at least three samples were collected. Analyses were performed by the Montana Bureau of Mines and Geology (MBMG) Analytical Laboratory. The laboratory data were supplemented by dissolved-constituent concentrations estimated from field measurements of specific conductance at an additional 18 wells.

**MAJOR IONS AND SUITABILITY FOR WATER USE**

Table 1 summarizes the results of the 87 ground-water analyses performed for this study. For reference, the U.S. Environmental Protection Agency's recommended maximum contaminant levels (MCL) and secondary maximum contaminant levels (SMCL) for public water supplies are also presented. Constituents for which maximum levels have been set may pose a health risk at elevated concentrations. Secondary levels are set for aesthetic reasons—elevated concentrations of these constituents may be a nuisance (bad taste or odor, or staining) but do not normally pose a health risk to public water supplies. For reference, the U.S. Environmental Protection Agency's recommended maximum contaminant levels (MCL) and secondary maximum contaminant levels (SMCL) for public water supplies are also presented. Constituents for which maximum levels have been set may pose a health risk at elevated concentrations. Secondary levels are set for aesthetic reasons—elevated concentrations of these constituents may be a nuisance (bad taste or odor, or staining) but do not normally pose a health risk to public water supplies. For reference, the U.S. Environmental Protection Agency's recommended maximum contaminant levels (MCL) and secondary maximum contaminant levels (SMCL) for public water supplies are also presented. Constituents for which maximum levels have been set may pose a health risk at elevated concentrations. Secondary levels are set for aesthetic reasons—elevated concentrations of these constituents may be a nuisance (bad taste or odor, or staining) but do not normally pose a health risk to public water supplies.

Radon is a colorless, odorless gas produced from the radioactive decay of uranium that occurs naturally in rocks and soil, and has been linked to lung cancer in humans (EPA, 1999). Radon in indoor air poses the greatest health risk. Most of the radon in indoor air seeps inside from the soil and rock beneath a dwelling. Water that contains radon can also be a minor source of radon in indoor air. The EPA estimates that radon released from drinking water accounts for less than 2 percent of radon in indoor air. Currently, there is no drinking water standard for radon, however the EPA is proposing a MCL of 500 pCi/L for radon in water for community water systems, and an alternative MCL of 4,000 pCi/L for states or community systems that have EPA-approved Maximum Contaminant Level Goal (MCLG) of 500 pCi/L. For radon, well water will not apply to private wells.

Radon was sampled in 10 wells for this study. In addition, the results from 8 other wells sampled between 1992 and 1995 are presented on the inset map. Of the 34 samples, only 1 had concentrations less than the proposed MCL of 500 pCi/L; those were greater than 300 pCi/L. On a state-wide basis, 73 percent of wells have been tested for radon but concentrations greater than 300 pCi/L (Miller and Coffey, 1998). The radon concentrations in the Kalispell valley show a strong correlation to aquifer materials. Those in water from wells completed in the Belt bedrock (average = 1,200 pCi/L) were twice as high as those in water from wells completed in the alluvium (average = 635 pCi/L).

For more information about radon and possible home treatment options the reader is referred to Miller and Coffey (1998) (EPA, 1999), and the U.S. Environmental Protection Agency web site (internet: www.epa.gov/owow/radon/).

**MAP CONSTRUCTION**

This map was constructed by plotting the "low" and "high" concentrations of dissolved constituents of 87 ground-water samples. The laboratory data were supplemented by dissolved-constituent concentrations derived from specific-conductance measurements made at an additional 18 wells. The specific-conductance (SC) measurements were according to the equation:  $DC = A \times SC$  (Elliott, 1992), where  $DC$  is the dissolved-constituent concentration in mg/L, and  $A$  is a conversion factor. The conversion factor for this study was 0.02, based on the relationship between SC and DC for the 87 samples analyzed in this study.

**ACKNOWLEDGMENTS**

Well owners who allowed collection of the data are gratefully acknowledged. Reviews of this report by Tom Farns, Wayne Wynn, Neilson, and Edmond Deal improved its clarity.

**SOURCES OF DATA**

Precipitation centers and roads are from 1:100,000-scale U.S. Geological Survey (USGS) Digital Line Graph files available from the National Resources Information System (NRIS) in the Montana State Library, Helena, Montana. Hydrography has been simplified from the 1:100,000-scale USGS Digital Line Graph files. Township boundaries are from U.S. Forest Service.

Point Data: Well location and water-quality data were obtained by Ground-Water Characterization Program and Department of Natural Resources and Conservation personnel. Well locations are accurate to the 2.5-acre resolution center interest. This map is available from the Montana Bureau of Mines and Geology, Montana Tech of The University of Montana, Helena, Montana.

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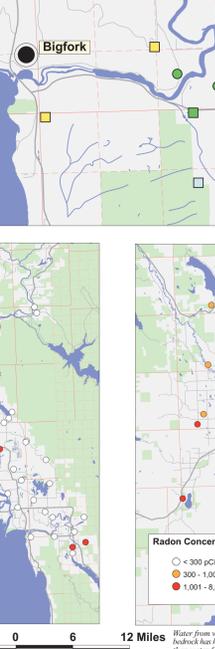
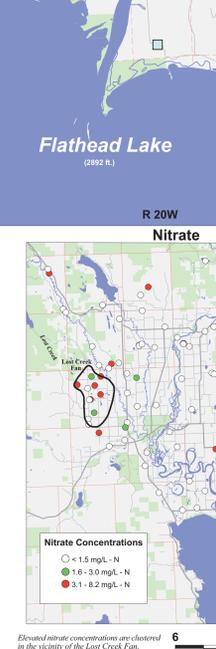
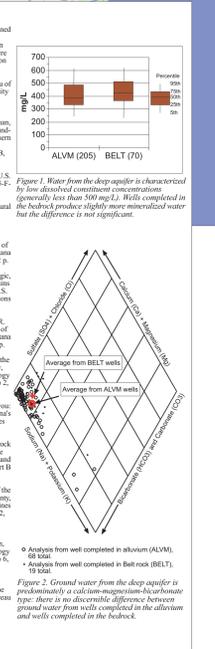
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Estimated nitrate concentrations are clustered in the vicinity of the Lost Creek Fan.

Water from wells completed in the Belt bedrock has higher radon concentrations than water from the deep alluvium.