

SURFICIAL HYDROGEOLOGY

By
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INTRODUCTION

The Upper Minnesota River Basin Regional Hydrogeologic Assessment focuses on ground-water resources, movement, and chemistry on a regional scale. The study area spans a large portion of the Minnesota River headwaters. The Minnesota River bisects the study area from northwest to southeast. The aquifers within and beneath the fine-grained sediments provide most domestic water supplies. The significant fraction of readily available water in these sediments is due to the high dissolved mineral content in ground water found in the aquifers in the area. Data collected from 84 wells included general water chemistry; radioactive isotopes of hydrogen and carbon; and stable isotopes of sulfur, hydrogen, and oxygen.

GROUND-WATER MOVEMENT AND MOVEMENT

Water Table

Water infiltrating the land surface moves generally downward through unsaturated soil and geologic materials. The water eventually reaches the water table, which is defined as that separates saturated sediments from overlying unsaturated sediments. The water table is commonly referred to as an unconfined surface; this means the pressure exerted on this surface is equal to atmospheric pressure. Most wells in the region, however, are completed in buried aquifers that generally have water that is under greater than atmospheric pressure. These buried aquifers are referred to as confined aquifers.

Contour lines on the map provide a regional depiction of the water-table surface. Delineation of the water-table contours relied on information available in the County Well Index (CWI) data base maintained by the Minnesota Geological Survey, including depth to water measurements taken when wells were drilled. Since only 6 percent of the wells in the data base for the study area are completed in surficial aquifers, and only a few wells are geographically limited, additional information was needed to determine depth to the water table. Water-table elevations were inferred where the water table is expressed at some lakes, streams, and wetlands. The water elevations for these features were obtained from the USGS 1:24,000-topographic maps. Depth to water-table was also determined using seismic refraction, which measures differences in the physical properties of saturated and unsaturated geologic materials to locate the water-table surface. Results obtained from eight locations show areas with low-relief topography generally have a shallow (less than 10 feet below land surface) water table, while areas with higher relief have deeper water tables, ranging from 30 feet below land surface. In most of the study area, the water table approximates a subsurface topography. Contours of surface elevations generated from the USGS 1:24,000-scale Digital Elevation Model (DEM) were used to guide placement of the water-table contours.

Ground-water residence time. All 84 wells were sampled for tritium, an isotope of hydrogen. Tritium is a naturally occurring radioactive isotope of hydrogen with a 12.3-year half-life that is useful for estimating ground-water residence time. Before 1954, tritium was not present in the environment (U.S. Environmental Protection Agency, 1997). Atmospheric testing of nuclear weapons during the 1950s and 1960s increased the tritium in precipitation more than a thousand-fold. Water that recharged before 1954 has lost most of its tritium by decay over several half-lives. The presence of more than about 8 TU in ground water indicates that recharge occurred since 1953 (modeling from Alexander, 1989).

Ground-water movement can be interpreted by relating the presence of tritium to well depth. Approximately 25 percent of the sampled wells contained detectable levels of tritium. Ground water less than 50 feet below the land surface will likely have measurable tritium, although two of the three wells sampled in the depth range did not have tritium. Less than half of the sampled wells contained tritium at depths greater than 100 feet below land surface. This depth range probably represents the maximum that tritium has penetrated since 1953. Only 16 percent of the sampled wells greater than 100 feet deep had tritium. This percentage likely overestimates the probability of tritium being present in wells greater than 100 feet deep. Well construction problems and geologic controls may account for wells being deeper than expected. For example, two wells completed at 150 and 215 feet deep had low but detectable levels of tritium. Two wells are close to each other and located southeast of the City of Appleton where some of the thickest sand and gravel aquifers in the study area are found. In the Appleton area (Figures 4 and 5 on Plate 2, Part A), some surficial sand and gravel deposits may be in close proximity with buried sand and gravel deposits. In these areas, water containing tritium could travel deeper into the aquifer system.

Wells completed deeper than 100 feet generally had water with no tritium. For these wells, the radioactive isotope of carbon was used to estimate ground-water residence time. Nine samples were analyzed for carbon-14 (^{14}C) age dating in wells ranging from 109 feet to 453 feet deep (Table 2). In addition, a 61-foot-deep well containing tritium was also sampled to calibrate the carbon-14 model.

The age-dating results show that water more than 100 feet below the land surface generally has a residence time from 1,000 to 9,000 years before present. The relatively young waters found in aquifers within 100 feet of the land surface suggest the presence of local and intermediate flow systems that recharge and discharge over shorter distances and times. The much older waters in aquifers below 100 feet are more likely to be associated with regional flow systems that can discharge many miles from where they receive recharge.

Water chemistry. The 10 samples listed in Table 2 were also analyzed for ratios of stable isotopes of hydrogen and oxygen. The results indicate that ground water in the deep regional flow systems largely originates from precipitation. The $\delta^{18}\text{O}$ values reflect the temperatures of the precipitation. Results show small variations that might be related to small climatic fluctuations. There is no indication of water originating from glacial meltwater, which is consistent with the calculated postglacial ground-water ages.

Wetlands are major sources of recharge to surficial aquifers and cause water levels to rise significantly. During the summer, groundwater movement is limited by the water table. When recharge does occur in the summer, it likely coincides with significant rainfall events. Recharge can also occur in the fall, depending on rainfall, runoff, and evapotranspiration rates. Water levels decline in the winter, when precipitation is stored on the land surface as snow, and typically a low point before spring thaw.

The chemistries of water samples from Cretaceous aquifers completely overlap the range of chemistries in Quaternary aquifers (see Figure 1). This observation is not surprising since large amounts of Cretaceous materials are incorporated into the glacial drift. The wide scattering of points on the diagram illustrates the wide variation in major ion water chemistry for both Cretaceous and Quaternary aquifers. The amount of dissolved minerals in ground water in the study area apparently evolves in a fairly sharp, linear fashion from the Quaternary to the Cretaceous. A relationship between the total dissolved mineral content and the observed residence time (as determined by tritium). Residence time refers to the time that ground water has resided below the land surface. This means that most of the dissolved mineral content of ground water results from chemical evolution that occurs in less than 45 years.

The cation exchange with sodium observed in some samples is associated with the Cretaceous aquifer. No samples were observed to have evidence of cation exchange. One possible explanation is that in sediments presently containing water with less than a 45-year residence time, any available sodium was exchanged and removed at an earlier time. It is also possible that these sediments never had sodium available for exchange.

Aquifer Water Chemistry

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Environmental Isotopes

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WATER CHEMISTRY

The chemical evolution of ground water begins as surface water and precipitation infiltrate below the land surface. The chemistry of the water changes as it percolates through soil and geologic material. Factors affecting ground-water chemistry include land use, first water chemistry, length of flow path, chemical reactions, and residence time.

Water samples were collected for chemical analysis in 84 wells from autumn 1997 to autumn 1998. Seventy-eight of these wells are completed in Quaternary sand and gravel, and the remaining six are completed in Cretaceous sandstones and fractured Precambrian bedrock. Precambrian igneous and metamorphic rocks underlie the entire study area. Wells are completed in Precambrian rocks because the yields are generally poor. They are used only when overlying Quaternary or Cretaceous aquifers are either absent or do not yield sufficient water. Cretaceous aquifers are common in areas where the underlying Quaternary aquifers are absent or lack sufficient yield. Magner and Anderson (1986) reported that yields for Cretaceous aquifers are generally from a few gpm to several tens of gpm.

During the past 800 years, several glacial advances have deposited a complex series of glacial sediments that are more than 600 feet thick in the study area (see Figure 3 on Plate 2, Part A) and are generally found southwest of the Minnesota River. Cretaceous sediments consist of interbedded shale, siltstone, and sandstone. Thicknesses of these sediments range from 100 to 300 feet, exceeding 600 feet southwest of the City of Canby in Yellow Medicine County. Approximately 12 percent of the wells in CWI are screened in Cretaceous sandstones; these wells range in depth from less than 50 feet to more than 400 feet below land surface. Wells completed in Cretaceous aquifers are common in areas where the underlying Quaternary aquifers are absent or lack sufficient yield. Magner and Anderson (1986) reported that yields for Cretaceous aquifers are generally from a few gpm to several tens of gpm.

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Water Quality Indicators

As water infiltrates into the subsurface, it accumulates additional carbon dioxide (CO_2) gas from decaying organic material. Dissolved carbon dioxide in sediments is limited by the amount of dissolved carbon dioxide in the water and the amount of dissolved calcium, magnesium, and bicarbonate. Water hardness is the sum of dissolved calcium and magnesium. Ground water from most sampled wells was very hard.

Another measure of water quality is the total dissolved mineral concentration in water samples. The TDS values ranged from 200 milligrams per liter (mg/L) to 2,599 mg/L. Most samples exceeded the U.S. Environmental Protection Agency's (EPA's) secondary standard for TDS of 500 mg/L. Other chemical constituents commonly exceeding EPA's secondary standards include sulfate (SO_4^{2-}), iron (Fe^{2+}), and manganese (Mn^{2+}). Excessive amounts of these ions may give water an objectionable taste or color and may damage pipes and equipment used for well screens. The lowest TDS and sulfate concentrations in the sampled wells are found in the northeast, primarily eastern Swift County. This water chemistry likely represents a difference in the mineralogy of subsurface glacial sediment in that area. In the northeastern part of the study area, Des Moines lobes sediments originating from the Des Moines River and the Mississippi River from the northeast (C. Patterson, oral commun., 1998). The northeast-southeast glacial sediments are generally not as calcareous and lack gypsum. Glacial deposits of the Des Moines lobe incorporated materials that are more calcareous and are commonly associated with gypsum.

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Major Ion Water Chemistry

The Piper trilinear diagram (Figure 1) shows the water chemistry results graphically. The sample points on each triangle (Figure 1) reflect the percentages in milligram equivalents per liter (mg/L) of the major cations and anions in each sample.

The lower left ternary diagram of Figure 1 compares the major ions for calcium and magnesium. There is a fair amount of calcium to magnesium throughout the study area. The study area superimposed on a trend toward sodium-rich waters. As waters containing calcium and magnesium pass through clays and shales, adsorbed sodium is exchanged into the water. This process is similar to how a household water softener works. Higher levels of magnesium (Mg^{2+}) associated with higher proportions of magnesium (Mg^{2+}) are an important indicator of human impact on ground water. Principal sources include septic systems, feedlot, and agricultural chemicals. EPA's primary public water supply standard for nitrate, 10 mg/L, was exceeded in only one of the 84 sampled wells, and two other wells had elevated levels (greater than 10 mg/L). The levels of nitrate elevated indicate concentrations of nitrate in the water are too high. Nitrate (NO_3^-) is usually found in near-surface ground water, but most of the wells sampled in this study are screened below the water table in aquifers overlying low-permeability glacial sediments. Second, biologically mediated nitrate removal (denitrification) may have occurred. The oxygen-poor (anoxic) ground water commonly found in glacial sediments may occur in bands within the aquifer. Nitrate is often found in near-surface ground water, but most of the wells sampled in this study are screened below the water table in aquifers overlying low-permeability glacial sediments. Second, biologically mediated nitrate removal (denitrification) may have occurred. The oxygen-poor (anoxic) ground water commonly found in glacial sediments may occur in bands within the aquifer.

Chloride (Cl^-) is another parameter that may indicate human impacts on ground water. Artificial sources of chloride include road salt; fertilizers; and industrial, human, and animal wastes. Several of the sampled wells in this study may have chloride from precipitation and Cretaceous rocks. Interpretations of chloride sources for individual wells are inconclusive because elevated natural sources of chloride levels were also encountered in some Cretaceous and buried glacial aquifers.

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Recharge and Discharge

Recharge to the water table occurs throughout the study area by infiltration of precipitation. If the areas are subject to high rainfall, recharge and surface runoff may be the dominant process for recharge areas. Sources of recharge include some lakes and wetlands and short reaches along stream segments. Water-table elevations fluctuate in response to seasonal variations in recharge and discharge from the ground-water system. Spring rain and

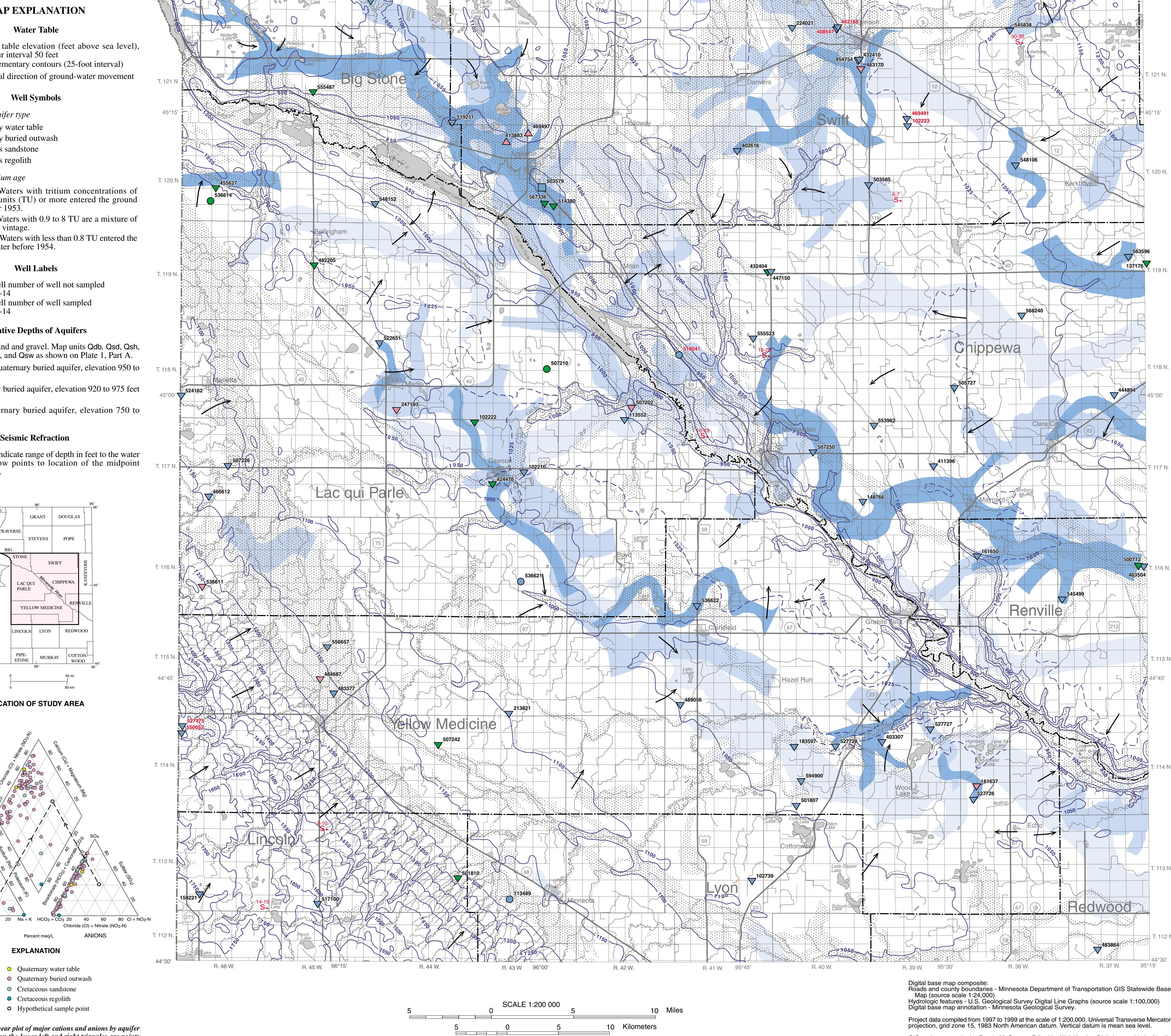


FIGURE 1. Piper trilinear plot of major cations and anions by aquifer classification. Points on the lower left and right triangles are points representing the positively charged ions (cations) and negatively charged ions (anions), respectively. The diamond-shaped field combines the components in the triangular fields as shown by the hypothetical sample point.

TABLE 1. Characteristics of natural waters by aquifer in the upper Minnesota River basin study area.

Well depth (feet)	TDS (mg/L)	pH (mV)	Dissolved oxygen (mg/L)	Eh (mV)	Cations												Anions												Total phosphorus (mg/L)
Na	Mg	Ca	K	Ca	Mg																								