Occurrence of Uranium and ²²²Radon in Glacial and Bedrock Aquifers in the Northern United States, 1993–2003

By Joseph D. Ayotte, Sarah M. Flanagan, and William S. Morrow

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Foreword

The U.S. Geological Survey (USGS) is committed to providing the Nation with credible scientific information that helps to enhance and protect the overall quality of life and that facilitates effective management of water, biological, energy, and mineral resources (http://www.usgs.gov/). Information on the Nation's water resources is critical to ensuring long-term availability of water that is safe for drinking and recreation and is suitable for industry, irrigation, and fish and wildlife. Population growth and increasing demands for water make the availability of that water, now measured in terms of quantity and quality, even more essential to the long-term sustainability of our communities and ecosystems.

The USGS implemented the National Water-Quality Assessment (NAWQA) Program in 1991 to support national, regional, State, and local information needs and decisions related to water-quality management and policy (http://water.usgs.gov/nawqa). The NAWQA Program is designed to answer: What is the condition of our Nation's streams and ground water? How are conditions changing over time? How do natural features and human activities affect the quality of streams and ground water, and where are those effects most pronounced? By combining information on water chemistry, physical characteristics, stream habitat, and aquatic life, the NAWQA Program aims to provide science-based insights for current and emerging water issues and priorities. From 1991-2001, the NAWQA Program completed interdisciplinary assessments and established a baseline understanding of water-quality conditions in 51 of the Nation's river basins and aquifers, referred to as Study Units (http://water.usgs.gov/nawqa/studyu.html).

In the second decade of the Program (2001–2012), a major focus is on regional assessments of water-quality conditions and trends. These regional assessments are based on major river basins and principal aquifers, which encompass larger regions of the country than the Study Units. Regional assessments extend the findings in the Study Units by filling critical gaps in characterizing the quality of surface water and ground water, and by determining status and trends at sites that have been consistently monitored for more than a decade. In addition, the regional assessments continue to build an understanding of how natural features and human activities affect water quality. Many of the regional assessments employ modeling and other scientific tools, developed on the basis of data collected at individual sites, to help extend knowledge of water quality to unmonitored, yet comparable areas within the regions. The models thereby enhance the value of our existing data and our understanding of the hydrologic system. In addition, the models are useful in evaluating various resource-management scenarios and in predicting how our actions, such as reducing or managing nonpoint and point sources of contamination, land conversion, and altering flow and (or) pumping regimes, are likely to affect water conditions within a region.

Other activities planned during the second decade include continuing national syntheses of information on pesticides, volatile organic compounds (VOCs), nutrients, selected trace elements, and aquatic ecology; and continuing national topical studies on the fate of agricultural chemicals, effects of urbanization on stream ecosystems, bioaccumulation of mercury in stream ecosystems, effects of nutrient enrichment on stream ecosystems, and transport of contaminants to public-supply wells.

The USGS aims to disseminate credible, timely, and relevant science information to address practical and effective water-resource management and strategies that protect and restore water quality. We hope this NAWQA publication will provide you with insights and information to meet your needs, and will foster increased citizen awareness and involvement in the protection and restoration of our Nation's waters.

The USGS recognizes that a national assessment by a single program cannot address all water-resource issues of interest. External coordination at all levels is critical for cost-effective management, regulation, and conservation of our Nation's water resources. The NAWQA Program, therefore, depends on advice and information from other agencies—Federal, State, regional, interstate, Tribal, and local—as well as nongovernmental organizations, industry, academia, and other stakeholder groups. Your assistance and suggestions are greatly appreciated.

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Conversion Factors, Water-Quality Units, and Abbreviations

Multiply	Ву	To obtain
	Length	
inch (in.)	2.54	centimeter (cm)
foot (ft)	0.3048	meter (m)
mile (mi)	1.609	kilometer (km)
acre	0.4047	hectare (ha)
square mile (mi²)	2.590	square kilometer (km²)
	Volume	
million gallons (Mgal)	3,785	cubic meter (m³)
gallon per minute (gal/min)	0.06309	liter per second (L/s)
million gallons per day (Mgal/d)	0.04381	cubic meter per second (m³/s)
	Radioactivity	
picocurie per liter (pCi/L)	0.037	becquerel per liter (Bq/L)

Temperature in degrees Celsius (°C) may be converted to degrees Fahrenheit (°F) as follows: °F=(1.8×°C)+32

Horizontal coordinate information is referenced to the North American Datum of 1983 (NAD 83).

ABBREVIATED WATER-QUALITY UNITS

μg/L micrograms per liter mg/L milligrams per liter pCi/L picocuries per liter

milligram per gram; equivalent to 1 part per million mg/g

parts per million; equivalent to 1 mg/g ppm

MISCELLANEOUS ABBREVIATIONS

redox oxidation-reduction potential MCL Maximum Contaminant Level AMCL Alternative Maximum Contaminant Level NAWQA National Water-Quality Assessment NURE National Uranium Resource Evaluation program **USEPA** U.S. Environmental Protection Agency

USGS U.S. Geological Survey TDS total dissolved solids

AQUIFER GROUP CODES

CPG	Columbia Plateau glacial aquifer group
COS	Cambrian-Ordovician aquifer group
ECG	East-Central glacial aquifer group
NEC	New York and New England crystalline aquifer group
NEG	Northeastern glacial aquifer group
NCG	North-Central glacial aquifer group
WCG	West-Central glacial aquifer group
NPG	Northern Pacific glacial aquifer group
PWG	Pre-Wisconsinan glacial aquifer group

Occurrence of Uranium and ²²²Radon in Glacial and Bedrock Aquifers in the Northern United States, 1993–2003

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Abstract

Water-quality data collected from 1,426 wells during 1993–2003 as part of the U.S. Geological Survey National Water-Quality Assessment (NAWQA) program were evaluated to characterize the water quality in glacial and bedrock aquifers of the northern United States. One of the goals of the NAWOA program is to synthesize data from individual studies across the United States to gain regional- and national-scale information about the behavior of contaminants. This study focused on the regional occurrence and distribution of uranium and ²²²radon in ground water in the glacial aquifer system of the United States as well as in the Cambrian-Ordovician and the New York and New England crystalline aquifer systems that underlie the glacial aquifer system. The occurrence of uranium and 222 radon in ground water has long been a concern throughout the United States. In the glacial aquifers, as well as the Cambrian-Ordovician and the New York and New England crystalline aquifer systems of the United States, concentrations of uranium and 222 radon were highly variable. High concentrations of uranium and ²²²radon affect ground water used for drinking water and for agriculture.

A combination of information or data on (1) nationalscale ground-water regions, (2) regional-scale glacial depositional models, (3) regional-scale geology, and (4) national-scale terrestrial gamma-ray emissions were used to confirm and(or) refine the regions used in the analysis of the water-chemistry data. Significant differences in the occurrence of uranium and ²²²radon, based primarily on geologic information were observed and used in this report. In general, uranium was highest in the Columbia Plateau glacial, West-Central glacial, and the New York and New England crystalline aquifer groups (75th percentile concentrations of 22.3, 7.7, and 2.9 micrograms per liter (µg/L), respectively). In the Columbia Plateau glacial and the West-Central glacial aquifer groups, about 10 percent of wells sampled had concentrations of uranium that exceeded the U.S. Environmental Protection Agency (USEPA) Maximum Contaminant Level of 30 µg/L; in the New York and New England crystalline aquifer group, 4 percent exceeded 30 μg/L.

Ground-water samples with high concentrations of uranium were commonly linked to geologic sources rich in

uranium. In seven of nine aquifer groups defined for this study, concentrations of uranium correlated significantly with concentrations of sulfate in ground water (Spearman's rho = 0.20 to 0.56; p < 0.05). In the Columbia Plateau, glacial aquifers were derived in part from basaltic lava flows, some felsic volcanic rocks, and some paleo-lake bed materials that may be rich in uranium. In the Columbia Plateau and West-Central glacial aquifer groups, uranium correlated with total dissolved solids, bicarbonate, boron, lithium, selenium, and strontium. In the West-Central glacial aquifer group, rocks such as Cretaceous marine shales, which are abundant in uranium, probably contribute to the high concentrations in ground water; in the southern part of this group, which extends into Nebraska, the glacial or glacial-related sediment may be interbedded with uranium-rich materials that originated to the north and west and in the Rocky Mountains. In New England, crystalline bedrock that is granitic, such as two-mica granites, as well as other high-grade metamorphic rocks, has abundant uranium that is soluble in the predominantly oxic to sub-oxic geochemical conditions. This appears to contribute to high uranium concentrations in ground water.

The highest ²²²radon concentrations were present in samples from wells completed in the New York and New England crystalline aquifer group; the median value (2,122 picocuries per liter (pCi/L)) was about 10 times the median value of the glacial and Cambrian-Ordovician aquifer groups. More than 25 percent of the samples from the New York and New England crystalline aquifer group wells had ²²²radon concentrations that exceeded the proposed USEPA Alternative Maximum Contaminant Level (AMCL) of 4,000 pCi/L. ²²²Radon concentrations in samples from the New York and New England crystalline aquifer group were similar to those found in other countries with similar geologic histories and(or) similar bedrock geology. In the New York and New England crystalline aquifer group, concentrations of ²²²radon were correlated with uranium (Spearman's rho = 0.60; p < 0.0001) and with gross alpha particle activity (Spearman's rho = 0.72; p < 0.0001). Also, in some parts of the New York and New England crystalline aquifer group, 222 radon correlated with lead (Spearman's rho = 0.52; p = 0.004) and some of the lead concentrations (greater than 2 µg/L) were associated with high ²²²radon concentrations (greater than 10,000 pCi/L).

Introduction

The U.S. Geological Survey National Water-Quality Assessment (NAWQA) Program conducts regional assessments of ground-water quality conditions (Lapham and others, 2005). These assessments target principal aquifers, many of which underlie multiple States, (fig. 1) that are major sources for drinking water, irrigation, and other uses. Regional assessments complement and extend the findings of the NAWQA Study Units (fig. 2) and expand our understanding of groundwater quality over broad regions. These assessments are designed to increase our understanding of how natural features and human activities affect ground-water quality and will help with extrapolating results to unmonitored, comparable areas in other parts of the regions (Lapham and others, 2005). This study focused on uranium and 222 radon occurrence in three principal aquifers in the glaciated northern United States (the glacial, Cambrian-Ordovician, and the New York and New England crystalline aquifer systems).

Uranium occurs naturally in the Earth's crust, in plants and animals, and in surface and ground waters. Uranium is among the heaviest and largest elements forming the Earth's crust, and ranks about 48th in abundance. Uranium has 16 isotopes, all of which are radioactive. Natural uranium has three primary isotopes: ²³⁸uranium (99.2 percent by weight), ²³⁵uranium (0.7 percent), and ²³⁴uranium (0.1 percent).

Uranium can be highly concentrated in intrusive igneous rocks, primarily granitic rocks (table 1). In felsic igneous rocks, uranium is concentrated in the micas and amphiboles. The large ionic radius and high valence state of uranium make it incompatible with the minerals created during the early stages of crystallization; therefore, uranium remains in the liquid phase of magma until the final stages of crystallization (Adams and others, 1959). Minerals formed during later stages of crystallization (such as biotite, muscovite, potassium feldspar, sodium feldspar, and quartz) can incorporate up to several percent of uranium in their crystal structure (Churchill, 1991). Uranium also is found in flat-lying bedded deposits in sedimentary rocks. Black shales, coals, asphalt, clays, silts, soils above limestone, and phosphate rocks also can contain large amounts of natural uranium (table 1).

²²²Radon is an inert gas and is not directly affected by geochemical conditions, yet is highly soluble in ground water. Although radon is formed by the radioactive decay process of uranium, radon and uranium do not necessarily co-occur in ground water. Concentrations of radon in ground water can be many orders of magnitude larger than the concentrations of its parent source. The decay of ²³⁸uranium through ²²⁶radium is the primary source of ²²²radon (fig. 3). Geologic sources that might lead to high concentrations of ²²²radon in soils and ground water are mainly (1) uranium-bearing metamorphosed rocks; (2) silica-rich volcanic rocks; (3) granite-intrusive (igneous) rocks that are highly deformed or sheared; and (4) sedimentary rocks including marine black shales, high-iron soils derived from carbonate, uranium-bearing fluvial

sediments, glacial deposits (such as deltaic, marine, and lacustrine deposits), and phosphate rocks (Gundersen, 1991; National Academy of Sciences, 1999; Schumann, 1993a). Because it is the immediate daughter product of ²²⁶radium and has a short half life (3.82 days), the presence of ²²⁷radon in ground water depends on a readily available source of ²²⁶radium and the hydraulic properties of the aquifer materials (Hall and others, 1987).

Concentrations of uranium in ground water also can differ as a result of the natural variability in the geology and in the depositional environment of the materials that compose these aquifers. Local factors, such as the availability of uranium minerals, ground-water flow characteristics, and geochemistry can affect uranium concentrations. Thus, the variation of uranium and ²²²radon concentrations in ground water in the glacial aquifer system depends, in part, on the source of the aquifer materials. The source of glacial materials depends on (1) the direction of glacial ice lobe movement, (2) the previous glacial deposits, and (3) the underlying bedrock geology. Ice that flowed into the central United States from Canada came from various directions and carried materials from along its path. Glacial-aquifer material is a mix of existing glacial deposits (which may have originated elsewhere) and materials from the underlying bedrock, creating a composite mixture that can affect concentrations of natural contaminants, such as uranium and ²²²radon.

Uranium and 222 radon can have adverse health effects on humans. In 2003, uranium became a regulated contaminant when the U.S. Environmental Protection Agency (USEPA) set a Maximum Contaminant Level (MCL) of 30 micrograms per liter (µg/L) for uranium in drinking water (U.S. Environmental Protection Agency, 2005). Regulating ²²²radon is complicated because 222 radon is an inert gas that is widely detected in ground water, but its primary hazard is through inhalation from indoor air in living spaces (U.S. Environmental Protection Agency, 1999). Currently (2007), the USEPA has not established an MCL for radon in drinking water, but has proposed an MCL of 300 pCi/L and an Alternative Maximum Contaminant Level (AMCL) of 4,000 pCi/L for public-water supplies (U.S. Environmental Protection Agency, 1999). The AMCL would be in effect in states that have established remediation guidelines for radon in indoor air.

In Canada, the drinking-water standard for uranium is $20~\mu g/L$ (Health Canada, 2001). The World Health Organization (WHO) has established the lowest drinking-water standard for uranium, at 15 $\mu g/L$ (World Health Organization, 2004). Two New England states have standards or recommendations for uranium that are lower than the USEPA MCL. The Vermont Department of Environmental Conservation, Water Supply Division, established an MCL for uranium in public water-supply system wells of $20~\mu g/L$, which is based on health risk information from the State of Vermont Department of Health (Vermont Department of Health, 2006). The Maine Bureau of Health established a uranium MCL of $30~\mu g/L$, but advises homeowners to "use less tap water for drinking" if uranium concentrations are between $20~\text{and}~30~\mu g/L$ (Maine

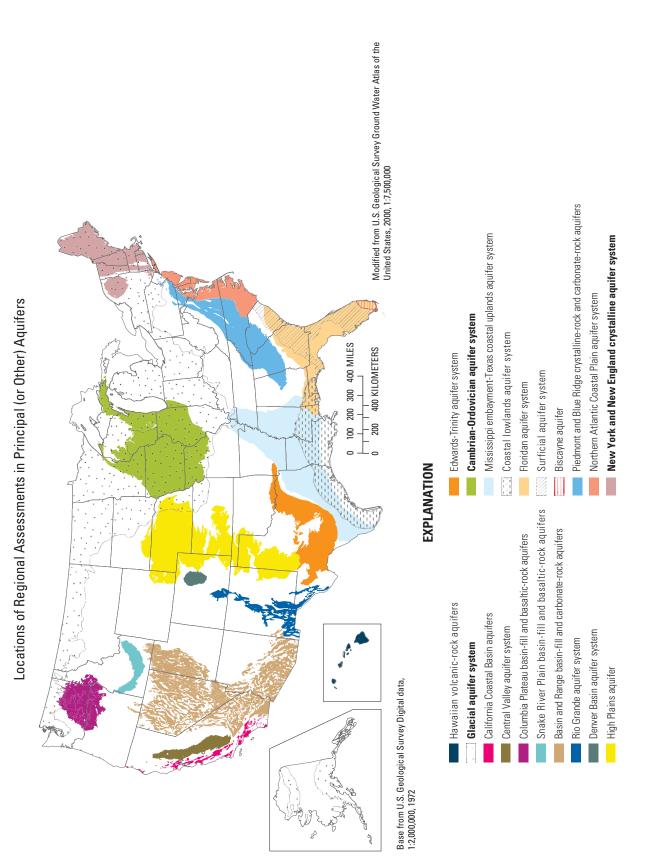
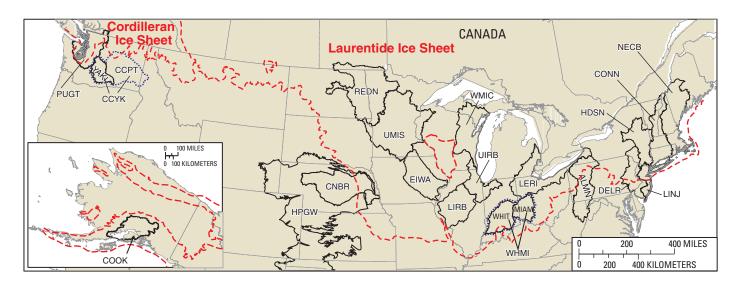


Figure 1. Locations of National Water-Quality Assessment (NAWQA) regional assessments in principal aquifers.

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EXPLANATION

NAWQA STUDY UNIT boundary

COMBINED STUDY UNIT boundaries

GLACIAL EXTENT (approximate)

NAWQA STUDY UNIT

ALMN	Allegneny & Mononganela River Basins	LIRB	Lower Illinois River Basin
CNBR	Central Nebraska Basins	LERI	Lake Erie-Lake St. Clair Drainage
CONN	Connecticut, Housatonic and Thames River Basins	LINJ	Long Island & New Jersey Coastal Drainages
COOK	Cook Inlet Basin	NECB	New England Coastal Basins
DELR	Delaware River Basin	PUGT	Puget Sound Basin
EIWA	Eastern Iowa Basins	REDN	Red River of the North Basin
HDSN	Hudson River Basin	UIRB	Upper Illinois River Basin
HPGW	High Plains Regional Ground-Water Study	UMIS	Upper Mississippi River Basin
		WMIC	Western Lake Michigan Drainage

COMBINED STUDY UNITS

CCYK Central Columbia Plateau (CCPT) and Yakima River Basin (YAKI)

WHMI White River Basin (WHIT) and Great and Little Miami River Basins (MIAM)

Figure 2. Locations of National Water-Quality Assessment (NAWQA) program study units in the northern United States.

Bureau of Health, 2005). Other states in the glaciated areas of the U.S. have not adopted guidelines for uranium.

Some state health and regulatory agencies have established their own recommendations concerning acceptable levels of ²²²radon in drinking water. For example, recommendations regarding treatment for the removal of ²²²radon from domestic drinking-water supplies take effect at 2,000 pCi/L in New Hampshire (New Hampshire Department of Environmental Services, 2005), at 4,000 pCi/L in Vermont, at 10,000 pCi/L in Massachusetts, and at 5,000 pCi/L in Connecticut (Connecticut Department of Health, 2006). The Maine Department of Human Services recommends that remedial actions should be taken when ²²²radon concentrations are greater than 20,000 pCi/L (University of Maine, 1996). Other states within the glaciated regions of the United States have not adopted guidelines for ²²²radon.

The glacial aguifers of the northern United States are important sources of water supply. Estimates of water use in the glacial aquifer system indicated that about 3.56 billion gallons per day (Bgal/d) were withdrawn for all uses during 2000 with about 1.95 Bgal/d for public supply (Maupin and Barber, 2005). Irrigation accounted for 29 percent (1.02 Bgal/d) of all withdrawals (Maupin and Barber, 2005). Irrigation can affect contaminant concentrations found in glaciofluvial aquifers. Irrigation water can contain high concentrations of naturally occurring dissolved salts, major ions, and trace elements, as well as fertilizers and pesticides, that are undesirable for crop growth and(or) human consumption. These chemical inputs can leach past the crop root zone and contaminate the underlying aquifer or subsurface drainage systems in what is called irrigation return flow (National Academy of Sciences, 1989).

Withdrawals from the Cambrian-Ordovician and the New York and New England crystalline aquifer systems totaled 932 million gallons per day (Mgal/d) and 96.3 Mgal/d, respectively in 2000. Public supply accounted for about 63 percent of total withdrawals from the Cambrian-Ordovician aquifer system and 76 percent of total withdrawals from the New York and New England crystalline aquifer system (Maupin and Barber, 2005).

Purpose and Scope

This report presents uranium and ²²²radon concentrations in three principal aquifers in the northern United States and describes possible sources. The three principal aquifers are (1) the glacial aquifer system in the northern United States, (2) the Cambrian-Ordovician aquifer system, primarily in Minnesota, Wisconsin, Iowa, and Michigan, and (3) the New York and New England crystalline aquifer system in New England and New York (fig. 1). The water-chemistry data used in the assessment were collected during 1993–2003 from 1,426 wells, which are part of the networks used in the NAWQA program. The distribution and concentrations of uranium and ²²²radon in glacial and bedrock aquifers of the

Table 1. Typical range of uranium concentrations in various rock types.

[<, less than; >, greater than; ~, approximately. Table 1 is modified from Ivanovich and Harmon (1992), table 2.1, p. 36]

Rock type	Name	Uranium (parts per million)			
Igneous	Two-mica granite	3–300			
	Calc-alkaline granites Alkalic plutonic rocks				
	Granites	2.2-6.1			
	Granodiorites				
	Rhyolites				
	Dacites				
	Gabbros	0.8			
	Basalts	0.1-1			
	Ultramafics	< 0.015			
Metamorphic	Eclogites	0.3–3			
	Granulites	4.9			
	Gneiss	2.0			
	Schist	2.5			
	Phyllite	1.9			
	Slate	2.7			
Sedimentary	Orthoquartzite	0.45-3.2			
	Greywackes	0.5 - 2.0			
	Shales: grey-green,	2–4			
	red-yellow, black	2–4			
	.	3–1, 250			
	Bauxite	11.4			
	Limestones	~2			
	Dolomites	0.03-2			
	Phosphates	50–300			
	Evaporites	< 0.1			
	Speleothem	< 0.03 – 100			
	Living molluses	<0.01-0.5			
	Fossil molluscs	0.15-8			
	Coral	2–4			
	Manganese nodules	2–8			
	Oceanic sands and clays	0.7–4			
	Peat	1–12			
	Lignite	<50-80			
	Coal	<10-<6,000			
	Asphalt	10–3,760			
	Oil	4–77			



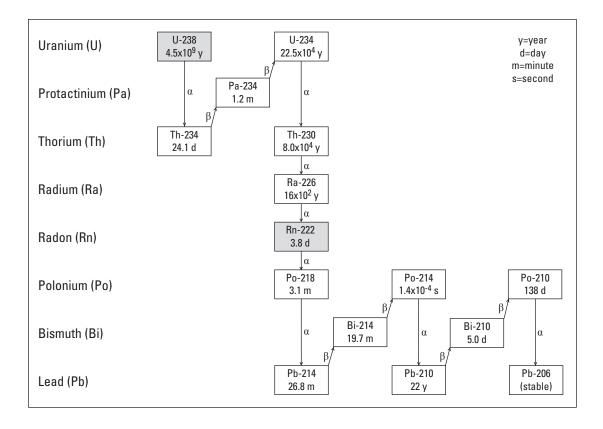


Figure 3. Uranium-238 radioactive decay series.

northern United States are shown on maps, and the relations of uranium and 222 radon to each other and to selected waterchemistry constituents, gross alpha particle radioactivity, and physical properties are illustrated in graphs. Statistical summaries of uranium and 222 radon concentrations and significant statistical correlations are listed in tables.

Health Effects of Uranium and ²²²Radon

Uranium enters the human body primarily through ingestion of food and water or by inhalation of airborne uranium-bearing dust particles or aerosols. Little is known about the long-term health effects on humans of exposure to low-level environmental uranium. Studies of occupationally exposed persons, like uranium miners, have shown that the major health effect of uranium in the body is renal (kidney) toxicity (Leggett, 1989; Taylor and Taylor, 1997). Because uranium tends to concentrate in specific locations in the body, such as in the skeletal structure and the liver, risk of bone and liver cancer and blood diseases (such as leukemia) also are increased.

A recent study of uranium concentrations in drinkingwater wells in Finland and in urine samples from individuals using these wells found an association between increased uranium exposure through drinking water and increased excretion levels of several solutes, including calcium, in urine (Kurttio and others, 2002). The Finnish study also suggests that a safe concentration for uranium in drinking water is less than 30 µg/L. In addition, recent epidemiology studies have shown that many heavy metals (cadmium, selenium, arsenic, cobalt, copper, nickel, chromium, lead, mercury, tin, uranium, and vanadium) possess estrogenic properties and may act as endocrine disrupters in mammals exposed to these metals in the environment (Choe and others, 2003; Martin and others, 2003).

The Surgeon General of the United States has recognized exposure to ²²²radon gas as being second only to cigarette smoking as a cause of lung cancer (U.S. Environmental Protection Agency, 1992). ²²²Radon itself is a relatively inert gas and almost all of the gas that is inhaled is exhaled (Kendall and Smith, 2002). One study found a significant correlation between lung cancer rates and radon levels in Maine, a state known for high concentrations of 222 radon in ground water and soils (Hess and others, 1979). An analysis of several epidemiologic studies, however, found a relation between lung cancer and inhalation of radon decay products only in underground miners (Lubin and Boice, 1997).

Previous Investigations

Previous investigations of uranium and 222 radon in ground water range from national to local scale. Several national studies provided a comprehensive assessment of the general occurrence and distribution of uranium and 222 radon in ground water throughout the United States (Aieta and others, 1987; Cothern and Lappenbusch, 1982; Focazio and others, 2001; Hess and others, 1985; Longtin, 1988; Michael and Cothern, 1986; Michael and Jordana, 1987; National Academy of Sciences, 1999; Otton and others, 1992; U.S. Environmental Protection Agency, 2005; Wanty and Nordstrom, 1993; Zapecza and Szabo, 1988). One study in particular that examined radionuclide occurrence by geologic provinces of the United States (Hess and others, 1985) found that uranium concentrations in ground water were highest in the Colorado Plateau, West Central Platform, and Rocky Mountains, whereas ²²²radon concentrations in ground water were highest in the New England and Appalachian Highlands/Piedmont. A study on the water-quality data for public-supply wells determined that high concentrations of 222 radon in ground water occurred primarily in the New England states, followed by the Appalachian states, the Rocky Mountain states, and small areas of the Southwest and the Great Plains (National Academy of Sciences, 1999). Most of these studies recognized the importance of geologic and geochemical controls on the occurrence of all radionuclides. Numerous regional and statewide studies pertinent to the northern United States also have been done and are described below by region.

Radionuclides in the Northeast

The occurrence of ²²²radon in the northeast United States has been documented in many studies. These studies found that ²²²radon concentrations were highest in granitic and high-grade metamorphic bedrock aquifers and were related to hydrologic and hydraulic factors including well pumping (Brutsaert and others, 1981; Hall and others, 1987; Hall and others, 1985; Hess and others, 1979; Maloney, 1988; Moore, 2004; Robinson and others, 2004; Thomas, 1987).

Studies on the occurrence of uranium in ground water in the northeast is limited except for several studies in New Jersey (Szabo and dePaul, 1998; Szabo and others, 2005; Szabo and Zapecza, 1991; Wanty and others, 1992a) and one in Connecticut (Brown and Zielinski, 2004). In these studies, concentrations of uranium and ²²²radon were related to pH, redox, geology, and other factors (Brown and Zielinski, 2004; Szabo and dePaul, 1998; Szabo and others, 2005; Szabo and Zapecza, 1991; Wanty and others, 1992a).

Radionuclides in the Midwest

Concentrations of ²²²radon in the glacial aquifer system in the Midwest and the Cambrian-Ordovician aquifer system generally were low but exceeded 100,000 pCi/L in northcentral Wisconsin (Mudrey and Bradbury, 1993). ²²²Radon was

present in ground water from crystalline bedrock and glacial aquifers on some Indian reservations in Wisconsin, but radon concentrations were higher in water from bedrock wells than from glacial wells (DeWild and Krohelski, 1995). Results from a study of ²²²radon in glacial till in southwestern Ohio and southeastern Indiana indicated that the original ²²⁶radium source (parent) material was shale in the till but that phosphate fertilizers could be a source (Hair and Baldwin, 1995). In Illinois, uranium concentrations in ground water correlated strongly with sulfate in oxic environments, and in general, ²²²radon concentrations were less than 1,000 pCi/L (Morrow, 2001). In the Cambrian-Ordovician aquifer system, uranium and ²²²radon concentrations were relatively low but, ²²⁶radium concentrations commonly exceeded the USEPA MCL of 5 pCi/L (Gilkeson and Cowart, 1987). Other studies in this region determined that ²²²radon concentrations were less than 1,000 pCi/L in glacial and bedrock aquifers (Field and Kross, 1998; Kelley and Mehrhoff, 1993; Stark and others, 2000).

Radionuclides in the West

Parts of the alluvial aquifers in eastern Nebraska and parts of the eastern Platte River valley are derived from glacial outwash (Jennifer Stanton, U.S. Geological Survey, written commun., 2006). In the South Platte River Basin of Nebraska and Colorado, concentrations of uranium from shallow, unconsolidated aquifers commonly exceeded the drinking-water standard of 30 µg/L (Bruce and McMahon, 1998; Dennehy and others, 1998; Parnell, 2000). Uranium concentrations were higher in ground-water samples from alluvial aquifers along the South Platte River than in those from bedrock aquifers in the Rocky Mountains. Background concentrations of ²²²radon in the glacial aquifer system were substantially higher than the national average; however, median radon concentrations were higher in water samples from the crystalline aguifer than in water samples from the alluvial aguifer (Dennehy and others, 1998). In Alaska, concentrations of uranium in ground water from glacial aquifers in the Cook Inlet area were all less than 1 μg/L (Glass, 2001).

Process Studies and Other Data Related to Radionuclides in Ground Water

Studies relating agricultural irrigation practices to concentrations of uranium in surface and ground waters in the western United States indicated that uranium concentrations were high (greater than 30 µg/L) in waters draining agricultural lands and where streamflow was predominantly from irrigation return flows (Naftz, 1996; Seiler, 2004; Zielinski and others, 1995). High uranium concentrations in surface waters from irrigated areas of the western United States were associated with high evaporation rates, as indicated by high total dissolved solids concentrations (greater than or equal to 500 mg/L), and with fine-grained marine sediments (Cretaceous shale) (Naftz, 1996). High concentrations of uranium in samples from aquifers in an

irrigated area in Nebraska, similar to deposits of the glacial aquifer system, likely resulted from dissolution of volcanic ash or other minerals in underlying bedrock (Verstraeten and others, 2000). Also, induced flow of pumped ground water to deeper parts of aquifers may be an important mechanism of moving uranium-enriched shallow, oxic ground water to public-supply wells (Craig Brown, U.S. Geological Survey, written commun., 2006).

Studies that use process-based models to explain ²²²radon occurrence provide insight into factors that control ²²²radon concentrations, although the conclusions derived from these studies sometimes differ. Researchers acknowledge that uranium concentrations in aguifer minerals, porosity, and density of bedrock are factors related to high 222 radon concentrations in ground water (Fleischer, 1983; Wanty and others, 1992b). The effect of the presence of ²²⁶radium along fractures, flow velocity, and fracture size also have been examined (Folger and others, 1996). More recent research proposes a chemically based diffusion/exchange model to explain elevated ²²²radon concentrations in fractured rock and the relation of ²²²radon to ²²⁶radium (Wood and others, 2004). A study of ²²²radon and uranium concentrations in ground water from glacial and fractured-rock aquifers found that porosity and location of the uranium on the fracture were the controlling factors for ²²²radon concentrations (Veeger and Ruderman, 1998). Other studies have identified areas of high 222 radon potential based on geologic characteristics for igneous, metamorphic, sedimentary, and unconsolidated deposits (Gundersen, 1989; Gundersen, 1991; Gundersen, 1993; Gundersen and others, 1992; Moreland and others, 1998).

Other regional data related to radionuclides in the environment were useful in explaining the occurrence of radionuclides in ground water. Data on uranium and other trace elements from the National Uranium Resource Evaluation (NURE) Hydrogeochemical and Stream Sediment Reconnaissance Program (HSSR), a program tasked with identifying uranium resources within the United States, are available for thousands of stream-sediment samples (Smith, 2001a; Smith, 2001b). These data were used as one indicator of areas where sources of uranium may be available to the ground-water system. These sources, however, may not affect concentrations in the ground water because other factors, such as geochemical controls may have greater effect on the solubility and mobility of uranium.

An aerial survey of terrestrial gamma-ray emissions of near-surface sediments, also part of the NURE program, resulted in maps that show estimated isotope concentrations of ⁴⁰potassium, equivalent ²³⁸uranium, and ²³²thorium in the United States (Duval, 1990). Most of the gamma-ray emissions were from the upper 25 centimeters (cm) of surficial materials (primarily rock or soil). Although ground water may not be in direct contact with the sediments that produce the gamma-ray response, these can be regional scale indicators of the potential for the presence of uranium and ²²²radon in ground water.

Study Design and Methods

This study focused on three principal aguifers within the glaciated terrain of the northern United States (including the Cook Inlet Basin in Alaska)—(1) the glacial (and glaciofluvial) aguifer system, (2) the Cambrian-Ordovician aguifer system of the upper Midwest and (3) the New York and New England crystalline aquifer system (Lapham and others, 2005) (fig. 1). The largest of the principal aquifers is the glacial aquifer system. In a generalized scheme, the glacial aquifer system has been divided into four areas as a framework to describe general variation in ground-water quality (Warner and Arnold, 2005). This generalized framework was based on hydraulic properties (susceptibility) and features related to contaminant sources (vulnerability), but because mineralogical and(or) geochemical differences among the areas have not been documented, these divisions are subject to revision (Warner and Arnold, 2005). Revisions to the NAWQA glacial aquifer system framework, based in part on information related to mineralogy and geochemistry, were made for the purposes of this report and are described below.

For this study, the glacial aguifer system framework has been modified to account for the geologic source material of the glacial aquifers as a proxy for uranium or other radionuclides in the aquifer matrix. The modified framework is generalized and differentiates the glacial aquifer system on the basis of information on surficial geology, bedrock geology, and late Wisconsinan glacial lobe positions (fig. 4) and flow directions (Duval, 1990; Heath, 1984; King and Beikman, 1974; Meinzer, 1923; Mickelson and others, 1983; Mickelson and Colgan, 2003; Schumann, 1993b). Data representing late Wisconsinan glacial lobe positions were grouped by considering (1) the direction of the glacial lobe movement, which accounts for geologic source materials in the up-ice direction and, (2) the underlying bedrock geology, which accounts for local geologic materials. In addition, national data on equivalent ²³⁸uranium (eU) and ²³²thorium (eTh) content from terrestrial gamma-ray surveys of near-surface sediments (Duval, 1990) were used to refine areas of potentially high radionuclide concentrations in ground water (fig. 5).

Based on the above criteria and for the purposes of this report, the study design for comparing uranium and ²²²radon concentrations in the glacial, Cambrian-Ordovician, and New York and New England crystalline aquifer systems includes nine aquifer groups: (1) Northern Pacific glacial aquifers, (2) Columbia Plateau glacial aquifers, (3) West-Central glacial aquifers from the James, Des Moines, and Red River (western part) glacial lobes, (4) North-Central glacial aquifers from the Late Wisconsinan Red River (eastern part), Wadena, Rainy, Lake Superior, and Green Bay glacial lobes, (5) East-Central glacial aquifers from the Late Wisconsinan Lake Michigan, Huron, and Huron-Erie glacial lobes, (6) Pre-Wisconsinan glacial aquifers, (7) Northeastern glacial aquifers, (8) Cambrian-Ordovician aquifers, and (9) New York and New England crystalline aquifers (fig. 6, table 2).

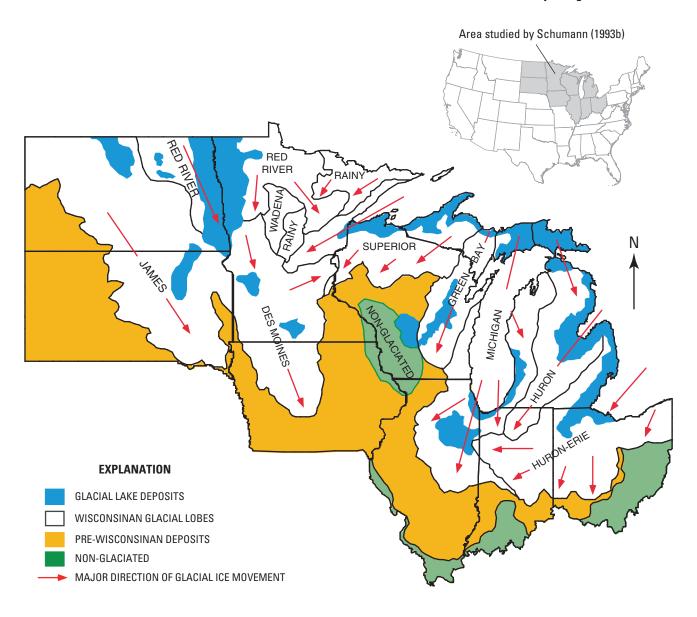


Figure 4. Generalized glacial geologic map of the upper Midwestern United States and names of major glacial lobes that were used to modify the glacial aquifer framework.

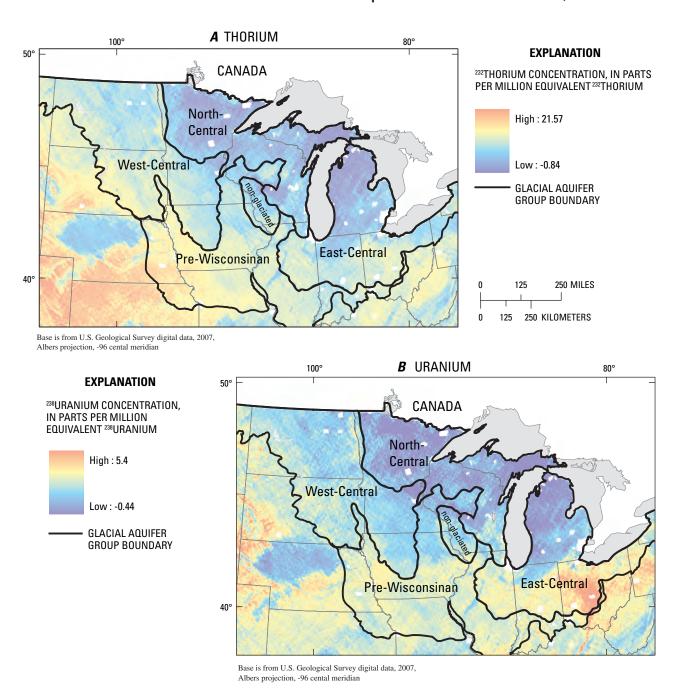
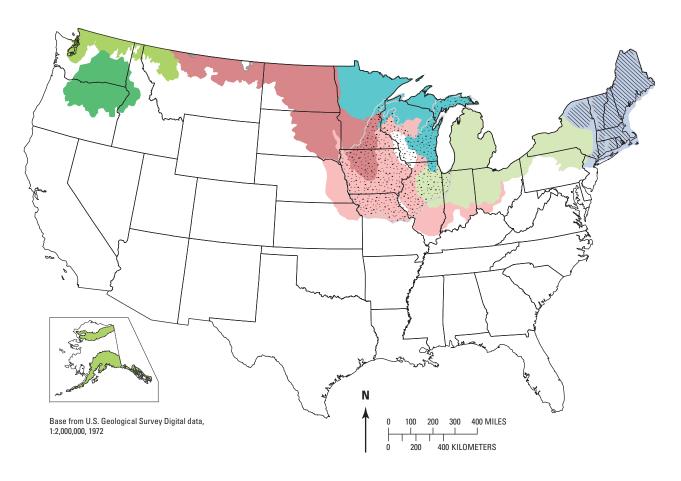


Figure 5. Distributions of (A) equivalent ²³²thorium and (B) equivalent ²³⁸uranium concentrations of near-surface sediments from aerometric gamma-ray surveys in the upper Midwestern United States used to modify the glacial aquifer framework.



EXPLANATION

AQUIFER GROUP CODE	AQUIFER GROUP
NPG	Northern Pacific glacial (includes Cook Inlet basin, Alaska, not shown)
CPG	Columbia Plateau glacial
WCG	West-Central glacial (James, Des Moines, and western part of the Red River glacial lobes)
NCG	North-Central glacial (eastern part of the Red River, Wadena, Rainy, Lake Superior, and Green Bay glacial lobes)
PWG	Pre-Wisconsinan glacial
ECG	East-Central glacial (Lake Michigan, Huron, and Huron-Erie glacial lobes)
COS	Cambrian-Ordovician (sandstones and dolomite)
NEG	Northeastern glacial
NEC	New York and New England crystalline (fractured bedrock)

Figure 6. Locations of generalized aquifer groups in the northern United States.

The boundaries of the Cambrian-Ordovician and the New York and New England crystalline aquifer systems were not modified but are referred to as groups, rather than systems, in this report, to be consistent with the terminology for the glacial aquifers. Additionally, the sediments in the Nebraska portion of the Pre-Wisconsinan glacial aquifers were derived in part from Cretaceous shale deposits transported within the Dakota lobe (which extended further south than the James lobe, into eastern Nebraska) (Colgan, 1999; Meinzer, 1923; Srhoba and others, 2001; Wayne, 1985) and associated loess (Muhs and Bettis, 2000) and may contain substantial amounts of uranium. This also was supported by the eTh and eU data (fig. 5) from aerometric surveys. Because these deposits are genetically similar to the deposits associated with the James lobe (Schumann, 1993b) and the Dakota lobe (Meinzer, 1923; Wayne, 1985), the wells in Nebraska were included in the West-Central glacial aquifers group. The eTh and eU data also were used to differentiate between the western part of the Red River lobe deposits (composed in part of Cretaceous shale) and the eastern part (composed of sediments derived more from granitic materials) (figs. 4 and 5).

Water-Quality Data Sources and Well Selection

NAWQA water-quality data are stored permanently in the USGS National Water Information System (NWIS). In 1999, the NAWQA program developed a data warehouse to better facilitate national and regional analysis of water-quality data from 51 study units started between 1991 and 1997 (Bell and Williamson, 2006). Water-quality data analyzed in this study were obtained from the NAWQA data warehouse in 2004.

The following criteria were used for selection of wells monitored as part of the NAWQA program: (1) wells were installed in glacial, Cambrian-Ordovician, or the New York and New England crystalline aquifers and were sampled as part of the NAWQA program (fig. 2) and (2) the wells had one uranium and(or) 222 radon water-quality data value. A total of 1,426 wells met the above criteria. Nearly 60 percent (817) of these wells had a water-quality value for both uranium and ²²²radon. In the dataset, 1,106 wells have uranium data and 1,137 have ²²²radon data. Within the NAWQA program, these wells also were assigned to a NAWQA well network. These well networks were designed to represent (1) the ground-water quality of major aquifer systems or (2) the immediate effects of a particular land-use setting on groundwater quality in an individual NAWQA study unit and are explained in appendix 1. The major aquifer networks were largely domestic and public-supply drinking-water wells and were selected without consideration of the surrounding land uses. Wells in the land-use networks were typically shallow, 2-in.-diameter monitoring wells, screened at or near the water table, and were selected to target predominantly agricultural or urban land uses within a 500-meter radius. In a few cases, the wells sampled for land-use networks were public-supply

or domestic wells (appendixes 1 and 2). A total of 56 NAWQA well networks were used in this study. On average, each NAWQA well network had about 30 wells, and the water-quality data were collected over an 11-year period, from 1993 to 2003. Statistical summaries for concentrations of uranium and ²²²radon for each NAWQA well network are presented in appendix 2.

Sampling and Analysis

Ground-water samples were collected and analyzed for major ions, nutrients, trace elements, chlorofluorocarbons (CFCs), and other constituents from multiple study units, in accordance with NAWQA protocols (Koterba and others, 1995; Lapham and others, 1995). Isotope data for radium species, ¹⁸oxygen, deuterium, and tritium also were available for some of the study units. Prior to collection of the samples, wells were pumped until the water temperature, dissolved oxygen, specific conductance, and pH were stable (Koterba and others, 1995). Cleaned fluorinated ethylene polypropylene tubing with stainless steel fittings was connected to either a submersible pump or to a water line close to the wellhead. Samples were collected at a flow rate of 0.05 liters per minute (L/min) in a portable sampling chamber. Samples for trace elements (including uranium), nutrients, and major ions were processed through a 0.45-µm disposable polypropylene capsule filter; the trace-element and cation samples were then acidified with ultra pure nitric acid to a pH of less than 2. The analyses for uranium and other trace elements were made at the USGS National Water Quality Laboratory (NWQL) in Denver, Colo., using inductively-coupled plasma mass spectrometry (ICP/MS) methods; laboratory reporting levels (LRL) ranged from 0.2-1 µg/L (Faires, 1993; Ivahnenko and others, 1996). Analyses for ²²²radon were done by liquid scintillation methods; LRLs ranged from 52 to 80 pCi/L (Mullin and Wanty, 1991; Prichard and Gessell, 1977).

Analysis of Ancillary Data and Explanatory Variables

A variety of other data were compiled to analyze the variation of radionuclides within all the aquifer groups. Many of these data were obtained from the USGS NAWQA data warehouse (U.S. Geological Survey, 2005). Ancillary data from geographic information systems (GIS) were also compiled for the analysis of uranium and ²²²radon occurrence. The information compiled included (1) aquifer type—bedrock or glacial unconsolidated, aquifer texture, geographic location and source material; (2) well construction information such as well depth, and well type; (3) population statistics (U.S. Bureau of the Census, 2000); (4) the lithology and lithochemical character of near surface bedrock in New England (Robinson and Kapo, 2003; Robinson, 1997); (5) National Uranium Resource Evaluation Stream Sediment Chemistry

Table 2. Description of aquifer groups within the glaciated northern United States investigated for the occurrence and distribution of radionuclides.

[Groups were compiled and modified from previous studies (Heath, 1984; Meinzer, 1923; Michael and Cothern, 1986; Michael and Jordana, 1987; Schumann, 1993; U.S. Geological Survey, 2000; U.S. Geological Survey, 2003; Wayne, 1985)]

Principal aquifer	l aquifer Aquifer group		Geologic framework	Hydrologic system			
Glacial aquifer system	Northern Pacific glacial aquifers (deposits of the Cordilleran ice sheet)	NPG	Variably mountainous terrain with narrow alluvial and glaciofluvial valleys	Ground-water withdrawals largely from alluvial and glaciofluvial sediments in the valleys; lesser amounts from underlying crystalline-bedrock aquifers.			
Glacial aquifer system	Columbia Plateau glacial aquifers	CPG	Volcanic rocks, thick sequence of lava flows interbedded with unconsolidated deposits, covered in places by sediment of glaciofluvial origin	Semiarid with large irrigation withdrawals from streams and the lava interflow zones; irrigation water moves downward into the ground-water system and then discharges to rivers.			
Glacial aquifer system	West-central glacial aquifers (deposits of the James, Des Moines, and the Red River (western part) glacial lobes)	WCG	Glacial sediments derived mostly from Cretaceous age sedimen- tary rocks consisting of shale, limestone, and sandstone	Largest ground-water withdrawals are from thick glacial deposits that include coarse- grained aquifers and from underlying sedimentary bedrock.			
Glacial aquifer system	North-central glacial aquifers (deposits of the Red River (eastern part) and Wadena, Rainy, Lake Superior, and Green Bay glacial lobes)	NCG	Glacial sediments overlying predominantly Precambrian granite, metamorphic, and volcanic rocks. Also, some areas of limestone	Largest ground-water withdrawals are from coarse-grained glacial deposits and from underlying Precambrian igneous and metamorphic rocks in most of the area. Some sedimentary bedrock in the south and east.			
Glacial aquifer system	East-central glacial aquifers (deposits of the Lake Michigan, Huron, and Huron-Erie glacial lobes)	ECG	Glacial sediments overlying pre- dominantly dolomites, shale, and sandstone	Largest ground-water withdrawals are from coarse-grained glacial deposits.			
Glacial aquifer system	Pre-Wisconsinan glacial aquifers	PWG	Glacial sediments overlying Paleozoic dolomite, carbonate, shale, and sandstone	Largest ground-water withdrawals are from coarse-grained glacial deposits.			
Glacial aquifer system	Northeastern glacial aquifers	NEG	Glacial sediments derived mostly from fractured crystalline rocks	Largest ground-water withdrawals are from coarse-grained glacial deposits and from underlying crystalline bedrock.			
Cambrian-Ordo- vician aquifer system	Cambrian–Ordovician aquifers	COS	Sandstone and dolomite inter- layered with confining units	Ground-water withdrawals primarily from sandstone units but also derive water from vertical leakage through confining layers.			
New York and New England crystalline aquifer system	New York and New England crystalline aquifers	NEC	Predominantly fractured igneous and metamorphic bedrock	Ground-water withdrawals for domestic and public supply from secondary brittle fractures in crystalline rock.			

data (Smith, 2001a; Smith, 2001b); (6) terrestrial gamma mapping data (Duval, 1990); (7) index of recharge to ground water and hydrologic landscape models (Wolock, 2003a; Wolock, 2003b); (8) water-sample temperature, pH, specific conductance, and dissolved oxygen data, and a variety of related water-chemistry data from samples collected concurrently with the radionuclide samples.

Statistical Analyses

A large proportion of the samples had uranium and other water-chemistry data that were reported as "less than the LRL." For each constituent with concentrations below multiple LRLs, the data were examined in order to determine the highest LRL. Reported concentrations above the lowest LRL, but below the highest LRL, were censored to the highest LRL. This enables the use of all the data in nonparametric statistical analyses without making assumptions about the distribution of the data below the LRL (Helsel and Hirsch, 1992). The highest LRL for uranium was 1 µg/L and the highest LRL for ²²²radon was 80 pCi/L. For this study, 62 percent of the uranium data and less than 1 percent of the 222 radon data were below their respective highest LRLs. Nonparametric statistics, which do not require distributional assumptions about data less than LRLs, were used to describe distributions or compare data among groups. Statistical analyses were performed using all data values, including censored data (data reported as below the LRL).

Contingency table analysis (Pearson's Chi Square test for independence) was used to measure the association of uranium and 222 radon (presence or absence) with a grouping variable of two or more categories (Helsel and Hirsch, 1992). Group-comparison tests such as the Wilcoxon and the Kruskal-Wallis tests were used to determine whether the distributions of the data from two or more groups were significantly different (Helsel, 2005; Helsel and Hirsch, 1992; SAS Institute, 1999). Where group-comparison tests indicated significant differences, the Tukey test was used on the ranks of the data to identify which group means were significantly different (Helsel, 2005; Helsel and Hirsch, 1992; SAS Institute, 1999). Although other methods exist for estimating the distribution of missing data and comparing groups, such as substitution and distributional methods (Helsel, 2005), nonparametric methods offer several benefits including (1) no assumption of normally distributed data; (2) good statistical power; and (3) data below the LRL are used without fabrication, resulting in an accurate portrayal of the "less than the LRL" information (Helsel, 1990; Helsel, 2005; Helsel and Cohn. 1988: Helsel and Hirsch, 1992).

Spearman correlation coefficients (rho) were computed to measure the strength of the relation between environmental factors and concentrations of uranium and 222 radon (Helsel and Hirsch, 1992). The significance level (α) for all statistical tests was equal to 0.05.

Evaluations of Quality-Assurance and Quality-Control Data

Water-quality data are subject to bias and variability during sample collection, processing, and analysis. The type and magnitude of bias and variability can be determined by analysis of quality-control samples (Mueller and others, 1997). Field-blank samples can verify that sample collection, processing, and analytical protocols were sufficient to prevent the introduction of contaminants into samples. Replicate environmental samples allow assessment of measurement variability due to sample collection, processing, and laboratory analysis protocols. Quality-control samples, including field blanks and replicates, were collected according to NAWQA protocols (Koterba and others, 1995).

For trace elements including uranium, 130 field blanks (12 percent) were collected for the 1,106 wells sampled. Five of 130 field-blank samples (4 percent) contained a measurable amount of uranium. The highest uranium concentration among those was 0.12 μ g/L—well below the highest LRL of 1 μ g/L used in this study.

Replicate samples for uranium analysis were collected at 65 wells (6 percent of the samples). Replicate samples for ²²²radon analysis were collected at 88 of the 1,137 wells sampled (8 percent). The relative percent difference (RPD) between the environmental sample and corresponding replicate sample was calculated as the absolute value of the difference in replicate concentrations divided by the environmental concentration, multiplied by 100. The RPD of the concentrations between the environmental samples and the corresponding replicate samples was less than 7 percent for uranium and 15 percent for ²²²radon. It was concluded that sample processing and analysis did not introduce enough variation in the environmental data to affect the interpretation of results.

Uranium and ²²²Radon in Glacial and Bedrock Aquifers

Uranium concentrations differ significantly by principal aquifer. The median uranium concentrations for each of the three principal aquifers were less than 1 μ g/L (table 3). Uranium concentrations were highest in the New York and New England crystalline aquifer systems and lowest in the Cambrian-Ordovician aquifer system (table 3). In the glacial aquifer system, uranium concentrations differed between specific geographic regions, and these differences are discussed by aquifer group in the following section. Concentrations of uranium in ground water exceeded 10 μ g/L in about 10 percent of the samples from wells in this study; however, only 2.5 percent of samples exceeded the USEPA MCL of 30 μ g/L for uranium. Thirty-eight percent of all samples had uranium concentrations greater than 1 μ g/L.

Table 3. Summary statistics for concentrations of uranium and ²²²radon, by principal aquifer in the glaciated northern United States, 1993–2003.

[Concentrations of uranium and ²²²radon for principal aquifers with the same Tukey group letter do not differ significantly at the 95-percent confidence level (alpha = 0.05). Max, maximum; Min, minimum; P, percentile (90, 75, 25, 10); <, less than]

Principal aquifer	Number of samples	Max	P90	P75	Median	P25	P10	Min	Tukey group
	Uraniun	n concentration	n, in microgra	ıms per liter	-				
New York and New England crystalline aquifer system	117	428.9	12.8	2.9	< 1	< 1	< 1	< 1	A
Glacial aquifer system	913	162.3	7.5	2.1	< 1	< 1	< 1	< 1	A
Cambrian-Ordovician aquifer system	76	5	3	1	< 1	< 1	< 1	< 1	В
	²²² Rado	n concentratio	n, in picocuri	es per liter					
New York and New England crystalline aquifer system	110	215,210	10,950	4,738	2,122	1,200	565	< 80	A
Glacial aquifer system	950	8,056	899	557	340	219	156	< 80	В
Cambrian-Ordovician aquifer system	77	2,700	940	648	370	230	184	< 80	В

Similarly, ²²²radon differed significantly by principal aquifer. The median ²²²radon concentration was highest in the New York and New England crystalline aquifer system. This concentration was seven times the proposed MCL (2,122 pCi/L). The 75th percentile for ²²²radon (4,738 pCi/L) exceeded the USEPA proposed AMCL of 4,000 pCi/L (table 3). The median ²²²radon concentration was slightly greater than the USEPA proposed MCL of 300 pCi/L in samples from the glacial (340 pCi/L) and Cambrian-Ordovician (370 pCi/L) aquifer systems.

Uranium by Aquifer Group

Concentrations of uranium in ground water differed significantly among the nine aquifer groups analyzed in this study (table 4). The highest concentrations of uranium in ground water were present in samples from the Columbia Plateau glacial aquifer group (CPG) (fig. 7A), the West-Central glacial aquifer group (WCG) (fig. 7A), and the New York and New England crystalline aquifer group (NEC) (fig. 8A).

Table 4. Summary statistics for concentrations of uranium, by aquifer group, for the glaciated northern United States, 1993–2003.

[See table 2 for descriptions of aquifer groups. For aquifer groups with the same Tukey group letter, concentrations of uranium do not differ significantly at the 95-percent confidence level (alpha = 0.05). Max, maximum; Min, minimum; P, percentile (90, 75, 25, 10); <, less than]

	Nhawaf	Concentration of uranium, in micrograms per liter									
Aquifer group (code)	Number of — samples	Max	P90	P75	Median	P25	P10	Min	Tukey group		
Columbia Plateau glacial (CPG)	46	67.9	29.9	22.3	11.9	6.0	3.8	< 1	A		
West-Central glacial (WCG)	177	162.3	31.9	7.7	2.9	1	< 1	< 1	В		
New York and New England crystalline (NEC)	117	428.9	12.8	2.9	< 1	< 1	< 1	< 1	С		
East-Central glacial (ECG)	283	21.1	3.3	1.4	< 1	< 1	< 1	< 1	CD		
Cambrian-Ordovician (COS)	76	5	3	1	< 1	< 1	< 1	< 1	CD		
North-Central glacial (NCG)	171	11.9	2.2	< 1	< 1	< 1	< 1	< 1	D		
Pre-Wisconsinan glacial (PWG)	71	17	2	< 1	< 1	< 1	< 1	< 1	D		
Northeastern glacial (NEG)	126	15.8	1.3	< 1	< 1	< 1	< 1	< 1	D		
Northern Pacific glacial (NPG)	39	< 1	< 1	< 1	< 1	< 1	< 1	< 1	D		

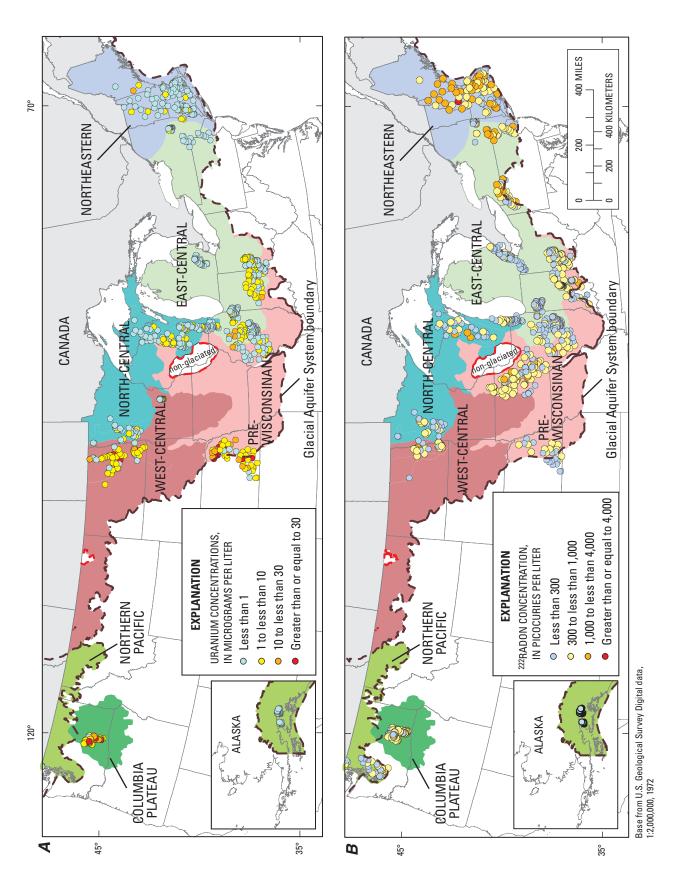


Figure 7. Distribution of (A) uranium and (B) 22radon concentrations in sampled wells in the glacial aquifer groups in the northern United States, 1993–2003.

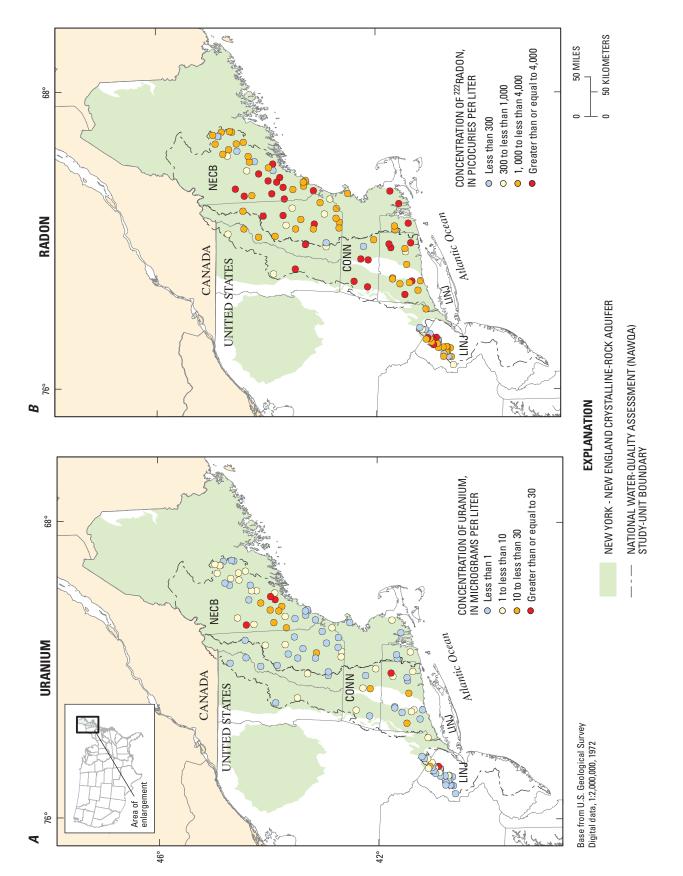


Figure 8. Distribution of (A) uranium and (B) 22 radon concentrations in sampled wells in the New York and New England crystalline aquifer group in the northern United States, 1993-2003.

Concentrations of uranium exceeded 10 μ g/L in more than 50 percent of ground-water samples from wells in the CPG aquifers (table 4), and about 10 percent of the samples from the CPG and WCG aquifers exceeded the USEPA MCL of 30 μ g/L. More than 10 percent of the wells sampled in the NEC aquifers had concentrations greater than 10 μ g/L. The greatest concentration, 429 μ g/L, was measured in a sample from a domestic bedrock well completed in the NEC aquifer group.

Concentrations of uranium in ground-water samples from North-Central glacial aquifer group (NCG), which are underlain mostly by Precambrian crystalline rocks, were less than 12 µg/L. The sediments that comprise the WCG aquifer group, immediately to the west of the NCG aquifers, were derived in part from the northwest and contain appreciable amounts of Cretaceous shale that is uraniferous. This difference may affect uranium source and(or) redox (Bohlke and others, 2002) and may contribute to the high concentrations of uranium in wells in the northern part of the WCG aquifers and the low concentrations of uranium in the wells in the NCG aquifers (figs. 5 and 7A). None of the samples from wells in Northern Pacific glacial aquifer group (NPG) (5 wells in Washington and 34 in Alaska) had uranium concentrations greater than 1 µg/L. The highest value reported for 76 wells in the Cambrian-Ordovician (COS) aquifer group was 5 μg/L (table 4, fig. 9A).

Results from a multiple comparison test showed that ground-water samples from wells in CPG aquifers had significantly higher uranium concentrations than samples from all other aquifer groups (table 4). Water samples from wells in WCG aquifers had significantly higher uranium concentrations than samples from all other aquifer groups, except for CPG aquifers. Whereas the NEC aquifer group had some individual samples with high uranium concentrations (greater than 30 $\mu g/L$), as a group, uranium concentrations in the NEC aquifer group were only significantly higher than those for NCG, Pre-Wisconsinan glacial (PWG), NPG, and NEG aquifer groups.

²²²Radon by Aquifer Group

Concentrations of ²²²radon in ground water from some of the nine aquifer groups also differed significantly. The highest ²²²radon concentrations were present in samples from wells completed in NEC aquifers; the median value (2,122 pCi/L) was about 10 times the median values of all other aquifer groups (table 3; fig. 8B). More than 30 percent of the 110 samples from wells in NEC aquifers had ²²²radon concentrations that exceeded the proposed USEPA AMCL of 4,000 pCi/L. The second highest median concentration (814 pCi/L) was in samples from the NEG aquifers, which were predominantly derived from and overlie the NEC aquifer. The NEG aquifer group was the only glacial aquifer group with a water sample having a ²²²radon concentration exceeding the proposed USEPA AMCL of 4,000 pCi/L. In

CPG aquifers, the median and 90th percentile concentrations of ²²²radon were 530 and 1,100 pCi/L, respectively. For all other aquifer groups (WCG, ECG, COS, NCG, NPG, and the PWG), the median value of ²²²radon was about 300 pCi/L and the 90th percentile values did not exceed 1,000 pCi/L (figs. 7B and 9B); these groups had similar ²²²radon concentrations.

Comparison of Uranium and ²²²Radon in Drinking-Water Supply and Monitoring Wells

Concentrations of uranium and 222 radon were evaluated for differences between monitoring wells representing particular land uses, and public- and domestic-supply wells used for drinking water (termed drinking-water wells). In this study, uranium concentrations were significantly higher in samples from monitoring wells than from drinking-water wells (p < 0.0001) in the ECG aguifers. However, uranium concentrations were significantly higher in drinking-water wells in the NEG (p = 0.0374) and the WCG (p = 0.0210) aquifer groups than in monitoring wells for these groups. Uranium concentrations did not differ significantly between samples from drinking-water and monitoring wells in the NCG, NPG, or PWG aquifer groups. This information is useful when evaluating uranium concentrations between aquifer groups that include a combination of drinking-water and monitoring-well data because it indicates that well type may also play a role in differences in uranium concentrations.

Concentrations of ²²²radon also differed by well type within the aquifer groups. ²²²Radon concentrations were significantly higher in samples from drinking-water wells than those from monitoring wells in the CPG aquifers and NEG aquifers (p = 0.0007 and 0.0078, respectively). ²²²Radon concentrations were significantly higher in monitoring wells than in drinking-water wells in the PWG and WCG aquifers (p = 0.0001 and 0.0257, respectively). For the ECG, NCG, and NPG glacial aquifer groups, no significant difference was observed between concentrations of ²²²radon in samples from drinking-water wells and monitoring wells.

Concentrations of uranium exceeded the USEPA maximum enforceable contaminant levels in some of the aquifer groups (table 6). Twelve percent of water samples from drinking-water wells in WCG aquifers and 4 percent of water samples from the wells in NEC aquifers exceeded the USEPA MCL of 30 μ g/L for uranium. Eleven percent of water samples from monitoring wells in WCG aquifers and 9 percent of samples from the monitoring wells in CPG aquifers exceeded the USEPA MCL of 30 μ g/L for uranium. Data for drinking-water wells were not available for the CPG aquifers and monitoring-well data were not available for the NEC or COS aquifers.

²²²Radon was measured at concentrations near or above the proposed USEPA MCL of 300 pCi/L in nearly 50 percent or more of all water samples, regardless of well type or aquifer group. However, 33 of the 34 water samples with ²²²radon concentrations greater than the proposed USEPA

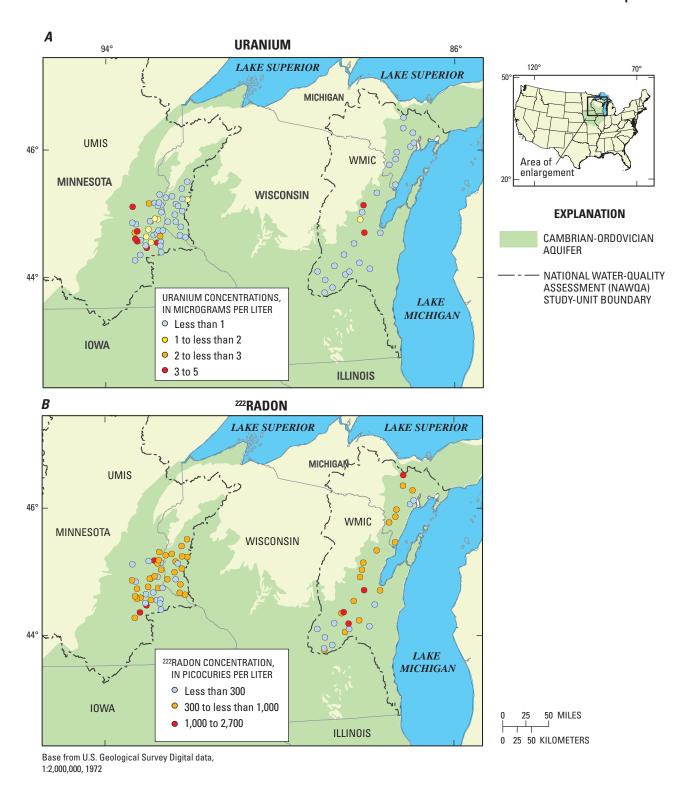


Figure 9. Distribution of (A) uranium and (B) 222 radon concentrations in sampled wells in the Cambrian-Ordovician aquifer group in the northern United States, 1993–2003.

Table 5. Summary statistics for concentrations of 222 radon, by aquifer group, in the glaciated northern United States, 1993–2003.

[See table 2 for descriptions of aquifer groups. Aquifer groups with the same Tukey group letter do not differ significantly at the 95-percent confidence level. N, number of samples; Max, maximum; Min, minimum; P, percentile (90, 75, 25, 10); <, less than]

Aifa	Nhanaf	Concentration of ²²² radon, in picocuries per liter									
Aquifer group (code)	Number of samples	Max	P90	P75	Median	P25	P10	Min	Tukey group (alpha = 0.05)		
New York and New England crystalline (NEC)	110	215,210	10,950	4,738	2,122	1,200	565	< 80	A		
Northeastern glacial (NEG)	93	8,056	2,087	1,269	814	508	344	< 80	В		
Columbia Plateau glacial (CPG)	85	1,900	1,100	710	530	320	265	110	C		
Cambrian-Ordovician (COS)	77	2,700	940	648	370	230	184	< 80	CD		
West-Central glacial (WCG)	76	1,100	626	394	313	245	217	83	DE		
Pre-Wisconsinan glacial (PWG)	175	2,146	767	454	322	210	150	< 80	DE		
East-Central glacial (ECG)	337	2,800	810	522	302	187	138	< 80	E		
Northern Pacific glacial (NPG)	77	860	540	439	290	200	146	100	E		
North-Central glacial (NCG)	107	2,436	463	360	252	184	152	< 80	Е		

Table 6. Percentage of drinking water and monitoring wells with concentrations of uranium and 222 radon greater than the U.S. Environmental Protection Agency Maximum Contaminant Level (MCL) for uranium and proposed MCL and AMCL for 222 radon, by aquifer group and well type, in the northern United States.

[See table 2 for descriptions of aquifer groups. N, number; >, greater than; µg/L, micrograms per liter; pCi/L, picocuries per liter; –, no data]

		Drinking wate	r (public	and domestic	Monitoring (unused) wells						
Aquifer group (code)	ı	Jranium		²²² Radon			Jranium		²²² Radon		
	N	Percentage >30 µg/L	N	Percentage >4,000 pCi/L	Percentage >300 pCi/L	N	Percentage >30 µg/L	N	Percentage >4,000 pCi/L	Percentage >300 pCi/L	
Northern Pacific glacial (NPG)	31	0	57	0	46	5	0	17	0	65	
Columbia Plateau glacial (CPG)	-	-	44	0	89	46	9	39	0	72	
West-Central gla- cial (WCG)	52	12	35	0	40	116	11	41	0	73	
North-Central glacial (NCG)	74	0	72	0	33	93	0	31	0	32	
East-Central glacial (ECG)	70	0	123	0	45	209	0	209	0	53	
Pre-Wisconsinan glacial (PWG)	59	0	88	0	41	8	0	83	0	66	
Cambrian-Ordovician (COS)	73	0	74	0	66	-	_	_	_	_	
Northeastern glacial (NEG)	67	0	61	2	92	55	0	29	0	93	
New York and New England crystal- line (NEC)	113	4	107	30	94	-	-	-	-	-	

AMCL of 4,000 pCi/L were from drinking-water wells in the northeastern United States. Thirty-two of the 33 samples were from domestic supply wells in the NEC aquifer group. The remaining water sample with high radon (8,056 pCi/L) was from a drinking-water well in the NEG aquifer group.

Data were available for five aquifer groups (ECG, WCG, NPG, COS, NEG) to evaluate differences in uranium and 222 radon concentrations in water samples between public and domestic wells used to supply drinking water. Concentrations of uranium did not differ significantly between public and domestic wells (p = 0.1784 to 0.7235) in any of the five aquifer groups. Similarly, 222 radon concentrations in public and domestic wells in the aquifer groups COS, ECG, NEG, and NPG were not significantly different (p = 0.0670 to 0.7344).

Relation of Uranium and ²²²Radon in Ground Water to Terrestrial Gamma-Ray Emissions

National-scale data on the equivalent uranium (eU) and thorium (eTh) content of near-surface sediments, determined from gamma-ray emissions measured during an aerometric survey (Duval, 1990), were compared to uranium and 222 radon concentrations measured in the associated ground water. More than 50 percent of the uranium concentrations measured in ground water were below the reporting level; therefore, the 75th percentiles of uranium and ²²²radon concentrations in ground water from each of the NAWQA well networks (appendix 2) were plotted against the 75th percentiles of eU and eTh concentrations from the aerometric gamma-ray surveys that were associated with each network (fig. 10). The eTh values may provide better correlations with uranium and(or) 222 radon concentrations in ground water than eU values because thorium is much less mobile than uranium in the environment and, thus, less susceptible to error related to secular disequilibrium in the sediment environment (Osmond and Ivanovich, 1992). Concentrations of uranium in ground water are somewhat related to eU and eTh concentrations in near-surface sediments. Where eU concentrations were greater than 1.25 ppm (parts per million) (fig. 10A), uranium concentrations in water varied and were often high (75th percentile > 10 µg/L). Similarly, where eTh concentrations were greater than 5 ppm (fig. 10B), uranium concentrations in ground water were high (75th percentile > $10 \mu g/L$); where eTh concentrations were less than 5 ppm, uranium concentrations in ground water were consistently low. The concentrations of ²²²radon generally increased with increasing eU and eTh concentrations, although ²²²radon concentrations were consistently low where eU concentrations were less than about 2.0 ppm. Where eU concentrations were greater than 2.0 ppm, ²²²radon concentrations increased sharply. This response is due largely to samples from New England in both the glacial (NEG) and crystalline (NEC) aquifer groups and from samples in the ECG (figs. 10C and 10D). Samples from the PWG aquifers had high eU and eTh values but consistently low concentrations of 222 radon, indicating that processes such

as redox may affect intermediate radionuclides (for example, ²²⁶radium) from which ²²²radon is derived. Thus, the relation between uranium and ²²²radon in ground water and terrestrial gamma-ray emissions is not consistent but provides an indication of potential for the presence of ²²²radon and, less so, uranium, in ground water.

Correlations of Uranium and ²²²Radon with Ancillary Data

Uranium and ²²²radon concentrations were correlated with ancillary data, including water chemistry, streambed-sediment chemistry, terrestrial gamma-ray emissions, and physical and hydrologic properties. Significant correlations of uranium and ²²²radon with ancillary data, by aquifer group, are listed in appendixes 3 and 4, respectively. Significant correlation coefficients for ground-water chemistry constituents that were commonly correlated with uranium, by aquifer group, are summarized in table 7. These correlations differed by aquifer group and may relate to large-scale hydrologic and geochemical processes that affect the concentrations of uranium and ²²²radon in ground water.

Uranium and ²²²radon concentrations in ground water correlated with concentrations of many other water-quality constituents; however, the constituents that correlated with uranium and ²²²radon differed among the aquifer groups. Some of the correlations were similar among aquifer groups, whereas others were unique to a specific aquifer group. The differences in the types of ancillary data that correlated could be due to differences in geology or the source of the geologic materials but also may be affected by the geochemical behavior of uranium and ²²⁶radium. Uranium readily forms complexes with several anions, including carbonate, sulfate, nitrate, chloride, and phosphate (Sorg, 1991), and correlations between uranium and anions may provide insights to the processes that affect uranium concentrations in the various aquifer groups. Major ions and trace elements were commonly correlated with uranium. Calcium correlated with uranium (p < 0.05) in eight of the nine aquifer groups, and sulfate correlated with uranium in seven of the nine aguifer groups. Total dissolved solids (TDS), acid neutralizing capacity (ANC), and bicarbonate (HCO₂) also correlated with uranium in most of the aquifer groups (table 7).

In the CPG and WCG glacial aquifer groups, uranium concentrations correlated with boron, lithium, selenium, and strontium (table 7). These correlations were strikingly similar to ones found in studies of uranium in surface waters in the western United States affected by irrigation (Seiler and others, 2003; Zielinski and others, 1995). Both studies cite the combination of geologic sources of uranium or selenium and heavy irrigation of agricultural land in a semiarid climate as possible explanations for mobilization of these trace elements. In the CPG and in the WCG aquifer groups, there are abundant natural sources of uranium in the aquifer sediments, and the re-use of irrigation waters from either ground water or surface water sources is a common practice.

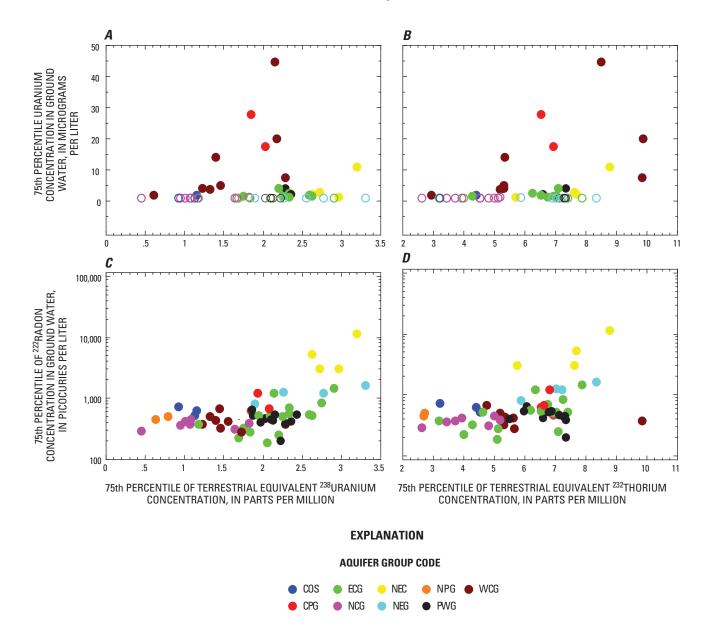


Figure 10. Relation of the 75th percentile of concentrations of uranium in ground water to the 75th percentile concentrations of terrestrial (A) equivalent ²³⁸uranium and (B) equivalent ²³²thorium; and 75th percentile of concentrations of ²²²radon in ground water to 75th percentile concentrations of terrestrial (C) equivalent ²³⁸uranium and (D) equivalent ²³²thorium in ground-water samples from wells in glacial, Cambrian-Ordovician, and New York and New England crystalline aquifer groups in the northern United States. Open circles indicate that 75th percentile for uranium was <1 microgram per liter.

Table 7. Spearman correlation coefficients between uranium and selected water-chemistry constituents, by aquifer group, in the northern United States.

[Aquifer group codes are defined in table 2; Values in bold have *p*-values less than 0.05; –, no data; ANC, Acid neutralizing capacity; B, boron; Ca, calcium; HCO₃, bicarbonate; Li, lithium; Mn, manganese; Ni, Nickel; Se, selenium; SO₄, sulfate; Sr, strontium; TDS, total dissolved solids; ²²²Rn, ²²²radon; alpha, gross alpha concentration]

Aquifer group code	ANC	В	Ca	HCO ₃	Li	Mn	Ni	Se	SO ₄	Sr	TDS	²²² Rn	alpha
NPG	_	_	_	_	_	_	_	_	_	_	_	_	_
CPG	0.68	0.42	0.31	0.77	0.47	0.22	0.31	0.37	0.51	0.52	0.68	0.18	_
WCG	0.40	0.36	0.37	0.37	0.43	-0.31	-0.01	0.60	0.27	0.32	0.39	0.06	_
NCG	0.18	0.04	0.19	0.24	0.20	0.18	0.42	0.19	0.20	0.14	0.17	0.10	0.06
ECG	0.35	0.18	0.40	0.20	0.11	0.46	0.43	0.16	0.41	0.23	0.36	-0.19	_
PWG	-0.16	-0.45	0.47	_	_	-0.04	0.45	0.13	0.56	0.61	0.29	0.41	_
NEG	0.31	-0.03	0.29	0.30	0.16	-0.01	-0.01	_	0.37	0.22	0.19	0.01	0.54
COS	0.48	-0.14	0.42	_	-0.22	0.36	0.47	0.06	0.32	0.02	0.27	0.09	_
NEC	0.18	-0.05	0.25	-0.03	0.24	-0.18	-0.01	-0.13	-0.03	0.10	0.14	0.60	0.72

Uranium correlated with nickel in glacial aquifer groups in the central United States (NCG, ECG, and PWG). Uranium also correlated with nickel in the COS aquifer group, although uranium concentrations were generally low in this aquifer group. The low uranium concentrations should be considered, however, in conjunction with information on other radio-nuclides, particularly ²²⁶radium, which is known to be high in parts of this aquifer system (Gilkeson and Cowart, 1987; Michael and Jordana, 1987; Saad, 1996).

²²²Radon correlated with uranium in the NEC and PWG aquifer groups, but not in any of the other aquifer groups. These correlations may be related to hydraulic properties of the NEC and PWG aquifers, such as the proximity of the sources to fractures that transmit water or by the geochemical behavior of uranium and ²²⁶radium.

Many of the variables that did not correlate significantly, or did not correlate consistently, by aquifer group, with uranium or ²²²radon are listed in appendixes 3 and 4. Examples of these include well depth, nitrate, dissolved oxygen, trace elements, and some of the NURE stream sediment data. There are a variety of possible explanations why significant correlations did not occur between uranium and 222 radon and other data. One reason may be that some of the ancillary data were not complete for the study area, such as the NURE streambed sediment chemistry data (Smith, 2001a; Smith, 2001b). For example, there were 283 uranium samples for the ECG aquifer group, but only 43 of these samples were from wells that intersected with areas containing NURE data. Another is that some trace elements, such as beryllium and cadmium, were not detected above laboratory reporting levels in 99 percent of the water samples. Also, some areas had no uranium or radon data or, as was the case in the Northern

Pacific aquifer group, none of water samples had uranium concentrations above 1 $\mu g/L$.

Anomalously High Uranium and ²²²Radon Concentrations and Related Factors

Uranium and 222 radon concentrations were anomalously high in samples from some aquifers in the northern United States. In this section, aquifer groups with large percentages of wells with uranium concentrations that exceeded the USEPA MCL of 30 μ g/L and 222 radon concentrations that exceeded the proposed USEPA AMCL of 4,000 pCi/L are examined in more detail. Three areas in particular were examined—the NEC, WCG, and the CPG aquifers. The distributions of the concentrations, as well as correlation to other factors, are presented.

New York and New England Crystalline Aquifer Group

Wells in the New York and New England crystalline (NEC) aquifer group (figs. 6 and 8) are predominantly domestic wells that intersect water-bearing fractures at various depths in the bedrock (Flanagan and others, 1999). Land use near the sampled wells is predominantly rural, forested land with minor amounts of residential and agricultural lands. The NEC aquifer group followed only the CPG and WCG aquifer groups for highest uranium concentrations (table 4) and ranked the highest among all groups for ²²²radon concentrations (table 5).

In the New England part of the NEC aquifer group, the uranium and 222 radon concentrations correlated with each other and with other radionuclide data such as gross alpha and gross beta particle activities, but not with most other constituent concentrations. Most notably, uranium concentrations correlated strongly with gross alpha (rho = 0.72, p < 0.0001) (appendix 3) and with 222 radon concentrations (rho = 0.60, p < 0.0001) (appendix 3). This relation was not found in other parts of the United States, and the primary reason is that ²²²radon is not subject to the same geochemical controls as uranium. In a Finnish study, however, ²²²radon concentrations in wells in similar bedrock formations, primarily granites or granitoid crystalline rocks, correlated with ²³⁸uranium concentrations (rho = 0.57, p < 0.0001) (Vesterbacka and others, 2005).

Results of a recent study for the mobilization of ²²²radon in fractured crystalline aquifers indicated that a diffusion/ion exchange model may help explain the anomalous concentrations of ²²²radon in ground water; in this model, ²²⁶radium diffuses toward the water-bearing fracture and decays to ²²²radon (Wood and others, 2004). They noted that results from a brine injection test showed that the activity of ²²⁶radium was 34 times greater than the activity of uranium at the fracture face. The correlation between uranium and ²²²radon in ground water determined in this study, especially in the ground waters in granitoid rocks (appendix 5, necbsus2 NAWQA well network), indicates that at least some of the ²²⁶radium, if present, may be derived from uranium at or near the fracture surface. Alternatively, ²²⁶radium may be less soluble than uranium under slightly oxic or mixed geochemical conditions typical of these aquifers. Uranium in samples from wells in the NEC aquifer group also correlated with gross beta particle activity (0.33, p = 0.0120) and with concentrations of molybdenum (rho = 0.35, p = 0.0001) (appendix 3).

²²²Radon also correlated with gross alpha particle activity (rho = 0.54, p < 0.0001; fig. 11B) and with total lead (rho = 0.54, p < 0.0001; fig. 11B)0.38, p = 0.0005; fig. 11D). The correlation between 222 radon and total lead was strongest (rho = 0.52, p = 0.004) in samples from the granitic and metamorphic crystalline aguifers of eastern New England, part of the NEC aquifer group (fig. 11D; appendix 6). This result is similar to a study in Finland that found a strong correlation (rho = 0.74, p < 0.0001) between ²²²radon and ²¹⁰lead in wells completed in crystalline aquifers (Vesterbacka, 2005; Vesterbacka and others, 2005). Lead can be present in ground water in two ways: it can be mobilized by the dissolution of the rock matrix or it can be formed in the water from the radioactive decay of 222 radon. Because 222 radon concentrations were anomalously high in the water from wells in this aguifer, the total lead concentrations measured in this study may be related, in part, to radiogenic lead (²⁰⁶Pb and ²⁰⁸Pb) dissolved, desorbed, or leached from labile phases in fractures or pore spaces in the rock, or alternatively, these rocks might simply be enriched in lead. In areas where ²²²radon in ground water is particularly high, it is possible that a minor amount of the lead (210Pb) associated with 222radon decay may remain dissolved in the water (Focazio and others,

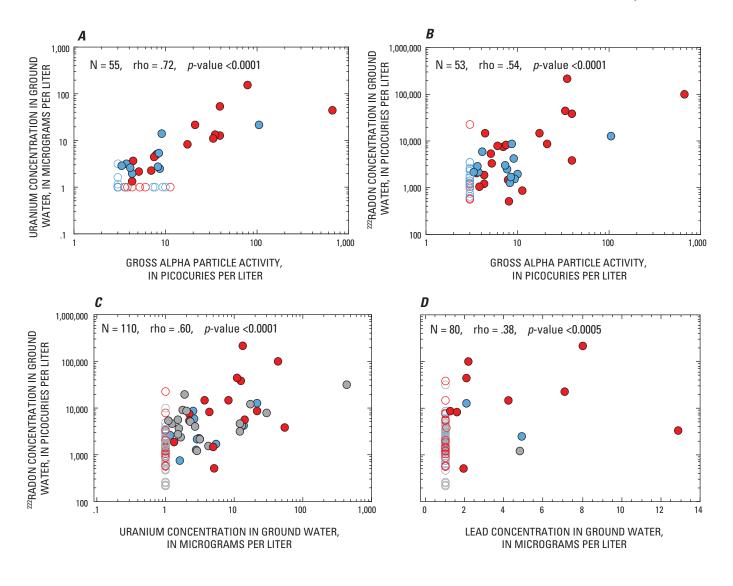
2001) although isotopic measurement, not simple mass measurement, is needed to confirm this hypothesis. Disequilibrium ratios could not be determined for this study because isotopic data for uranium and lead were not collected.

In New England, crystalline bedrock that is granitic, such as two-mica granites, as well as other high-grade metamorphic rocks, appears to contain abundant uranium that is soluble in the predominantly sub-oxic to oxic and high-pH geochemical conditions (Boudette, 1977). In New Hampshire, a study of concentration data from self-selected domestic well samples sent to the New Hampshire Department of Environmental Services laboratory for radon analysis showed results similar to those found in this study for the NEC aquifer group (Moore, 2004). The New Hampshire study found that 40 percent of the domestic well-water samples, primarily from the southern part of the state, exceeded the 4,000 pCi/L proposed USEPA AMCL for ²²²radon; 30 percent of the public and domestic well-water samples from the NEC aquifer group exceeded the proposed AMCL (table 8).

Concentrations of uranium and 222 radon in drinking water from public and domestic wells in the NEC aguifer group were similar to those from domestic wells completed in bedrock in parts of several northern European countries, including Norway, Sweden, and Finland (Vesterbacka, 2005). In these countries, bedrock formations are similar to those found in the NEC. A comparison of radionuclide data from the NEC aquifer in this study with data from studies in these European countries is shown in table 9. In the Norwegian study, 3,494 domestic bedrock wells were sampled. The wells are completed in rock formations that include uranium-rich granites in the southeast part of Norway where the portion of the population obtaining drinking water from domestic bedrock wells has increased (Strand and others, 1998). In southeastern Sweden, the highest concentrations of 222 radon and uranium were in samples from drinking-water wells completed in primarily granite bedrock formations and granite-pegmatite mixes (Isam Salih and others, 2002).

West-Central Glacial Aguifer Group

Ground water in the West-Central glacial (WCG) aquifer group generally had high concentrations of uranium, and these high uranium concentrations may be affected by the source material of the glacial sediments. Land use near the sampled wells included urban, agricultural, and mixed land uses. The WCG aquifers are derived from Cretaceous-age rocks composed predominantly of shale and some sandstone and limestone. Parts of North and South Dakota, western Minnesota, and northeastern Nebraska are underlain by glacial materials deposited in association with the Des Moines, James, and Red River glacial lobes (Meinzer, 1923; Schumann, 1993b; Wayne, 1985). Despite the complexity of these glacial deposits, much of the material contained within them is derived from Cretaceous-age, uraniferous shale. The Pierre Shale, for example, contains higher amounts of uranium than the average crustal abundance of 2.5 mg/kg (Schumann,



EXPLANATION

LITHOLOGY OF NEW YORK AND NEW ENGLAND CRYSTALLINE AQUIFER GROUP

- METASEDIMENTARY, VARIABLY CALCAREOUS
- FELSIC IGNEOUS AND METASEDIMENTARY, UNDIFFERENTIATED
- CRYSTALLINE ROCK, UNDIFFERENTIATED

Figure 11. Relation of concentrations of (A) uranium and (B) ²²²radon to gross alpha particle activity; and concentrations of ²²²radon to (C) uranium and (D) lead in ground-water samples from wells in the New York and New England crystalline aquifer group in the northern United States, 1993–2003. Open circles indicate that the concentration is less than the laboratory reporting limit.

Table 8. Percentage of wells with uranium and ²²²radon concentrations exceeding selected standards and proposed standards, by lithology, for the New York and New England crystalline aquifer group.

[See appendix 1 for description of National Water-Quality Assessment Program (NAWQA) networks. NA, not applicable]

Lithology	NAWQA well	Number of	Percentage of wells with uranium greater tha (micrograms per liter)				
	network code	samples –	30	20	10	1	
Felsic igneous and undifferentiated metasedimentary rocks	necbsus2	30	10	13	27	57	
Metasedimentary rocks, variably calcareous	necbsus1	28	0	4	7	50	
Crystalline, undifferentiated	connsus1, linjsus1	59	3	3	10	37	
Total	NA	117	4	6	13	45	

Lithology	NAWQA well	Number of	Percenta	eater than		
	network code	samples -	20,000	4,000	300	80
Felsic igneous and undifferentiated metasedimentary rocks	necbsus2	28	18	46	96	96
Metasedimentary rocks, variably calcareous	necbsus1	26	0	15	100	100
Crystalline, undifferentiated	connsus1, linjsus1	56	4	29	91	100
Total	NA	110	6	30	94	99

Table 9. Comparison of maximum and mean concentrations of uranium and ²²²radon in ground water from areas dominated by crystalline aquifers in New England, United States; Norway; Sweden; and Finland.

[–, no data; N, number of samples; $\mu g/L$, micrograms per liter; pCi/L, picocuries per liter]

Country	N	Maximum	Mean	Reference
	l	Jranium (μg/L)		
Finland	288	800	20.9	(Vesterbacka and others, 2005)
Sweden	328	427	14.3	(Isam Salih and others, 2002)
Norway	_	_	_	_
New York and New England crystalline aquifers group, United States	113	4,291	4.51	This study
	2	²² Radon (pCi/L)		
Finland	288	232,432	12,432	(Vesterbacka and others, 2005)
Sweden	328	219,054	6,946	(Isam Salih and others, 2002)
Norway	3,494	864,864	10,541	(Strand and others, 1998)
New York and New England crystalline aquifers group, United States	107	215,210	7,360	This study

¹ Value represents total uranium rather than ²³⁸uranium and the mean is estimated on the basis of a log-normal distribution because of censored data (Helsel, 1990).

1993b). Thus, glacial aquifers in this region may contain appreciable amounts of uranium that is available for dissolution in ground water.

In addition, much of eastern Nebraska was occupied by pre-Wisconsinan ice of the Dakota and Minnesota lobes (Colgan, 1999; Meinzer, 1923; Srhoba and others, 2001; Wayne, 1985). The Dakota lobe in particular may have carried and deposited uraniferous shale to eastern Nebraska and northeastern Kansas. Loess deposits from the western United States (Muhs and Bettis, 2000) may also have contributed uranium to aquifers that resulted in high concentrations of uranium in ground water in the Nebraska wells (fig. 7A).

Ten percent (17 of 177) of samples in the WCG aquifer group contained uranium concentrations greater than the USEPA MCL of 30 μ g/L; the highest uranium concentration was 162.3 μ g/L. Although this aquifer group had significantly higher uranium concentrations than other aquifer groups studied (table 4), it had one of the lowest ²²²radon distributions (table 5). The highest ²²²radon concentration measured in this aquifer group was 1,100 pCi/L.

The strongest correlations for uranium were with selenium (rho = 0.60), carbonate (CO $_3$) ion (rho = 0.52), potassium, lithium, and calcium (rho = 0.52, 0.43, and 0.37, respectively), acid neutralizing capacity (rho = 0.40), and total dissolved solids (rho = 0.39) (appendix 3). Uranium correlated somewhat with sodium (rho = 0.40) and nitrate (rho = 0.34). Uranium was inversely correlated with total iron (rho = -0.46), indicating redox controls. Uranium also correlated with terrestrial gamma-ray emissions: ⁴⁰potassium (rho = 0.46) and eTh (rho = 0.43).

²²²Radon in the WCG aquifer group inversely correlated with well depth, depth to top of open interval, and depth to bottom of open interval (rho = -0.35, -0.36, and -0.35, respectively) (appendix 4). Also, ²²²radon correlated positively with tritium (rho = 0.35), and inversely with 18 oxygen (^{18}O) (rho = -0.34). These correlations indicate that water in these aquifers may be young. Young ground water may be an indication of high transmissivities (high hydraulic conductivity) and may indicate that there is a pathway for ²²²radon, thus increasing concentrations in ground water. Alternatively, young ground water may have the redox properties necessary to mobilize uranium, or the water may be affected by agricultural practices such as irrigation. Inverse correlations were observed between 222 radon and terrestrial gamma-ray measurements of 40 potassium and eTh (-0.32 and -0.34, respectively).

Within the WCG aquifer group, concentrations of uranium in ground water generally were higher in samples from the wells in Nebraska (NAWQA well networks cnbrluscr1 and hpgwdwgs1; appendix 2) than in the wells in eastern North Dakota and western Minnesota (NAWQA well networks rednlusag2, rednsus1, rednsus3, and rednsus5; appendix 2). This may be a result of available uranium in aquifer materials in combination with enhanced mobility from evaporative concentration associated with agricultural irrigation. In general, high uranium concentrations correlated with high

total dissolved solids concentrations. This correlation was strongest for the ground water in shallow wells in an agricultural cropland setting in northeastern Nebraska (NAWOA well network cnbrluscr1; appendix 5). These wells tap shallow ground water in predominantly corn and soybeans croplands (appendix 1). Uranium in ground water from these wells correlated most strongly with major ions, including lithium (rho = 0.81), boron (rho = 0.79), potassium (rho = 0.71), sodium (rho = 0.57), strontium (rho = 0.51), and selenium (rho = 0.50) (appendix 5). In general, the samples from wells in eastern North Dakota, South Dakota, and Minnesota had lower uranium concentrations than those from the northeastern Nebraska wells, but several of the wells had samples with concentrations exceeding or approaching the USEPA MCL of 30 µg/L. Uranium concentrations in these wells also correlated significantly with total dissolved solids, magnesium, potassium, and other major ions (appendix 5).

The relations of uranium to other dissolved constituents in the samples from the WCG aquifers were similar to those in surface water in the Arkansas River in eastern Colorado (Zielinski and others, 1995). In this study, the ground water discharging to the river (as irrigation return flow) was affected by geologic sources of uranium in shale bedrock, enhanced uranium leaching from irrigation water, and evaporative concentration from the semiarid climate. For the WCG aquifer samples, graphs of uranium with calcium, lithium, boron, sodium, strontium, and selenium illustrate significant correlations (fig. 12) among those constituents and, except for sulfate and chloride, the correlations were similar to those for surface waters in the Arkansas River study.

In the Arkansas River study, uranium concentrations increased in the river with increasing distance from its source in the Rocky Mountains and with increasing inputs of ground-water discharge from aquifers that underlie irrigated agricultural lands (Zielinski and others, 1995). It is possible that one factor related to the high uranium concentrations in ground water in the WCG aquifer group also is recharge from irrigated agricultural lands.

Columbia Plateau Glacial Aquifer Group

Columbia Plateau glacial aquifers (CPG) (fig. 6) are composed of shallow to deep glaciofluvial sediments overlying sequences of lava flows interbedded with unconsolidated paleo-lakebed deposits (U.S. Geological Survey, 2000). Wells in the CPG aquifer group are located in predominantly agricultural land. Shallow monitoring wells in this group have associated uranium data (appendixes 1 and 2). Uranium data are not available for the deep domestic wells (appendix 2). CPG aquifers are affected by irrigation for agriculture where surface-water irrigation is a main source of recharge to underlying aquifers (Embrey, 1988). This may indirectly affect the concentrations of uranium in shallow ground water because agricultural irrigation usually results in increased concentrations of dissolved solids, nitrate, and salts in the underlying aquifer.

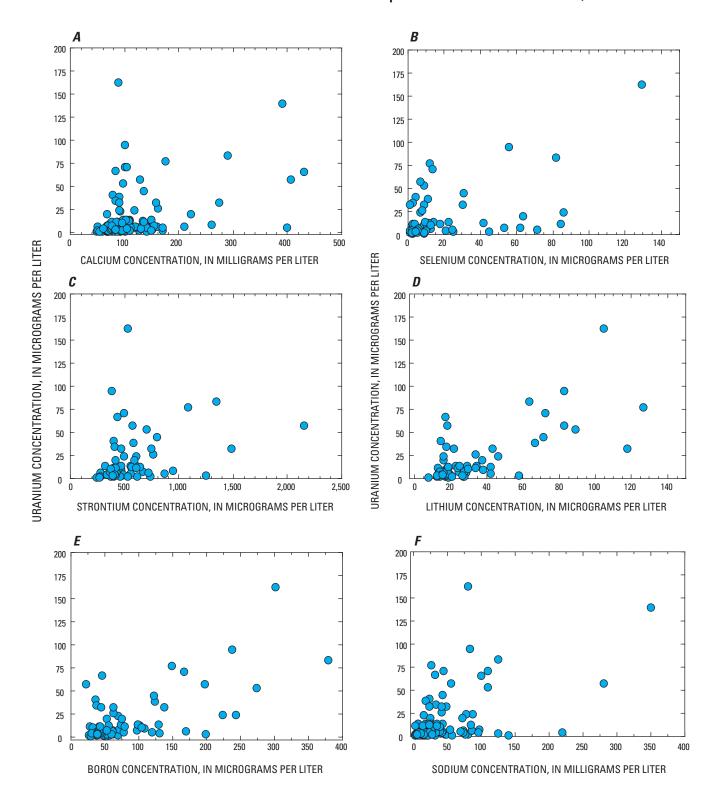


Figure 12. Relation of concentrations of uranium to concentrations of (*A*) calcium, (*B*) selenium, (*C*) strontium, (*D*) lithium, (*E*) boron, and (*F*) sodium in ground-water samples from the West-Central glacial aquifer group in the northern United States, 1993–2003.

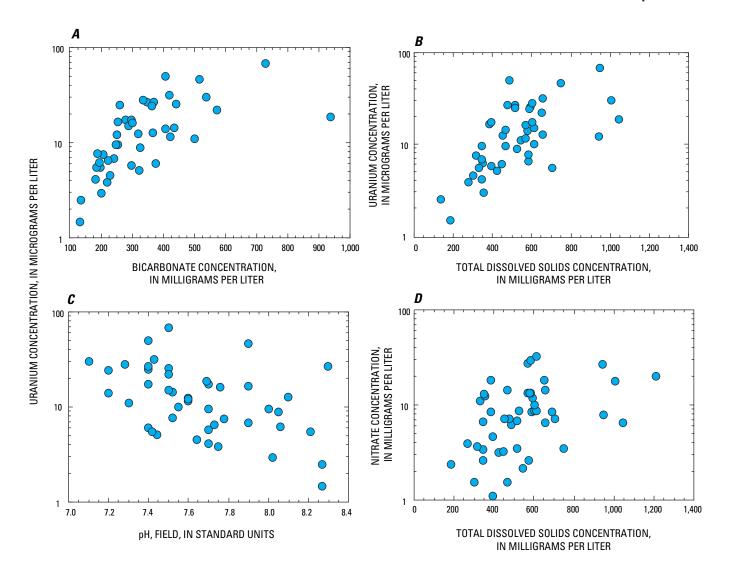


Figure 13. Relation of concentrations of uranium to concentrations of (*A*) bicarbonate, (*B*) total dissolved solids, and (*C*) pH, and (*D*) relation of concentrations of nitrate to concentrations of total dissolved solids in ground-water samples from wells in the Columbia Plateau glacial aquifer group in the northern United States, 1993–2003.

Nine percent of the samples from the CPG aquifers contained uranium concentrations that exceeded the USEPA MCL of 30 μ g/L (table 6). The maximum uranium concentration was 67.9 μ g/L (table 4). The concentrations of uranium in ground water from this aquifer group were higher than in any of the other aquifer groups studied (table 4). The maximum 222 radon concentration for 85 samples in the CPG aquifer group was 1,900 pCi/L (table 5).

The strongest uranium correlations in CPG aquifers were with bicarbonate (HCO $_3$) ion (rho = 0.77), field specific conductance (rho = 0.71), sodium (rho = 0.69), total dissolved solids (rho = 0.68), and acid neutralizing capacity (rho = 0.68) (appendix 3). Other strong correlations include sulfate (rho = 0.51) and pH (rho = -0.48). Uranium also correlated with some trace elements in this aquifer, including copper (rho = 0.47), lithium (rho = 0.47), magnesium (rho = 0.44), and boron (rho = 0.42). Significant uranium correlations are listed by individual well networks in the CPG aquifer group in appendix 5.

These correlations are consistent with the complexation of uranium with bicarbonate and sulfate ions (Gascoyne, 1992; Hem, 1985), and indicate that this process may be a major control on uranium occurrence in ground water in the CPG aquifer group. Concentrations of TDS that exceed 500 mg/L in shallow aquifers are indicative of irrigation (Whitehead, 2000). Wells in CPG aquifers had a median TDS concentration of 441 mg/L and a mean of 494 mg/L. TDS also correlated strongly with nitrate concentrations, further indicating that TDS is related to irrigated agriculture in this aquifer group (fig. 13). The inverse correlation of uranium with pH may be related to the age of the water, indicating that less chemically evolved water had correspondingly high uranium concentrations (fig. 13).

Lithium, bromide, and nickel were the only elements that correlated positively with 222 radon in CPG aquifers (rho = 0.51, 0.35, and 0.32, respectively) (appendix 4). Vanadium and phosphorus were inversely correlated to 222 radon (rho = -0.31 and -0.33, respectively). Well depth and radon were positively correlated (rho = 0.37).

Summary and Conclusions

Uranium and ²²²radon in ground water have long been a concern throughout the United States. Radionuclide concentrations were compared among the glacial, Cambrian-Ordovician, and the New York and New England crystalline aquifer systems of the northern United States. In these aquifer systems, uranium and ²²²radon concentrations differed greatly. In certain aquifers, concentrations also exceeded human-health standards in ground water used for drinking water and for agriculture.

In this study, water-quality data collected during 1993 to 2003 from 56 well networks in the U.S. Geological Survey National Water-Quality Assessment (NAWQA) program

were evaluated to determine the occurrence of uranium and ²²²radon in water in aquifers in the northern United States. The well networks, generally consisting of about 30 wells in similar aquifers and(or) land-use settings, were used in this investigation if the wells were completed in one of the three aquifer systems described above and had a measurement for uranium and(or) ²²²radon. A total of 1,426 wells sampled as part of the NAWQA program met the above criteria. About 60 percent of these wells had water-quality data for both uranium and ²²²radon.

The glacial aquifer system is geographically extensive and the composition of aquifer materials differs across the Unites States. For this reason, the glacial aguifer system was further divided into seven aquifer groups on the basis of the origin of the glacial deposits from which they were formed. Comparisons were made among the seven glacial aquifer groups and the Cambrian-Ordovician and New York and New England Crystalline aquifer systems. The chemical composition of samples from the resulting nine aguifer groups was compared, and concentrations of uranium and ²²²radon were found to differ significantly. In general, uranium concentrations were highest in the glacial aquifer group of the Columbia Plateau, the West-Central glacial aquifer group, and the New York and New England crystalline aquifer group. About 10 percent of the wells in the Columbia Plateau glacial and the West-Central glacial aquifer groups had water with concentrations of uranium that exceeded the U.S. Environmental Protection Agency (USEPA) Maximum Contaminant Level (MCL) of 30 µg/L. For the New York and New England crystalline aquifer group, samples from 4 percent of the wells sampled exceeded the MCL of 30 μg/L for uranium.

In areas with high concentrations of uranium, the link to uranium-enriched geologic material is a common feature. In the Columbia Plateau glacial aquifer group, the deposits were derived in part from basaltic lava flows, with some felsic volcanic rocks and probably some paleo lakebeds that may be rich in uranium. In the West-Central glacial aguifer group, rocks such as Cretaceous marine shale that contain abundant uranium probably contribute to the high concentrations of uranium in ground water; in the southern part of this area, including northern Nebraska, the glacial or glacial-related sediment may be interbedded with uraniumrich materials that originated in the Rocky Mountains. These areas coincide with some of the highest terrestrial gamma-ray measurements in the United States, further indicating that a geologic source of uranium is a major factor controlling ground-water concentrations. In New England, crystalline bedrock that is granitic, such as two-mica granites, as well as other high-grade metamorphic rocks, appears to contain abundant uranium that is soluble in the predominant sub-oxic to oxic and high-pH geochemical conditions. In the Columbia Plateau and West-Central glacial aquifers, uranium correlated with total dissolved solids and bicarbonate as well as with boron, lithium, selenium, and strontium. In the New York and New England crystalline aquifer group, uranium correlated

primarily with other radionuclides, such as gross alpha particle activity, and with ²²²radon. In addition to having a geologic source of uranium, factors that affect oxidation-reduction potential (redox) or ground-water flow could have an effect on uranium concentrations in ground water. For example, irrigation agriculture is particularly prevalent in the glacial aquifers of the Columbia Plateau and West-Central glacial aquifer groups, and irrigation return flows might enhance the mobility of uranium.

Median concentrations of ²²²radon were 10 times higher in the New York and New England crystalline aquifer group than in the glacial aquifer group and the Cambrian-Ordovician aquifer group. Concentrations of ²²²radon in New England were similar to those found in other countries (mainly Scandinavia) with similar geologic histories and(or) similar bedrock geology. In the New York and New England crystalline aquifer group, uranium and ²²²radon concentrations strongly correlated with each other and with gross alpha particle activity. This was not found in any other aquifers studied in this investigation.

Also, 222 radon correlated with lead in water from wells completed in felsic igneous and undifferentiated metasedimentary rocks in the New York and New England crystalline aquifer group—a finding that is similar to findings in studies in Finland. Some of the lead concentrations associated with high 222 radon concentrations (greater than 10,000 pCi/L) in these wells are at levels near the USEPA action level for lead of 15 µg/L.

Although water-quality data for 1,426 wells were evaluated in this investigation, spatial coverage of areas where water is used for drinking-water supply was limited. Additional data from the state drinking-water programs or other sources could provide the necessary spatial coverage to validate findings presented in this report as well as to refine or develop new models of uranium and ²²²radon occurrence.

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Appendixes 1–6

Appendix 1. Abbreviations and descriptions of the National Water Quality Assessment (NAWQA) program well networks used to study water quality in glacial, Cambrian-Ordovician, and New York and New England crystalline aquifer systems in the northern United States, 1993–2003.

[NAWQA, National Water Quality Assessment Program; SUID, Study-Unit Identifier. NAWQA summary reports provide more detailed information on the individual study units and their well networks used in this study. Access to these reports can be found online at http://water.usgs.gov/nawqa/nawqa_sumr.html]

NAWOA Study Unit Identifiers

ALMN	Allegheny and Monongahela River Basins (Pennsylvania, West Virginia, New York, and Maryland)
CCYK	Central Columbia Plateau (Washington and Idaho) combined with the Yakima River Basin (Washington)
CNBR	Central Nebraska Basins (Nebraska)
CONN	Connecticut, Housatonic, and Thames Rivers Basins (Connecticut, Massachusetts, New Hampshire, and Vermont)
COOK	Cook Inlet Basin (Alaska)
DELR	Delaware River Basin (Delaware, Maryland, New Jersey, Pennsylvania, and Virginia)
EIWA	Eastern Iowa Basins (Iowa and Minnesota)
HDSN	Hudson River Basin (New York and adjacent states)
HPGW	High Plains Regional ground-water study (Colorado, Kansas, Nebraska, New Mexico, Oklahoma, South Dakota, and Texas)
LERI	Lake Erie-Lake Saint Clair Drainages (Michigan, Ohio, Indiana, New York, and Pennsylvania)
LINJ	Long Island-New Jersey Coastal Drainages (New Jersey and New York)
LIRB	Lower Illinois River Basin (Illinois)
NECB	New England Coastal Basins (New Hampshire, Massachusetts, Maine, and Rhode Island)
PUGT	Puget Sound Basin (Washington and British Columbia, Canada)
REDN	Red River of the North Basin (Minnesota, North Dakota, and South Dakota)
UIRB	Upper Illinois River Basin (Illinois, Indiana, and Wisconsin)
UMIS	Upper Mississippi River Basin (Minnesota, Wisconsin, South Dakota, Iowa, and North Dakota)
WHMI	White River Basin (Indiana) combined with the Great and Little Miami River Basins (Ohio and Indiana)
WMIC	Western Lake Michigan Drainages (Michigan and Wisconsin)

Aquifer Group Codes

CPG	Columbia Plateau glacial
COS	Cambrian-Ordovician (sandstone and dolomite)
ECG	East-Central glacial
NEC	New York and New England crystalline (fractured bedrock)
NEG	Northeastern glacial
NCG	North-Central glacial (deposits of the Red River and Wadena, Rainy, Lake Superior, and Green Bay glacial lobes)
WCG	West-Central glacial (deposits of the Dakota, James, Des Moines, and eastern Red River glacial lobes)
NPG	Northern Pacific glacial
PWG	Pre-Wisconsinan glacial

Appendix 1. Abbreviations and descriptions of the National Water Quality Assessment (NAWQA) program well networks used to study water quality in glacial, Cambrian-Ordovician, and New York and New England crystalline aquifer systems in the northern United States, 1993–2003.—Continued

NAWQA SUID (see figure 2)	NAWQA well network	Aquifer group code	Predominant land-use setting	Aquifer setting	NAWOA summary report reference
			ຶ່ງ	Cambrian-Ordovician aquifer system	
UMIS	umissus3	COS	Mixed	Unconfined (dolomite and sandstone) aquifers in Minnesota and Wisconsin	Stark and others, 2000
UMIS	umissus4	COS	Mixed	Confined (dolomite and sandstone) aquifers in Minnesota and Wisconsin	Do.
WMIC	wmicsus1	COS	Mixed	(Sandstone) aquifers, west of Maquoketa shale in Michigan and Wisconsin	Peters and others, 1998
			New York 6	New York and New England crystalline aquifer system	
CONN	connsus1	NEC	Mixed	Fractured, crystalline-bedrock (undifferentiated) aquifers in Connecticut, Vermont, Massachusetts, New Hampshire	Garabedian and others, 1998
LINJ	linjsus1	NEC	Mixed	Fractured, crystalline-bedrock (undifferentiated) aquifers in New Jersey and New York	Ayers and others, 2000
NECB	necbsus1	NEC	Mixed	Fractured, crystalline-bedrock (metasedimentary, variably calcareous) aquifers in New Hampshire and Maine	Robinson and others, 2004
NECB	necbsus2	NEC	Mixed	Fractured, crystalline-bedrock (felsic igneous and metasedimentary, undifferentiated) aquifers in New Hampshire, Maine, and Massachusetts	Do.
				Glacial aquifer system	
ALMN	almnsus2	ECG	Mixed	Glacial (coarse-grained glaciofluvial sediments) aquifers in northwestern Pennsylvania and western New York	Anderson and others, 2000
CCYK	ccptlusag2a	CPG	Potatoes and corn	Deep, basin-fill glacial aquifers in eastern Washington State	Williamson and others, 1998
CCYK	ccptlusag2b	CPG	Potatoes and corn	Shallow, basin-fill glacial aquifers in eastern Washington State	Do.
CCYK	ccptlusor1a	CPG	Orchards and vineyards	Deep, basin-fill glacial aquifers in eastern Washington	Do.
CCYK	ccptlusor1b	CPG	Orchards and vineyards	Shallow, basin-fill glacial aquifers in eastern Washington	Do.
CNBR	cnbrluscr1	MCG	Corn and soybeans	Loess-mantled glacial till aquifers in eastern Nebraska	Unpublished data ¹
CONN	conndwgs1 connlusrc1 connsus2	NEG NEG NEG	Mixed New residential and commercial Mixed	Glacial (sand and gravel) aquifers in Connecticut Glacial (sand and gravel) aquifers in Connecticut and Massachusetts Unconsolidated glacial aquifers in western New England	Do. Do.
COOK	cooksus1	NPG	Mixed	Glacial aquifers in Cook Inlet area, Alaska	Glass and others, 2004

Appendix 1. Abbreviations and descriptions of the National Water Quality Assessment (NAWQA) program well networks used to study water quality in glacial, Cambrian-Ordovician, and New York and New England crystalline aquifer systems in the northern United States, 1993–2003.—Continued

NAWQA SUID (see figure 2)	NAWOA well network	Aquifer group code	Predominant land-use setting	Aquifer setting	NAWOA summary report reference
			9	Glacial aquifer system—Continued	
DELR	delrsus3	ECG	Mixed	Coarse-grained, glacial outwash aquifers in New York, New Jersey, and Pennsylvania	Fischer and others, 2004
EIWA	eiwaluscr1 eiwalusrc1	PWG, WCG PWG, WCG	Corn and soybeans Residential, commercial, and	Unconsolidated glacial aquifers in Iowa and Minnesota Unconsolidated glacial aquifers in Iowa and Minnesota	Kalkhoff and others, 2000 Do.
EIWA	eiwasus2	PWG, WCG	ındustrial Mixed	Alluvial glacial aquifers in Iowa	Do.
HDSN HDSN	hdsnlusag1 hdsnlusur1	ECG	Unknown Urban	Unconsolidated glacial aquifers in New York Unconsolidated glacial aquifers in New York	Wall and others, 1998 Do.
HPGW HPGW	hpgwdwgs1 hpgwsus4	WCG WCG	Mixed Mixed	Quaternary (glacial) aquifers in southeastern Nebraska Unconfined glacial aquifers in Nebraska (northern high plains)	Unpublished data ¹ Do.
LERI LERI	lerilusrc1 lerisus1	ECG	Urban Mixed	Unconfined glacial aquifers outside Detroit, Michigan Glacial aquifers overlain by fractured clay-till in Michigan and Ohio	Myers and others, 2000 Do.
LIRB	lirbluscr1	ECG	Corn and soybeans	Shallow glacial aquifers, Bloomington Ridged Till Plains not overlying buried bedrock valley in Illinois	Groschen and others, 2000
LIRB	lirbluscr2	ECG	Corn and soybeans	Shallow glacial aquifers, Bloomington Ridged Till Plains, overlying Mahomet-Teays buried bedrock valley, in Illinois	Do.
LIRB	lirbsus1	PWG	Mixed	Deep glacial aquifers, Bloomington Ridged Till Plains, overlying Mahomet-Teays buried bedrock valley, in Illinois	Do.
LIRB	lirbsus2	PWG	Mixed	Shallow glacial aquifers, Galesburg/Springfield Till Plains, not overlying buried bedrock valley, in Illinois	Do.
WHMI WHMI WHMI	miamluscr1 miamlusrc1 miamsus1	ECG, PWG ECG, PWG ECG, PWG	Corn and soybeans Urban Mixed	Glacial (outwash) aquifers in Indiana and Ohio Glacial (outwash) aquifers in Ohio Confined glacial (buried valley) aquifers in Indiana and Ohio	Rowe and others, 2004 Do. Do.
NECB NECB	necblusrc1 necbsus3	NEG NEG	New residential and commercial Mixed	Coarse-grained, glacial aquifers in Massachusetts and New Hampshire Shallow and deep glacial aquifers in eastern New England	Robinson and others, 2004 Do.

Appendix 1. Abbreviations and descriptions of the National Water Quality Assessment (NAWQA) program well networks used to study water quality in glacial, Cambrian-Ordovician, and New York and New England crystalline aquifer systems in the northern United States, 1993–2003.—Continued

NAWQA SUID (see figure 2)	NAWQA well network	Aquifer group code	Predominant land-use setting	Aquifer setting	NAWOA summary report reference
				Glacial aquifer system—Continued	
PUGT PUGT PUGT	pugduscr1 pugdusrs1 pugtsus1	NPG NPG NPG	Raspberries cropland New residential Mixed	Coarse-grained, glacial aquifers in western Washington State Coarse-grained, glacial aquifers in western Washington State Unconfined (Fraser) glacial aquifers in western Washington State	Ebbert and others, 2000 Do. Do.
REDN REDN	rednlusag1 rednlusag2	NCG WCG	Irrigated cropland Agriculture	Glacial outwash (Otter Tail) aquifer in Minnesota Glacial (Sheyenne Delta) aquifer in South Dakota	Stoner and others, 1998 Do.
REDN REDN REDN	rednsus1 rednsus2 rednsus3 rednsus5	WCG, NCG WCG, NCG WCG, NCG	Mixed Mixed Mixed Mixed	Contined and unconfined glactal aquifers in North Dakota and Minnesota Unconfined glacial aquifers in North Dakota Unconfined glacial aquifers in North Dakota Buried glacial aquifers in North Dakota	Бо. Оо.
UIRB UIRB UIRB	uirbluscr1 uirblusrc1 uirbsus1 uirbsus2	BCG NCG NCG BCG	Irrigated rowcrops Hi-density suburban Mixed Mixed	Coarse-grained, glacial aquifers in Illinois and Indiana Thin sand and gravel lenses in till, overlying Silurian-Devonian aquifers, in Illinois and Wisconsin Shallow, coarse-grained, glacial aquifers in Illinois and Wisconsin Deep, coarse-grained glacial aquifers in Illinois and Indiana	Groschen and others, 2004 Do. Do.
UMIS	umislusrc1	WCG	New residential	Coarse-grained, unconfined, glacial aquifers in Minnesota	Stark and others, 2000
WHMI WHMI WHMI	whitluscr1 whitluscr3 whitlusur1	ECG PWG, ECG ECG, PWG	Croplands Croplands Urban lands	Unconfined sand and gravel, and glacial till, aquifers in Indiana Coarse-grained glacial aquifers in Indiana Coarse-grained, glacial aquifers in Indiana	Fenelon, 1998 Do. Do.
WMIC WMIC	wmiclusag1a wmiclusag2 wmicsus2	NCG NCG NCG	Corn and alfalfa Corn and alfalfa Mixed	Glacial till (sand, gravel, and clay) aquifers in Wisconsin Coarse-grained glacial (sand and gravel) aquifers in Wisconsin Glacial aquifers in Wisconsin and Michigan	Peters and others, 1998 Do. Unpublished data ¹

¹ Unpublished data collected during the second assessment period of the National Water Quality Assessment Program. Public access to this data is available from the NAWQA Data Warehouse (Bell and Williamson, 2006; Williamson, 2006; Williamson, 2006).

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Statistical summary of uranium and 22 radon concentrations in ground water in the NAWQA well networks used to study water quality in glacial, Cambrian-Ordovician, and New York and New England crystalline aquifer systems in the northern United States, 1993—2003. Appendix 2.

available; e, estimated concentration; Domestic, drinking-water well used for private (or household) supply; Monitor, observation well installed in a specific land-use setting for monitoring water quality; Public [N, number of samples; Min, minimum; 10th, tenth percentile; 25th, twenty-fifth percentile; Med, median; 75th, seventy-fifth percentile; 90th, ninetieth percentile; Max, maximum; <, less than; -, no data supply, drinking-water well used for municipal supply]

			N.				Lani	Concentra	Ilranium concentrations in micrograms ner liter	rodrame ne	ii				222 Ra r	Jon concer	in ations in	222Radon concentrations in nicocuries ner liter	or liter	
NAWQA well network	Water use	Sample years	ber of wells in net- work	Median well depth, in feet	z	Min	10th	25th	Med	75th	90th	Мах	z	Min	10th	25th	Med	75th	90th	Мах
								Camb	Cambrian-Ordovician aquifer system	ian aquifer	system									
umissus3	Domestic	1996	25	180	25		1 ^	^		-	-	5	24	100	190	230	340	530	730	2,700
umissus4	Domestic	1996	25	200	25	× -	^	^	^ 1	2	3	4	23	74	190	250	420	630	0006	2,600
wmicsus1	Domestic/pub-lic supply	2002	30	155	26	e .01	> .01	0.1	0.28	0.73	1.8	4.7	30	140	180	230	523	730	1,072	1,957
							Nev	v York and	New York and New England crystalline aquifer system	d crystalline	aquifer sys	tem								
connsus1	Domestic	1995	30	235	30	4. ^	4. ^	4. ^	0.75	2.3	12	30	30	220	730	1,300	2,150	5,400	9,500	20,000
linjsus1	Domestic	1997	29	201	29	^	^	^	^ _	1.24	6.4	429	26	218	245	554	1,547	3,124	4,738	31,820
necbsus1	Domestic	1999–00	28	188	28	^ _	^ 	^ _	^	2.8	6.1	21.8	26	663	947	1,404	2,185	3,027	5,920	12,776
necbsus2	Domestic	2000	30	303	30	e .01		^	1.9	11	32.4	151.5	28	75		1,115	3,567	11,740	45,290	215,210
									Glacial aqu	Glacial aquifer system										
almnsus2	Domestic	1996	30	73	0	ı	1	1	1	ı	1	ı	30	180	195	330	999	1,200	1,700	2,800
ccptlusag2a	Domestic	1993–94	30	133	0	ı	ı	ı	ı	ı	I	ı	28	210	240	388	555	069	885	1,250
ccptlusag2b	Monitor	1993–94	32	38	27	0.92	2.5	5.7	10.1	17.5	26.6	50.5	19	110	230	270	395	476	831	086
ccptlusor1a	Domestic	1994–95	18	165	0	ı	1	ı	I	ı	1	ı	18	310	310	009	915	1,200	1,800	1,900
ccptlusor1b	Monitor	1994–95	22	29.5	19	3.8	4.1	9	14.5	27.9	46.2	89	20	210	250	340	505	620	705	850
cnbrluscr1	Monitor	2003	30	I	29	0.76	2.9	10.9	19.3	8.44	83.8	162	0	ı	ı	I	I	I	I	I
conndwgs1	Public supply	2002-03	15	70	10	e .01	.01	0.07	0.24	0.38	76.	1.4	v	577	I	762	1,103	1,208	1	1,292
connlusrc1	Monitor	2003	28	22	28	e .01	< .02	0.02	0.024	0.44	.92	13.6	0	I	I	ı	I	I	I	I
connsus2	Domestic	2002	30	110	30	e .01	< .02	0.07	0.31	0.97	2.9	6.2	29	166	270	484	758	1,247	2,244	8,056
cooksus1	Domestic	1999	34	5.76	34	^	^ _	^	^	^	^ 	^	31	139	151	197	256	382	454	609
delrsus3	Domestic	2001	16	114	16	e .01	< .02	< .02	< 0.02	0.2	0.59	6.0	16	373	497	771	1,011	1,483	1,644	2,282
eiwaluscr1	Monitor	1997	31	I	0	I	ı	ı	1	1	1	ı	31	187	221	266	396	488	861	1,500
eiwalusrc1	Monitor	1997	30	I	0	I	I	ı	I	1	I	ı	30	172	236	281	378	657	186	1,221
eiwasus2	Domestic	1998	32	52	0	I	I	I	I	1	I	ı	32	53	196	245	340	515	692	2,146

Appendix 2. Statistical summary of uranium and 22 radon concentrations in ground water in the NAWQA well networks used to study water quality in glacial, Cambrian-Ordovician, and New York and New England crystalline aquifer systems in the northern United States, 1993–2003.—Continued

[N, number of samples; Min, minimum; 10th, tenth percentile; 25th, twenty-fifth percentile; Med, median; 75th, seventy-fifth percentile; 90th, ninetieth percentile; Max, maximum; <, less than; -, no data available; e, estimated concentration; Domestic, drinking-water well used for private (or household) supply; Monitor, observation well installed in a specific land-use setting for monitoring water quality; Public supply, drinking-water well used for municipal supply]

							=								- 4666				100	
NAWOA			Num- ber of	Median			Oranium	concentra	Uranium concentrations, in micrograms per liter	crograms pe	er liter				Rad	***Kadon concentrations, in picocuries per liter	ations, in pic	cocuries per	Iter	
well	Water use	Sample years	wells in net- work	well depth, in feet	z	Min	10th	25th	Med	75th	90th	Мах	z	Min	10th	25th P	Med	75th	90th	Мах
								Glaci	Glacial aquifer system—Continued	rstem—Con	tinued									
hdsnlusag1	Monitor	1993–94	41	19.5	0	ı	ı	ı	1	ı	1	ı	8	110	ı	ı	672	ı	1	841
hdsnlusur1	Unused/public supply	1993–94	28	27.5	27	<u>^</u>	, 1	^	^	<u>^</u>	71	71	18	69	82	102	169	281	531	1,086
hpgwdwgs1	Public supply	2002	15	292	15	0.19	0.5	2.21	6.4	19.9	34.4	9.99	0	I	I	I	I	I	I	I
hpgwsus4	Domestic	2003	30	187.5	27	0.09	0.43	1.8	4.2	7.56	12.2	57.4	26	162	187	233	276	372	979	006
lerilusrc1	Monitor	1996–97	30	25.5	29		<u>~</u>	^	<u>^</u>	<u>^</u>	4	7	21	93	140	170	310	380	520	069
lerisus1	Domestic	1998	27	06	0	I		I	I	ı	I	ı	27	104	128	145	171	188	245	302
lirbluscr1 lirbluscr2	Monitor Monitor	1997 1997	28	27.5	28	^ ^ 	^	^ ^ 	1.7	4 1.6	10.2	15	28	08 08	< 80 143	158 200	216 348	251 472	605	677 1,301
lirbsus1 lirbsus2	Domestic Domestic	1996 1996	30	245 50	30	^ ^ ^	^ ^ 	^ ^ 	^ ^ 	^	< 13.5	1>	30	110 < 52	120	140 210	160 340	200	260	730
miamluscr1	Monitor	2000	34	27.5	34		<u>~</u>	^	<u>^</u>	1.75	4.2	41	34	86	215	308	384	909	651	791
miamlusrc1	Monitor	2001	27	38	25	0.23	.42	0.5	8.0	1.1	1.24	1.8	25	199	210	392	449	619	867	1,013
miamsus1	Domestic	1999	30	09	30		<u>~</u>	^	^	1.7	2.5	3.7	30	163	181	222	359	529	029	1,386
necblusrc1	Monitor	1999	29	23	29	^	× 	^	^ 	^	^ 1	> 2	29	165	344	454	683	822	1,253	1,289
necbsus3	Public supply	2001	30	99	30	<.02	<.02	0.026	0.13	0.39	4.1	15.8	30	311	442	671	1,165	1,627	2,407	3,745
pugtluscr1	Monitor	I	22	I	ς.	^	× 	^	× .	^ 	^	^ 	0	ı	ı	I	I	ı	I	I
pugtlusrs1	Monitor	1996	27	ı	0	1	I	I	I	1	1	I	16	160	180	240	385	510	550	630
pugtsus 1	Domestic	1996	30	ı	0	1	I	I	I	1	1	ı	30	100	125	180	290	450	625	098
rednlusag1 rednlusag2	Monitor Monitor	1994	29	25.5 15	3 25	^ ^ 4. 4.	1 A 4.	0.7	4. 4. 4. 4.	1 4	_ 11	<.4 71	3 21	< 80 217	_ 222	264	250 341	383	- 469	290

Statistical summary of uranium and 22 radon concentrations in ground water in the NAWQA well networks used to study water quality in glacial, Cambrian-Ordovician, and New York and New England crystalline aquifer systems in the northern United States, 1993–2003.—Continued Appendix 2.

available; e, estimated concentration; Domestic, drinking-water well used for private (or household) supply; Monitor, observation well installed in a specific land-use setting for monitoring water quality; Public [N, number of samples; Min, minimum; 10th, tenth percentile; 25th, twenty-fifth percentile; Med, median; 75th, seventy-fifth percentile; 90th, ninetieth percentile; Max, maximum; <, less than; -, no data supply, drinking-water well used for municipal supply]

			Num	1017			Uranium	concentra	Uranium concentrations, in micrograms per liter	crograms p	er liter				222 Ra u	don concen	²²² Radon concentrations, in picocuries per liter	icocuries pe	ır liter	
NAWOA well network	Water use	Sample years	ber of wells in net- work	weulall well depth, in feet	z	Min	10th	25th	Med	75th	90th	Мах	z	Min	10th	25th	Med	75th	90th	Мах
								Glac	Glacial aquifer system—Continued	ystem—Cor	tinued									
rednsus1	Monitor/ domestic	1993–95	29	35	17	4.	^ 4.	0.7	1.9	6.7	23	99	14	190	217	230	325	640	029	1,100
rednsus2	Monitor/ domestic	1993	25	58	22	^ 4.	^ 4.	^ 4.	< 0.4	0.7	2.1	5.5	20	130	160	189	267	361	452	594
rednsus3	Monitor/ domestic	1993–95	24	41	24	^ 4.	^ 4.	1.2	2.1	3.5	6.7	140	5	290	I	290	469	200	ı	820
rednsus5	Domestic	1994	34	121	19	^ 4.	^ 4.	^ 4.	2.7	5	9	7.5	∞	83	83	220	305	380	390	390
uirbluscr1	Monitor	1999	59	41	29	×	^	^	<u>~</u>	1.7	7.8	21.1	29	137	143	182	260	323	569	969
uirblusrc1	Monitor	2000	27	59	26	^		^	<u>~</u>	1.3	4.4	5.9	21	149	182	210	265	388	456	287
uirbsus 1 uirbsus 2	Domestic Domestic	2001	27	72	27	e .02 e .01	< .02	0.11	0.27	0.42	.98	2.5	27	145	153 128	171	221 166	310	377 259	418
umislusrc1	Monitor	1996	30	19	30	× 	^	^	<u>~</u>	2	5.5	13	0	I	I	ı	I	I	I	I
whitluscr1 whitluscr3	Monitor Monitor	1994	30	28 22.5	23	0.5	.46	0.68	1.7	2.4	6.4	12.1	0 29	160	170	210	310	540	790	1,600
whitlusur1	Monitor	1995	28	59	0	I	I	I	I	I	I	ı	28	140	190	300	405	505	069	920
wmiclusag1a wmiclusag2	Monitor Monitor	1993–94 1994	26 32	28	25	< .4 e .01	< 1< .02	0.06	0.7	0.83	2.1	5.5	0	1 1	1 1	1 1	1 1	1 1	1 1	1 1
wmicsus2	Domestic	2002	28	ı	28	e .01	<.01	90.0	0.28	0.75	1.9	3.3	28	132	4	173	249	424	1,115	2,436

Appendix 3. Selected rho and probability values from Spearman correlations of selected trace elements, major ions, and physical properties in ground water, National Uranium Resource Evaluation program streambed sediment, terrestrial gamma-ray emissions, and ancillary data with uranium concentrations in ground water, by aquifer group, in the northern United States, 1993–2003.

For each aquifer group, the order of the correlation variable is listed by the highest correlation (either negative or positive) value to the lowest correlation value. Spearman's coefficients (rho) are significant when the *P*-values are less than or equal to 0.05; a negative rho value indicates an inverse relation. The following variables consist of geochemical data for the United States based primarily on streambed sediments and analyzed for using a consistent set of methods as part of the USGS National Geochemical Survey (U.S. Geological Survey Open-File Report 2004–1001). Most of the original samples were collected as part of the U.S. Geological Survey's National Uranium Resource Evaluation (NURE) Program from 1975 to 1983 and re-analyzed (data were accessed online May 2005, at http://tin.er.usgs.gov/geochem/doc/home.htm).

The variables are expressed in parts per million, unless stated otherwise:

Calcium (NURE) in percent weight

Molybdenum (NURE)

Selenium (NURE)

Sodium (NURE) in percent weight

Lithium (NURE)

Lead (NURE)

Potassium (NURE)

Iron (NURE)

Uranium (NURE)

< less than
U uranium

Runoff mean annual runoff, for 1951 to 1980, in inches (Wolock, D.M., 2003)

precip90 mean annual precipitation, for 1961 through 1990 (Source of data is at Oregon State University in

Corvallis, OR, accessed online December 2006, at http://www.ocs.oregonstate.edu/prism/index.phtml)

Population

population density from the 2000 Census, in persons per square kilometer within 500 meters of sampled wells

Density

¹⁸Oxygen a stable isotope, reported as the ratio per mill (%o, or per thousand) of ¹⁸O/¹⁶O (δ ¹⁸O) Deuterium a stable isotope, reported as the ratio per mill (%o, or per thousand) of ²H/¹H (δ D)

⁴⁰Potassium aerial gamma-ray survey across the conterminous United States that measured the gamma-ray flux produced

by the radioactive decay of the naturally-occurring element 40Potassium in the top few centimeters of rock and

soil, expressed in units of percent potassium (Phillips and others, 1993)

²³²Thorium aerial gamma-ray survey across the conterminous United States that measure the gamma-ray flux produced

by the radioactive decay of the naturally-occurring element ²³²Thorium in the top few centimeters of rock and

soil, expressed as parts per million equivalent thorium (eTh) (Phillips and others, 1993).

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Radionuclide	Correlation variable	Number of samples	Spearman's coefficient (rho)	<i>P</i> -value
	Cambria	ın-Ordovician (COS) aquifer		
U	Sodium (NURE)	11	0.501	0.116
U	Acid neutralizing capacity (water)	74	0.478	< 0.0001
U	Nickel (water)	76	0.472	< 0.0001
U	Molybdenum (NURE)	19	-0.443	0.058
U	Calcium (water)	75	0.415	< 0.0001
U	Calcium (NURE)	11	-0.400	0.223
U	Magnesium (water)	75	0.392	< 0.0001
U	Specific conductance, field (water)	76	0.358	0.001
U	Manganese (water)	76	0.355	0.002
U	Runoff	75	-0.336	0.003
U	Selenium (NURE)	19	-0.327	0.171
U	Sulfate (water)	75	0.320	0.005
U	Molybdenum (water)	76	0.308	0.007
U	precip90	75	-0.306	0.008
U	Dissolved oxygen (water)	76	-0.280	0.014
U	Thallium (water)	26	0.277	0.171
U	Aluminum (water)	74	0.273	0.019
U	Total dissolved solids (water)	75	0.271	0.019
U	Ammonia + organic nitrogen (water)	38	0.243	0.142
U	Potassium (water)	75	0.226	0.051
U	Turbidity (water)	74	0.223	0.056
U	Tritium (water)	49	-0.219	0.130
U	Zinc (water)	76	0.218	0.058
U	Lithium (water)	26	-0.215	0.291
U	²³² Thorium (gamma-ray)	73	0.201	0.089
U	Cobalt (water)	76	0.190	0.100
U	Population density	76	0.187	0.106
U	Silica (water)	75	0.183	0.116
U	ortho-phosphate (water)	75	-0.182	0.117
U	Water temperature	76	0.175	0.131
U	Barium (water)	76	0.148	0.202
U	Sodium (water)	75	0.147	0.207
U	Age of water (time of recharge)	62	0.140	0.278
U	Copper (water)	76	-0.139	0.232
U	Boron	26	-0.135	0.510
U	Depth to water table	75	-0.128	0.272
U	⁴⁰ Potassium (gamma-ray)	76	0.119	0.303
U	Chloride (water)	75	-0.117	0.318
U	Arsenic (water)	76	-0.114	0.327
U	Fluoride (water)	75	0.110	0.346
U	Radon (water)	73	0.087	0.463
U	Depth to top of open interval	75	0.081	0.490
U	Phosphorus (water)	74	-0.069	0.561
U	Nitrate (water)	75	0.064	0.586

Appendix 3. Selected rho and probability values from Spearman correlations of selected trace elements, major ions, and physical properties in ground water, National Uranium Resource Evaluation program streambed sediment, terrestrial gamma-ray emissions, and ancillary data with uranium concentrations in ground water, by aquifer group, in the northern United States, 1993–2003.

—Continued

Radionuclide	Correlation variable	Number of samples	Spearman's coefficient (rho)	<i>P</i> -value
	Cambrian-Ord	ovician (COS) aquifer—Contir		
U	Dissolved organic carbon (water)	73	0.063	0.596
U	Selenium (water)	76	0.062	0.594
U	pH, field (water)	76	-0.061	0.598
U	Well depth	76	-0.029	0.808
U	Lead (water)	76	-0.020	0.866
U	Strontium (water)	26	0.015	0.940
	Columbia	Plateau Glacial (CPG) aquifer		
U	Bicarbonate (water)	45	0.765	< 0.0001
U	Calcium (NURE)	10	0.745	0.013
U	Specific conductance, field (water)	46	0.711	< 0.0001
U	Sodium (water)	46	0.692	< 0.0001
U	Total dissolved solids (water)	46	0.683	< 0.0001
U	Acid neutralizing capacity (water)	35	0.680	< 0.0001
U	Ammonia + organic nitrogen (water)	46	0.658	< 0.0001
U	Molybdenum (NURE)	12	-0.544	0.068
U	Strontium (water)	46	0.518	< 0.0001
U	Sulfate (water)	46	0.513	< 0.0001
U	Field pH (water)	46	-0.485	< 0.0001
U	Dissolved organic carbon (water)	46	0.473	< 0.0002
U	Copper (water)	46	0.472	< 0.0002
U	Lithium (water)	46	0.469	0.001
U	Sodium (NURE)	12	-0.452	0.140
U	Magnesium (water)	46	0.442	0.002
U	Iron (NURE)	12	0.424	0.170
U	Boron (water)	46	0.424	0.170
U	Chloride (water)	46	0.418	0.004
U	Selenium (water)	46	0.369	0.010
U	* * *	12	0.354	0.012
	Uranium (NURE)			
U	Nickel (water)	46	0.310	0.036
U	Calcium (water)	46	0.309	0.037
U	Bromide (water)	46	0.300	0.043
U	Nitrate (water)	46	0.291	0.050
U	Ortho-phosphate (water)	46	0.284	0.056
U	Depth to water table	35	-0.269	0.119
U	Phosphorus (water)	35	0.255	0.139
U	Iron (water)	46	0.253	0.090
U	Molybdenum (water)	46	0.247	0.098
U	Water temperature	44	-0.232	0.130
U	Ammonia + organic nitrogen (water)	46	-0.230	0.124
U	Manganese (water)	46	0.215	0.151
U	Zinc (water)	46	0.212	0.157
U	Thallium (water)	46	-0.203	0.175
U	Aluminum (water)	46	-0.186	0.215
U	Radon (water)	33	0.176	0.327

Appendix 3. Selected rho and probability values from Spearman correlations of selected trace elements, major ions, and physical properties in ground water, National Uranium Resource Evaluation program streambed sediment, terrestrial gamma-ray emissions, and ancillary data with uranium concentrations in ground water, by aquifer group, in the northern United States, 1993–2003.

—Continued

Radionuclide	Correlation variable	Number of samples	Spearman's coefficient (rho)	<i>P</i> -value
	Columbia Plate	au Glacial (CPG) aquifer—Cont	· · · · · · · · · · · · · · · · · · ·	
U	Population density	46	0.141	0.348
U	Dissolved oxygen (water)	46	-0.140	0.354
U	Depth to top of open interval	46	0.139	0.356
U	Well depth	46	0.126	0.404
U	Depth to bottom of open interval	46	0.125	0.406
U	Fluoride (water)	46	0.099	0.511
U	Vanadium (water)	46	-0.077	0.609
U	²³² Thorium (gamma-ray)	46	-0.076	0.617
U	Potassium (water)	46	0.063	0.680
U	Turbidity (water)	31	-0.044	0.813
U	Barium (water)	46	0.043	0.777
U	Silica (water)	46	-0.042	0.780
U	precip90	35	0.042	0.812
U	⁴⁰ Potassium (gamma-ray)	46	-0.036	0.813
U	Arsenic (water)	46	-0.012	0.937
		Central Glacial (ECG) aquifer		
U	Manganese (water)	283	0.460	< 0.0001
U	Magnesium (water)	282	0.452	< 0.0001
U	Nickel (water)	283	0.425	< 0.0001
U	Sulfate (water)	279	0.414	< 0.0001
U	Calcium (water)	282	0.400	< 0.0001
U	Molybdenum (water)	283	0.380	< 0.0001
U	Sodium (NURE)	43	0.375	0.013
U	Turbidity (water)	179	0.371	< 0.0001
U	Total dissolved solids (water)	280	0.361	< 0.0001
U	Acid neutralizing capacity (water)	131	0.353	< 0.0001
U	Thallium (water)	68	0.342	0.004
U	Carbonate (water)	45	0.329	0.027
U	Cobalt (water)	283	0.316	< 0.0001
U	Fluoride (water)	282	0.297	< 0.0001
U	Calcium (NURE)	39	0.288	0.076
U	Dissolved organic carbon (water)	277	0.278	< 0.0001
U	Specific conductance, field (water)	282	0.271	< 0.0001
U	Phosphorus (water)	259	-0.264	< 0.0001
U	Depth to water table	280	-0.262	< 0.0001
U	Depth to bottom of open interval	275	-0.260	< 0.0001
U	Well depth	283	-0.259	< 0.0001
U	Depth to top of open interval	279	-0.258	< 0.0001
U	Strontium (water)	157	0.227	0.004
U	Nitrite (water)	282	0.224	0.004
U	²³² Thorium (gamma-ray)	266	0.213	0.000
U	Ortho-phosphate (water)	282	-0.211	0.001
U	Barium (water)	283	0.209	0.000
U	Potassium (NURE)	43	0.209	0.000

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Appendix 3. Selected rho and probability values from Spearman correlations of selected trace elements, major ions, and physical properties in ground water, National Uranium Resource Evaluation program streambed sediment, terrestrial gamma-ray emissions, and ancillary data with uranium concentrations in ground water, by aquifer group, in the northern United States, 1993–2003. —Continued

Radionuclide	Correlation variable	Number of samples	Spearman's coefficient (rho)	<i>P</i> -value
	East-Central 0	Glacial (ECG) aquifer—Continu		
U	Runoff	280	-0.205	0.001
U	Iron (NURE)	43	0.202	0.193
U	Bicarbonate (water)	73	0.197	0.095
U	Radon (water)	241	-0.191	0.003
U	⁴⁰ Potassium (gamma-ray)	283	0.189	0.001
U	Lithium (NURE)	43	-0.185	0.235
U	Boron (water)	167	0.184	0.017
U	Selenium (NURE)	30	0.176	0.353
U	Potassium (water)	282	0.174	0.003
U	Dissolved oxygen (water)	277	-0.166	0.006
U	Uranium (NURE)	43	-0.158	0.311
U	Selenium (water)	282	0.157	0.008
U	Molybdenum (NURE)	43	-0.158	0.317
U	Population density	283	-0.155	0.009
U	Water temperature	280	0.137	0.021
U	Lead (water)	283	-0.132	0.026
U	Antimony (water)	282	0.129	0.030
U	²²⁸ Radium (water)	26	0.118	0.566
U	¹⁸ Oxygen (water)	36	0.115	0.504
U	Arsenic (water)	282	0.114	0.055
U	Lithium (water)	157	0.113	0.159
U	Deuterium (water)	36	0.109	0.528
U	Copper (water)	283	0.088	0.139
U	Bromide (water)	280	0.085	0.155
U	Ammonia (water)	282	0.068	0.257
U	Vanadium (water)	157	-0.060	0.456
U	Chloride (water)	282	-0.059	0.325
U	Aluminum (water)	282	0.045	0.454
U	pH, field (water)	281	-0.044	0.461
U	Zinc (water)	282	0.043	0.469
U	Ammonia + organic nitrogen (water)	209	0.025	0.715
U	Sodium (water)	282	-0.024	0.693
U	Tritium (water)	116	-0.024	0.810
U	precip90	280	-0.022	0.810
U	Silica (water)	282	-0.013	0.822
U		282	-0.011	0.830
	Nitrate (water)			
U	Iron (water)	282	0.002	0.973
U	Cobalt (water)	entral Glacial (NCG) aquifer 116	0.463	<0.0001
U	Nickel (water) Nitrite (water)	114	0.420	<0.0001
U		167	0.330	<0.0001
U	²²⁴ Radium (water)	27	0.313	0.112
U	²²⁸ Radium (water)	27	-0.313	0.112

Appendix 3. Selected rho and probability values from Spearman correlations of selected trace elements, major ions, and physical properties in ground water, National Uranium Resource Evaluation program streambed sediment, terrestrial gamma-ray emissions, and ancillary data with uranium concentrations in ground water, by aquifer group, in the northern United States, 1993–2003.

—Continued

Radionuclide	Correlation variable	Number of samples	Spearman's coefficient (rho)	<i>P</i> -value
	North-Centra	l Glacial (NCG) aquifer—Continue		
U	Magnesium (water)	171	0.259	< 0.0001
U	Fluoride (water)	170	0.250	0.001
U	Potassium (water)	166	0.237	0.002
U	Bicarbonate (water)	74	0.235	0.044
U	Specific conductance, field (water)	170	0.207	0.007
U	Gross beta particle activity (water)	45	0.199	0.190
U	Lithium (water)	119	0.198	0.031
U	Sulfate (water)	166	0.200	0.011
U	Tritium (water)	50	0.190	0.186
U	Calcium (water)	171	0.188	0.014
U	Selenium (water)	124	0.186	0.039
U	Acid neutralizing capacity (water)	87	0.183	0.090
U	Manganese (water)	171	0.175	0.022
U	Total dissolved solids (water)	169	0.174	0.023
U	Chloride (water)	170	0.170	0.027
U	Copper (water)	115	0.162	0.084
U	²²⁶ Radium (water)	77	-0.161	0.161
U	Calcium (NURE)	26	0.158	0.440
U	Thallium (water)	83	-0.156	0.159
U	Molybdenum (water)	114	0.152	0.107
U	Strontium (water)	119	0.143	0.121
U	Barium (water)	124	0.141	0.117
U	Potassium (NURE)	26	-0.141	0.492
U	Well depth	143	-0.137	0.103
U	Molybdenum (NURE)	57	-0.136	0.311
U	¹⁸ Oxygen (water)	55	-0.131	0.339
U	Depth to top of open interval	143	-0.127	0.129
U	Depth to bottom of open interval	143	-0.125	0.135
U	Dissolved organic carbon (water)	163	0.120	0.126
U	Deuterium (water)	55	-0.120	0.383
U	Carbonate (water)	73	0.117	0.325
U	Zinc (water)	107	0.111	0.252
U	Nitrate (water)	152	0.109	0.181
U	Sodium (water)	171	0.107	0.165
U	Radon (water)	107	0.102	0.297
U	Phosphorus (water)	137	-0.099	0.249
U	pH, field (water)	170	-0.095	0.216
U	Ortho-phosphate (water)	166	-0.093	0.210
U	Iron (water)	171	-0.091 -0.091	0.241
U	Aluminum (water)	74	0.089	0.230
U	Water temperature	170	0.085	0.268
U	232Thorium (gamma-ray)	171	0.077	0.318
U	Bromide (water) Lead (water)	170 114	0.074 0.069	0.341 0.465

Appendix 3. Selected rho and probability values from Spearman correlations of selected trace elements, major ions, and physical properties in ground water, National Uranium Resource Evaluation program streambed sediment, terrestrial gamma-ray emissions, and ancillary data with uranium concentrations in ground water, by aquifer group, in the northern United States, 1993–2003.

—Continued

Radionuclide	Correlation variable	Number of samples	Spearman's coefficient (rho)	<i>P</i> -value
	North-Central			
U	Turbidity (water)	108	0.063	0.515
U	Gross alpha particle activity (water)	49	0.058	0.693
U	Ammonia (water)	167	-0.053	0.491
U	Selenium (NURE)	57	0.051	0.705
U	Arsenic (water)	116	-0.050	0.594
U	Population density	143	0.047	0.573
U	Cadmium (water)	124	-0.045	0.620
U	Age of water (time of recharge)	54	-0.045	0.749
U	Dissolved oxygen (water)	162	-0.044	0.580
U	Boron (water)	110	0.044	0.650
U	Silica (water)	171	-0.043	0.578
U	precip90	141	0.026	0.767
U	Runoff	141	0.025	0.767
U	Ammonia + organic nitrogen (water)	146	-0.024	0.776
U	⁴⁰ Potassium (gamma-ray)	171	-0.020	0.800
U	Depth to water table	141	-0.014	0.872
U	Lithium (NURE)	57	-0.007	0.960
		ew England Crystalline (NEC) a		
U	Gross alpha particle activity (water)	55	0.719	< 0.0001
U	Radon (water)	110	0.598	< 0.0001
U	Molybdenum (water)	87	0.350	< 0.0001
U	Gross beta particle activity (water)	56	0.333	0.012
U	Calcium (NURE)	52	0.281	0.044
U	Iron (water)	117	-0.266	0.004
U	Lead (water)	87	0.257	0.016
U	Calcium (water)	117	0.254	0.006
U	²²⁶ Radium (water)	56	0.243	0.071
U	Lithium (water)	58	0.237	0.073
U	Barium (water)	87	-0.233	0.030
U	Antimony (water)	86	0.211	0.050
U	Arsenic (water)	110	0.188	0.049
U	Manganese (water)	117	-0.181	0.050
U	Iron (NURE)	117	-0.178	0.054
U	²²⁴ Radium (water)	53	-0.177	0.203
U	Acid neutralizing capacity (water)	116	0.176	0.058
U	Depth to water table	117	-0.170	0.066
U	Deuterium (water)	58	-0.167	0.210
U	Cadmium (water)	87	0.165	0.127
U	Potassium (water)	117	0.162	0.081
U	Specific conductance, field (water)	117	0.160	0.085
U	Population density	117	-0.156	0.093
U	Selenium (NURE)	29	-0.154	0.425
U	Well depth	117	0.153	0.100
C	¹⁸ Oxygen (water)	58	-0.150	0.260

Appendix 3. Selected rho and probability values from Spearman correlations of selected trace elements, major ions, and physical properties in ground water, National Uranium Resource Evaluation program streambed sediment, terrestrial gamma-ray emissions, and ancillary data with uranium concentrations in ground water, by aquifer group, in the northern United States, 1993–2003.

—Continued

Radionuclide	Correlation variable	Number of samples	Spearman's coefficient (rho)	<i>P</i> -value
	New York and New Eng	land Crystalline (NEC) aquifer	Continued	
U	Fluoride (water)	117	0.144	0.120
U	Total dissolved solids (water)	117	0.143	0.125
U	Potassium (NURE)	116	-0.141	0.132
U	Sodium (water)	117	0.138	0.138
U	Water temperature	117	0.137	0.141
U	Selenium (water)	76	-0.134	0.250
U	Aluminum (water)	86	-0.130	0.231
U	Silica (water)	117	-0.113	0.227
U	Sodium (NURE)	117	0.112	0.230
U	Beryllium (water)	87	0.109	0.314
U	pH, field (water)	117	0.107	0.251
U	²²⁸ Radium (water)	58	-0.100	0.453
U	Uranium (NURE)	117	0.100	0.284
U	Ammonia (water)	117	-0.098	0.292
U	Strontium (water)	58	0.097	0.467
U	Runoff	117	0.089	0.336
U	⁴⁰ Potassium (gamma-ray)	117	0.088	0.343
U	Ammonia + organic nitrogen (water)	65	-0.080	0.526
U	Magnesium (water)	117	0.078	0.405
U	Zinc (water)	86	0.066	0.548
U	Dissolved organic carbon (water)	81	0.064	0.569
U	Ortho-phosphate (water)	117	-0.060	0.521
U	precip90	117	-0.059	0.528
U	Dissolved oxygen (water)	115	-0.054	0.566
U	Boron (water)	53	-0.049	0.730
U	Age of water (time of recharge)	55	0.042	0.760
U	Molybdenum (NURE)	116	-0.039	0.673
U	Lithium (NURE)	116	-0.036	0.697
U	Bromide (water)	117	-0.032	0.729
U	Bicarbonate (water)	88	-0.029	0.786
U	Sulfate (water)	116	-0.025	0.789
U	Phosphorus (water)	117	-0.024	0.800
U	Depth to top of open interval	117	-0.022	0.814
U	Nitrate (water)	117	-0.015	0.868
U	Copper (water)	87	-0.012	0.911
U	Nickel (water)	87	-0.007	0.948
U	Chloride (water)	117	0.001	0.990
	Northea	stern Glacial (NEG) aquifer		
U	Gross alpha particle activity (water)	63	0.543	< 0.0001
U	Sulfate (water)	126	0.368	< 0.0001
U	Field pH (water)	126	0.345	< 0.0001
U	Acid neutralizing capacity (water)	124	0.310	< 0.0001
U	Molybdenum (water)	126	0.306	< 0.0001

Appendix 3. Selected rho and probability values from Spearman correlations of selected trace elements, major ions, and physical properties in ground water, National Uranium Resource Evaluation program streambed sediment, terrestrial gamma-ray emissions, and ancillary data with uranium concentrations in ground water, by aquifer group, in the northern United States, 1993–2003.

—Continued

Radionuclide	Correlation variable	Number of samples	Spearman's coefficient (rho)	<i>P</i> -value
	Northeastern	Glacial (NEG) aquifer—Contin		
U	Well depth	125	0.303	< 0.0001
U	Bicarbonate (water)	126	0.302	< 0.0001
U	Depth to bottom of open interval	116	0.291	0.002
U	Calcium (water)	126	0.287	0.001
U	Depth to top of open interval	116	0.286	0.002
U	¹⁸ Oxygen (water)	38	-0.262	0.111
U	Deuterium (water)	38	-0.262	0.112
U	Aluminum (water)	104	-0.249	0.011
U	Arsenic (water)	126	0.248	0.005
U	Strontium (water)	125	0.220	0.014
U	Chloride (water)	126	-0.207	0.020
U	Dissolved oxygen (water)	126	-0.204	0.022
U	precip90	58	-0.200	0.131
U	Gross beta particle activity (water)	63	0.196	0.124
U	Potassium (water)	126	0.190	0.033
U	Total dissolved solids (water)	123	0.187	0.039
U	²²⁶ Radium (water)	50	0.176	0.220
U	Specific conductance, field (water)	126	0.167	0.061
U	Calcium (NURE)	61	0.164	0.207
U	Lithium (water)	125	0.160	0.074
U	Magnesium (water)	126	0.155	0.082
U	Population density	125	-0.139	0.120
U	Cobalt (water)	126	-0.124	0.166
U	Depth to water table	58	-0.124	0.355
U	Selenium (NURE)	55	0.120	0.382
U	Lithium (NURE)	119	0.116	0.208
U	Sodium (water)	126	-0.111	0.217
U	Iron (water)	126	0.108	0.229
U	Sodium (NURE)	125	-0.106	0.241
U	⁴⁰ Potassium (gamma-ray)	124	0.095	0.292
U	Iron (NURE)	125	0.087	0.337
U	Bromide (water)	126	0.086	0.341
U	Dissolved organic carbon (water)	120	-0.082	0.373
U	Molybdenum (NURE)	119	0.080	0.385
U	²²⁸ Radium (water)	53	0.074	0.597
U	Uranium (NURE)	125	-0.073	0.419
U	Ortho-phosphate (water)	126	0.071	0.429
U	Barium (water)	126	-0.054	0.748
U	²²⁴ Radium (water)	42	0.051	0.748
U	Lead (water)	126	-0.042	0.639
U	Water temperature	126	-0.042	0.641
U	Ammonia (water)	126	-0.040	0.659
U	²³² Thorium (gamma-ray)	108	0.038	0.695
U	Runoff	58	-0.038	0.776

Appendix 3. Selected rho and probability values from Spearman correlations of selected trace elements, major ions, and physical properties in ground water, National Uranium Resource Evaluation program streambed sediment, terrestrial gamma-ray emissions, and ancillary data with uranium concentrations in ground water, by aquifer group, in the northern United States, 1993–2003.

—Continued

Radionuclide	Correlation variable	Number of samples	Spearman's coefficient (rho)	<i>P</i> -value
	Northeastern (Glacial (NEG) aquifer—Contin		
U	Zinc (water)	126	-0.034	0.704
U	Vanadium (water)	125	-0.031	0.735
U	Beryllium (water)	126	-0.030	0.736
U	Boron (water)	113	-0.028	0.771
U	Copper (water)	126	-0.028	0.759
U	Ammonia + organic nitrogen (water)	126	-0.027	0.763
U	Silica (water)	126	0.016	0.861
U	Radon (water)	92	0.014	0.890
U	Potassium (NURE)	119	-0.014	0.883
U	Phosphorus (water)	58	0.008	0.952
U	Nitrate (water)	116	-0.008	0.933
U	Turbidity (water)	38	0.008	0.964
U	Thallium (water)	98	0.007	0.944
U	Nickel (water)	126	-0.007	0.941
U	Manganese (water)	126	-0.005	0.952
U	Fluoride (water)	126	-0.009	0.991
		Pacific Glacial (NPG) aquifer		
U	No correlations between uranium in ground	water and other variables.		
	Pre-Wisco	onsinan glacial (PWG) aquifer		
U	Strontium (water)	11	0.607	0.048
U	Sulfate (water)	71	0.560	< 0.0001
U	Field pH (water)	71	-0.519	< 0.0001
U	Calcium (water)	70	0.467	< 0.0001
U	Nickel (water)	71	0.453	< 0.0001
U	Phosphorus (water)	66	-0.449	< 0.0001
U	Dissolved oxygen (water)	71	0.448	< 0.0001
U	Ammonia (water)	71	-0.442	< 0.0001
U	Depth to top of open interval	71	-0.441	< 0.0001
U	Nitrate (water)	71	0.434	< 0.0001
U	Ortho-phosphate (water)	71	-0.434	< 0.0001
U	Iron (water)	71	-0.424	< 0.0001
U	Well depth	71	-0.422	< 0.0001
U	Depth to bottom of open interval	71	-0.421	< 0.0001
U	Barium (water)	71	-0.410	< 0.0001
U	Radon (water)	70	0.406	< 0.0001
U	Copper (water)	71	0.390	< 0.0001
U	Bromide (water)	69	0.376	0.001
U	Magnesium (water)	71	0.358	0.002
U	Runoff	71	-0.356	0.002
U	Molybdenum (water)	71	-0.347	0.002
U	Potassium (water)	71	-0.327	0.005
U	Tritium (water)	34	0.326	0.060
U	Aluminum (water)	71	0.320	0.007
U	Cobalt (water)	71	0.314	0.008

Appendix 3. Selected rho and probability values from Spearman correlations of selected trace elements, major ions, and physical properties in ground water, National Uranium Resource Evaluation program streambed sediment, terrestrial gamma-ray emissions, and ancillary data with uranium concentrations in ground water, by aquifer group, in the northern United States, 1993–2003.

—Continued

Radionuclide	Correlation variable	Number of samples	Spearman's coefficient (rho)	<i>P</i> -value
	Pre-Wisconsina	n glacial (PWG) aquifer—Con	tinued	
U	Turbidity (water)	63	-0.312	0.013
U	Sodium (water)	71	-0.307	0.009
U	Total dissolved solids (water)	70	0.288	0.015
U	Antimony (water)	71	0.285	0.016
U	Ammonia + organic nitrogen (water)	53	-0.276	0.046
U	Arsenic (water)	71	-0.275	0.020
U	Dissolved organic carbon (water)	70	-0.242	0.044
U	Zinc (water)	71	0.240	0.044
U	Fluoride (water)	71	-0.226	0.057
U	Nitrite (water)	71	0.188	0.116
U	precip90	71	-0.176	0.141
U	Acid neutralizing capacity (water)	60	-0.163	0.213
U	Water temperature	71	0.149	0.214
U	Selenium (water)	71	0.131	0.277
U	Specific conductance, field (water)	71	0.122	0.311
U	⁴⁰ Potassium (gamma-ray)	71	-0.080	0.507
U	²³² Thorium (gamma-ray)	69	-0.073	0.553
U	Depth to water table	71	-0.067	0.580
U	Lead (water)	71	-0.056	0.641
U	Manganese (water)	71	-0.042	0.730
U	Chloride (water)	70	0.029	0.809
U	Population density	71	-0.023	0.852
U	Silica (water)	71	0.010	0.937
		entral Glacial (WCG) aquifer	0.010	0.507
U	Selenium (water)	116	0.598	< 0.0001
U	Carbonate (water)	135	0.524	< 0.0001
U	Potassium (water)	169	0.521	< 0.0001
U	Aluminum (water)	52	-0.483	< 0.0001
U	⁴⁰ Potassium (gamma-ray)	175	0.462	< 0.0001
U	Iron (water)	174	-0.458	< 0.0001
U	Lithium (water)	72	0.431	< 0.0001
U	Fluoride (water)	176	0.428	< 0.0001
U	²³² Thorium (gamma-ray)	175	0.425	< 0.0001
U	Lithium (NURE)	132	0.416	< 0.0001
U	Acid neutralizing capacity (water)	96	0.410	< 0.0001
U	Sodium (water)	176	0.398	< 0.0001
U	Total dissolved solids (water)	176	0.390	<0.0001
U	Bicarbonate (water)	176	0.373	<0.0001
U	Calcium (water)	176	0.373	<0.0001
	Potassium (NURE)	132	0.368	< 0.0001
		132	0.500	<0.0001
U		132	-0.363	∠0 0001
U U	Ammonia + organic nitrogen (water)	133	-0.363 0.357	< 0.0001
U		133 78 161	-0.363 0.357 0.335	<0.0001 0.001 <0.0001

Appendix 3. Selected rho and probability values from Spearman correlations of selected trace elements, major ions, and physical properties in ground water, National Uranium Resource Evaluation program streambed sediment, terrestrial gamma-ray emissions, and ancillary data with uranium concentrations in ground water, by aquifer group, in the northern United States, 1993–2003.

—Continued

Radionuclide	Correlation variable	Number of samples	Spearman's coefficient (rho)	<i>P</i> -value
	West-Central	Glacial (WCG) aquifer—Contir	nued	
U	Specific conductance, field (water)	175	0.328	< 0.0001
U	Strontium (water)	72	0.315	0.007
U	Manganese (water)	176	-0.311	< 0.0001
U	Calcium (NURE)	132	-0.304	< 0.0001
U	Vanadium (water)	72	-0.303	0.010
U	Ammonia (water)	172	-0.301	< 0.0001
U	Depth to bottom of open interval	124	0.300	< 0.0001
U	Turbidity (water)	30	-0.292	0.118
U	Well depth	131	0.280	0.001
U	Sulfate (water)	175	0.265	< 0.0001
U	Depth to top of open interval	125	0.258	0.004
U	Phosphorus (water)	100	-0.250	0.012
U	Runoff	105	-0.248	0.011
U	Copper (water)	116	0.243	0.009
U	Bromide (water)	176	0.229	0.002
U	Dissolved oxygen (water)	175	0.218	0.004
U	Selenium (NURE)	132	0.214	0.014
U	Zinc (water)	116	-0.195	0.036
U	Barium (water)	116	0.194	0.037
U	Magnesium (water)	176	0.193	0.010
U	Silica (water)	176	0.181	0.016
U	Molybdenum (water)	116	0.177	0.057
U	Uranium (NURE)	132	0.176	0.043
U	precip90	105	-0.176	0.073
U	Age of water (time of recharge)	28	-0.169	0.390
U	Arsenic (water)	122	0.157	0.084
U	Thallium (water)	72	-0.155	0.193
U	Cobalt (water)	115	-0.150	0.110
U	Water temperature	175	0.127	0.094
U	Ortho-phosphate (water)	169	0.126	0.103
U	Molybdenum (NURE)	132	0.116	0.185
U	Sodium (NURE)	132	-0.113	0.195
U	Dissolved organic carbon (water)	162	-0.111	0.160
U	Radon (water)	60	0.064	0.627
U	pH, field (water)	177	-0.060	0.424
U	Chloride (water)	175	0.058	0.444
U	Nitrite (water)	172	0.050	0.516
U	Depth to water table	105	-0.050	0.613
U	Iron (NURE)	132	0.041	0.640
U	Population density	177	0.014	0.851
U	Lead (water)	115	0.013	0.892
U	Antimony (water)	116	0.012	0.901
U	Nickel (water)	116	-0.010	0.916

References

Phillips, J.D., Duval, J.S., and Ambroziak, R.A., 1993, National Geophysical Data Grids: Gamma-ray, gravity, magnetic, and topographic data for the conterminous United States: U.S. Geological Survey Digital Data Series DDS-9, accessed online May 2005, at http://ftpext.usgs.gov/pub/cr/co/denver/musette/pub/GEOPHYSICAL_DATA/cdrom_DDS-9/

Wolock, D.M., 2003, Estimated mean annual natural ground-water recharge in the conterminous United States: U.S. Geological Survey Open-File Report 03–311, online at http://water.usgs.gov/lookup/getspatial?rech48grd.

Appendix 4. Selected rho and probability values from Spearman correlations of selected trace elements, major ions, and physical properties in ground water, National Uranium Resource Evaluation program streambed sediment, terrestrial gamma-ray emissions, and other ancillary data with ²²²radon concentrations in ground water, by aquifer group, in the northern United States, 1993–2003.

For each aguifer group, the order of the correlation variable is listed by the highest correlation (either negative or positive) value to the lowest correlation value. Spearman's coefficients (rho) are significant when the P-values are less than or equal to 0.05; a negative rho value indicates an inverse relation. The following variables consist of geochemical data for the United States based primarily on streambed sediments and analyzed for using a consistent set of methods as part of the USGS National Geochemical Survey (U.S. Geological Survey Open-File Report 2004–1001). Most of the original samples were collected as part of the U.S. Geological Survey's National Uranium Resource Evaluation (NURE) Program from 1975 to 1983 and re-analyzed (data were accessed online May 2005, at http://tin.er.usgs.gov/geochem/doc/home.htm).

The variables are expressed in parts per million, unless stated otherwise:

Calcium (NURE) in percent weight

Molybdenum (NURE)

Selenium (NURE)

Sodium (NURE) in percent weight

Lithium (NURE)

Lead (NURE)

Potassium (NURE)

Iron (NURE)

Uranium (NURE)

< less than ²²²Rn radon

Runoff mean annual runoff, for 1951 to 1980, in inches (Wolock, D.M., 2003)

mean annual precipitation, for 1961 through 1990 (Source of data is at Oregon State University in precip90

Corvallis, OR, accessed online December 2006, at http://www.ocs.oregonstate.edu/prism/index.phtml)

Population

Density

population density from the 2000 Census, in persons per square kilometer within 500 meters of sampled wells

¹⁸Oxygen a stable isotope, reported as the ratio per mill (%o, or per thousand) of $^{18}O/^{16}O$ ($\delta^{18}O$) Deuterium a stable isotope, reported as the ratio per mill (%o, or per thousand) of ${}^{2}H/{}^{1}H$ (δD)

40Potassium aerial gamma-ray survey across the conterminous United States that measured the gamma-ray flux produced

by the radioactive decay of the naturally-occurring element 40 Potassium in the top few centimeters of rock and

soil, expressed in units of percent potassium (Phillips and others, 1993)

²³²Thorium aerial gamma-ray survey across the conterminous United States that measure the gamma-ray flux produced

by the radioactive decay of the naturally-occurring element ²³²Thorium in the top few centimeters of rock and

soil, expressed as parts per million equivalent thorium (eTh) (Phillips and others, 1993).

Appendix 4. Selected rho and probability values from Spearman correlations of selected trace elements, major ions, and physical properties in ground water, National Uranium Resource Evaluation program streambed sediment, terrestrial gamma-ray emissions, and other ancillary data with ²²²radon concentrations in ground water, by aquifer group, in the northern United States, 1993–2003.

—Continued

Radionuclide	Correlation variable	Number of samples	Spearman's coefficient (rho)	<i>P</i> -value
	Cambrian-O	rdovician (COS) aquifer		
²²² Rn	Sodium (NURE)	11	0.474	0.141
²²² Rn	Thallium (water)	26	0.393	0.047
222 Rn	Depth to water table	76	-0.278	0.015
222 Rn	Dissolved organic carbon (water)	74	0.215	0.066
222 Rn	precip90	76	0.200	0.083
222 Rn	Nickel (water)	73	-0.179	0.129
222 Rn	Runoff	76	0.175	0.130
222 Rn	Nitrite (water)	76	0.174	0.133
222 Rn	Aluminum (water)	71	-0.170	0.155
222 Rn	Age of water (recharge date)	64	0.167	0.186
²²² Rn	Water temperature	77	-0.167	0.146
222 Rn	Lead (water)	73	-0.166	0.161
²²² Rn	Strontium (water)	26	0.164	0.424
²²² Rn	Silica (water)	76	0.162	0.162
²²² Rn	Zinc (water)	73	-0.149	0.210
²²² Rn	Well depth	77	-0.123	0.288
²²² Rn	Depth to top of open interval	76	-0.114	0.328
²²² Rn	pH, field (water)	77	-0.108	0.348
²²² Rn	Potassium (water)	76	0.107	0.357
²²² Rn	⁴⁰ Potassium (gamma-ray)	77	0.105	0.365
²²² Rn	Arsenic (water)	73	-0.103	0.384
²²² Rn	Iron (water)	76	-0.093	0.424
²²² Rn	Barium (water)	73	0.087	0.463
²²² Rn	Uranium (water)	73	0.087	0.463
²²² Rn	Acid neutralizing capacity (water)	75	0.080	0.498
²²² Rn	Cobalt (water)	73	-0.078	0.512
²²² Rn	Nitrate (water)	76	0.073	0.529
²²² Rn	Turbidity (water)	71	0.066	0.582
²²² Rn	Dissolved oxygen (water)	76	-0.057	0.623
²²² Rn	Lithium (water)	26	-0.056	0.784
²²² Rn	Calcium (water)	76	-0.052	0.654
²²² Rn	Fluoride (water)	76	0.051	0.663
²²² Rn	Molybdenum (water)	73	-0.047	0.691
²²² Rn	Total dissolved solids (water)	76	0.045	0.697
²²² Rn	Sodium (water)	76	0.045	0.699
²²² Rn	Ammonia + organic nitrogen (water)	39	0.042	0.802
²²² Rn	Selenium (water)	73	0.035	0.772
²²² Rn	²³² Thorium (gamma-ray)	73	0.033	0.772
²²² Rn	Ortho-phosphate (water)	75 76	-0.032	0.784
²²² Rn	Magnesium (water)	76 76	0.031	0.784
²²² Rn	Ammonia (water)	76 76	0.031	0.790
²²² Rn		76		
	Copper (water)		-0.022	0.855
²²² Rn	Tritium (water)	50	0.021	0.887

Appendix 4. Selected rho and probability values from Spearman correlations of selected trace elements, major ions, and physical properties in ground water, National Uranium Resource Evaluation program streambed sediment, terrestrial gamma-ray emissions, and other ancillary data with ²²²radon concentrations in ground water, by aquifer group, in the northern United States, 1993–2003.

—Continued

Radionuclide	Correlation variable	Number of samples	Spearman's coefficient (rho)	<i>P</i> -value
	Cambrian-Ordovic	ian (COS) aquifer—Contin	ued	
²²² Rn	Population density	77	-0.016	0.888
²²² Rn	Chloride (water)	76	-0.013	0.914
²²² Rn	Specific conductance, field (water)	77	-0.006	0.957
²²² Rn	Boron (water)	26	0.004	0.982
²²² Rn	Phosphorus (water)	75	-0.003	0.982
²²² Rn	Sulfate (water)	76	0.002	0.984
²²² Rn	Manganese (water)	77	0.001	0.991
	Columbia Plat	eau Glacial (CPG) aquifer		
²²² Rn	Lithium (water)	33	0.510	0.002
²²² Rn	Lead (NURE)	21	-0.417	0.060
²²² Rn	Depth to bottom of screen interval	83	0.370	0.001
²²² Rn	Well depth	85	0.368	0.001
²²² Rn	Bromide (water)	85	0.348	0.001
²²² Rn	Phosphorus (water)	85	-0.326	0.002
²²² Rn	Nickel (water)	33	0.323	0.067
²²² Rn	Vanadium (water)	33	-0.311	0.078
²²² Rn	Thallium (water)	33	-0.301	0.089
²²² Rn	Iron (water)	85	0.298	0.006
²²² Rn	Calcium (NURE)	21	-0.279	0.220
²²² Rn	Zinc (water)	33	0.274	0.123
²²² Rn	Arsenic (water)	33	-0.258	0.147
²²² Rn	Ortho-phosphate (water)	85	-0.247	0.023
²²² Rn	Molybdenum (water)	33	0.231	0.197
²²² Rn	Depth to top of open interval	84	0.225	0.040
²²² Rn	Selenium (water)	33	0.216	0.228
²²² Rn	Boron (water)	33	-0.202	0.256
²²² Rn	Bicarbonate (water)	76	-0.188	0.104
²²² Rn	Depth to water table	85	0.187	0.087
²²² Rn	⁴⁰ Potassium (gamma-ray)	85	0.186	0.088
²²² Rn	Uranium (water)	33	0.176	0.327
²²² Rn	Sulfate (water)	85	0.158	0.148
²²² Rn	Chloride (water)	85	0.156	0.155
²²² Rn	Manganese (water)	85	0.142	0.196
²²² Rn	Nitrate (water)	85	-0.132	0.227
²²² Rn	Copper (water)	33	0.122	0.498
²²² Rn	Ammonia + organic nitrogen (water)	44	0.100	0.518
²²² Rn	Water temperature	81	0.097	0.389
²²² Rn	Magnesium (water)	85	-0.095	0.387
²²² Rn	Potassium (water)	85	0.076	0.387
²²² Rn	Calcium (water)	85	-0.065	0.491
²²² Rn	Specific conductance, field (water)	85	-0.065	0.556
²²² Rn	Strontium (water)	33	-0.053	0.336
²²² Rn		85		
KII	Total dissolved solids (water)	83	-0.051	0.641

Appendix 4. Selected rho and probability values from Spearman correlations of selected trace elements, major ions, and physical properties in ground water, National Uranium Resource Evaluation program streambed sediment, terrestrial gamma-ray emissions, and other ancillary data with ²²²radon concentrations in ground water, by aquifer group, in the northern United States, 1993–2003.

—Continued

Radionuclide	Correlation variable	Number of samples	Spearman's coefficient (rho)	<i>P</i> -value
	Columbia Plateau Gl	acial (CPG) aquifer—Cont	inued	
²²² Rn	Fluoride (water)	85	0.046	0.673
²²² Rn	Barium (water)	33	-0.037	0.836
²²² Rn	Nitrite (water)	85	0.037	0.737
²²² Rn	Silica (water)	85	0.029	0.790
²²² Rn	Dissolved organic carbon (water)	61	-0.029	0.827
²²² Rn	Aluminum (water)	33	-0.017	0.927
²²² Rn	Acid neutralizing capacity (water)	61	-0.015	0.907
²²² Rn	Population density	85	-0.015	0.888
²²² Rn	Ammonia (water)	85	0.012	0.914
²²² Rn	precip90	85	0.012	0.916
²²² Rn	Sodium (water)	85	-0.012	0.917
²²² Rn	Dissolved oxygen (water)	85	-0.010	0.926
²²² Rn	pH, field (water)	85	-0.003	0.979
74.1		ol Glacial (ECG) aquifer	0.000	0.,,,
²²² Rn	Calcium (NURE)	35	-0.707	< 0.0001
²²² Rn	Sodium (NURE)	39	-0.672	< 0.0001
²²² Rn	Lithium (NURE)	39	0.583	0.000
²²² Rn	Uranium (NURE)	39	0.570	0.000
²²² Rn	Nitrate (water)	336	0.554	< 0.0001
²²² Rn	Iron (water)	336	-0.507	< 0.0001
²²² Rn	precip90	337	0.500	< 0.0001
²²² Rn	Ammonia (water)	336	-0.486	< 0.0001
²²² Rn	Selenium (NURE)	23	-0.464	0.026
²²² Rn	Dissolved oxygen (water)	331	0.440	< 0.0001
²²² Rn	Arsenic (water)	241	-0.430	< 0.0001
²²² Rn	Manganese (water)	337	-0.428	< 0.0001
²²² Rn	Ammonia + organic nitrogen (water)	216	-0.398	< 0.0001
²²² Rn	Runoff	337	0.381	< 0.0001
²²² Rn	²²⁸ Radium (water)	26	-0.357	0.074
²²² Rn	Sulfate (water)	333	-0.349	< 0.0001
222Rn		39	0.348	0.030
222Rn	Molybdenum (NURE) Molybdenum (water)	241		< 0.0001
222Rn	²³² Thorium (gamma-ray)	296	-0.341 0.337	< 0.0001
²²² Rn		39		
	Potassium (NURE)		-0.334	0.038
²²² Rn	Silica (water)	336	-0.327	< 0.0001
²²² Rn	Dissolved organic carbon (water)	330	-0.301	<0.0001
²²² Rn	Iron (NURE)	39	-0.301	0.063
²²² Rn	Bicarbonate (water)	70	0.282	0.018
²²² Rn	Fluoride (water)	336	-0.278	< 0.0001
²²² Rn	Deuterium (water)	36	0.264	0.120
²²² Rn	Thallium (water)	43	-0.264	0.088
²²² Rn	¹⁸ Oxygen (water)	36	0.249	0.143
²²² Rn	Copper (water)	241	0.248	0.000
²²² Rn	Acid neutralizing capacity (water)	189	-0.239	0.001

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—Continued

Radionuclide	Correlation variable	Number of samples	Spearman's coefficient (rho)	<i>P</i> -value
	East-Central Glac	ial (ECG) aquifer—Continu	ied	
²²² Rn	Lithium (water)	132	-0.239	0.006
²²² Rn	pH, field (water)	335	-0.225	< 0.0001
²²² Rn	Total dissolved solids (water)	334	-0.199	0.000
²²² Rn	Boron (water)	139	-0.198	0.019
²²² Rn	Uranium (water)	241	-0.191	0.003
²²² Rn	Nickel (water)	242	-0.188	0.003
²²² Rn	Magnesium (water)	336	-0.172	0.002
²²² Rn	Cobalt (water)	241	-0.161	0.013
²²² Rn	Selenium (water)	241	0.141	0.029
²²² Rn	Strontium (water)	132	-0.139	0.113
²²² Rn	Calcium (water)	336	-0.136	0.012
²²² Rn	Specific conductance, field (water)	336	-0.134	0.014
²²² Rn	Aluminum (water)	241	-0.132	0.041
²²² Rn	Nitrite (water)	336	-0.131	0.016
²²² Rn	Potassium (water)	336	-0.124	0.023
²²² Rn	Chloride (water)	336	0.102	0.062
²²² Rn	Depth to water table	337	0.097	0.076
²²² Rn	Antimony (water)	240	-0.087	0.178
²²² Rn	Turbidity (water)	182	0.085	0.256
²²² Rn	Water temperature	334	-0.068	0.213
²²² Rn	Zinc (water)	240	-0.046	0.477
²²² Rn	Ortho-phosphate (water)	336	-0.043	0.427
²²² Rn	Well depth	337	-0.041	0.449
²²² Rn	Tritium (water)	120	0.037	0.686
²²² Rn	Depth to bottom of open interval	329	-0.034	0.535
²²² Rn	Sodium (water)	336	-0.031	0.575
²²² Rn	Vanadium (water)	132	-0.026	0.764
²²² Rn	²³² Thorium (gamma-ray)	337	-0.025	0.651
²²² Rn	Barium (water)	242	-0.024	0.708
²²² Rn	Population density	337	0.023	0.669
²²² Rn	Lead (water)	241	0.017	0.788
²²² Rn	Total phosphorus (water)	316	-0.006	0.910
	North-Centr	al Glacial (NCG) aquifer		
²²² Rn	Sodium (NURE)	22	0.348	0.113
²²² Rn	Gross alpha particle activity (water)	48	-0.344	0.017
²²² Rn	Nitrate (water)	88	0.342	0.001
²²² Rn	Iron (water)	107	-0.329	0.001
²²² Rn	precip90	79	-0.303	0.007
²²² Rn	Bicarbonate (water)	30	0.284	0.128
²²² Rn	²²⁴ Radium (water)	27	-0.282	0.153
²²² Rn	Gross beta particle activity (water)	44	-0.279	0.067
²²² Rn	Manganese (water)	107	-0.270	0.005
²²² Rn	Total phosphorus (water)	75	0.256	0.027
²²² Rn	Runoff	79	-0.249	0.027

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—Continued

Radionuclide	Correlation variable	Number of samples	Spearman's coefficient (rho)	<i>P</i> -value
	North-Central Glac	ial (NCG) aquifer—Contin	nued	
²²² Rn	²²⁸ Radium (water)	27	-0.233	0.242
²²² Rn	²²⁶ Radium (water)	76	-0.226	0.050
²²² Rn	Selenium (water)	91	0.220	0.036
²²² Rn	Zinc (water)	76	0.215	0.062
²²² Rn	Nickel (water)	81	0.212	0.058
²²² Rn	²³² Thorium (gamma-ray)	107	0.211	0.029
²²² Rn	Ortho-phosphate (water)	102	0.209	0.035
²²² Rn	Turbidity (water)	76	0.207	0.072
²²² Rn	Depth to top of open interval	79	-0.200	0.077
²²² Rn	Lithium (NURE)	29	0.195	0.311
²²² Rn	Copper (water)	82	0.188	0.090
²²² Rn	Acid neutralizing capacity (water)	31	0.185	0.320
²²² Rn	Ammonia (water)	103	-0.178	0.072
²²² Rn	Depth to bottom of open interval	79	-0.173	0.128
²²² Rn	Well depth	79	-0.170	0.133
²²² Rn	Thallium (water)	55	-0.170	0.215
²²² Rn	Molybdenum (NURE)	29	0.169	0.380
²²² Rn	Cobalt (water)	83	0.169	0.127
²²² Rn	Depth to water table	79	-0.155	0.173
²²² Rn	Sodium (water)	107	0.141	0.149
²²² Rn	Lithium (water)	86	-0.133	0.223
²²² Rn	Molybdenum (water)	81	-0.127	0.260
²²² Rn	Potassium (water)	102	0.121	0.227
²²² Rn	⁴⁰ Potassium (gamma-ray)	107	0.110	0.261
²²² Rn	pH, field (water)	107	-0.109	0.263
²²² Rn	Dissolved organic carbon (water)	100	0.102	0.313
²²² Rn	Uranium (water)	107	0.102	0.297
²²² Rn	Specific conductance, field (water)	107	0.096	0.326
²²² Rn	Lead (water)	81	0.094	0.403
²²² Rn	Strontium (water)	86	-0.086	0.430
²²² Rn	Dissolved oxygen (water)	103	0.078	0.436
²²² Rn	Barium (water)	91	-0.076	0.473
²²² Rn	Total dissolved solids (water)	105	0.071	0.472
²²² Rn	Chloride (water)	106	0.069	0.484
²²² Rn	Selenium (NURE)	29	-0.062	0.751
²²² Rn	Ammonia + organic nitrogen (water)	97	-0.058	0.574
²²² Rn	Calcium (water)	107	0.042	0.669
²²² Rn	Magnesium (water)	107	0.040	0.679
²²² Rn	Arsenic (water)	83	0.040	0.721
²²² Rn	Silica (water)	107	0.039	0.689
²²² Rn	Cadmium (water)	91	0.036	0.734
²²² Rn	Nitrite (water)	103	-0.032	0.749
²²² Rn	Water temperature	107	-0.024	0.804
²²² Rn	Fluoride (water)	106	-0.024	0.855

Appendix 4. Selected rho and probability values from Spearman correlations of selected trace elements, major ions, and physical properties in ground water, National Uranium Resource Evaluation program streambed sediment, terrestrial gamma-ray emissions, and other ancillary data with ²²²radon concentrations in ground water, by aquifer group, in the northern United States, 1993–2003.

—Continued

Radionuclide	Correlation variable	Number of samples	Spearman's coefficient (rho)	<i>P</i> -value	
	North-Central Glad	cial (NCG) aquifer—Contin	ued		
²²² Rn	Uranium (NURE)	29	0.016	0.935	
²²² Rn	Population density	79	0.012	0.916	
²²² Rn	Sulfate (water)	102	0.008	0.933	
²²² Rn	Aluminum (water)	43	-0.007	0.965	
²²² Rn	Boron (water)	77	0.001	0.100	
	New York and New E	England Crystalline (NEC) a	quifer		
²²² Rn	Uranium (water)	110	0.598	< 0.0001	
²²² Rn	Gross alpha particle activity (water)	53	0.537	< 0.0001	
²²² Rn	Gross beta particle activity (water)	54	0.424	0.001	
²²² Rn	Lead (water)	80	0.378	0.001	
²²² Rn	Strontium (water)	54	-0.347	0.010	
²²² Rn	Barium (water)	80	-0.347	0.002	
²²² Rn	Sulfate (water)	109	-0.328	0.001	
²²² Rn	Calcium (NURE)	45	0.325	0.029	
²²² Rn	Bicarbonate (water)	84	-0.324	0.003	
²²² Rn	²²⁶ Radium (water)	54	0.257	0.060	
²²² Rn	Beryllium (water)	47	0.245	0.028	
²²² Rn	Boron (water)	47	-0.242	0.102	
²²² Rn	²³² Thorium (gamma-ray)	106	0.228	0.019	
²²² Rn	Iron (NURE)	110	-0.222	0.020	
²²² Rn	⁴⁰ Potassium (gamma-ray)	110	0.224	0.020	
²²² Rn	Lithium (water)	54	0.200	0.148	
²²² Rn	Cadmium (water)	80	0.192	0.087	
²²² Rn	Fluoride (water)	110	0.188	0.049	
²²² Rn	Molybdenum (water)	80	0.187	0.097	
²²² Rn	Sodium (NURE)	110	0.184	0.054	
²²² Rn	¹⁸ Oxygen (water)	54	0.171	0.217	
²²² Rn	Antimony (water)	79	-0.158	0.164	
²²² Rn	Age of water (recharge date)	51	0.154	0.282	
²²² Rn	Acid neutralizing capacity (water)	109	-0.139	0.148	
²²² Rn	Deuterium (water)	54	0.138	0.320	
²²² Rn	Zinc (water)	79	0.133	0.244	
²²² Rn	Water temperature	110	0.126	0.190	
²²² Rn	Total dissolved solids (water)	110	-0.122	0.203	
²²² Rn	Iron (water)	110	-0.122	0.203	
²²² Rn	Phosphorus (water)	110	0.118	0.219	
²²² Rn	Copper (water)	80	0.118	0.219	
222Rn	Runoff	110	0.113	0.219	
222Rn	Dissolved oxygen (water)	108	0.113	0.239	
²²² Rn	Dissolved oxygen (water) Dissolved organic carbon (water)	75			
²²² Rn			-0.101	0.386	
²²² Rn ²²² Rn	Potassium (water)	110	0.097	0.315	
	Specific conductance, field (water)	110	-0.096	0.317	
²²² Rn	²²⁴ Radium (water)	51	0.089	0.536	

Appendix 4. Selected rho and probability values from Spearman correlations of selected trace elements, major ions, and physical properties in ground water, National Uranium Resource Evaluation program streambed sediment, terrestrial gamma-ray emissions, and other ancillary data with ²²²radon concentrations in ground water, by aquifer group, in the northern United States, 1993–2003. —Continued

Radionuclide	Correlation variable	Number of samples	Spearman's coefficient (rho)	<i>P</i> -value
	New York and New England	d Crystalline (NEC) aquifer	—Continued	
²²² Rn	Nickel (water)	80	-0.080	0.481
²²² Rn	Nitrite (water)	110	-0.080	0.409
²²² Rn	Molybdenum (NURE)	110	-0.069	0.473
²²² Rn	Uranium (NURE)	110	0.068	0.483
²²² Rn	Manganese (water)	110	-0.064	0.507
²²² Rn	Magnesium (water)	110	-0.060	0.530
²²² Rn	Well depth	110	-0.059	0.537
²²² Rn	Population density	110	-0.059	0.537
²²² Rn	Ammonia (water)	110	-0.058	0.548
²²² Rn	Silica (water)	110	-0.058	0.550
²²² Rn	Ortho-phosphate (water)	110	0.045	0.639
²²² Rn	Calcium (water)	110	-0.042	0.665
²²² Rn	Chloride (water)	110	-0.038	0.690
²²² Rn	precip90	110	0.037	0.703
²²² Rn	pH, field (water)	110	-0.037	0.705
²²² Rn	Ammonia + organic nitrogen (water)	59	-0.021	0.874
²²² Rn	Arsenic (water)	104	-0.019	0.846
²²² Rn	Sodium (water)	110	0.018	0.853
222 Rn	²²⁸ Radium (water)	56	0.017	0.901
²²² Rn	Potassium (NURE)	109	0.016	0.872
222 Rn	Nitrate (water)	110	-0.013	0.893
²²² Rn	Aluminum (water)	79	0.008	0.943
222 Rn	Depth to top of open interval	110	-0.008	0.936
222 Rn	Depth to water table	110	-0.003	0.976
²²² Rn	Selenium (water)	71	-0.001	0.994
		n Glacial (NEG) aquifer		
²²² Rn	Water temperature, in deg. C	93	-0.461	< 0.0001
²²² Rn	Barium (water)	93	-0.414	< 0.0001
²²² Rn	Deuterium (water)	39	-0.286	0.078
²²² Rn	Nickel (water)	93	-0.285	0.006
²²² Rn	Fluoride (water)	93	0.265	0.010
²²² Rn	¹⁸ Oxygen (water)	39	-0.263	0.105
²²² Rn	²²⁴ Radium (water)	43	-0.258	0.095
222 Rn	Nitrite (water)	93	-0.248	0.017
²²² Rn	²²⁸ Radium (water)	54	-0.221	0.108
²²² Rn	Dissolved organic carbon (water)	89	-0.209	0.049
²²² Rn	Silica (water)	93	0.208	0.045
²²² Rn	Potassium (NURE)	86	0.198	0.043
²²² Rn	Sodium (NURE)	92	0.198	0.067
222Rn	Well depth	93	0.192	0.066
222Rn	Ammonia + organic nitrogen (water)	93	-0.191	0.067
		93	-0.191	0.087
²²² Rn Cobalt (water)		73	-0.101	0.063
	²²² Rn Aluminum (water)		-0.176	0.108

Appendix 4. Selected rho and probability values from Spearman correlations of selected trace elements, major ions, and physical properties in ground water, National Uranium Resource Evaluation program streambed sediment, terrestrial gamma-ray emissions, and other ancillary data with ²²²radon concentrations in ground water, by aquifer group, in the northern United States, 1993–2003.

—Continued

Radionuclide	Correlation variable	Number of samples	Spearman's coefficient (rho)	<i>P</i> -value	
	Northeastern Glad	ial (NEG) aquifer—Continued			
²²² Rn	Iron (water)	93	-0.168	0.107	
²²² Rn	Population density	93	-0.168	0.106	
²²² Rn	pH, field (water)	93	0.167	0.111	
²²² Rn	Boron (water)	80	-0.161	0.153	
²²² Rn	⁴⁰ Potassium (gamma-ray)	91	0.161	0.128	
²²² Rn	Manganese (water)	93	-0.161	0.124	
²²² Rn	Specific conductance, field (water)	93	-0.159	0.128	
²²² Rn	Sodium (water)	93	-0.158	0.130	
²²² Rn	Molybdenum (water)	92	0.150	0.154	
²²² Rn	Potassium (water)	93	0.148	0.156	
²²² Rn	Gross alpha particle activity (water)	63	-0.143	0.265	
²²² Rn	Thallium (water)	64	-0.142	0.262	
²²² Rn	Depth to water table	59	-0.138	0.299	
²²² Rn	Lithium (water)	92	0.136	0.197	
²²² Rn	²³² Thorium (gamma-ray)	75	0.135	0.248	
²²² Rn	Age of water (recharge date)	33	0.129	0.476	
²²² Rn	Total dissolved solids (water)	90	-0.125	0.240	
²²² Rn	Zinc (water)	93	-0.122	0.242	
²²² Rn	Strontium (water)	92	0.122	0.247	
²²² Rn	Dissolved oxygen (water)	93	0.122	0.246	
²²² Rn	Phosphorus (water)	59	-0.120	0.363	
²²² Rn	Vanadium (water)	91	-0.116	0.272	
²²² Rn	Magnesium (water)	93	0.112	0.286	
²²² Rn	Turbidity (water)	39	-0.104	0.527	
²²² Rn	Bicarbonate (water)	93	0.102	0.330	
²²² Rn	Arsenic (water)	93	-0.102	0.333	
²²² Rn	Depth to top of open interval	84	0.097	0.382	
²²² Rn	Beryllium (water)	92	0.093	0.379	
²²² Rn	Chloride (water)	93	-0.086	0.411	
²²² Rn	Ammonia (water)	93	-0.083	0.429	
²²² Rn	²²⁶ Radium (water)	51	0.080	0.576	
²²² Rn	Acid neutralizing capacity (water)	93	0.059	0.572	
²²² Rn	Selenium (NURE)	47	-0.058	0.698	
²²² Rn	Molybdenum (NURE)	86	0.055	0.614	
²²² Rn	Lithium (NURE)	86	0.049	0.655	
²²² Rn	Nitrate (water)	87	-0.046	0.675	
²²² Rn	precip90	59	-0.036	0.789	
²²² Rn	Sulfate (water)	93	-0.035	0.738	
²²² Rn	Iron (NURE)	92	0.028	0.790	
²²² Rn	Gross beta particle activity (water)	63	-0.025	0.847	
²²² Rn	Calcium (NURE)	50	0.020	0.893	
²²² Rn	Lead (water)	92	0.017	0.871	
²²² Rn	Ortho-phosphate (water)	93	0.016	0.882	
²²² Rn	Copper (water)	93	0.015	0.887	

Appendix 4. Selected rho and probability values from Spearman correlations of selected trace elements, major ions, and physical properties in ground water, National Uranium Resource Evaluation program streambed sediment, terrestrial gamma-ray emissions, and other ancillary data with ²²²radon concentrations in ground water, by aquifer group, in the northern United States, 1993–2003.

—Continued

Radionuclide	Correlation variable	Number of samples	Spearman's coefficient (rho)	<i>P</i> -value	
	Northeastern Glad	cial (NEG) aquifer—Contin	ued		
²²² Rn	Uranium (water)	92	0.014	0.892	
²²² Rn	Runoff	59	0.013	0.921	
²²² Rn	Calcium (water)	93	0.012	0.908	
	Northern Pac	cific Glacial (NPG) aquifer			
²²² Rn	²³² Thorium (gamma-ray)	42	0.535	0.000	
222 Rn	⁴⁰ Potassium (gamma-ray)	43	0.501	0.001	
²²² Rn	Aluminum (water)	31	-0.421	0.018	
²²² Rn	Potassium (NURE)	24	0.420	0.041	
²²² Rn	Arsenic (water)	30	-0.401	0.028	
²²² Rn	Field pH (water)	77	-0.326	0.004	
²²² Rn	Iron (water)	77	-0.323	0.004	
²²² Rn	Depth to bottom of screen interval	50	-0.305	0.031	
²²² Rn	Depth to water table	77	0.304	0.007	
²²² Rn	Molybdenum (water)	31	-0.291	0.112	
²²² Rn	Nitrate (water)	77	0.266	0.019	
²²² Rn	Barium (water)	31	-0.263	0.152	
²²² Rn	Calcium (NURE)	31	-0.241	0.191	
²²² Rn	precip90	46	-0.239	0.110	
²²² Rn	²²⁶ Radium (water)	31	-0.218	0.240	
²²² Rn	Manganese (water)	77	-0.208	0.069	
²²² Rn	Gross alpha particle activity (water)	31	0.191	0.303	
²²² Rn	Runoff	45	-0.169	0.141	
²²² Rn	Bicarbonate (water)	62	-0.155	0.230	
²²² Rn	Population density	46	0.151	0.314	
²²² Rn	Gross beta particle activity (water)	31	0.148	0.425	
²²² Rn	Cobalt	31	0.143	0.443	
²²² Rn	Water temperature	77	0.142	0.218	
²²² Rn	Iron (NURE)	31	-0.142	0.448	
²²² Rn	Chloride	76	0.137	0.239	
²²² Rn	Well depth	77	-0.136	0.238	
²²² Rn	Sodium (NURE)	31	-0.135	0.467	
²²² Rn	Phosphorus (water)	77	-0.130	0.467	
²²² Rn	Silica (water)	77	0.128	0.269	
222Rn	Magnesium (water)	77	-0.125	0.209	
222 R n	Depth to top of open interval	77	-0.123	0.277	
222Rn	Zinc (water)	31	0.115	0.237	
222Rn	, ,	77			
²²² Rn	Sodium (water)	31	0.114 0.099	0.322 0.595	
²²² Rn	Copper (water)	77		0.393	
	Ortho-phosphate		-0.098		
²²² Rn	Acid neutralizing capacity (water)	46	-0.097	0.523	
²²² Rn	Lithium (NURE)	30	0.095	0.617	
²²² Rn	Nitrite (water)	77	0.088	0.447	
²²² Rn	Nickel (water)	31	-0.083	0.655	
²²² Rn	Potassium (water)	77	0.081	0.486	

Appendix 4. Selected rho and probability values from Spearman correlations of selected trace elements, major ions, and physical properties in ground water, National Uranium Resource Evaluation program streambed sediment, terrestrial gamma-ray emissions, and other ancillary data with 222 radon concentrations in ground water, by aquifer group, in the northern United States, 1993–2003. —Continued

Radionuclide	Correlation variable	Number of samples	Spearman's coefficient (rho)	<i>P</i> -value	
	Northern Pacific Gla	cial (NPG) aquifer—Cont	inued		
²²² Rn	Specific conductance (water)	77	-0.074	0.521	
²²² Rn	Uranium (NURE)	31	-0.073	0.696	
²²² Rn	Ammonia (water)	77	-0.047	0.682	
²²² Rn	Dissolved oxygen (water)	77	0.046	0.694	
²²² Rn	Lead (water)	31	0.041	0.827	
²²² Rn	Sulfate (water)	76	0.036	0.757	
²²² Rn	¹⁸ Oxygen (water)	28	-0.036	0.857	
²²² Rn	Deuterium (water)	28	-0.029	0.883	
²²² Rn	Dissolved organic carbon (water)	77	-0.025	0.831	
²²² Rn	Total dissolved soilds (water)	76	-0.022	0.850	
²²² Rn	²²⁴ Radium (water)	21	-0.020	0.933	
²²² Rn	²²⁸ Radium (water)	29	0.019	0.923	
²²² Rn	Ammonia + organic nitrogen (water)	34	-0.018	0.918	
²²² Rn	Calcium (water)	77	0.012	0.919	
²²² Rn	Fluoride (water)	76	-0.012	0.921	
²²² Rn	Selenium (water)	29	-0.008	0.968	
	Pre-Wisconsin	nan Glacial (PWG) aquifer			
²²² Rn	Copper (water)	70	0.558	< 0.0001	
²²² Rn	Dissolved oxygen (water)	173	0.541	< 0.0001	
²²² Rn	Depth to bottom of screen interval	157	-0.535	< 0.0001	
²²² Rn	Barium (water)	70	-0.519	< 0.0001	
²²² Rn	Depth to top of screen interval	157	-0.503	< 0.0001	
²²² Rn	Molybdenum (water)	70	-0.477	< 0.0001	
²²² Rn	Ammonia (water)	175	-0.473	< 0.0001	
²²² Rn	Well depth	175	-0.449	< 0.0001	
²²² Rn	Iron (water)	175	-0.408	< 0.0001	
²²² Rn	Uranium (water)	70	0.406	0.000	
²²² Rn	Ammonia + organic nitrogen (water)	95	-0.404	< 0.0001	
²²² Rn	²²⁸ Radium (water)	4	0.400	0.600	
²²² Rn	Arsenic (water)	70	-0.344	0.004	
²²² Rn	Ortho-phosphate (water)	175	-0.330	< 0.0001	
²²² Rn	Runoff	175	-0.328	< 0.0001	
²²² Rn	Selenium (water)	70	0.319	0.007	
²²² Rn	Nitrate (water)	175	0.308	< 0.0001	
²²² Rn	⁴⁰ Potassium (gamma-ray)	175	-0.300	0.000	
²²² Rn	Sulfate (water)	175	0.295	0.000	
²²² Rn	precip90	175	-0.293	0.000	
²²² Rn	Acid neutralizing capacity (water)	164	-0.290	0.000	
²²² Rn	Field pH (water)	175	-0.289	0.000	
²²² Rn	Fluoride (water)	175	-0.287	0.000	
²²² Rn	Phosphorus (water)	170	-0.281	0.000	
²²² Rn	Sodium (water)	175	-0.281	0.000	
²²² Rn	Dissolved organic carbon (DOC)	174	-0.274	0.000	
222 R n	Dissolved organic carbon (DOC)	83	-0.274	0.000	

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—Continued

Radionuclide	Correlation variable	Number of samples	Spearman's coefficient (rho)	<i>P</i> -value	
	Pre-Wisconsinan G	lacial (PWG) aquifer—Con	tinued		
²²² Rn	Deuterium (water)	69	-0.228	0.059	
²²² Rn	Population density	175	0.190	0.012	
²²² Rn	Calcium (water)	174	0.164	0.031	
²²² Rn	Specific conductance (water)	175	-0.160	0.034	
²²² Rn	Potassium (water)	175	-0.159	0.035	
²²² Rn	Magnesium (water)	175	-0.152	0.044	
²²² Rn	²³² Thorium (gamma-ray)	173	-0.145	0.057	
²²² Rn	Cobalt (water)	70	0.144	0.233	
²²² Rn	Bicarbonate (water)	90	0.130	0.222	
²²² Rn	Lead (water)	70	0.128	0.290	
²²² Rn	Manganese (water)	166	-0.117	0.132	
²²² Rn	Turbidity (water)	161	0.111	0.162	
²²² Rn	Water temperature	173	-0.100	0.189	
²²² Rn	Nickel (water)	70	0.091	0.455	
²²² Rn	Total dissolved solids (water)	174	-0.087	0.255	
²²² Rn	Zinc (water)	70	-0.075	0.539	
²²² Rn	Tritium (water)	133	-0.072	0.411	
²²² Rn	Antimony (water)	70	0.057	0.641	
²²² Rn	Chloride (water)	174	-0.045	0.558	
²²² Rn	Aluminum (water)	70	-0.032	0.791	
²²² Rn	Carbonate (water)	90	-0.026	0.805	
²²² Rn	Nitrite (water)	175	-0.020	0.862	
²²² Rn	Silica (water)	175	-0.013	0.959	
²²² Rn	Depth to water table	175	-0.004	0.980	
TO		ral Glacial (WCG) aquifer	0.002	0.700	
²²² Rn	Depth to top of screen interval	63	-0.364	0.003	
²²² Rn	Tritium (water)	34	0.352	0.041	
²²² Rn	Depth to bottom of screen interval	62	-0.348	0.006	
²²² Rn	Well depth	70	-0.345	0.003	
²²² Rn	¹⁸ Oxygen (water)	17	-0.338	0.184	
²²² Rn	²³² Thorium (gamma-ray)	76	-0.337	0.003	
²²² Rn	⁴⁰ Potassium (gamma-ray)	76	-0.322	0.005	
222 R n	Depth to water table	50	0.321	0.003	
²²² Rn	Thallium (water)	26	0.307	0.127	
²²² Rn	Potassium (NURE)	64	-0.298	0.127	
222 R n	Iron (water)	68	-0.249	0.017	
222 R n	Bromide (water)	70		0.046	
²²² Rn	Boron (water)	32	-0.267 0.234	0.026	
²²² Rn	Phosphorus (water)	32 44	-0.221	0.198	
²²² Rn	-	64			
	Molybdenum (NURE)		-0.206	0.102	
²²² Rn	Lead (water)	30	0.204	0.280	
²²² Rn	Nickel (water)	31 31	0.188 0.179	0.312 0.334	
²²² Rn Selenium (water)		4.1	11 1 /9	0.554	

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Radionuclide	Correlation variable	Number of samples	Spearman's coefficient (rho)	<i>P</i> -value
	West-Central Glac			
²²² Rn	Lithium (NURE)	64	-0.165	0.191
²²² Rn	Ammonia (water)	70	-0.160	0.187
²²² Rn	Dissolved organic carbon (water)	69	0.159	0.193
²²² Rn	Water temperature	70	-0.153	0.207
²²² Rn	Carbonate (water)	69	-0.147	0.227
²²² Rn	Nitrite (water)	70	0.142	0.241
²²² Rn	Nitrate (water)	66	0.134	0.282
²²² Rn	Molybdenum (water)	31	0.132	0.479
²²² Rn	Magnesium (water)	69	0.125	0.306
²²² Rn	Selenium (NURE)	64	-0.121	0.341
²²² Rn	Fluoride (water)	70	-0.112	0.356
²²² Rn	Ortho-phosphate (water)	70	-0.110	0.366
²²² Rn	Copper (water)	31	0.106	0.569
²²² Rn	Vanadium (water)	26	0.106	0.608
²²² Rn	Chloride (water)	70	-0.102	0.400
²²² Rn	Population density	76	-0.097	0.405
²²² Rn	Silica (water)	69	-0.096	0.434
²²² Rn	Potassium (water)	70	-0.092	0.449
²²² Rn	Arsenic (water)	37	0.089	0.602
²²² Rn	Runoff	50	0.089	0.54
²²² Rn	Strontium (water)	26	0.074	0.719
²²² Rn	Iron (NURE)	64	-0.071	0.577
²²² Rn	Lithium (water)	26	0.070	0.735
²²² Rn	Sodium (NURE)	64	-0.067	0.597
²²² Rn	Barium (water)	31	0.064	0.731
²²² Rn	Uranium (water)	60	0.064	0.627
²²² Rn	Specific conductance (water)	68	-0.059	0.634
²²² Rn	precip90	50	-0.057	0.693
²²² Rn	Turbidity (water)	10	0.055	0.881
²²² Rn	Ammonia + organic nitrogen (water)	54	0.053	0.703
²²² Rn	Sodium (water)	69	-0.051	0.680
²²² Rn	Dissolved oxygen (water)	69	-0.050	0.683
²²² Rn	pH, field (water)	70	0.050	0.683
²²² Rn	Sulfate (water)	70	-0.045	0.709
²²² Rn	Manganese (water)	69	0.044	0.719
²²² Rn	Bicarbonate (water)	70	0.030	0.803
²²² Rn	Zinc (water)	31	-0.014	0.942
²²² Rn	Acid neutralizing capacity (water)	43	0.012	0.938
²²² Rn	Calcium (water)	69	-0.011	0.929
²²² Rn	Total dissolved solids (water)	70	-0.010	0.932
²²² Rn	Uranium (NURE)	64	-0.005	0.970

References

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Appendix 5. Significant rho and probability values from Spearman correlations of selected trace elements, major ions, and physical properties in ground water, National Uranium Resource Evaluation program streambed sediment, terrestrial gamma-ray emissions, and other ancillary data with uranium concentrations in selected NAWQA well networks in the Columbia Plateau glacial, West-Central glacial, and New York and New England crystalline aquifer groups, in the northern United States, 1993–2003.

For each aquifer group, the order of the correlation variable is listed by the highest correlation (either negative or positive) value to the lowest correlation value. Spearman's coefficients (rho) are significant when the *P*-values are less than or equal to 0.05; a negative rho value indicates an inverse relation. The following variables consist of geochemical data for the United States based primarily on streambed sediments and analyzed for using a consistent set of methods as part of the USGS National Geochemical Survey (U.S. Geological Survey Open-File Report 2004–1001). Most of the original samples were collected as part of the U.S. Geological Survey's National Uranium Resource Evaluation (NURE) Program from 1975 to 1983 and re-analyzed (data were accessed online May 2005, at http://tin.er.usgs.gov/geochem/doc/home.htm).

The variables are expressed in parts per million, unless stated otherwise:

Calcium (NURE) in percent weight

Molybdenum (NURE)

Selenium (NURE)

Sodium (NURE) in percent weight

Lithium (NURE)

Lead (NURE)

Potassium (NURE)

Iron (NURE)

Uranium (NURE)

< less than
U uranium

deg. C degrees Celsius

precip90 mean annual precipitation, for 1961 through 1990 (Source of data is at Oregon State University in

Corvallis, OR, accessed online December 2006, at http://www.ocs.oregonstate.edu/prism/index.phtml)

¹⁸Oxygen a stable isotope, reported as the ratio per mill (%o, or per thousand) of ¹⁸O/¹⁶O (δ ¹⁸O) Deuterium a stable isotope, reported as the ratio per mill (%o, or per thousand) of ²H/¹H (δ D)

⁴⁰Potassium aerial gamma-ray survey across the conterminous United States that measured the gamma-ray flux produced

by the radioactive decay of the naturally-occurring element 40 Potassium in the top few centimeters of rock and

soil, expressed in units of percent potassium (Phillips and others, 1993)

²³²Thorium aerial gamma-ray survey across the conterminous United States that measure the gamma-ray flux produced

by the radioactive decay of the naturally-occurring element 232Thorium in the top few centimeters of rock and

soil, expressed as parts per million equivalent thorium (eTh) (Phillips and others, 1993).

Appendix 5. Significant rho and probability values from Spearman correlations of selected trace elements, major ions, and physical properties in ground water, National Uranium Resource Evaluation program streambed sediment, terrestrial gamma-ray emissions, and other ancillary data with uranium concentrations in selected NAWQA well networks in the Columbia Plateau glacial, West-Central glacial, and New York and New England crystalline aquifer groups, in the northern United States, 1993–2003.—Continued

NAWQA well network (See appendix 1 for definition)	Aquifer group code	Radionuclide	Variable name	Number of samples	Spearman's coefficient (rho)	<i>P</i> -value
ccptlusag2b	CPG	U	Bicarbonate (water)	26	0.826	< 0.0001
ccptlusag2b	CPG	U	Calcium (NURE)	10	0.745	0.013
ccptlusag2b	CPG	U	Sodium (water)	27	0.737	< 0.0001
ccptlusag2b	CPG	U	Ammonia + organic nitrogen (water)	27	0.664	0.000
ccptlusag2b	CPG	U	Acid neutralizing capacity (water)	16	0.664	0.005
ccptlusag2b	CPG	U	Boron (water)	27	0.638	_
ccptlusag2b	CPG	U	Lithium (water)	27	0.616	0.001
ccptlusag2b	CPG	U	Specific conductance, field (water)	27	0.595	0.001
ccptlusag2b	CPG	U	Molybdenum (NURE)	12	-0.544	0.068
ccptlusag2b	CPG	U	Total dissolved solids (water)	27	0.531	0.004
ccptlusag2b	CPG	U	Copper (water)	27	0.527	0.005
ccptlusag2b	CPG	U	Field pH (water)	27	-0.500	0.008
ccptlusag2b	CPG	U	Dissolved organic carbon (water)	27	0.471	0.013
ccptlusag2b	CPG	U	Strontium (water)	27	0.463	0.015
ccptlusag2b	CPG	U	Sodium (NURE)	12	-0.452	0.140
ccptlusag2b	CPG	U	Iron (NURE)	12	0.424	0.170
ccptlusag2b	CPG	U	²²² Radon (water)	16	0.421	0.105
ccptlusag2b	CPG	U	Magnesium (water)	27	0.403	0.037
ccptlusag2b	CPG	U	Sulfate (water)	27	0.386	0.046
ccptlusag2b	CPG	U	Thallium (water)	27	-0.373	0.055
ccptlusag2b	CPG	U	Manganese (water)	27	0.371	0.057
ccptlusag2b	CPG	U	Uranium (NURE)	12	0.354	0.259
ccptlusag2b	CPG	U	Molybdenum (water)	27	0.344	0.079
ccptlusag2b	CPG	U	Fluoride (water)	27	0.326	0.097
ccptlusag2b	CPG	U	precip90	16	0.298	0.263
ccptlusag2b	CPG	U	Barium (water)	27	0.284	0.150
ccptlusag2b	CPG	U	⁴⁰ Potassium (gamma-ray)	27	0.279	0.158
ccptlusag2b	CPG	U	Ammonia (water)	27	-0.277	0.162
ccptlusag2b	CPG	U	Selenium (water)	27	0.264	0.184
ccptlusor1b	CPG	U	Specific conductance, field (water)	19	0.914	< 0.0001
ccptlusor1b	CPG	U	Total dissolved solids (water)	19	0.909	< 0.0001
ccptlusor1b	CPG	U	Acid neutralizing capacity (water)	19	0.719	0.001
ccptlusor1b	CPG	U	Bicarbonate (water)	19	0.705	0.001
ccptlusor1b	CPG	U	Sodium (water)	19	0.691	0.001
ccptlusor1b	CPG	U	Ammonia + organic nitrogen (water)	19	0.681	0.001
ccptlusor1b	CPG	U	Sulfate (water)	19	0.644	0.003
ccptlusor1b	CPG	U	Dissolved organic carbon (water)	19	0.558	0.013
ccptlusor1b	CPG	U	Strontium (water)	19	0.540	0.017
ccptlusor1b	CPG	U	Nitrate (water)	19	0.467	0.044
ccptlusor1b	CPG	U	Selenium (water)	19	0.455	0.050
ccptlusor1b	CPG	U	Field pH (water)	19	-0.441	0.059
ccptlusor1b	CPG	U	Depth to water table	19	-0.437	0.061

Appendix 5. Significant rho and probability values from Spearman correlations of selected trace elements, major ions, and physical properties in ground water, National Uranium Resource Evaluation program streambed sediment, terrestrial gamma-ray emissions, and other ancillary data with uranium concentrations in selected NAWQA well networks in the Columbia Plateau glacial, West-Central glacial, and New York and New England crystalline aquifer groups, in the northern United States, 1993–2003.—Continued

NAWQA well network (See appendix 1 for definition)	Aquifer group code	Radionuclide	Variable name	Number of samples	Spearman's coefficient (rho)	<i>P</i> -value
ccptlusor1b	CPG	U	Nickel (water)	19	0.434	0.063
ccptlusor1b	CPG	U	Chloride (water)	19	0.433	0.064
ccptlusor1b	CPG	U	Water temperature, in deg. C	19	-0.433	0.064
ccptlusor1b	CPG	U	Magnesium (water)	19	0.432	0.065
ccptlusor1b	CPG	U	Ortho phosphate (water)	19	0.404	0.086
ccptlusor1b	CPG	U	Phosphorus (water)	19	0.401	0.089
ccptlusor1b	CPG	U	Calcium (water)	19	0.396	0.093
ccptlusor1b	CPG	U	Iron (water)	19	0.387	0.101
ccptlusor1b	CPG	U	Copper (water)	19	0.346	0.147
ccptlusor1b	CPG	U	Thallium (water)	19	0.344	0.149
ccptlusor1b	CPG	U	Zinc (water)	19	0.295	0.221
cnbrluscr1	WCG	U	Lithium (water)	29	0.810	<0.0001
cnbrluser1	WCG	U	Boron (water)	29	0.790	_
cnbrluscr1	WCG	U	Potassium (water)	26	0.714	< 0.0001
cnbrluscr1	WCG	U	Sodium (water)	29	0.574	0.001
cnbrluscr1	WCG	U	Strontium (water)	29	0.510	0.005
cnbrluser1	WCG	U	Selenium (water)	29	0.504	0.005
cnbrluscr1	WCG	U	Total dissolved solids (water)	29	0.486	0.007
cnbrluser1	WCG	U	Uranium (NURE)	28	-0.471	0.011
cnbrluscr1	WCG	U	Fluoride (water)	29	0.459	0.012
cnbrluscr1	WCG	U	²³² Thorium (gamma-ray)	28	-0.449	0.016
cnbrluser1	WCG	U	Specific conductance, field (water)	29	0.437	0.018
cnbrluscr1	WCG	U	Sulfate (water)	28	0.425	0.024
cnbrluscr1	WCG	U	Bicarbonate (water)	28	0.420	0.026
cnbrluscr1	WCG	U	Calcium (water)	29	0.353	0.060
cnbrluscr1	WCG	U	Cabonate (water)	28	0.339	0.078
cnbrluscr1	WCG	U	Aluminum (water)	6	-0.338	0.512
cnbrluscr1	WCG	U	Iron (water)	28	-0.332	0.084
cnbrluscr1	WCG	U	Chloride (water)	29	0.324	0.086
cnbrluscr1	WCG	U	Silica (water)	29	0.304	0.109
cnbrluscr1	WCG	U	Calcium (NURE)	28	0.297	0.125
cnbrluscr1	WCG	U	⁴⁰ Potassium (gamma-ray)	28	-0.277	0.154
cnbrluscr1	WCG	U	Arsenic (water)	29	0.271	0.155
cnbrluscr1	WCG	U	Manganese (water)	29	-0.264	0.166
cnbrluscr1	WCG	U	Potassium (NURE)	28	0.263	0.176
cnbrluscr1	WCG	U	Lithium (NURE)	28	0.258	0.184
hpgwdwgs1	WCG	U	Ammonia + organic nitrogen (water)	15	-0.718	0.003
hpgwdwgs1	WCG	U	Ammonia (water)	15	-0.659	0.008
hpgwdwgs1	WCG	U	Iron (water)	15	-0.643	0.010
hpgwdwgs1	WCG	U	Copper (water)	15	0.620	0.014
hpgwdwgs1	WCG	U	Iron (NURE)	15	0.532	0.041

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NAWQA well network (See appendix 1 for definition)	Aquifer group code	Radionuclide	Variable name	Number of samples	Spearman's coefficient (rho)	<i>P</i> -value
hpgwdwgs1	WCG	U	Sodium (NURE)	15	-0.509	0.053
hpgwdwgs1	WCG	U	Manganese (water)	15	-0.464	0.082
hpgwdwgs1	WCG	U	Arsenic (water)	15	-0.452	0.091
hpgwdwgs1	WCG	U	Barium (water)	15	0.398	0.142
hpgwdwgs1	WCG	U	Ortho phosphate (water)	15	0.384	0.158
hpgwdwgs1	WCG	U	Nitrite (water)	15	0.379	0.163
hpgwdwgs1	WCG	U	Molybdenum (NURE)	15	-0.376	0.168
hpgwdwgs1	WCG	U	Thallium (water)	15	-0.372	0.172
hpgwdwgs1	WCG	U	Fluoride (water)	15	-0.370	0.175
hpgwdwgs1	WCG	U	Selenium (water)	15	0.359	0.189
hpgwdwgs1	WCG	U	Boron (water)	15	-0.330	_
hpgwdwgs1	WCG	U	Selenium (NURE)	15	-0.321	0.243
hpgwdwgs1	WCG	U	Calcium (NURE)	15	-0.316	0.252
hpgwdwgs1	WCG	U	Sodium (water)	15	0.315	0.252
hpgwdwgs1	WCG	U	Dissolved organic carbon (water)	15	0.297	0.282
hpgwdwgs1	WCG	U	Cabonate (water)	13	0.291	0.335
hpgwsus4	WCG	U	Selenium (water)	27	0.589	0.001
hpgwsus4	WCG	U	Iron (water)	27	-0.487	0.010
hpgwsus4	WCG	U	Ammonia (water)	27	-0.464	0.015
hpgwsus4	WCG	U	Nitrite (water)	27	0.455	0.017
hpgwsus4	WCG	U	Tritium (water)	12	0.366	0.241
hpgwsus4	WCG	U	Sulfate (water)	27	0.364	0.062
hpgwsus4	WCG	U	Cabonate (water)	27	0.349	0.075
hpgwsus4	WCG	U	Ammonia + organic nitrogen (water)	27	-0.344	0.079
hpgwsus4	WCG	U	Depth to bottom of screen interval	27	-0.330	0.093
hpgwsus4	WCG	U	Well depth	27	-0.324	0.099
hpgwsus4	WCG	U	Calcium (water)	27	0.305	0.122
hpgwsus4	WCG	U	Copper (water)	27	0.300	0.129
hpgwsus4	WCG	U	Selenium (NURE)	27	-0.262	0.187
hpgwsus4	WCG	U	²²² Radon (water)	26	0.258	0.202
hpgwsus4	WCG	U	Depth to top of screen interval	27	-0.256	0.198
rednlusag2	WCG	U	Tritium (water)	9	-0.586	0.097
rednlusag2	WCG	U	Total dissolved solids (water)	25	0.553	0.004
rednlusag2	WCG	U	Specific conductance, field (water)	25	0.523	0.007
rednlusag2	WCG	U	Magnesium (water)	24	0.506	0.012
rednlusag2	WCG	U	Bicarbonate (water)	25	0.501	0.011
rednlusag2	WCG	U	Acid neutralizing capacity (water)	25	0.460	0.021
rednlusag2	WCG	U	Well depth	17	-0.408	0.104
rednlusag2	WCG	U	Silica (water)	24	-0.394	0.057
rednlusag2	WCG	U	Sodium (water)	24	0.385	0.063
rednlusag2	WCG	U	Sulfate (water)	25	0.384	0.058

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NAWQA well network (See appendix 1 for definition)	Aquifer group code	Radionuclide	Variable name	Number of samples	Spearman's coefficient (rho)	<i>P</i> -value
rednlusag2	WCG	U	Calcium (water)	24	0.377	0.069
rednlusag2	WCG	U	Depth to bottom of screen interval	17	-0.348	0.171
rednlusag2	WCG	U	Chloride (water)	24	0.330	0.115
rednlusag2	WCG	U	Depth to top of screen interval	17	-0.317	0.216
rednlusag2	WCG	U	Calcium (NURE)	25	-0.308	0.134
rednlusag2	WCG	U	Deuterium (water)	9	-0.285	0.458
rednlusag2	WCG	U	¹⁸ Oxygen (water)	9	-0.285	0.458
rednlusag2	WCG	U	Depth to water table	25	-0.272	0.189
rednlusag2	WCG	U	Ortho phosphate (water)	25	-0.269	0.193
rednlusag2	WCG	U	Molybdenum (NURE)	25	0.254	0.220
rednlusag2	WCG	U	Potassium (NURE)	25	0.254	0.220
rednsus1	WCG	U	Magnesium (water)	11	0.890	0.000
rednsus1	WCG	U	Total dissolved solids (water)	11	0.793	0.004
rednsus1	WCG	U	Specific conductance, field (water)	10	0.736	0.015
rednsus1	WCG	U	Acid neutralizing capacity (water)	11	0.724	0.012
rednsus1	WCG	U	⁴⁰ Potassium (gamma-ray)	11	0.724	0.012
rednsus1	WCG	U	Arsenic (water)	5	-0.700	0.188
rednsus1	WCG	U	Dissolved oxygen (water)	11	0.676	0.022
rednsus1	WCG	U	Bicarbonate (water)	11	0.661	0.027
rednsus1	WCG	U	Sulfate (water)	11	0.648	0.031
rednsus1	WCG	U	Calcium (water)	11	0.600	0.051
rednsus1	WCG	U	Potassium (water)	11	0.533	0.091
rednsus1	WCG	U	²³² Thorium (gamma-ray)	11	0.492	0.124
rednsus1	WCG	U	Iron (NURE)	11	-0.442	0.174
rednsus1	WCG	U	Iron (water)	10	-0.427	0.219
rednsus1	WCG	U	Silica (water)	11	-0.425	0.193
rednsus1	WCG	U	Selenium (NURE)	11	-0.401	0.222
rednsus1	WCG	U	Sodium (water)	11	0.392	0.233
rednsus1	WCG	U	Calcium (NURE)	11	0.351	0.290
rednsus1	WCG	U	Uranium (NURE)	11	0.287	0.392
rednsus1	WCG	U	Depth to water table	11	-0.279	0.406
rednsus1	WCG	U	Chloride (water)	11	0.269	0.424
rednsus3	WCG	U	Potassium (water)	26	0.794	<0.0001
rednsus3	WCG	U	Acid neutralizing capacity (water)	17	0.741	0.001
rednsus3	WCG	U	Sodium (water)	26	0.725	< 0.0001
rednsus3	WCG	U	Specific conductance, field (water)	25	0.715	0.000
rednsus3	WCG	U	Magnesium (water)	26	0.701	0.000
rednsus3	WCG	U	Total dissolved solids (water)	25	0.701	0.000
rednsus3	WCG	U	Calcium (water)	26	0.694	0.000
rednsus3	WCG	U	Sulfate (water)	25	0.681	0.000
rednsus3	WCG	U	Fluoride (water)	25	0.665	0.000

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NAWQA well network (See appendix 1 for definition)	Aquifer group code	Radionuclide	Variable name	Number of samples	Spearman's coefficient (rho)	<i>P</i> -value
rednsus3	WCG	U	Dissolved organic carbon (water)	17	0.650	0.005
rednsus3	WCG	U	Lithium (NURE)	19	0.641	0.003
rednsus3	WCG	U	Bicarbonate (water)	26	0.544	0.004
rednsus3	WCG	U	Chloride (water)	25	0.464	0.019
rednsus3	WCG	U	Ortho phosphate (water)	26	-0.452	0.020
rednsus3	WCG	U	Ammonia + organic nitrogen (water)	15	0.429	0.111
rednsus3	WCG	U	Uranium (NURE)	19	0.415	0.077
rednsus3	WCG	U	Selenium (NURE)	19	0.366	0.124
rednsus3	WCG	U	Depth to water table	26	0.321	0.110
rednsus3	WCG	U	Phosphorus (water)	26	-0.313	0.119
rednsus3	WCG	U	Potassium (NURE)	19	0.281	0.243
rednsus3	WCG	U	Sodium (NURE)	19	-0.253	0.296
rednsus5	WCG	U	Sodium (NURE)	7	0.800	0.031
rednsus5	WCG	U	Fluoride (water)	13	-0.794	0.001
rednsus5	WCG	U	Iron (water)	13	-0.646	0.017
rednsus5	WCG	U	Dissolved organic carbon (water)	8	0.634	0.092
rednsus5	WCG	U	Selenium (NURE)	7	-0.600	0.154
rednsus5	WCG	U	Phosphorus (water)	8	-0.584	0.129
rednsus5	WCG	U	Lithium (NURE)	7	-0.564	0.187
rednsus5	WCG	U	Ammonia (water)	8	-0.503	0.204
rednsus5	WCG	U	Ortho phosphate (water)	8	-0.500	0.207
rednsus5	WCG	U	Potassium (water)	8	0.491	0.217
rednsus5	WCG	U	Zinc (water)	13	0.468	0.107
rednsus5	WCG	U	Copper (water)	13	0.455	0.118
rednsus5	WCG	U	Ammonia + organic nitrogen (water)	7	-0.411	0.359
rednsus5	WCG	U	Arsenic (water)	13	0.404	0.172
rednsus5	WCG	U	Depth to top of screen interval	13	-0.382	0.198
rednsus5	WCG	U	Depth to bottom of screen interval	13	-0.362	0.224
rednsus5	WCG	U	Well depth	13	-0.362	0.224
rednsus5	WCG	U	Molybdenum (water)	13	0.347	0.246
rednsus5	WCG	U	Potassium (NURE)	7	0.346	0.448
rednsus5	WCG	U	Uranium (NURE)	7	0.346	0.448
rednsus5	WCG	U	Manganese (water)	13	0.296	0.326
rednsus5	WCG	U	Molybdenum (NURE)	7	0.295	0.521
rednsus5	WCG	U	⁴⁰ Potassium (gamma-ray)	13	0.294	0.329
rednsus5	WCG	U	precip90	13	0.275	0.363
rednsus5	WCG	U	Calcium (water)	13	0.269	0.374
rednsus5	WCG	U	Nickel (water)	13	0.262	0.387
rednsus5	WCG	U	Nitrite (water)	8	0.260	0.534
umislusrc1	WCG	U	Nitrite (water)	30	0.516	0.003
umislusrc1	WCG	U	Aluminum (water)	30	-0.448	0.013

Appendix 5. Significant rho and probability values from Spearman correlations of selected trace elements, major ions, and physical properties in ground water, National Uranium Resource Evaluation program streambed sediment, terrestrial gamma-ray emissions, and other ancillary data with uranium concentrations in selected NAWQA well networks in the Columbia Plateau glacial, West-Central glacial, and New York and New England crystalline aquifer groups, in the northern United States, 1993–2003.—Continued

NAWQA well network (See appendix 1 for definition)	Aquifer group code	Radionuclide	Variable name	Number of samples	Spearman's coefficient (rho)	<i>P</i> -value
umislusrc1	WCG	U	Potassium (water)	30	0.434	0.017
umislusrc1	WCG	U	Cobalt (water)	30	0.409	0.025
umislusrc1	WCG	U	Antimony (water)	30	0.376	0.041
umislusrc1	WCG	U	Manganese (water)	30	0.369	0.045
umislusrc1	WCG	U	Sodium (water)	30	0.348	0.059
umislusrc1	WCG	U	Specific conductance, field (water)	30	0.345	0.062
umislusrc1	WCG	U	Phosphorus (water)	30	-0.327	0.078
umislusrc1	WCG	U	Dissolved oxygen (water)	30	-0.317	0.088
umislusrc1	WCG	U	Sulfate (water)	30	0.315	0.090
umislusrc1	WCG	U	Magnesium (water)	30	0.313	0.093
umislusrc1	WCG	U	Total dissolved solids (water)	30	0.305	0.101
umislusrc1	WCG	U	Chloride (water)	30	0.303	0.103
umislusrc1	WCG	U	Nickel (water)	30	0.295	0.114
umislusrc1	WCG	U	Turbidity (water)	30	-0.292	0.118
umislusrc1	WCG	U	Depth to water table	30	0.286	0.125
umislusrc1	WCG	U	Barium (water)	30	0.270	0.149
umislusrc1	WCG	U	Calcium (water)	30	0.263	0.160
umislusrc1	WCG	U	Tritium (water)	30	0.261	0.164
umislusrc1	WCG	U	Bicarbonate (water)	30	0.252	0.179
connsus1	NEC	U	Calcium (NURE)	7	-0.709	0.074
connsus1	NEC	U	²²² Radon (water)	30	0.621	0.000
connsus1	NEC	U	Selenium (NURE)	6	0.522	0.288
connsus1	NEC	U	Iron (water)	30	-0.491	0.006
connsus1	NEC	U	⁴⁰ Potassium (gamma-ray)	30	0.378	0.039
connsus1	NEC	U	²³² Thorium (gamma-ray)	29	0.370	0.048
connsus1	NEC	U	Calcium (water)	30	0.364	0.048
connsus1	NEC	U	Specific conductance, field (water)	30	0.361	0.050
connsus1	NEC	U	Depth to top of screen interval	30	-0.349	0.058
connsus1	NEC	U	Total dissolved solids (water)	30	0.342	0.064
connsus1	NEC	U	Phosphorus (water)	30	-0.257	0.171
linjsus1	NEC	U	Acid neutralizing capacity (water)	28	0.701	0.000
linjsus1	NEC	U	Calcium (NURE)	16	0.609	0.012
linjsus1	NEC	U	Molybdenum (water)	29	0.601	0.001
linjsus1	NEC	U	Calcium (water)	29	0.565	0.001
linjsus1	NEC	U	Field pH (water)	29	0.561	0.002
linjsus1	NEC	U	²²² Radon (water)	26	0.535	0.005
linjsus1	NEC	U	Boron (water)	29	0.505	_
linjsus1	NEC	U	Specific conductance, field (water)	29	0.472	0.010
linjsus1	NEC	U	Total dissolved solids (water)	29	0.446	0.015
linjsus1	NEC	U	Dissolved oxygen (water)	27	-0.436	0.023
linjsus1	NEC	U	Iron (water)	29	-0.398	0.032

Appendix 5. Significant rho and probability values from Spearman correlations of selected trace elements, major ions, and physical properties in ground water, National Uranium Resource Evaluation program streambed sediment, terrestrial gamma-ray emissions, and other ancillary data with uranium concentrations in selected NAWQA well networks in the Columbia Plateau glacial, West-Central glacial, and New York and New England crystalline aquifer groups, in the northern United States, 1993–2003.—Continued

NAWQA well network (See appendix 1 for definition)	Aquifer group code	Radionuclide	Variable name	Number of samples	Spearman's coefficient (rho)	<i>P</i> -value
linjsus1	NEC	U	Potassium (NURE)	29	-0.390	0.037
linjsus1	NEC	U	Fluoride (water)	29	0.378	0.043
linjsus1	NEC	U	Phosphorus (water)	29	0.357	0.057
linjsus1	NEC	U	Barium (water)	29	-0.349	0.064
linjsus1	NEC	U	Magnesium (water)	29	0.346	0.066
linjsus1	NEC	U	Manganese (water)	29	-0.337	0.074
linjsus1	NEC	U	Nickel (water)	29	-0.303	0.110
linjsus1	NEC	U	Depth to top of screen interval	29	0.295	0.120
linjsus1	NEC	U	⁴⁰ Potassium (gamma-ray)	29	-0.269	0.158
necbsus1	NEC	U	Gross alpha particle activity (water)	27	0.520	0.005
necbsus1	NEC	U	Calcium (NURE)	19	0.441	0.059
necbsus1	NEC	U	Depth to water table	28	-0.437	0.020
necbsus1	NEC	U	Strontium (water)	28	0.408	0.031
necbsus1	NEC	U	Selenium (NURE)	15	-0.382	0.160
necbsus1	NEC	U	Deutritium (water)	28	-0.364	0.057
necbsus1	NEC	U	Antimony (water)	27	0.350	0.073
necbsus1	NEC	U	Molybdenum (NURE)	28	0.350	0.068
necbsus1	NEC	U	¹⁸ Oxygen (water)	28	-0.325	0.092
necbsus1	NEC	U	²²² Radon (water)	26	0.308	0.126
necbsus1	NEC	U	Sodium (water)	28	0.252	0.195
necbsus2	NEC	U	Gross alpha particle activity (water)	28	0.875	< 0.0001
necbsus2	NEC	U	²²² Radon (water)	28	0.620	0.000
necbsus2	NEC	U	Boron (water)	18	-0.528	_
necbsus2	NEC	U	Gross beta particle activity (water)	29	0.525	0.003
necbsus2	NEC	U	Arsenic (water)	29	0.465	0.011
necbsus2	NEC	U	Dissolved oxygen (water)	30	0.414	0.023
necbsus2	NEC	U	Uranium (NURE)	30	0.397	0.030
necbsus2	NEC	U	Manganese (water)	30	-0.376	0.041
necbsus2	NEC	U	Iron (water)	30	-0.347	0.060
necbsus2	NEC	U	Lead (water)	30	0.330	0.075
necbsus2	NEC	U	Barium (water)	30	-0.304	0.102
necbsus2	NEC	U	²²⁶ Radium (water)	29	0.295	0.120
necbsus2	NEC	U	Copper (water)	30	0.284	0.128
necbsus2	NEC	U	Lithium (water)	30	0.279	0.136
necbsus2	NEC	U	²²⁴ Radium (water)	27	-0.268	0.177

Reference

Phillips, J.D., Duval, J.S., and Ambroziak, R.A., 1993, National Geophysical Data Grids: Gamma-Ray, Gravity, Magnetic, and Topographic Data for the Conterminous United States: U.S. Geological Survey Digital Data Series DDS-9, accessed online May 2005 at https://ftpext.usgs.gov/pub/cr/co/denver/musette/pub/GEOPHYSICAL_DATA/cdrom_DDS-9/

Appendix 6. Significant rho and probability values from Spearman correlations of selected trace elements, major ions, and physical properties in ground water, National Uranium Resource Evaluation program streambed sediment, terrestrial gamma-ray emissions, and other ancillary data with 222 radon concentrations in the NAWQA well networks in the New York and New England crystalline aquifer group, in the northern United States, 1993-2003.

For each well network, the order of the correlation variable is listed by the highest correlation (either negative or positive) value to the lowest correlation value. Spearman's coefficients (rho) are significant when the P-values are less than or equal to 0.05; a negative rho value indicates an inverse relation. The following variables consist of geochemical data for the United States based primarily on streambed sediments and analyzed for using a consistent set of methods as part of the USGS National Geochemical Survey (U.S. Geological Survey Open-File Report 2004–1001). Most of the original samples were collected as part of the U.S. Geological Survey's National Uranium Resource Evaluation (NURE) Program from 1975 to 1983 and re-analyzed (data were accessed online May 2005, at http://tin.er.usgs.gov/geochem/doc/home.htm).

The variables are expressed in parts per million, unless stated otherwise:

Calcium (NURE) in percent weight

Molybdenum (NURE)

Selenium (NURE)

Sodium (NURE) in percent weight

Lithium (NURE)

Lead (NURE)

Potassium (NURE)

Iron (NURE)

Uranium (NURE)

< less than ²²²Rn radon

deg. C degrees Celsius

mean annual precipitation, for 1961 through 1990 (Source of data is at Oregon State University in precip90

Corvallis, OR, accessed online December 2006, at http://www.ocs.oregonstate.edu/prism/index.phtml)

¹⁸Oxygen a stable isotope, reported as the ratio per mill (%o, or per thousand) of $^{18}O/^{16}O$ ($\delta^{18}O$) Deuterium a stable isotope, reported as the ratio per mill (%o, or per thousand) of ${}^{2}H/{}^{1}H$ (δD)

aerial gamma-ray survey across the conterminous United States that measured the gamma-ray flux produced 40Potassium

by the radioactive decay of the naturally-occurring element 40 Potassium in the top few centimeters of rock and

soil, expressed in units of percent potassium (Phillips and others, 1993)

aerial gamma-ray survey across the conterminous United States that measure the gamma-ray flux produced ²³²Thorium

by the radioactive decay of the naturally-occurring element ²³²Thorium in the top few centimeters of rock and

soil, expressed as parts per million equivalent thorium (eTh) (Phillips and others, 1993).

Appendix 6. Significant rho and probability values from Spearman correlations of selected trace elements, major ions, and physical properties in ground water, National Uranium Resource Evaluation program streambed sediment, terrestrial gamma-ray emissions, and other ancillary data with ²²²radon concentrations in the NAWQA well networks in the New York and New England crystalline aquifer group, in the northern United States, 1993–2003.—Continued

NAWQA well network See appendix 1 for definition)	Radionuclide	Correlation variable	Number of samples	Spearman's coefficient (rho)	<i>P</i> -value
connsus1	²²² Rn	Selenium (NURE)	6	0.943	0.005
connsus1	²²² Rn	Uranium (water)	30	0.621	0.000
connsus1	²²² Rn	Lithium (NURE)	30	-0.293	0.116
connsus1	²²² Rn	Bromide (water)	30	-0.271	0.148
connsus1	²²² Rn	Depth to top of open interval	30	-0.262	0.162
connsus1	²²² Rn	Sulfate (water)	30	-0.256	0.173
linjsus1	²²² Rn	Calcium (NURE)	13	0.674	0.012
linjsus1	²²² Rn	Uranium (water)	26	0.535	0.005
linjsus1	²²² Rn	Barium (water)	26	-0.518	0.007
linjsus1	²²² Rn	Depth to top of open interval	26	0.461	0.018
linjsus1	²²² Rn	Lead (NURE)	26	-0.460	0.018
linjsus1	²²² Rn	Field pH (water)	26	0.452	0.021
linjsus1	²²² Rn	Silica (water)	26	-0.426	0.030
linjsus1	²²² Rn	Fluoride (water)	26	0.389	0.049
linjsus1	²²² Rn	Phosphorus (water)	26	0.375	0.059
linjsus1	²²² Rn	Molybdenum (water)	26	0.354	0.076
linjsus1	²²² Rn	Potassium (water)	26	-0.353	0.077
linjsus1	²²² Rn	Manganese (water)	26	-0.343	0.086
linjsus1	²²² Rn	Uranium (NURE)	26	0.331	0.099
linjsus1	²²² Rn	Sulfate (water)	26	-0.327	0.103
linjsus1	²²² Rn	Dissolved oxygen (water)	24	-0.305	0.147
linjsus1	²²² Rn	Sodium (NURE)	26	0.302	0.133
linjsus1	²²² Rn	Iron (NURE)	26	0.300	0.137
linjsus1	²²² Rn	Iron (water)	26	-0.289	0.153
linjsus1	²²² Rn	Depth to water table	26	-0.282	0.162
necbsus1	²²² Rn	Dissolved organic carbon (water)	22	-0.476	0.025
necbsus1	²²² Rn	Water temperature, in deg. C	26	0.400	0.043
necbsus1	²²² Rn	Gross alpha particle activity (water)	26	0.397	0.045
necbsus1	²²² Rn	Sulfate (water)	26	-0.355	0.075
necbsus1	²²² Rn	Iron (NURE)	26	-0.325	0.105
necbsus1	²²² Rn	Uranium (water)	26	0.308	0.126
necbsus1	²²² Rn	Lead (water)	26	0.301	0.135
necbsus1	²²² Rn	Calcium (NURE)	17	0.290	0.260
necbsus1	²²² Rn	Barium (water)	26	-0.272	0.179
necbsus1	²²² Rn	Acid neutralizing capacity (water)	26	-0.268	0.185
necbsus1	²²² Rn	Dissolved oxygen (water)	26	0.256	0.207
necbsus1	²²² Rn	precip90	26	0.255	0.208
necbsus1	²²² Rn	Uranium (NURE)	26	0.252	0.214
necbsus1	²²² Rn	²³² Thorium (gamma-ray)	25	0.251	0.227
necbsus1	²²² Rn	Lead (NURE)	26	-0.251	0.217

Appendix 6. Significant rho and probability values from Spearman correlations of selected trace elements, major ions, and physical properties in ground water, National Uranium Resource Evaluation program streambed sediment, terrestrial gamma-ray emissions, and other ancillary data with ²²²radon concentrations in the NAWQA well networks in the New York and New England crystalline aquifer group, in the northern United States, 1993–2003.—Continued

NAWQA well network (See appendix 1 for definition)		Correlation variable	Number of samples	Spearman's coefficient (rho)	<i>P</i> -value	
necbsus2	²²² Rn	Uranium (water)	28	0.620	0.000	
necbsus2	²²² Rn	Gross alpha particle activity (water)	27	0.608	0.001	
necbsus2	²²² Rn	Acid neutralizing capacity (water)	28	-0.580	0.001	
necbsus2	²²² Rn	Gross beta particle activity (water)	28	0.576	0.001	
necbsus2	²²² Rn	Bicarbonate (water)	28	-0.570	0.002	
necbsus2	²²² Rn	Lead (water)	28	0.521	0.004	
necbsus2	²²² Rn	Dissolved oxygen (water)	28	0.503	0.006	
necbsus2	²²² Rn	Boron (water)	16	-0.487	_	
necbsus2	²²² Rn	Copper (water)	28	0.469	0.012	
necbsus2	²²² Rn	Strontium (water)	28	-0.442	0.019	
necbsus2	²²² Rn	Barium (water)	28	-0.405	0.032	
necbsus2	²²² Rn	Depth to bottom of open interval	28	-0.384	0.044	
necbsus2	²²² Rn	Well depth	28	-0.384	0.044	
necbsus2	²²² Rn	Field pH (water)	28	-0.383	0.044	
necbsus2	²²² Rn	Potassium (NURE)	27	0.360	0.065	
necbsus2	²²² Rn	Total dissolved solids (water)	28	-0.359	0.060	
necbsus2	²²² Rn	Specific conductance, field (water)	28	-0.349	0.068	
necbsus2	²²² Rn	Zinc (water)	28	0.347	0.071	
necbsus2	²²² Rn	²²⁶ Radium (water)	28	0.340	0.077	
necbsus2	²²² Rn	precip90	28	0.336	0.080	
necbsus2	²²² Rn	Depth to water table	28	0.322	0.095	
necbsus2	²²² Rn	Cobalt (water)	28	0.322	0.095	
necbsus2	²²² Rn	¹⁸ Oxygen (water)	28	0.319	0.098	
necbsus2	²²² Rn	⁴⁰ Potassium (gamma-ray)	28	0.308	0.111	
necbsus2	²²² Rn	Beryllium (water)	28	0.307	0.112	
necbsus2	²²² Rn	Antimony (water)	28	-0.298	0.124	
necbsus2	²²² Rn	Aluminum (water)	28	0.292	0.132	
necbsus2	²²² Rn	Time of recharge (age date)	26	0.288	0.153	
necbsus2	²²² Rn	Thallium (water)	11	-0.272	0.419	
necbsus2	²²² Rn	Iron (NURE)	28	-0.256	0.188	
necbsus2	²²² Rn	Deuterium (water)	28	0.255	0.191	
necbsus2	²²² Rn	Ortho-phosphate (water)	28	0.252	0.195	

Reference

Phillips, J.D., Duval, J.S., and Ambroziak, R.A., 1993, National Geophysical Data Grids: Gamma-ray, gravity, magnetic, and topographic data for the conterminous United States: U.S. Geological Survey Digital Data Series DDS-9, accessed online May 2005, at http://ftpext.usgs.gov/pub/cr/co/denver/musette/pub/GEOPHYSICAL_DATA/cdrom_DDS-9/

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