

**Establishment of Threshold Effects for the
Forest County Potawatomi Community (FCPC)
Class I Air Quality Related Values (AQRV)**

Final Report

**Timothy J. Sullivan
E&S Environmental Chemistry, Inc.
P.O. Box 609
Corvallis, OR 97339**

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EXECUTIVE SUMMARY

At the request of the Forest County Potawatomi Community (FCPC), the U.S. Environmental Protection Agency (EPA) redesignated the FCPC Reservation as a non-federal Class I area in April, 2008. The FCPC agreed, in the Class I Agreement with the State of Wisconsin, to list and protect air quality related values (AQRVs).

The FCPC has designated Water Quality and Aquatic Systems as AQRVs to be protected against air pollution degradation. The FCPC selected these as AQRVs for a number of reasons, including the following:

- tribal members rely on fish as part of their traditional subsistence way of life,
- pure water is important to the tribe's cultural, medicinal, and religious practices, and
- recreational fishing is important to the tourism economy in and around the FCPC reservation, upon which the FCPC depends.

These AQRVs can be adversely impacted by atmospheric deposition of Hg, S, and N. In particular, adverse effects include Hg methylation (which is related to both Hg and S deposition), acidification (which is related to both S and N deposition), and fertilization of water and soils (which is related to N deposition). The primary concern for protection of these AQRVs is the potential Hg methylation in Devils Lake and other water bodies in the 393 km² study area (which includes all land area from which surface water drains onto or within FCPC reservation lands).

To protect the AQRVs from adverse impacts, the FCPC has agreed to establish threshold effects levels. Threshold effects levels generally represent the amount of additional (above ambient) deposition from a proposed new or modified source below which an adverse effect will not occur. In addition, once atmospheric deposition of Hg, S, or N is high enough that a deposition concern threshold has been reached, which is true for Hg levels in Devils Lake and other water bodies in the study area, any further increase in emissions that would contribute additional deposition is considered to adversely impact the AQRV.

The threshold effects levels identified in this report are shown in Table ES-1. These levels are based on and consistent with extensive studies of FCPC surface waters and surface waters elsewhere in Wisconsin and the broader northern Great Lakes region, Federal Land Managers Air Quality Related Values Work Group (FLAG) guidance (as amended), U.S. Forest Service (USFS) designated concern thresholds, and USFS and National Park Service deposition analysis thresholds (DATs).

Table ES-1. Deposition analysis thresholds (DATs) for FCPC lands.

Atmospherically Deposited Pollutant	Purpose of DAT	DAT Value
N or S	Protect against water acidification or eutrophication	0.01 kg/ha/yr
S	Protect against Hg methylation	0.0012 kg/ha/yr
Hg	Protect against Hg methylation	0.0035 $\mu\text{g}/\text{m}^2/\text{yr}$

The foregoing DAT levels satisfy FCPC's obligations under the 1999 Class I Agreement and, more importantly, help to protect FCPC's essential resources for current members and future generations.

1. BACKGROUND

1.1. Introduction

The Forest County Potawatomi Community (FCPC) is a federally recognized tribe with headquarters near Crandon, Wisconsin. At the request of the FCPC, the U.S. Environmental Protection Agency (EPA) redesignated the FCPC Reservation as a non-federal Class I area in April, 2008. The FCPC agreed, in the Class I Agreement with the State of Wisconsin, to list and protect air quality related values (AQRVs).

This report identifies appropriate threshold effects levels for water quality and aquatic systems AQRVs on FCPC lands. The primary concern for protection of these AQRVs is the potential for increases in methylmercury (MeHg) in Devils Lake and other water bodies on FCPC Class I lands that could be caused by increased atmospheric deposition of either sulfur (S) or mercury (Hg). The most important route of entry of Hg into remote ecosystems in the Great Lakes region is atmospheric deposition of inorganic Hg (Fitzgerald et al. 1998). However, the most important source of the neurotoxic MeHg found in fish and other biota is via microbially mediated methylation of inorganic Hg (Ullrich et al. 2001).

Other important concerns for AQRV protection on FCPC lands include acidification of soil and surface water caused by atmospheric deposition of S and nitrogen (N) and nutrient enrichment of aquatic and terrestrial systems caused by atmospheric deposition of N.

In accordance with the Prevention of Significant Deterioration (PSD) permit application process and the Class I Agreement, the Wisconsin Department of Natural Resources (WDNR) will notify FCPC of all major emissions source permit applications located within a 62-mile radius of the reservation. As part of the permit application process, an applicant may be required by WDNR to conduct an air quality impact analysis that will assess the impacts of its proposed emissions on the FCPC Class I area. FCPC will then assess whether the proposed air emissions will adversely impact the designated AQRVs. If FCPC determines that an adverse impact is likely, FCPC will notify WDNR that it has made an adverse impact finding for the proposed source; this finding will be considered by WDNR in its permitting decisions.

1.2. Effects of Atmospheric Deposition in the Northern Great Lakes Region

1.2.1. Mercury

Based on published research findings and spatial patterns in pollutant load and resource sensitivity and effects nationwide, the air pollution effects of greatest concern in the northern Great Lakes region appear to be Hg methylation, lake acidification, and nutrient enrichment. The most thoroughly studied toxic agent that affects Class I lands in the northern Great Lakes region via atmospheric deposition pathways is Hg. It is a heavy metal that occurs naturally in several forms. Elemental Hg is released from sources in the earth's crust into the global environment through volcanic and geothermal activity and the weathering of rocks. It is also emitted by human emissions sources. Once released into the atmosphere, Hg can be deposited to the earth's surface and transformed through natural processes into a toxic form that can bioaccumulate in food webs. Contamination of aquatic ecosystems by Hg has reduced the benefits provided by fisheries resources in many inland and coastal waters throughout the United States.

Within the northern Great Lakes region, Hg is well known as constituting a risk to the health of wildlife and humans, including members of the FCPC, who consume large quantities of fish. Much of the landscape is Hg-sensitive. The major source of Hg in the watersheds of this region is atmospheric deposition (Mason and Sullivan 1997, Rolffhus et al. 2003, Wiener et al. 2006), most of which originates from human-caused sources (Swain et al. 1992, Lorey and Driscoll 1999, Lamborg et al. 2002).

Mercury emissions and deposition tend to be relatively high in some portions of the northern Great Lakes region (Figure 1). Initial scientific focus on Hg contamination in the Great Lakes region was directed at industrial point sources such as pulp and paper mills and chlor-alkali plants (Turner and Southworth 1999; Wiener et al. 2003, 2012a). However, point source release of Hg into surface waters decreased substantially in the 1980s, and atmospheric deposition is now the main source of Hg to watersheds in the Great Lakes region. The region contains a number of anthropogenic atmospheric emissions sources that collectively emit more than 50 kg per year of Hg (Wiener et al. 2012a). Watershed characteristics in portions of the region that contain abundant wetlands and forests facilitate Hg watershed inputs via dry deposition processes, methylation, transport, and biomagnification.

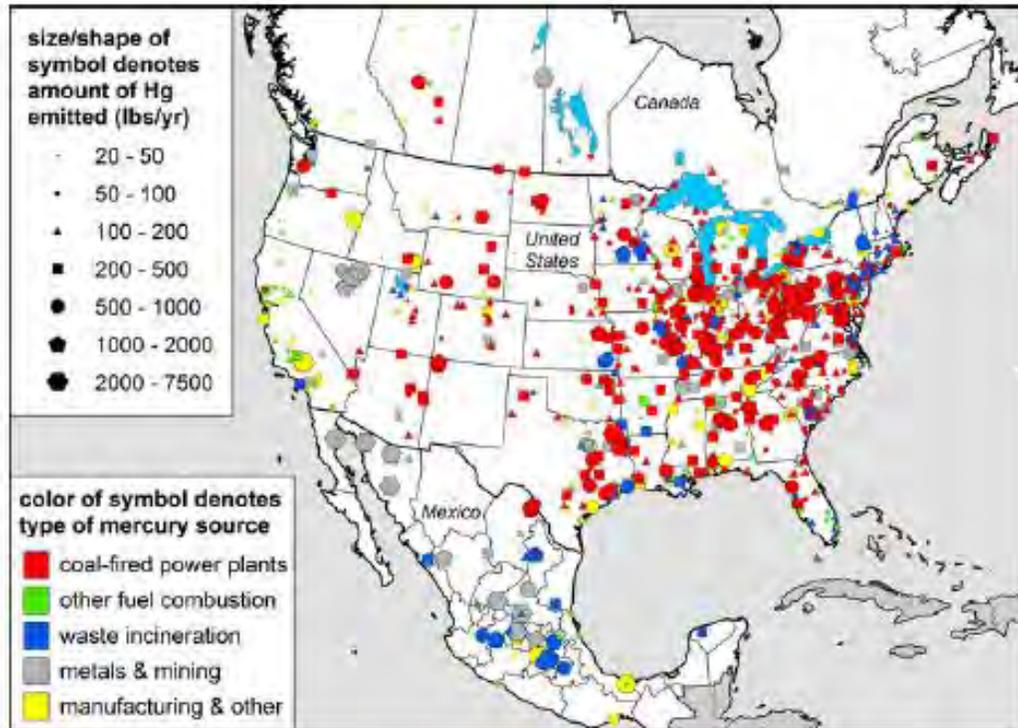


Figure 1. Distribution of total Hg annual air emissions for point sources in the United States, Canada, and Mexico. Total number of point sources in inventory is 15,892, which equals 155 tons/year. Point sources not Mapped are those under 20 lbs/year (n = 3,508 point sources which equals 1.75 tons/year). Point source emissions in Alaska and Hawaii are minor (under 20 lbs/year) and not depicted here. (Source: Schmeltz et al. 2011)

Risch et al. (2012a) reported wet Hg deposition throughout the Great Lakes region during the period 2002-2008, based on data from three Hg and precipitation monitoring networks. Areas in southern Indiana and Illinois, eastern Pennsylvania, and central Michigan had highest estimated wet Hg deposition, ranging from about 10 to 14 $\mu\text{g}/\text{m}^2$. Such high wet deposition levels were caused by high Hg concentration in precipitation ($> 12 \text{ ng}/\text{L}$) and/or high precipitation amount (120-140 cm/yr). There was not a change in deposition observed over the course of the monitoring period; any small decrease that may have occurred in Hg concentration in precipitation was apparently offset by increased precipitation amount (Risch et al. 2012a).

Dry Hg deposition estimates, derived by Zhang et al. (2012) using the Community Multiscale Air Quality (CMAQ) and Global/Regional Atmospheric Heavy Metals (GRAHM) models, suggested that total annual dry Hg deposition was generally $< 5 \mu\text{g}/\text{m}^2$ to the water

surface of the Great Lakes and between 5 and 40 $\mu\text{g}/\text{m}^2$ to land surfaces to the south of the U.S./Canada border.

Mercury is highly volatile and is easily transported in the atmosphere from human sources to remote regions. Most Hg in the atmosphere is gaseous elemental Hg (Hg^0). It can be photochemically oxidized to form reactive Hg ($\text{Hg}(\text{II})$; Schroeder and Munthe 1998). The reactive Hg is water soluble and can become dissolved in precipitation or deposited as dry deposition that is subsequently transported to the forest floor in throughfall or litterfall (St. Louis et al. 2001).

Although some Hg is released to the atmosphere from natural sources, a sizeable fraction of the total atmospheric Hg burden is a result of human activities (Driscoll et al. 2007). Anthropogenic Hg emissions are primarily due to the combustion of fossil fuels and secondarily to waste incineration. Depending on the source, Hg emissions to the atmosphere can be comprised of a combination of three inorganic forms: elemental, reactive gaseous (“oxidized”), and particulate. On average, half of the Hg emitted by coal-fired power plants in the United States is elemental, 40% is reactive gaseous, and 10% is particulate (Driscoll et al. 2007).

Although atmospheric Hg deposition has been well studied, it is difficult to establish total Hg deposition rates. Wet deposition in the United States is generally estimated to range from about 2 to 20 $\mu\text{g}/\text{m}^2\text{-yr}$, and tends to be highest in summer (MDN: <http://nadp.sws.uiuc.edu/MDN/why.aspx>). Wet Hg deposition tends to be especially high in Indiana, Ohio, Illinois, and portions of Pennsylvania, Michigan, and Wisconsin (Risch et al. 2012a, Wiener et al. 2012a). Dry deposition of Hg is poorly known, but is likely in the range of the wet values or slightly higher (Risch et al. 2012b).

Dry deposition of Hg is difficult to measure and the associated uncertainty is high (St. Louis et al. 2001, Lyman et al. 2007, Graydon et al. 2008, Zhang et al. 2009). Inferential methods are commonly used to estimate dry deposition, based on assumed dry deposition velocities and measured ambient concentrations of the principal Hg species found in the atmosphere: gaseous elemental Hg (GEM), gaseous oxidized Hg (GOM), and particulate bound Hg (PBM) at hourly time scales (Engle et al. 2010, Zhang et al. 2012).

The complexities of Hg deposition were illustrated by Graydon et al. (2006) who found that a portion of wet deposited $\text{Hg}(\text{II})$ to forest canopies may be photo-reduced to gaseous $\text{Hg}(0)$ that is then re-emitted to the atmosphere. Gustin et al. (2006) showed that re-emission of

deposited Hg from soil can be affected by environmental conditions such as soil moisture, temperature, light, the presence of atmospheric oxidants, and Hg concentrations in air (U. S. EPA 2009b).

The dry deposition velocity for each atmospheric Hg species can be two to five times higher to forest surfaces as compared with non-forested vegetated and unvegetated surfaces (Zhang et al. 2009, Risch et al. 2012a). Risch et al. (2012a) measured the concentration of Hg in autumn litterfall collected during three years from 23 Hg deposition monitoring stations across 15 eastern states. Forests were primarily hardwood. On average, the ratio of dry (estimated from litterfall) to measured wet Hg deposition was 1.3 to 1.0.

Deposited Hg(II) can revolatilize back to the atmosphere, be methylated in the soil, or be transported to a water body via runoff and leaching. Methylation can also occur within the water body, and either Hg(II) or methyl Hg can be reintroduced from the water back to the atmosphere. Mercury deposited to the O horizon of the soil can be strongly retained, and then re-emitted to the atmosphere with burning (Dicosty et al. 2006).

The importance of atmospheric deposition of Hg as a source of aquatic contamination in the upper Midwest was further established when a study of lake sediments illustrated that wet atmospheric Hg deposition could explain most of the observed sediment Hg loadings (Sorensen et al. 1990, Glass and Sorensen 1999). Analyses of lake sediment cores reported by Swain et al. (1992) suggested that the rate of Hg accumulation in lake sediments in the Upper Midwest had nearly tripled since preindustrial time.

Stratigraphic data from eight lakes in rural Minnesota indicated that atmospheric Hg deposition in this region peaked in the 1960s and 1970s, followed by decreases attributable to decreased emissions from regional sources (Engstrom and Swain 1997). Based on comparison with data from remote lakes in southeastern Alaska, the authors estimated that global Hg emissions in the Northern Hemisphere could account for about $7.4 \mu\text{g}/\text{m}^2$ of annual Hg deposition in rural Minnesota. The remaining deposition (about $5.1 \mu\text{g}/\text{m}^2$, or 40% of the total) was estimated to derive mainly from regional anthropogenic Hg sources in the upper Midwest and eastern United States. It has been estimated that about 60% of the Hg in lake sediments in Voyageurs National Park (VOYA) was derived from atmospheric sources (Sorensen et al. 1990, Wiener et al. 2006). Some of that deposited Hg probably originated in Asia (Pittman et al. 2011).

The Hg that has accumulated historically in lake sediments provides an index of changes over time in Hg inputs from atmospheric deposition and other sources. Drevnick et al. (2012) analyzed 91 sediment cores collected from relatively undisturbed inland lakes in the Great Lakes region. Inferred rates of Hg accumulation in these lakes increased about 7-fold from pre-industrial times to a peak in about the 1980s. Over the past two to three decades, inferred inputs declined about 20%. This recent decrease in sediment Hg is consistent with documented trends in Hg deposition (Drevnick et al. 2012).

Since 1990, anthropogenic emissions of Hg in the United States have decreased dramatically, from 246 tons in 1990 to 101 tons in 2005 (U.S. EPA 2000, 2005). This decrease has largely been attributed to controls on municipal and medical waste incinerators (Butler et al. 2008). The largest single type of emissions source in 2005 was coal-fired power plants (52.3 tons emitted), followed by other boilers, cement manufacturing, and electric arc furnaces (Schmeltz et al. 2011).

1.2.1.1. Bioaccumulation

Bioaccumulation is the progressive accumulation of chemicals with increasing trophic level (LeBlanc 1995). Organic Hg is the most likely metal to bioaccumulate, in part because organisms can efficiently assimilate MeHg and it is slowly eliminated (Reinfelder et al. 1998, Croteau et al. 2005).

Although any form of Hg can be considered toxic, the greatest threat to the environment, wildlife, and people is exposure to MeHg. MeHg is a persistent bioaccumulative toxin that is formed when certain bacteria convert inorganic Hg into MeHg. It can enter through the digestive systems of animals and bind to proteins. It bioaccumulates as each successive predator in the aquatic food chain consumes higher concentrations of MeHg. Low levels of MeHg in surface water (less than about 1 part per trillion) can bioaccumulate over a million-fold and reach toxic levels in fish (Driscoll et al. 2007; MDN: <http://nadp.sws.uiuc.edu/MDN/why.aspx>). Predatory fish, piscivorous birds, and mammals (including humans) that eat contaminated fish can be exposed to toxic levels of MeHg.

In anaerobic environments in wetlands and lake or stream sediments, Hg(II) can be methylated to MeHg (Wiener et al. 2003, Hall et al. 2005). Sulfate (SO_4^{2-})-reducing bacteria are responsible for production of MeHg from inorganic Hg (Compeau and Bartha 1985, Gilmour et

al. 1992). Relatively low concentrations of inorganic Hg in surface water can contribute to substantial Hg bioaccumulation in aquatic ecosystems when and where environmental conditions favor methylation, which is largely carried out by SO_4^{2-} -reducing microbes. Organic MeHg is among the most toxic and widespread environmental contaminants that affect aquatic ecosystems within remote areas (cf., Brumbaugh et al. 2001).

The bioavailability of MeHg in the environment is also of particular importance to humans who can accumulate high levels of MeHg from eating contaminated fish. Avoiding exposure to Hg is particularly important for pregnant women and nursing mothers. Mercury poisoning can cause neurological impairment in children, leading to effects on memory, visual and spatial ability, information processing, and general intelligence (Mahaffey 2005).

Because atmospheric Hg must be methylated in order to bioaccumulate in the food web, significant efforts have been made to understand the bacteria responsible for this chemical transformation. Sulfate-reducing bacteria have been shown to be the main agents of Hg methylation in sediments (Gilmour et al. 1992) and wetlands (St. Louis et al. 1994, Branfireun et al. 1999). In ecosystems that are SO_4^{2-} -poor, the total amount of biologically-available S controls the activity of these bacteria, and thereby the rate of MeHg production. Atmospheric SO_4^{2-} deposition, leading to the stimulation of SO_4^{2-} -reducing bacteria and increased Hg methylation may have caused or contributed to the observed post-industrial amplification of MeHg concentrations in fish (Jeremiason et al. 2006). Studies have demonstrated increased levels of MeHg production with experimental addition of SO_4^{2-} (Gilmour and Henry 1991, Gilmour et al. 1992, Branfireun et al. 1999, Jeremiason et al. 2006). In addition, fish have shown increased MeHg burdens in lakes acidified by atmospheric S deposition as compared with non-acidified lakes receiving similar atmospheric Hg deposition (Gilmour and Henry 1991). Thus, atmospheric S deposition is an important driver of Hg methylation and subsequent biomagnification, and reduced SO_4^{2-} emissions and deposition may be expected to cause a corresponding reduction in Hg methylation and bioavailability (Jeremiason et al. 2006).

In large part because of the common occurrence of wetlands, Hg methylation and bioaccumulation are important concerns within the northern Great Lakes region. This is also true on FCPC lands. Swackhamer and Hornbuckle (2004) assessed air quality and air pollutant impacts in two of the national parks within this region (Isle Royale [ISRO] and VOYA) in a report to the National Park Service (NPS). They concluded that Hg was one of the air pollutants

of most concern in these two parks. Exposure of wildlife to Hg occurs largely through fish consumption, and may be linked to reproductive, neurological, and developmental problems. Inputs of Hg to national park ecosystems are primarily atmospheric, attributable mostly to long range transport.

Mercury binds to organic matter. As a consequence, dissolved organic carbon (DOC) is an important parameter affecting Hg bioavailability and transport through watersheds (Grigal 2002). Methylation is critical to the effects of Hg on aquatic biota. Methylating bacteria require SO_4^{2-} to carry on their metabolic activities (Galloway and Branfireun 2004, ICF International 2006, Jeremiason et al. 2006). Methylation is also correlated with water acid-base chemistry (Wiener et al. 2006, Driscoll et al. 2007). The northern Great Lakes region, and eastward into northern New York, is especially sensitive to Hg bioaccumulation, due in part to relatively high Hg deposition and in particular due to watershed and lake characteristics that exacerbate Hg transport, methylation, and bioaccumulation (Evers et al. 2011a). Such characteristics also apply to Devils Lake within the FCPC study area.

The distribution and magnitude of the impacts of Hg contamination of natural ecosystems in the Great Lakes region is more substantial than previously recognized, with concentrations of Hg exceeding risk thresholds in many species of fish and wildlife across the region (Evers et al. 2011a). Many species of fish in the inland waters of the Great Lakes region, including FCPC's Devils Lake, have average Hg concentration in fillets above risk thresholds for piscivorous wildlife (0.27 ppm).

Much of the research that has been conducted in the United States on Hg methylation, the influence of SO_4^{2-} on methylation rates, and controls on Hg transport within watersheds has been conducted in the northern Great Lakes region (cf., Watras et al. 1998, 2000; Hrabik and Watras 2002; Wiener et al. 2006). Wetlands constitute important features of the landscape in this region. Wetlands act as important sources of MeHg to fresh water ecosystems. This is likely due in large part to two characteristics of wetlands: 1) high availability of DOC, and 2) anaerobic conditions in sediments. Both enhance methylation rates, and DOC also enhances the transport of MeHg to downstream receiving waters. As a consequence of these wetland influences on Hg methylation and transport, the percentage of wetland areas within watersheds is commonly correlated with MeHg flux (Grigal 2002).

Wetlands are important sources of organic matter (OM) and associated Hg and also are active sites for methylation. The MeHg produced in wetlands can bind with OM and be transported to hydrologically connected lakes (St. Louis et al. 1996, Sellers et al. 2001, Grigal 2002).

Hg in Fish

Wiener et al. (2012b) synthesized data on 6,400 yellow perch (*Perca flavescens*) from the Great Lakes and associated inland lakes and reservoirs in and around the northern Great Lakes region. Yellow perch from 6.5% of the waters examined had average whole fish concentrations of Hg high enough to cause adverse effects on fish. Concentrations of Hg were higher in fish at higher trophic levels.

Mercury loads in fish collected in the northern Great Lakes region tended to be higher in inland waters (e.g., median Hg in yellow perch fillets 0.14 ppm [ww¹]) than in the Great Lakes themselves (median Hg in yellow perch fillets 0.09 ppm (ww); Wiener et al. 2012b). Similarly, Hg concentrations in fish from the Great Lakes were about 55% lower for walleye (*Stizostedion vitreum*) and 25% lower for largemouth bass (*Micropterus salmoides*; Monson et al. 2011) as compared with concentrations in fish collected from inland waters within the region.

Gorski et al. (2003) investigated bioaccumulation of Hg in northern pike (*Esox lucius*) in two inland lakes in ISRO: one Hg advisory lake (Sargent Lake) and one non-advisory lake (Lake Richie). Concentrations of total and MeHg were analyzed in water, zooplankton, macroinvertebrates, and fish. Although concentrations of total Hg were significantly higher in pike collected from Sargent Lake, concentrations in open water were slightly higher in Lake Richie. Analyses of stable isotopes of C and N in biota indicated that pike from the two lakes were positioned at approximately the same trophic level (4.2 and 4.3). However, the food web in Sargent Lake was more pelagic-based, whereas in Lake Richie it was more benthic-based. Gorski et al. (2003) speculated that the pelagic food web in Sargent Lake may enhance bioaccumulation. For example, Campbell et al. (2000) showed by ¹³C analysis that organochlorines can bioaccumulate to a greater extent in pelagic food webs. It remains difficult

¹ ww refers to wet weight, reflecting the basis for reporting the concentration of Hg in fish tissue

to identify *a priori* lakes where Hg concentrations in fish will exceed health thresholds for humans or wildlife without direct measurement of Hg content in fish.

The concentration of Hg in fish tissue is often positively correlated with lake and/or watershed area, and negatively correlated with pH, acid neutralizing capacity (ANC), and zooplankton density (Chen et al. 2005, Driscoll et al. 2007). Lake types that are generally associated with the most Hg bioaccumulation are poorly buffered, low in pH and productivity, and have forested watersheds and little human development within the watershed (Chen et al. 2005). Such lakes and watersheds are common throughout the northern Great Lakes region. Lake pH, SO_4^{2-} concentration, and extent of wetland connectivity were the major factors identified by Wiener et al. (2006) controlling Hg bioaccumulation in predatory fish within VOYA. Low-ANC, low-pH lakes are common in VOYA. Such lakes are prone to having high levels of Hg in fish (Swain and Helwig 1989, Wiener and Spry 1996). The review of Evers (2005) classified Hg-sensitive surface waters as those having:

- high SO_4^{2-} concentrations,
- low pH and ANC,
- extensive wetlands,
- large watershed area relative to lake area,
- fluctuating water level, and
- low nutrient concentration

The USGS sampled 20 lakes in VOYA during the period 2000-2002 and analyzed the water and fish for Hg. Near-surface water samples had MeHg concentrations ranging from below the method detection level of 0.04 ng/L to concentrations that were more than an order of magnitude higher. In near-by Ontario, Scheuhammer and Blancher (1994) estimated that 30% of the studied lakes had prey-size fish that had Hg concentrations sufficiently high as to cause reproductive impairment of common loons (*Gavia immer*) that consumed those fish.

Changes in atmospheric Hg deposition can have an important influence on Hg concentrations in fish. Hrabik and Watras (2002) found approximately a 30% reduction in fish Hg concentrations between 1994 and 2000 in a Wisconsin seepage lake in response to decreased atmospheric Hg deposition of about 10% per year between 1995 and 1999 (Watras et al. 2000). It must be noted, however, that seepage lakes, such as the lake studied by Hrabik and Watras (2002), might be expected to respond more markedly and quickly to decreased atmospheric Hg

loading (Harris et al. 2007), compared with drainage lakes that typically receive a large portion of their water and Hg input via the watershed.

VOYA contains many lakes that vary in size, productivity, and fish Hg content (Sorensen et al. 2005b, Wiener et al. 2006). Within VOYA, lakes with extremely low and extremely high fish Hg content are situated within a few kilometers of each other. Fish collected from Ryan Lake and Mukooda Lake in VOYA had some of the highest and lowest Hg concentrations recorded in Minnesota for northern pike (Rolfhus et al. 2011). These two lakes receive nearly identical Hg inputs in atmospheric deposition (Wiener et al. 2006, Rolfhus et al. 2011). Thus, lake- and watershed-specific factors have large influence on Hg bioaccumulation.

Within the Great Lakes region, walleye and largemouth bass exhibited increasing Hg concentrations from south to north and from west to east. Portions of the region having highest concentrations were forested and contained substantial wetland area (Evers et al. 2011a).

Sorensen et al. (2005a) reported results of a monitoring study of Hg concentrations in young-of-the-year (YOY) yellow perch and water level fluctuations in 14 lakes in northeastern Minnesota, including 6 lakes in or adjacent to VOYA. Twelve years of monitoring data were collected for Sand Point Lake, and three years for each of the other lakes. Over the three-year period of record across all lakes, mean fish Hg concentrations varied in each lake by nearly a factor of 2, on average. For the 12-year monitoring period at Sand Point Lake, values ranged from 38 ng/g (ww) in 1998 to 200 ng/g (ww) in 2001. Annual water level fluctuation was correlated with Hg concentrations in YOY perch. Because the Hg concentration measurements were made on YOY fish at the end of each growing season, they represented water conditions and bioaccumulation for each year individually. The researchers concluded that annual water level fluctuations in the reservoir systems in VOYA have significant influence on Hg bioaccumulation in perch.

Bioaccumulation of Hg and the factors that control it have been topics of considerable research in VOYA. Results of a key study were reported by Wiener et al. (2006), who studied Hg concentrations and bioaccumulation in 17 lakes in the park. They assessed the importance of atmospheric and geologic sources of Hg to interior lakes and their watersheds and identified factors associated with lake-to-lake variation in MeHg contamination of lacustrine food webs. Geologic sources were judged to be small. Most Hg found in the upper O and A soil horizons was of atmospheric origin, primarily from anthropogenic sources. The most important factors

influencing MeHg concentrations in lake water and fish were identified as lake pH, SO_4^{2-} concentration in lake water, and total organic C in lake water, which largely reflects wetland influence. The area of connected wetlands adjoining the 17 study lake watersheds represented from 3.5 to 20% of total watershed area (Wiener et al. 2006).

Northern pike from some lakes in VOYA have contained the highest concentrations of Hg reported in Minnesota (Wiener et al. 2006). Concentrations of Hg in northern pike collected in VOYA varied almost 10-fold among the small lakes in the park (Wiener et al. 2006). This reflects the importance of lake and watershed characteristics, in addition to atmospheric deposition, as controlling factors in the cycling and bioaccumulation of Hg. There was no indication in the deposition measurements or distribution of local Hg emissions sources that the atmospheric loading of Hg would explain these spatial differences.

Mean concentrations of total Hg in one-year-old yellow perch ($n = 612$) varied more than 5-fold across the 17 study lakes of Wiener et al. (2006; 182 ng/g [dw]) in Mukooda Lake to 942 ng/g [dw] in Ryan Lake). A model with pH and lake SO_4^{2-} concentration explained 53% of the variation in total Hg concentration in yellow perch. A model with SO_4^{2-} concentration, TOC, and $[\text{TOC}]^2$ explained 60% of the variation in total Hg. The model judged best, based on the Akaike Information Criteria (AIC), for predicting MeHg concentration in lake water included SO_4^{2-} , TOC and $[\text{TOC}]^2$, and explained 79% of the variation in MeHg concentration in water (Wiener et al. 2006).

Ryan and Tooth lakes had both the highest concentrations of Hg in yellow perch and the highest concentrations of lake SO_4^{2-} (up to 74 $\mu\text{eq/L}$) among the studied lakes. Fish Hg concentrations were lowest in the three lakes (Little Trout, Mukooda, O'Leary) that had $\text{pH} > 7.0$ (Wiener et al. 2006), a pattern observed previously (Grieb et al. 1990, Spry and Wiener 1991).

Piscivorous Wildlife

Relatively high levels of Hg have been documented in many species of bird and mammal, in addition to fish, at multiple trophic positions in a variety of habitats across the Great Lakes region (Evers et al. 2011a). Fish constitute an important pathway for transferring MeHg to wild mammals and birds. Monitoring of fish for Hg in this region has often focused on both small prey fish and large, older piscivorous fish for evaluating trophic transfer of Hg. The former are expected to respond quickly to changes in Hg exposure and reflect trophic transport of

contaminants in food webs (Wiener et al. 2007). The latter respond more gradually to Hg bioavailability, and are influenced by such factors as fish age, fish size, nutrient input, and inter-species competition (Mason et al. 2005). Thus, fish monitoring is often targeted at both the mid-trophic level omnivorous prey fish and the higher-trophic level predatory fish in order to capture the full range of fish Hg conditions (Kamman et al. 2005).

In the southern portions of the Great Lakes region, agricultural land uses predominate and nutrient inputs from fertilizer use and livestock waste have contributed to increased algal populations. The abundant algae biodilute MeHg in the food web, resulting in lower concentrations in fish and wildlife at higher trophic levels (Evers et al. 2011a).

Analyses of soil, bedrock, and lake sediments indicated that atmospheric deposition was the primary source of Hg to 17 study lakes in VOYA (Wiener et al. 2006). Bedrock and lower soil horizon material contained much lower Hg concentrations than upper A and O soil horizons. Within the A horizon, Hg concentration was positively correlated with organic content ($r^2 = 0.82$). Lake sediment cores showed patterns of low background concentration in sediments deposited before 1860 (47 to 167 ng/g [dw²]) to maximum values in the late 1900s (102 to 364 ng/g [dw]). Sediment Hg concentrations were lowest in the lake cores that exhibited the highest sedimentation rates, suggesting dilution of sediment Hg concentration by sediment flux. The ratio of recent Hg concentration in sediment divided by preindustrial Hg concentration was relatively consistent among lakes (mean \pm SD: 2.1 \pm 0.1). Measured Hg fluxes represented in lake sediment cores collected from five lakes were not controlled mainly by atmospheric Hg deposition. Rather, they were strongly influenced by sedimentation patterns within the lakes and watershed characteristics that control runoff, including watershed-to-lake area ratio and the percent of the watershed covered by wetlands.

Analyses conducted by Wiener et al. (2006) confirmed that both watershed and in-lake processes affect the concentrations of Hg in lacustrine food webs in the northern Great Lakes region. The concentration of SO₄²⁻, TOC, and pH in water are robust indicators of the sensitivity of lake watersheds to Hg inputs in landscapes affected by atmospheric Hg deposition.

Elevated MeHg accumulation in fish-eating birds in both Wisconsin and in the northeastern United States has been linked to lake acidification (Meyer et al. 1995, Hrabik and Watras 2002, Evers et al. 2007). This form of Hg accumulates in top predators to levels of

² dw signifies that the concentration is reported on a dry weight basis

concern for both human health and the environment (Evers et al. 2007). Several piscivorous bird and mammal species have been suggested as biomonitors of Hg bioaccumulation in the Northeast and the northern Great Lakes region (Wolfe et al. 2007). The common loon and bald eagle (*Haliaeetus leucocephalus*) are especially good indicators of Hg risk to wildlife (Evers 2006). Increased concentrations of Hg have been found to be associated with behavioral, physiological, and reproductive effects on these bird species (Burgess and Meyer 2008, Evers et al. 2008). The common loon has become established as one of the main avian indicators for evaluating the exposure and effects of MeHg in the United States (Evers 2006; Evers et al. 2008, 2009, 2011b).

Acidic deposition contributes to Hg toxicity in fish-eating birds, including the common loon, in part because SO_4^{2-} addition to wetland and lake sediment environments stimulates the production of MeHg, thereby increasing lake water concentrations of MeHg (Jeremiason et al. 2006). Kramar et al. (2005) determined that the extent of wetland located in close proximity (less than 150 m) to loon territory was positively correlated with Hg concentrations in loon blood.

The Lowest Observed Adverse Effect Level (LOAEL) provides a benchmark for quantifying potential injury to wildlife from Hg exposure. For example, the LOAEL for the common loon has been established as 3.0 $\mu\text{g/g}$ of Hg in adult loon blood (Evers et al. 2007, 2008). This level of Hg is associated with reproductive effects such as reduced fledgling success (Burgess and Meyer 2008). However, loon Hg exposure data can include either sex; different ages, locations, and time periods; and can be derived from analysis of blood, tissue or eggs. It has therefore been difficult to standardize the data regarding MeHg availability. Evers et al. (2011b) developed linkages among loon Hg measurements in eggs, blood, and fish prey in the Great Lakes region. Data were normalized into standard loon tissue units. Use of a standard unit of measure that combines multiple tissues of a high profile species and its principal prey items facilitates examination of spatial gradients in pollution effects (Evers et al. 2011b). Based on analysis of over 8,000 male loon units (MLUs), seven biological Hg hotspots were identified in the region. The average MLU concentration across the Great Lakes region was 1.8 $\mu\text{g/g}$; 82% were above 1 $\mu\text{g/g}$ and 9.8% were above the LOAEL of 3 $\mu\text{g/g}$. At a given lake, MLUs tend to have higher Hg concentration than female loon units (FLUs). This is because male loons are typically 21% larger than female loons (Evers et al. 2010) and therefore eat larger fish (Barr 1996). These larger fish tend to have higher Hg concentration (Sandheinrich and Wiener 2011).

Evers et al. (2011b) identified two biological Hg hotspots within the northern Great Lakes region for which rationales were well described for the observed high concentrations of Hg in loons: northeastern Minnesota and northern Wisconsin/upper Michigan. Both were characterized by mixed deciduous and coniferous forest (79% and 61%, respectively) interspersed with scrub-shrub and emergent wetland (12% and 20%, respectively). In the northeastern Minnesota hotspot, high MLUs were further associated with reservoir impoundments characterized by fluctuating water levels and with small natural lakes having low pH and high SO_4^{2-} and DOC concentration. Some of these high MLU lakes were located in VOYA (Wiener et al. 2006) and nearby Superior National Forest. In northern Wisconsin/upper Michigan, high MLUs were associated with low lake ANC (Cope et al. 1990; Meyer et al. 1995, 1998) and interactions with SO_4^{2-} and DOC (Watras and Morrison 2008).

Although the common loon has been well studied as an example of Hg bioaccumulation in wildlife, this species is actually relatively insensitive (LC 50 of MeHg injected into eggs > 1 ppm [ww]) to adverse effects of Hg compared with other avian species. For example, American kestrel (*Falco sparverius*), white ibis (*Eudocimus albus*), snowy egret (*Egretta thula*), osprey (*Pandion haliaetus*), and tricolored heron (*Egretta tricolor*) are all thought to be much more sensitive (LC 50 of MeHg injected into eggs < 0.25 ppm [ww]), based on research by Heinz et al. (2009) and Kenow et al. (2011).

The NPS is monitoring environmental contaminants in bald eagle nestlings at three national park units (APIS, MISS, and SACN) in the upper Great Lakes region (Route et al. 2011). Monitoring includes heavy metals, agricultural pesticides, and industrial chemicals. Contaminant levels in 288 eaglets were associated with proximity to municipal and industrial waste. Mercury was highest where wetlands influenced water chemistry.

Total Hg concentrations in mink in the Great Lakes region have been in many cases high enough to suggest the likelihood of subclinical effects (Basu et al. 2007b). In one survey, Hg concentrations in mink were highest in wetlands along impounded rivers that had fluctuating water levels and that were downstream of large historical point sources (Hamilton et al. 2011). Similarly, bald eagles in this region accumulated sufficient Hg as to suggest subclinical neurological damage. An estimated 14% to 27% of the eagles studied had Hg tissue concentration above the proposed risk threshold for liver toxicity of 16.7 ppm (Zillioux et al. 1993, Rutkiewicz et al. 2011).

The concentration of Hg in fish and birds of the Great Lakes region have declined during recent decades. From 1967 to 2009, Hg concentrations in walleye, largemouth bass, and the eggs of herring gulls throughout the Great Lakes region decreased substantially (Figure 2; Evers et al. 2011a, Monson et al. 2011, Weseloh et al. 2011). This decrease can at least partly be attributed to decreases in atmospheric Hg deposition.

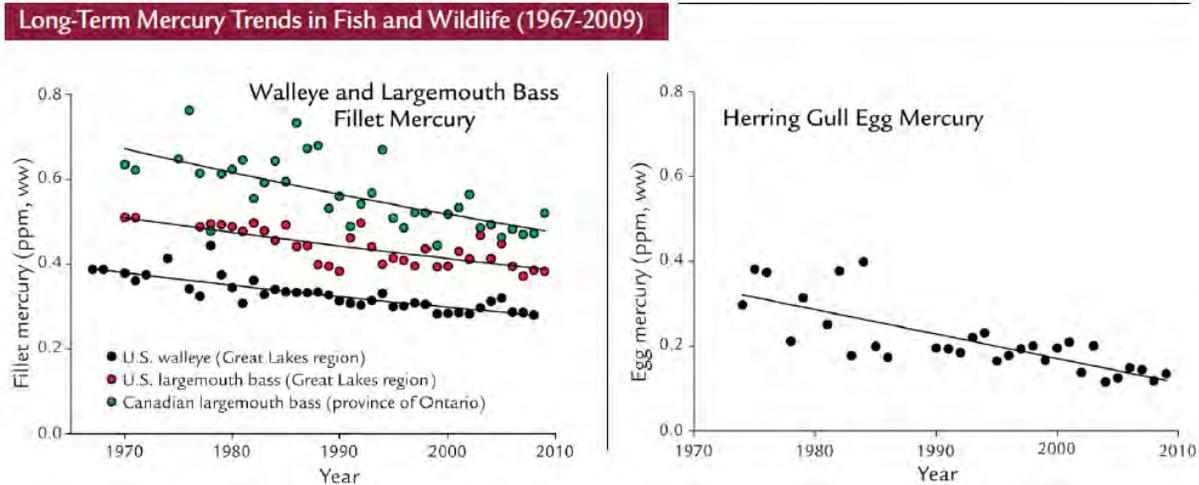


Figure 2. Temporal trends in herring gull egg mercury concentrations (averaged by year across multiple sites in the Great Lakes region; Weseloh et al. 2011) and fish fillet mercury concentrations (walleye and largemouth bass, averaged by year across multiple sites in the Great Lakes and inland water bodies in the U.S. Great Lakes states and the province of Ontario; Monson et al. 2011). These data are characteristic of the regional trend of decreasing mercury concentrations in fish and wildlife in recent decades. Much of this decrease has been attributed to reductions in regional mercury emissions, although there may be other contributing factors as well (Weseloh et al. 2011). (Source: Evers et al. 2011a)

In spite of the multidecadal decline in Hg concentrations in fish and wildlife of the Great Lakes region, there has also been evidence of a more recent increase in Hg concentrations for some species at some locations. This has been shown for walleye from Ontario, walleye and northern pike from Minnesota (Monson 2009, Monson et al. 2011), walleye from Lake Erie (Bhavsar et al. 2010, Zananski et al. 2011), loon blood from northern Wisconsin (Meyer et al. 2011), and bald eagles from VOYA (Pittman et al. 2011). The cause of this recent increase in Hg concentrations in fish and wildlife is not known, but may be related to lower water levels and increased exposed shoreline associated with changing climatic conditions (Meyer et al. 2011), effects of introduced exotic species (Monson et al. 2011), or decreased nutrient loading (Zananski et al. 2011).

Rolfhus et al. (2011) evaluated the results of ten studies in the western Great Lakes region to document the trophic transfer efficiency of MeHg in the pelagic food webs of lakes. The largest increases in MeHg (largest biomagnification) were found at the base of the food web, between water and suspended particles. The observed similarity in the efficiency of trophic transfer suggested that the aqueous supply of MeHg controls much of the bioaccumulation in pelagic food webs in this region.

Bald eagles in the United States have been found to contain substantial amounts of Hg in feathers, eggs, liver, and brain (cf., Wood et al. 1996, Bechard et al. 2007, Scheuhammer et al. 2008). Impacts of Hg toxicity on eagle reproduction and survival can affect individual and population condition. It is therefore important to identify markers of subclinical effects of Hg on the brains of eagles and other piscivorous bird species as early warning signals. Rutkiewicz et al. (2011) evaluated Hg exposure to bald eagles collected in Iowa, Michigan, Minnesota, Ohio, and Wisconsin. Levels of Hg in eagle brains were associated with neurochemical receptors and enzymes. Results suggested that bald eagles in this region are exposed to Hg at levels high enough to cause subclinical neurological damage (Rutkiewicz et al. 2011). The concentrations of total Hg in bald eagle brain and liver tissues were lowest in Ohio and Wisconsin, and highest in Michigan, Iowa, and Minnesota.

Feathers from nestling bald eagles in VOYA have been collected over the past two decades and analyzed for Hg content. This monitoring has focused on nesting sites near three impoundments: Rainy, Kabetogama, and Namakan (included together with Crane and Sand Pointe) lakes. The VOYA environment has high potential for Hg methylation because wetlands are abundant (Grim and Kallemeyn 1995). Since 1999, the water levels of these impoundments have been stabilized to more closely match natural conditions. The lake level stabilization was ordered by the International Joint Commission because research had shown that large changes in lake stage can increase MeHg concentrations in lake water (Rudd 1995, St. Louis et al. 2004, Sorensen et al. 2005a). Apparently in response to this control order, the annual geometric mean of Hg concentrations in bald eagle nestling feathers decreased 74.4% from 1989 to 2010 (Pittman et al. 2011).

1.2.1.2. Effects of Atmospherically Deposited Mercury

Increased body burdens of Hg in fish can lead to decreased reproductive success and behavioral alterations. Yellow perch are often studied as a primary indicator species for Hg contamination in the environment. Ecological effects thresholds for Hg concentration in fish prey have been proposed at levels lower than human health thresholds. Reproduction effects on fish-eating birds have been reported at fish Hg levels as low as 0.16 ppm (Fenn et al. 2011).

As more research has been conducted during the last few decades, high concentrations of Hg have been increasingly documented in more species of wildlife across the Great Lakes region and into the northeastern United States. Recent research has also decreased estimates of effects level, suggesting sublethal effects on wildlife (including effects on reproduction and biochemical processes) at whole-fish concentrations of 0.2 to 0.3 ppm (Beckvar et al. 2005, Dillon et al. 2010, Sandheinrich and Wiener 2011). All 50 states issued Hg advisories for human fish consumption in 2008 (Fenn et al. 2011). That represented an increase in lake area under advisory of 19% and an increase in river length under advisory of 42% compared with 2006. These increases are likely due mainly to increased availability of measured data, rather than increases in Hg deposition or methylation (Fenn et al. 2011). The U.S. EPA (2001) recommended a human health criterion of 0.3 ppm in fish and shellfish tissue to protect the general population. More stringent restrictions may be appropriate for women of child-bearing age and children. Some states, including Maine and Minnesota, use a more restrictive human health standard of 0.2 ppm.

The National Environmental Justice Advisory Council (NEJAC; 2002) issued a report on fish contamination and consumption and environmental justice that addressed the issue of improvement of the quality of aquatic ecosystems in order to protect the health and safety of people who consume fish, aquatic plants, and wildlife. Those affected by contaminated and depleted aquatic ecosystems are disproportionately communities of color, low income, tribes, and other indigenous peoples. The report recommended that EPA work expeditiously to prevent and reduce the generation and release of contaminants that pose the greatest risk of harm to human health and aquatic resources, including Hg. Aquatic ecosystem health is key to addressing issues of environmental justice. Tribes and other indigenous peoples depend to a greater extent than the general population on healthy aquatic ecosystems and the fish and wildlife that these ecosystems support. In addition, harm caused by degradation of aquatic habitats and depletion of fisheries affect not only the present tribal generations, but also future generations and the transfer

of knowledge from one generation to the next. This knowledge transfer includes ecological knowledge, customs and traditions surrounding the harvest, and preparation and consumption of fish and other aquatic resources (NEJAC 2002). Depletion of aquatic environments and contamination of aquatic resources threaten tribal subsistence, economic, traditional, cultural, and religious practices.

Methylmercury causes damage to the vertebrate central nervous system. Embryos are the most sensitive life forms of vertebrates (Scheuhammer 1991, Clarkson 1992). Low-level dietary Hg exposures that cause no measureable effects on adult birds can impair egg fertility, hatchling survival, and reproductive success (Scheuhammer 1991). In reproducing females of fish, birds, and piscivorous mammals, MeHg passes directly to developing egg or embryo (Evers et al. 2003, Hammerschmidt and Sandheinrich 2005, Heinz et al. 2010). These early life stages are more sensitive than adults to the adverse effects of MeHg exposure (Evers et al. 2003, Wiener et al. 2003, Scheuhammer et al. 2007).

In piscivorous birds, including loons and bald eagles, Hg poisoning can lead to brain lesions, reduced reproductive success, increased chick mortality, spinal cord collapse, and neuromuscular problems. The common loon has been widely used as a Hg bioaccumulation indicator for risk to piscivorous birds (Evers et al. 2008, 2011b). It is listed as threatened in Michigan and as a species of special concern in Wisconsin. Because loons feed almost exclusively on fish and crayfish and are relatively long-lived, they can bioaccumulate a substantial amount of Hg (Evers et al. 2011a). Nevertheless, common loons are considered less sensitive to Hg than some other piscivorous birds. They do, however, concentrate Hg in their blood to levels that are high enough to impair reproduction at some locations (Burgess and Meyer 2008, Evers et al. 2011a).

Mink (*Neovison vison*) has been proposed as a sentinel species indicating toxic contaminant exposure (Basu et al. 2007a, Martin et al. 2011). This species constitutes a good candidate for biomonitoring due to its wide distribution and abundance, trophic status, and availability of tissue samples from trappers (Mason and Wren 2001). Symptoms of Hg toxicity in mink include decreased coordination, loss of weight, and splaying of hind legs (Wobeser et al. 1976, Wren et al. 1987).

In its Risk and Exposure Assessment for Review of the Secondary National Ambient Air Quality Standards for Oxides of Nitrogen and Oxides of Sulfur, the U.S. EPA (2009a) concluded the following:

- The production of the large majority of methylmercury is mediated by sulfate-reducing bacteria (SRB), and changes in SO_4^{2-} deposition have resulted in changes in both mercury methylation and mercury concentration in fish (pg. 6-3).
- Experimental evidence has established that only inconsequential amounts of methylmercury can be produced in the absence of SO_4^{2-} (pg. 6-4).
- Elevated mercury levels in fish can only occur where substantial amounts of methylmercury are present (pg. 6-4).
- Evidence is sufficient to infer a causal relationship between sulfur deposition and increased mercury methylation in wetlands and aquatic environments (pg. 6-11).
- Decreases in SO_4^{2-} deposition have already shown promising reductions in methylmercury (pg. 6-13).
- Reducing SO_x emissions could have a beneficial effect on levels of methylmercury in many waters of the United States (pg. 6-13).

1.2.2. Acidification of Terrestrial Ecosystems

Data are generally not available regarding the sensitivity of terrestrial ecosystems to acidification in the northern Great Lakes region or the extent to which terrestrial acidification has occurred. Nevertheless, sugar maple occurs in the region and is known to be sensitive to acidification on base-poor soils. It is not known whether soil acidification has occurred sufficiently within the region to cause damage to this, or other, sensitive plant species.

Possible effects of atmospherically deposited S and toxic metals have been the focus of research interest at ISRO for several decades (cf., Bennett 1995). Subtle patterns of slightly elevated elements in two lichen species collected in ISRO suggested potential impacts from atmospheric deposition. Lichens collected from areas having higher exposures were found to have higher element concentrations of S, Mn, Se, and other heavy metals (Bennett 1995).

1.2.3. Acidification of Aquatic Ecosystems

Atmospheric emissions and deposition of S and N vary dramatically across the Upper Midwest, with relatively low values to the northwest and higher values to the southeast. The Upper Midwest contains numerous lakes created by glaciation. They have been the subject of extensive research on lake acidification (cf., Cook and Jager 1991, NAPAP 1991). The region

has little topographic relief and due to a deep glacial overburden, it also has little exposed bedrock.

Acid-sensitive surface waters in the Upper Midwest are mainly groundwater recharge seepage lakes and small dilute drainage lakes (Eilers et al. 1983). Most drainage lakes and some of the seepage lakes (the flow-through type) in the Upper Midwest region receive substantial inflow from groundwater, which is generally high in base cation concentrations from dissolution of carbonate and silicate minerals. Relatively high concentrations of base cations in many of these lakes make them insensitive to acidification from acidic deposition. The seepage lakes that have low base cation concentrations, and that are therefore acid-sensitive, generally receive most of their water input from precipitation directly on the lake surface (Baker et al. 1991). These groundwater recharge seepage lakes can be identified on the basis of having Si concentration greater than about 1 mg/L (Baker et al. 1991). The acid-sensitivity of streams within the region has not been well studied.

1.2.3.1. Current Status

Based on the EPA's Eastern Lakes Survey (ELS), the Upper Midwest region had a large population of low ANC lakes (Linthurst et al. 1986a, 1986b). Low-ANC lakes in this region are primarily small, shallow, seepage lakes that are hydrologically isolated from the surrounding terrain and that have low concentrations of base cations and Al and moderate SO_4^{2-} concentrations. Some are small drainage lakes. Groundwater flow-through seepage lakes in that region generally have high pH and ANC, due to groundwater inputs of base cations (e.g., Baker et al. 1991). Based on results from the ELS survey, only 6% of the groundwater flow-through seepage lakes had $\text{ANC} \leq 50 \mu\text{eq/L}$. In contrast, the more acid-sensitive groundwater recharge seepage lakes sampled by the ELS constituted 71% of the seepage lakes in the region and were frequently low pH and ANC. Nine percent had $\text{pH} \leq 5.5$. Such lakes tend to be susceptible to acidification from acidic deposition.

Lakewater concentrations of inorganic N reported by the ELS were low throughout the Upper Midwest. In addition, snowmelt would not be expected to provide any significant nitrate (NO_3^-) influx to lakes in the Upper Midwest because most snowmelt in this region infiltrates the soil before reaching the drainage lakes, and because snowmelt input of N into seepage lakes

would be limited mainly to the snow on the lake surface and immediate near-shore environment (Cook and Jager 1991).

1.2.3.2. Past Acidification

Space-for-time substitution analysis was used by Sullivan (2000) to infer the general levels of past change in lake water acid-base chemistry in the Great Lakes region. The concentration of lakewater ($\text{Ca}^{2+} + \text{Mg}^{2+}$) also decreased with increasing acidic deposition across the region, probably due to hydrologic differences and to lower levels of base cation deposition and greater amounts of precipitation in the eastern portion of the region. Atmospheric deposition is an important source of base cations for groundwater recharge seepage lakes because of minimal groundwater inputs. In the eastern portion of the region, such lakes are more sensitive to pH and ANC depression in response to either elevated SO_4^{2-} or DOC. The spatial patterns for low ANC groundwater recharge lakes in the region are consistent with the following hypotheses (Sullivan 1990, 2000):

1. Sensitivity to mineral and organic acidity increased from west to east because of decreasing lakewater base cation concentrations, and this may have been due, in part, to changes in base cation deposition and precipitation volume along this gradient.
2. High concentrations of DOC were responsible for the acidic conditions in some of the lakes, and DOC may have decreased in response to acidic deposition.
3. Many of the groundwater recharge seepage lakes in the eastern portion of the Upper Peninsula of Michigan, and some in Wisconsin, were acidic ($\text{ANC} \leq 0 \mu\text{eq/L}$) because of high SO_4^{2-} relative to base cation concentration, and had probably been acidified by acidic deposition.

1.2.3.3. Recent Trends

Regional trends analyses for long-term monitoring lakes in the Upper Midwest during the period 1990 to 2000 suggested that SO_4^{2-} declined in lakewater by $3.63 \mu\text{eq/L/yr}$, whereas lakewater NO_3^- concentrations were relatively constant (Stoddard et al. 2003). The large decrease in SO_4^{2-} concentration in lake water was mainly balanced by a combination of a large decrease in base cation concentrations ($-1.42 \mu\text{eq/L/yr}$) and an increase in ANC ($+1.07 \mu\text{eq/L/yr}$). All of these trends were significant at $p < 0.01$ (Stoddard et al. 2003). An estimated 80 of 251 lakes in the Upper Midwest region that were acidic in the mid-1980s were no longer acidic in 2000. This change was probably caused by decreased S deposition (Stoddard et al. 2003).

Whole lake experiments and artificial stream channel experiments have shown that acidification can lead to loss of fish species. Work at Little Rock Lake in Wisconsin suggested that rock bass suffered recruitment failure at pH 5.6 or below. Artificial channel studies showed poor survival and reproductive success for fathead minnow at pH 5.9 to 6.0.

One decade after cessation of the experimental acidification of Little Rock Lake in Wisconsin, recovery of the zooplankton community was complete (Frost et al. 2006). Recovery did not follow the same trajectory as the initial acidification, however, indicating a substantial hysteresis in zooplankton community recovery. About 40% of the zooplankton species in the lake exhibited a lag of 1 to 6 years to recover to levels noted in the neutral reference basin.

1.2.4. Nutrient Enrichment

1.2.4.1. Terrestrial Ecosystems

Reduced species richness and biodiversity of terrestrial and wetland ecosystems in response to N addition are of concern, in large part because of the potential interactions among diversity and ecosystem processes and functioning. A species-poor plant community may be more likely to exhibit compromised ecosystem functioning as compared with a species-rich community.

In terrestrial ecosystems, changes in plant species occurrence in response to increased N deposition should be most obvious in ecosystems that are naturally deficient in N. This is because species that are adapted to low N supply will often be more readily outcompeted by species that have higher N demand (Aerts 1990, Tilman and Wedin 1991, Krupa 2003). Plant species having high N-demand are often non-native and invasive (Dukes and Mooney 1999). As a consequence, some native species can be extirpated by N-enhanced eutrophication and potentially replaced by non-native, often invasive species. (Ellenberg 1985; Falkengren-Grerup 1986, 1989; Roelofs 1986; Stevens et al. 2004).

Experimental N addition at > 50 kg N/ha/yr to tallgrass prairie increased productivity and decreased species richness, probably as a consequence of reduced light penetration and increased litter biomass (Baer et al. 2003). Wedin and Tilman (1996) presented results of 12 years of experimental N addition to 162 grassland plots in Minnesota. N loading dramatically changed plant species composition, decreased species diversity, and increased aboveground productivity in the experimental plots. Species richness declined by more than 50% across the N-deposition

gradient, with the greatest losses at levels of N input of 10 to 50 kg N/ha/yr. This loss of richness was accompanied by large changes in plant species composition, with C₄ grasses declining and the weedy Eurasian C₃ quackgrass (*Agropyron repens*) becoming dominant at high N addition rates (Wedin and Tilman 1996). The authors concluded that N loading is a major threat to grassland ecosystems and causes loss of diversity, increased abundance of non-native species, and disruption of ecosystem functioning. A major uncertainty, however, is the rate of N loading at which such changes may be manifested. Total N loading to ecosystems in the northern Great Lakes region is lower than the loading rates typically used in experimental treatments, such as those of Wedin and Tilman (1996).

Effects of N fertilization on vegetation, in turn, can influence faunal populations. For example, fertilization in a Minnesota old-field at relatively low rates (≤ 20 kg N/ha/yr) for 14 years resulted in decreased species diversity of insects in response to decreased diversity of plant food sources. Plant species richness decreased, and quackgrass and Kentucky bluegrass (*Poa pratensis*) became dominant in response to the N addition (Haddad et al. 2000). Changes in the abundance of insect functional groups were also observed. Herbivores (especially the dominant species) increased in numbers; parasitoid insect species decreased. Over the long term, changes in plant species composition would be expected to either increase or decrease insect herbivore activity, depending on whether there is a shift toward or away from herbivore-preferred plant species (Throop et al. 2004, Clark 2011).

Forests can also be responsive to nutrient addition. Northern hardwood forests that predominate in the northern portions of Minnesota, Wisconsin, and Michigan are expected to show growth enhancement by some, but not all, tree species in response to N addition. Eastern hardwood forests from Wisconsin to Maine and south to Virginia were shown to have experienced increased growth with increasing N deposition across a depositional gradient from about 3 to 11 kg N/ha/yr (Thomas et al. 2010). Nevertheless, responses were species-specific. Growth increases were most pronounced for red maple (*Acer rubrum*), sugar maple (*Acer saccharum*), white ash (*Fraxinus americana*), yellow poplar (*Liriodendron tulipifera*), black cherry (*Prunus serotina*), balsam fir (*Abies balsamea*), pignut hickory (*Carya glabra*), eastern white pine (*Pinus strobus*), quaking aspen (*Populus tremuloides*), northern red oak (*Quercus rubra*), and scarlet oak (*Quercus coccinea*). Other tree species showed negative response to N addition. Growth decreased by a statistically significant amount with increasing deposition for

red pine (*Pinus resinosa*), red spruce (*Picea rubens*), and northern white cedar (*Thuja occidentalis*). Mortality increased with increasing deposition for yellow birch (*Betula alleghaniensis*), eastern white pine, basswood (*Tilia* spp.), quaking aspen, bigtooth aspen (*Populus grandidentata*), scarlet oak, chestnut oak (*Quercus prinus*), and northern red oak. At higher rates of N supply on nutrient-poor sites, tree growth might be expected to decline (Pardo et al. 2011a).

Eastern hardwood forests can also respond to increased N deposition with a decrease in herbaceous plant biodiversity (Gilliam et al. 2006). The response of the herbaceous layer of the forest can be pronounced, with initial increases in herbaceous plant cover, followed by decreases in species richness and species evenness, two components of biodiversity. The response time is typically shorter if ambient N deposition is relatively high (Gilliam et al. 2006, 2011; Fraterrigo et al. 2009; Royo et al. 2010). Because the plants in the herbaceous layer of eastern hardwood forests tend to have foliage with relatively high nutrient content, herbaceous plants in these forests influence N cycling to a level that is disproportionate to their biomass (Muller 2003, Moore et al. 2007, Welch et al. 2007, Gilliam et al. 2011).

Lichens have also been heavily impacted throughout the eastern hardwood forests. High levels of air pollution, including both S and N, has played an important role in causing such impacts (McCune 1988, Wetmore 1988).

1.2.4.2. Wetland Ecosystems

Wetlands are common throughout the northern Great Lakes region. They contribute to high production of organic matter which is reflected in high DOC concentrations (higher than about 5 mg/L) in many lakes. Wetlands considered sensitive to nutrient enrichment from N deposition typically contain plants species that have evolved under N-limited conditions (U.S. EPA 2008). It is believed that the competitive balance among plant species in some sensitive wetland ecosystems can be altered by N addition, with resulting displacement of some species by others that can utilize the excess N more efficiently (U.S. EPA 1993, 2008). This effect has been well documented in heathlands in The Netherlands, where heather (*Calluna vulgaris*) has been replaced by grass species (Heil and Bruggink 1987, Tomassen et al. 2003). It is important to note, however, that N deposition in The Netherlands has been much higher (about 20 to 60 kg

N/ha/yr) than levels commonly encountered in wetland areas in the Great Lakes region (usually less than 10 to 15 kg N/ha/y).

There is a great deal of diversity in terms of types of wetlands and how they respond to N addition. Ombotrophic bogs are typically acidic and are dominated by mosses. They are especially common in northern boreal forested regions and develop where precipitation is higher than evapotranspiration in areas that exhibit an impediment to downward drainage in the soil (Mitsch and Gosselink 2000). Freshwater marshes develop where water inputs in groundwater plus surface water inflow approximate precipitation input (Koerselman 1989). Vegetation primarily consists of tall graminoid plants. Freshwater swamps have hydrological conditions that are generally similar to marshes, but vegetation is forested (Greaver et al. 2011).

The response of freshwater wetlands to N addition varies with hydrological conditions. Bogs that receive much of their water input from precipitation are especially sensitive to the effects of N input (Morris 1991). Ombotrophic bogs receive most of their nutrient supply from precipitation; as a consequence plants are adapted to low inputs of N and other nutrients (Shaver and Melillo 1984, Bridgham et al. 1995). Peat-forming bog ecosystems are among the most sensitive wetland ecosystems to the effects of N deposition. In the conterminous United States, peat-forming bogs are most common in areas that were glaciated, especially in portions of the Northeast and Upper Midwest (U.S. EPA 1993). Bogs can host several federally listed rare and endangered plant species. These include multiple species of quillworts (*Isoetes* spp.), sphagnum mosses (*Sphagnum* spp.), and the green pitcher plant (*Sarracenia oreophila*; Greaver et al. 2011).

Increased N availability in nutrient-poor wetland environments may lead to a decrease in species diversity and increased risk of extinction for some of the more sensitive and rare species (Moore et al. 1989). As a consequence, peatlands and bogs are among the most vulnerable transitional ecosystems to adverse nutrient-enrichment effects of N deposition (Krupa 2003). The sensitivity of peatland *Sphagnum* species to elevated atmospheric N deposition is well documented in Europe (Berendse et al. 2001, Tomassen et al. 2004). *Sphagnum squarrosum* and *S. fallax* have been observed to be negatively affected by experimentally elevated atmospheric N and S inputs in Europe (Kooijman and Bakker 1994). Roundleaf sundew (*Drosera rotundifolia*) is also susceptible to elevated atmospheric N deposition (Redbo-Torstensson 1994).

In other studies, wetland species such as heather can successfully compete with grasses even at relatively high rates of N deposition, as long as the vegetative canopies are closed (Aerts et al. 1990). However, N deposition causes nutrient imbalances, including increase in the shoot-to-root ratio, and therefore increases in the sensitivity of shrubs to drought stress, frost stress, and attack by insect pests (Heil and Diemont 1983). These can result in gaps in the canopy of the shrub layer, which can then be readily invaded by grasses that are more efficient in using the additional N and therefore gain a competitive advantage (Krupa 2003).

Data are not available with which to evaluate the extent to which wetlands in the northern Great Lakes region have been affected by nutrient enrichment from N deposition. Wetlands are widely distributed, including within some areas that receive moderate levels of N deposition. This N deposition may or may not be sufficiently high to cause species shifts in wetland plants. If such effects do occur, they are most likely in wetlands such as bogs that normally receive most of their nutrients from atmospheric inputs. These wetlands have been shown in Europe to experience changes in plant species composition in response to high levels of atmospheric N deposition. Such ecosystems tend to occur in portions of the northern Great Lakes region that receive much lower levels of N deposition than do many of the affected wetlands in Europe (Bobbink et al. 2010). It is not clear to what extent such effects occur under ambient N deposition levels in Wisconsin. The risk of species composition change is important, in part because wetland ecosystems often contain relatively large numbers of rare plant species (U.S. EPA 2009a).

Greaver et al. (2011) estimated that the critical load (CL) of atmospheric N deposition to protect peatlands in the United States from increased productivity was in the range of 2.7 to 13 kg N/ha/yr. This estimate was based on consideration of the results of the studies of Aldous et al. (2002), Moore et al. (2004), Rochefort et al. (1990), and Vitt et al. (2003).

1.2.4.3. Surface Water Ecosystems

Atmospheric inputs of N can change the productivity of N-limited fresh water ecosystems, altering the make-up of phytoplankton communities. In lakes that are limited in their biological activity by N, increased atmospheric N deposition can cause phytoplankton species that use N most efficiently to dominate the lake. Species that use N less efficiently can be extirpated. However, many lakes and streams are limited by phosphorus (P), light, or some other

critical constituent. In such waters that are not N-limited, addition of atmospheric N is unlikely to cause nutrient enrichment (eutrophication) impacts. The degree to which lakes and streams in the Upper Midwest might be N-limited is not well known. However, Baron et al. (2011) estimated, based on measured N:P ratios in the U.S. EPA’s Eastern Lakes Survey conducted in 1984, that 37% of Upper Midwestern lakes were N-limited and an additional 31% were either co-limited by N and P or limited by something else other than P (Table 1).

Table 1. Proportion of lakes in three nutrient limitation classes (N limited, P limited, and N and P co-limited) based on ratios of dissolved inorganic nitrogen (DIN; NO₃⁻ + NH₄⁺) to total phosphorus (TP). Data are from the Eastern Lakes Survey (Linthurst et al. 1986b), conducted in the fall of 1984. (Source: Baron et al. 2011)

Region	Number of Lakes ^a	Number of N Limited Lakes (%) ^b	Number of P Limited Lakes (%) ^c	Number of Lakes with Co-limited N:P (%) ^d
Upper Midwest	8,755	3,142 (37%)	2,787 (32%)	2,646 (31%)

^a The Eastern Lakes Survey was a stratified random sample of lakes; estimates of the number of lakes in the region are based on the target population size for the survey.

^b Lakes with DIN:TP ratios (by weight) less than four were characterized as N limited, based on the work of Morris and Lewis (1988).

^c Lakes with DIN:TP ratios (by weight) greater than 12 were characterized as P limited, based on the work of Morris and Lewis (1988).

^d Lakes with DIN:TP ratios (by weight) between 4 and 12 could not be assigned to nutrient limitation class, and are characterized as either co-limited or limited by something other than N or P.

2. DESIGNATION OF AQRVS FOR FCPC CLASS I LANDS

Atmospheric pollutant thresholds are addressed in the Federal Land Managers Air Quality Related Values Work Group (FLAG) Phase I report, initially drafted in 2000 (U.S. Forest Service et al. 2000) and more recently revised (U.S. Forest Service et al. 2010, 2011). The purpose of the FLAG Phase I report is to assist permit applicants, state authorities, and land managers to develop more consistent and objective approaches to the evaluation of AQRVs. This is needed for evaluation of air pollution effects, caused by new or modified emissions sources, on AQRVs in Class I areas under the New Source Review (NSR) program, especially in the review of Prevention of Significant Deterioration (PSD) air quality permit applications. The FLAG Phase I report and its modifications provide background information on effects levels and screening tools to determine if effects are expected to be negligible based on the planned

quantity of emissions, the distance between the proposed source and the Class I area, and dose/response relationships. Among the primary issues of concern to Federal Land Managers (FLMs) and tribal land managers are the effects of pollutant deposition (including S, N, and Hg) on soils and surface waters.

As part of the review of a PSD application for a new or modified emissions source, an AQRV impact analysis is conducted to ensure that the identified AQRVs in the Class I area will not be adversely impacted by the proposed increased emissions. The AQRV impact analysis is conducted partly by the permit applicant and partly by the land manager. The applicant calculates the changes in pollutant concentrations, deposition amounts, or visibility extinction that will result from the proposed project emissions. The land manager assesses the extent to which these changes in pollution level will affect sensitive resources and have an adverse impact on an AQRV. As part of this process, the land manager will, to the extent allowable by available data:

- identify AQRVs
- identify data and studies that constitute the basis for identification of AQRVs
- identify sensitive receptors and pollutants of concern that can potentially impact those receptors
- identify pollutant concentrations or loads above which adverse impacts are expected
- recommend methods for predicting pollutant concentrations, loads, and related impacts
- suggest screening level values to assess whether the proposed emissions would have a *de minimis* impact on AQRVs

The FCPC has designated Water Quality and Aquatic Systems as AQRVs to be protected against air pollution degradation. These AQRVs can be adversely impacted by atmospheric deposition of Hg, S, and N. Important effects include Hg methylation (which is related to both Hg and S deposition), acidification (which is related to both S and N deposition), and fertilization of water and soils (which is related to N deposition). In order for the FCPC to determine whether any future emissions source would adversely impact the designated AQRVs, The FCPC must establish threshold effects levels, which represent the amount of additional (above ambient) deposition from a proposed new, or modified existing, source below which an adverse effect will not occur. Such thresholds are proposed herein. Once atmospheric deposition

of S, N, or Hg is high enough that a deposition concern threshold has been reached, any further increase in emissions that would contribute additional deposition and cause increased adverse effects is considered unlikely to be acceptable. This is especially true for a process such as Hg methylation because MeHg bioaccumulates in fish, sometimes to toxic levels.

2.1. Importance of Water Quality and Aquatic System AQRVs to the FCPC

Potawatomi tribal members have a close link to the natural environment, including its water quality and wildlife. Elements of the natural environment are important to religious rituals, ceremonies, and medicines. Contamination of tribal waters, including entry of Hg into the food web, jeopardizes the tribe's cultural and religious beliefs and practices and threatens public health. The FCPC designated water quality and aquatic systems as AQRVs because:

- tribal members rely on fish as part of their traditional subsistence way of life,
- pure water is important to the tribe's cultural and religious practices, and
- recreational fishing is important to the tourism economy in and around the FCPC reservation, upon which the FCPC depends.

The traditional subsistence way of life followed by many FCPC tribal members relies heavily on fish, clean water, and other natural resources. A Wisconsin state-wide fish advisory was issued by WDNR due to Hg contamination of fish

(<http://dnr.wi.gov/fish/consumption/FishAdvisoryweb2011.pdf>). Thus, Hg contamination threatens tribal public health. Because many tribal members consume large quantities of fish compared to the general population, they are at heightened risk of Hg contamination.

The FCPC is the largest employer in Forest County and is dependent on tourism focused on outdoor recreation. Contamination of fish and other aquatic resources with Hg discourages recreational fishing and threatens the economic well-being of the FCPC and its tribal members.

2.2. FCPC Study Area

The 393 km² FCPC study area designated for this AQRV evaluation is comprised of all land area from which surface water drains onto or within FCPC reservation lands (Figure 3). Primary drainages within the study area include Otter Creek, Rat River, and North Branch Oconto River. Wetlands, which are strongly associated with Hg methylation, are abundant in the study area and throughout the state of Wisconsin. More than 5 million acres (15.5% of total land area) of wetland exist within Wisconsin. The study area is located in the southern portion of

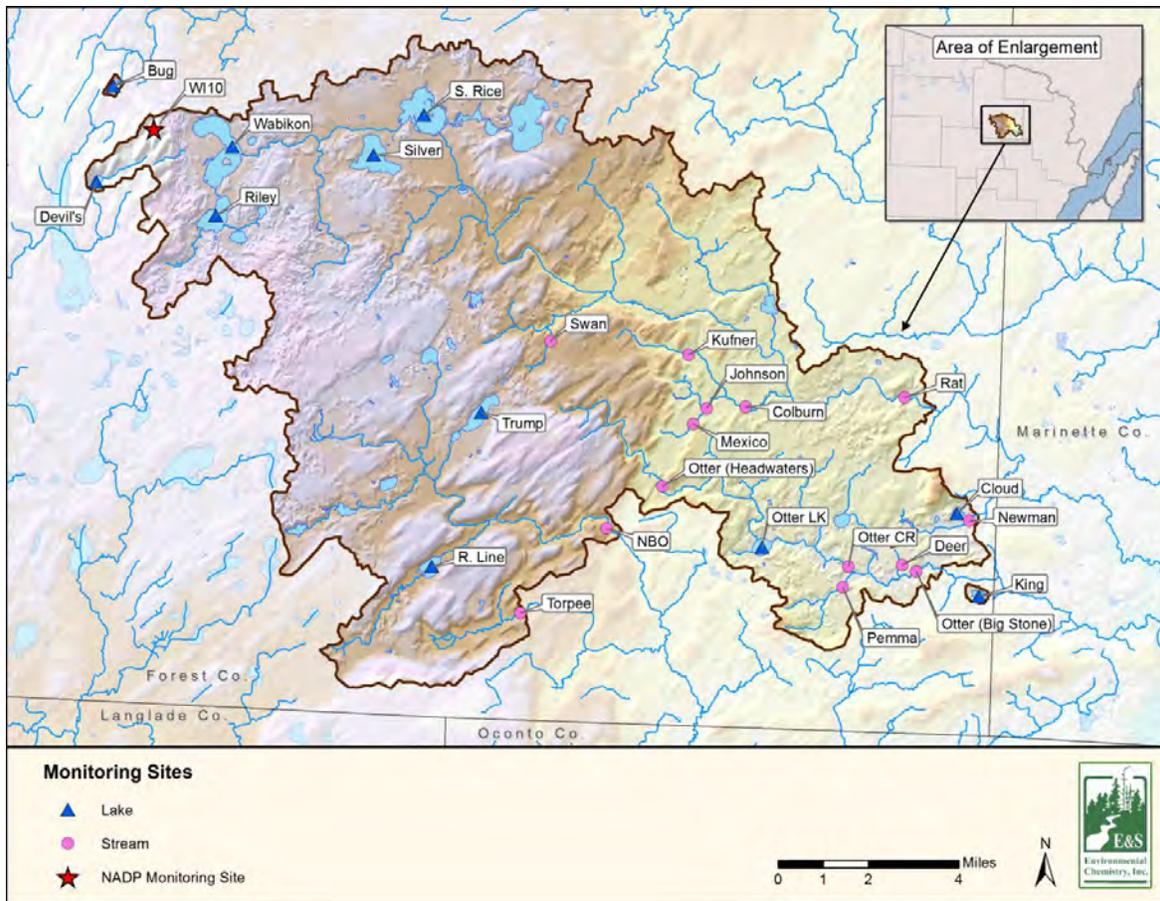


Figure 3. Study area for evaluation of FCPC air quality related values, as determined by mapping all drainage water contributing areas that drain into surface waters that occur on FCPC lands. Sampling locations for lakes, streams, and wet deposition monitoring are also shown.

Forest County, which is ranked 9th among all 72 Wisconsin counties in wetland percentage, with almost one-quarter of the land area mapped as wetland (data from WDNR; <http://dnr.wi.gov/wetlands/acreage.html>). The 2006 National Land Cover Database (NLCD; U.S. EPA 2011) estimates that 27.2% of the study area is classified as wetland (Figure 4). The abundance of wetlands within the study area increases sensitivity to Hg methylation, especially in surface waters such as Devils Lake that are high in DOC and low in ANC.

Estimates of annual wet and dry deposition of N, S, and Hg to the FCPC study area are summarized in Table 2. Measured wet N, S and Hg deposition data were derived from the NADP NTN/MDN deposition monitoring site located in the northwestern portion of the study area near Devils Lake (WI10; elevation 570 m). Dry N and S deposition were estimated using

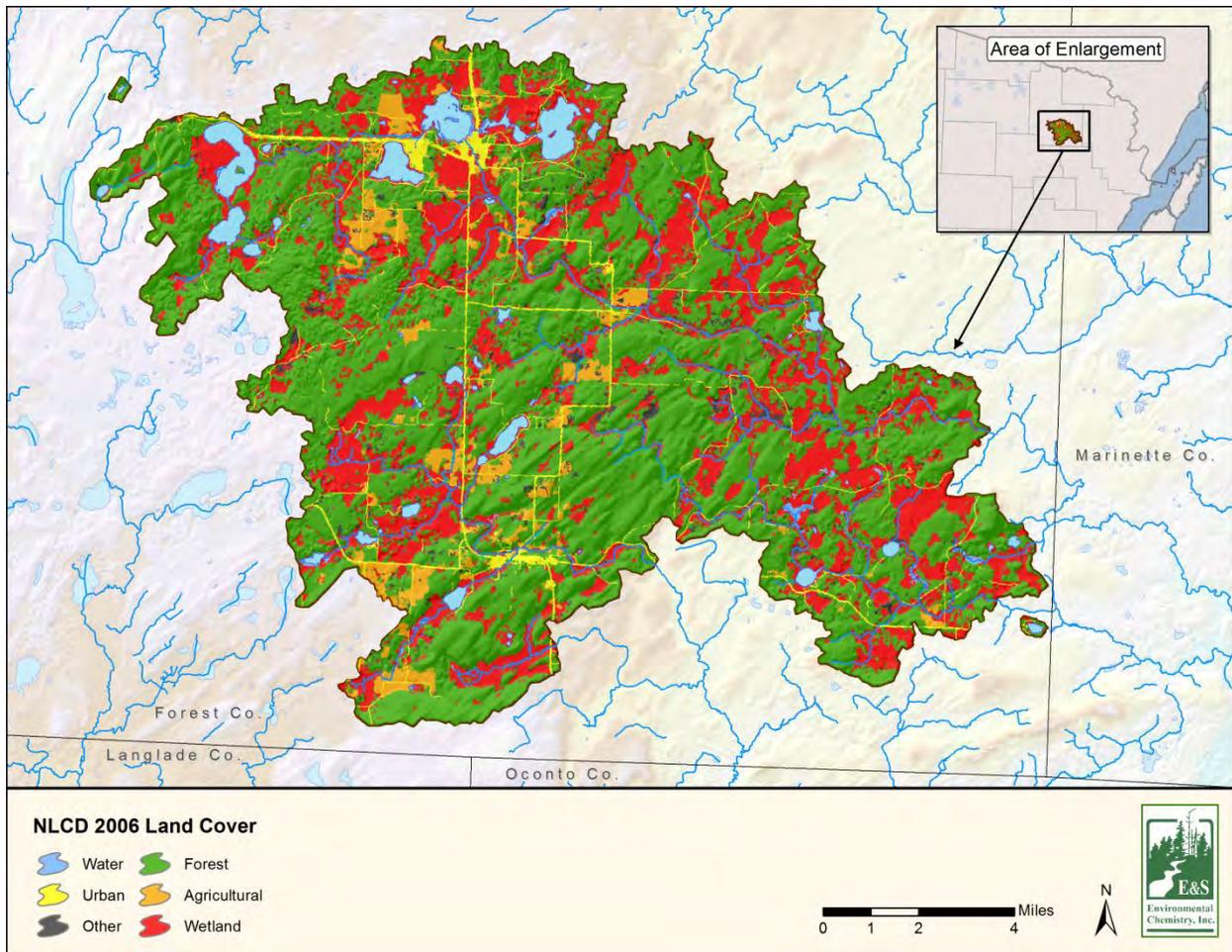


Figure 4. Land cover in the FCPC study area, based on National Land Cover Data. Note: “Other” includes: Barren land, Shrub/Scrub and Grassland/Herbaceous.

Table 2. Measured values of annual wet deposition at the NADP NTN/MDN monitoring site near Devils Lake and estimated dry deposition based on dry-to-wet deposition ratios at the CASTNET site PRK134 for the period 2006-2010.

Annual Wet Deposition			Dry-to-Wet Deposition Ratios			Total Deposition		
S (kg/ha/yr)	N (kg/ha/yr)	Hg ($\mu\text{g}/\text{m}^2/\text{yr}$)	S	N	Hg	S (kg/ha/yr)	N (kg/ha/yr)	Hg ($\mu\text{g}/\text{m}^2/\text{yr}$)
1.83	3.41	6.4	0.24	0.17	1.3 ¹	2.27	3.99	8.32

¹ Dry deposition of Hg is poorly known, but is estimated to range up to about 1.3 times wet Hg deposition (Risch et al. 2012a)

measurements of wet deposition combined with dry-to-wet ratios calculated at the nearest CASTNET dry deposition monitoring site (PRK134; elevation 472 m) located approximately 144 km ESE of the study area. Dry Hg deposition is less well known, but is probably equal to, or slightly higher than, wet Hg deposition.

Annual average wet S and N deposition amounts at NADP/NTN monitoring station WI10 (Devils Lake) for the five-year period 2006 through 2010 were 1.83 kg S/ha/yr and 3.41 kg N/ha/yr (Table 2). The dry-to-wet ratios for S and N averaged over the most recent available five-year period at the nearest CASTNET dry deposition monitoring station were 0.24 and 0.17, respectively. Thus, estimated total (wet plus dry) deposition on FCPC lands is currently approximately 2.27 kg S/ha/yr and 3.99 kg N/ha/yr. Measured annual wet Hg deposition, averaged across the same five-year period was 6.4 $\mu\text{g}/\text{m}^2/\text{yr}$. Dry Hg deposition is not commonly measured, but has been estimated to approximately equal or exceed wet deposition by about 30% (Risch et al. 2012a). Assuming that dry Hg deposition equals 1.3 times the wet deposition level, the estimated current annual average total Hg deposition on FCPC lands is 8.32 $\mu\text{g}/\text{m}^2/\text{yr}$.

2.3. Protection Against Methylation of Hg

Because of its cultural and recreational importance, Devils Lake has been sampled and tested for Hg contamination since 1995 by the FCPC (Horsley & Witten 2003). Devils Lake is heavily used by the Potawatomi Tribe as a fishing and recreation site. In 2006, Devils Lake plus three other lakes on the reservation (Bug, Cloud, and King lakes) were sampled to determine the concentrations of Hg in fish tissue. High concentrations of Hg were found in fish, especially in largemouth bass.

Published total Hg concentration in water for Devils Lake and its inlet stream suggest epilimnetic concentrations in the lake near 3 ng/L; both hypolimnetic concentrations in the lake and concentrations in the stream have been measured to be three- to four-fold higher (Eckley et al. 2005, Watras et al. 2005). Such conditions of relatively low pH and ANC and relatively high DOC and Hg concentrations are considered highly susceptible to Hg methylation and bioaccumulation.

Studies of FCPC surface waters and surface waters elsewhere in Wisconsin and the broader northern Great Lakes region have documented that many water bodies are contaminated with high levels of Hg. This is especially true for Devils Lake, which is of great cultural,

recreational, and spiritual importance to the Tribe. As a consequence of high levels of Hg found in fish tissue tested from Devils Lake, the Tribe issued a fish advisory to limit human consumption of fish from Devils Lake. Because Hg is already at unsafe levels in Devils Lake and other water bodies in and near the FCPC reservation, the collective CLs of S and Hg deposition for protecting these impacted watersheds against Hg methylation have already been exceeded. Additional sources of either S or Hg are considered unacceptable because of the likelihood of enhanced Hg methylation that this would cause. Additional inputs of either S or Hg to FCPC lakes would contribute to further diminishment of the water bodies' significance to the Tribe, impair the natural ecosystem, and impair the quality of Tribal members' and visitor's experiences. Based on guidance from the Federal Land Managers provided in the FLAG Phase I Report (U.S. Forest Service et al. 2010), such effects are unacceptable.

Devils Lake and other water bodies on and near the FCPC Class I area contain levels of Hg that are considered to be unsafe for protecting human health and fish-eating wildlife. Any increase in the atmospheric deposition of either S or Hg would likely increase MeHg production within the Class I area, thereby causing additional Hg contamination of fish and wildlife. For that reason, the threshold effects level for additional atmospheric deposition of both S and Hg should be set at zero in order to prevent further Hg methylation and bioaccumulation.

Lakes and streams sampled for water quality by the FCPC are mapped in Figure 3. These water quality data confirm that sampled lakes and streams on and around the FCPC reservation are wetland-influenced, with DOC concentrations typically higher than 10 mg/L. ANC values are relatively high in most waters, although Devils and Bug Lakes are both low in alkalinity, with values below 60 $\mu\text{eq/L}$ (Table 3). Box plots showing the distributions of DOC, SO_4^{2-} concentration, and alkalinity are shown for sampled lakes in Figure 5 and streams in Figure 6. Lake SO_4^{2-} concentrations are typically between about 20 and 50 $\mu\text{eq/L}$; SO_4^{2-} concentrations in streams are somewhat higher.

The FCPC and WDNR have analyzed nearly 300 fish samples for Hg concentrations. Analyzed fish have primarily been largemouth bass (n=94), bluegill (n=57), and yellow perch (n=30). Distributions of fish samples across lakes are given in Table 4. Among the lakes showing highest Hg concentrations in fish fillets are Otter, Trump, Silver, and Devils Lakes, all having median fish Hg concentration higher than 0.25 ppm and 75th percentile Hg concentrations near or above 0.4 ppm (Figure 7). Fish Hg concentrations averaged across all lakes were highest for walleye,

Table 3. List of lake and stream water quality sampling sites. Site ID and initial year of available water quality data are specified.

Site Type	Site Name	Site ID	Initial Year
Lake	Bug Lake	Bug	2004
	Cloud Lake	Cloud	2003
	Devil's Lake	Devils	2004
	King Lake	King	2004
Stream	Colburn Creek	Colburn	2002
	Unnamed	Deer	2008
	Johnson Creek	Johnson	2011
	Kufner Creek	Kufner	2008
	Mexico Creek	Mexico	2008
	Newman Creek	Newman	2002
	North Branch Oconto River	NBO	2002
	Otter Creek	Otter	2006
	Otter Creek (Big Stone)	Otter BS	2006
	Otter Creek (Headwaters)	Otter HW	2002
	Pemma Creek	Pemma	2006
	Rat River	Rat	2002
	Swan Creek	Swan	2006
	Torpee Creek	Torpee	2002

Table 4. Number of fish samples and date ranges for which skin-on fillet total Hg samples are available, by species. Data were compiled from the Forest County Potawatomi Community (Steinbach 2011) and the Wisconsin Department of Natural Resources databases.

Species ID	Species	Number of Samples	Years for Which Data Are Available
B. Trout	Brook trout	4	2006
Bluegill	Bluegill	57	2006-2010
Crappie	Black crappie	15	2006-2010
L. Bass	Largemouth bass	94	1998-2010
N. Pike	Northern pike	24	1998-2010
Pumpkin	Pumpkinseed	14	2006-2010
R. Bass	Rock bass	15	2006-2010
R. Trout	Rainbow trout	2	2010
S. Bass	Smallmouth bass	2	1998
W. Sucker	White sucker	9	2006-2010
Walleye	Walleye	11	1998-2010
Y. Bull	Yellow bullhead	1	2006
Y. Perch	Yellow perch	30	2006-2010

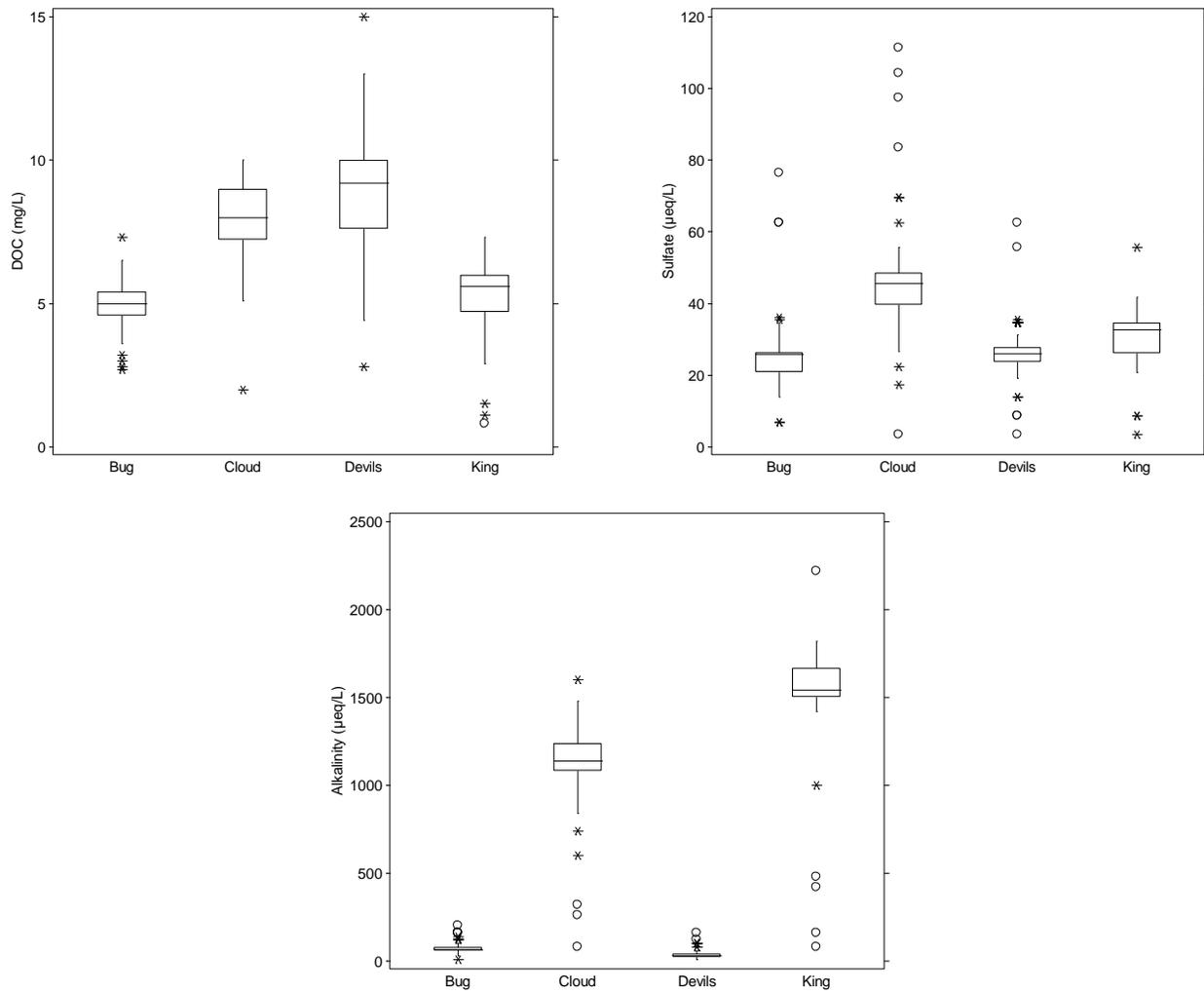


Figure 5. Box plots showing the distribution of lake water DOC, sulfate and alkalinity at FCPC lake monitoring sites. Each site is represented by at least 76 water samples.

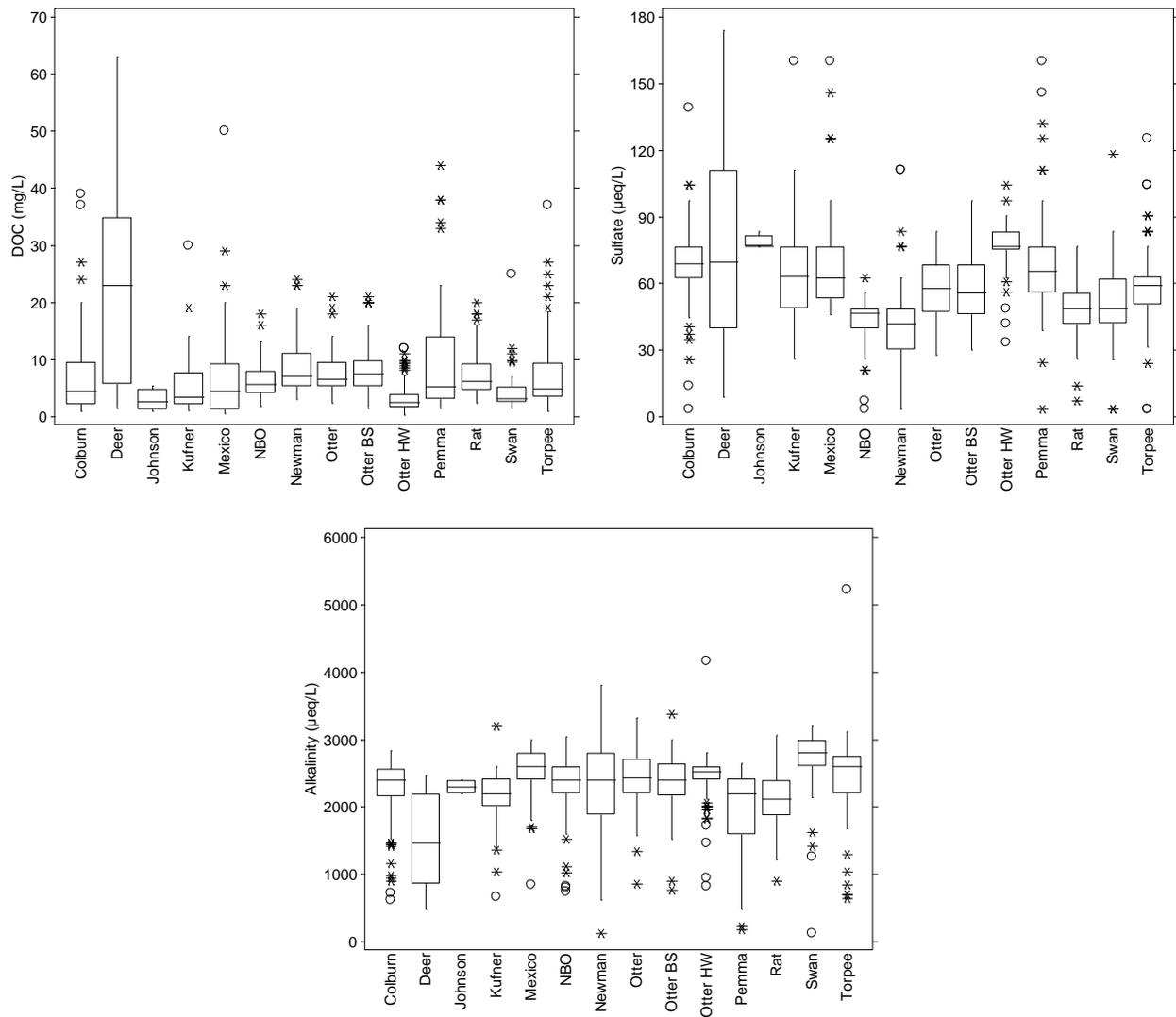


Figure 6. Box plots showing the distribution of stream water DOC, sulfate and alkalinity at FCPC stream monitoring sites. Each site is represented by at least 40 samples, except Johnson Creek which only had 4 recent samples.

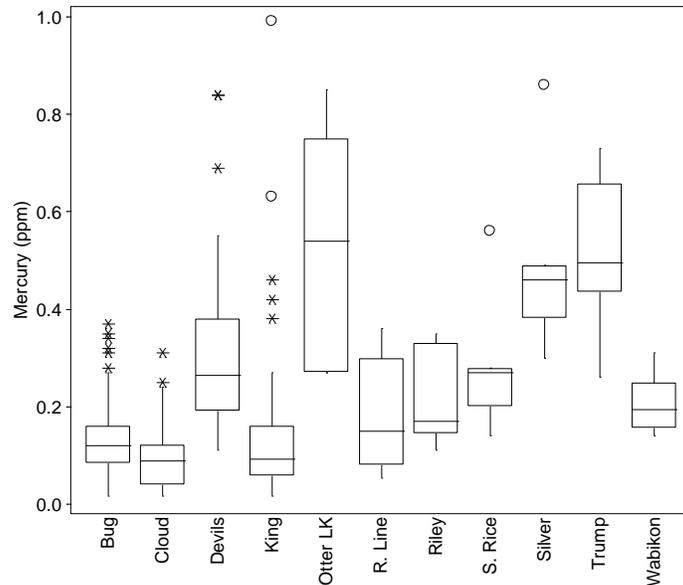


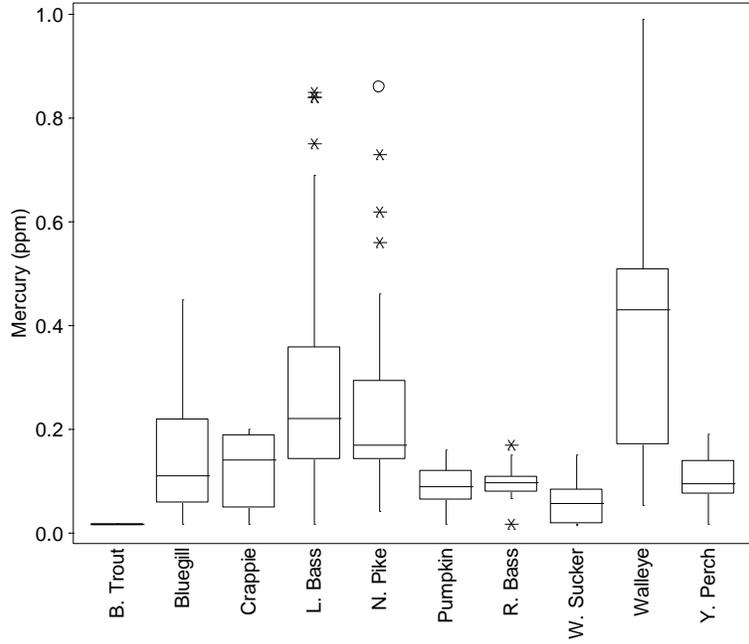
Figure 7. Total Hg concentrations in skin-on fillets of all sampled fish species, by lake.

largemouth bass, and northern pike (Figure 8a). Largemouth bass in Devils Lake showed a median Hg concentration higher than 0.4 ppm (Figure 8b).

Reported fish Hg content risk thresholds for protecting piscivorous wildlife against sublethal effects range between about 0.2 and 0.3 ppm (ww, reported in fish fillets [Evers et al. 2011a]) or whole fish (Beckvar et al. 2005, Dillon et al. 2010, Sandheinrich and Wiener 2011). Measured values in fish collected from several of the sampled lakes on FCPC lands and elsewhere within the study area (Figure 7) were substantially higher than these threshold effects levels. Furthermore, concentrations of Hg measured in fish from Devils Lake were substantially higher than the median Hg concentrations reported for yellow perch collected throughout the northern Great Lakes region by Wiener et al. (2012b; 0.14 ppm, ww).

The U.S. Forest Service designated concern thresholds in the Boundary Waters Canoe Area Wilderness and the Rainbow Lakes Wilderness for Hg in fish. The concern threshold for Hg in fish tissue is 0.05 ppm (ww, skin-on fillet; http://www.fs.fed.us/air/technical/class_1/wilds.php?recordID=6) in both wilderness areas. This level is exceeded in all monitored FCPC lakes (Figure 7).

A)



B)

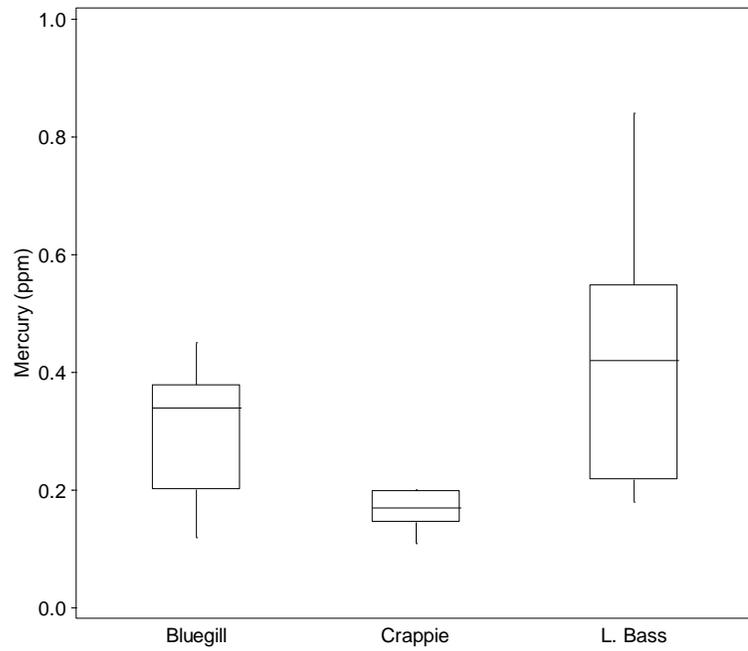


Figure 8. Total Hg concentrations in skin-on fillets of fish species sampled in A) all lakes located within the study area and B) Devils Lake only.

2.4. Protection Against Acidification

The U.S. Forest Service designated concern thresholds for surface water ANC in two federal Class I areas in reasonable proximity to the FCPC lands that are the subject of the analysis reported here: the Boundary Waters Canoe Area Wilderness, MN and Rainbow Lakes Wilderness, WI. The threshold for maintaining healthy biological functioning in lakes and streams requires chronic ANC ≥ 25 $\mu\text{eq/L}$. Thus, FLMs have adopted a screening level for lake ANC equal to 25 $\mu\text{eq/L}$ to evaluate the likelihood of adverse impacts of acidification on lake ecosystems. This screening level is applied here to measured data for lakes on FCPC lands.

Water chemistry has been measured by FCPC in 4 lakes and 12 streams over the last decade (Table 5). Published surface water ANC has been below 25 $\mu\text{eq/L}$ in Devils Lake (Eckley et al. 2005) and its inlet stream (Watras et al. 2005). ANC in Devils Lake measured by FCPC

Table 5. Lake and stream DOC, alkalinity and calculated ANC values for two sampling occasions represented by data necessary (major base cations and strong mineral acid anions) for determining the calculated ANC (see text for the definition of calculated ANC).

Site Type	SITE	July 2006 Sampling			October 2010 Sampling		
		DOC (mg/L)	Alkalinity ($\mu\text{eq/L}$)	Calculated ANC ($\mu\text{eq/L}$)	DOC (mg/L)	Alkalinity ($\mu\text{eq/L}$)	Calculated ANC ($\mu\text{eq/L}$) ¹
Lake	Bug	4.6	60	109	5.1	58	114
	Cloud	10.0	1,080	1,139	8.9	1,220	1,239
	Devils	10.0	20	129	10.0	30	118
	King	6.5	1,460	1,423	6.2	1,520	1,447
Stream	Colburn	39.0	900	1,343	37.0	940	2,345
	Deer		No Data		48.0	760	1,243
	Kufner		No Data		30.0	660	2,275
	Mexico		No Data		50.0	840	2,615
	NBO	13.0	1,680	1,832	18.0	1,600	2,365
	Newman	19.0	1,980	2,117	23.0	1,720	2,422
	Otter	14.0	1,920	2,043	19.0	1,680	2,302
	Otter BS	20.0	1,520	1,570	21.0	1,540	2,161
	Otter HW	11.0	1,960	2,135	12.0	2,000	2,551
	Pemma	38.0	580	913	44.0	600	1,825
	Rat	15.0	1,480	1,562	20.0	1,540	1,997
	Swan	6.2	120	2,549	25.0	1,620	2,880
Torpee	27.0	1,720	1,969	19.0	2,000	2,689	

¹ It was necessary to combine data from two sampling occasions (Oct 13th and 25th, 26th, or 27th) to calculate ANC values representative of 2010.

was 20 µeq/L in 2006 and 30 µeq/L in 2010 (Table 5). Published water chemistry data for the Devils Lake watershed are consistent with an interpretation of generally acid-sensitive, high-DOC water having pH below 6.0 and DOC near 10 mg/L (Eckley et al. 2005, Watras et al. 2005).

2.5. Protection Against Fertilization of Water and Soils

It is not known to what extent surface waters on FCPC lands are N-limited and therefore susceptible to nutrient enrichment impacts due to atmospheric N deposition. It is also not known whether atmospheric inputs of N have been sufficient to cause changes in plant species distributions or abundance due to nutrient enrichment. Current atmospheric N deposition to FCPC lands in the range of 4 kg N/ha/yr (Table 2) is at the low end of the range suggested by Greaver et al. (2011) to protect peatlands in the United States from effects associated with changes in productivity. Peatlands are among the plant community types expected to be most sensitive to nutrient enrichment. Thus, fertilization effects on FCPC lands are possible for the most sensitive plant communities under current N loading.

3. CRITICAL LOAD

The CL is the threshold of air pollution deposition that causes harm to sensitive resources. If harm has been caused, then by definition the CL has been exceeded in the short term. In other words, it is exceeded based on ambient conditions, rather than based on long-term steady-state conditions. The CL is usually calculated as a steady-state value, representing the deposition level that can be sustained over the long term without experiencing ecosystem harm.

3.1. Critical Load to Protect Against Hg Methylation

The CL approach has not yet been applied in the scientific literature to the topic of atmospheric Hg deposition or for protecting ecosystems against Hg methylation caused by atmospheric S deposition. Nevertheless, the basic principles remain the same. The observation that fish Hg levels in FCPC lakes exceed thresholds for protection of wildlife and human health indicate that short-term CLs of Hg and S deposition for protection against Hg methylation have been exceeded. In other words, methylation of Hg is already so high that the health of wildlife

and humans who consume fish is threatened. Any further increases in the air pollutants that contribute to methylation is considered to be unacceptable.

3.2. Critical Load to Protect Waters Against Acidification

The CL to protect against acidification can be calculated using an acid-base chemistry model such as the Model of Acidification of Groundwater in Catchments (MAGIC) or the Photosynthesis and EvapoTranspiration-Biogeochemical (PnET-BGC) model, or using a simple mass balance model such as the Steady State Water Chemistry (SSWC) model. The CL can also be derived from empirical dose-response data. The CL can be calculated to protect particular sensitive resources from harm, and these can include, for example, fish presence, fish species richness, concentration of Hg in fish or wildlife, lake zooplankton, stream benthic invertebrates, terrestrial plant biodiversity, etc. Nevertheless, the fact that lakewater ANC in Devils Lake has been reported below 25 $\mu\text{eq/L}$ indicates that the CL to protect against water acidification has been exceeded in the Devils Lake watershed, at least in the short term. Long-term steady-state CL values might be higher or lower. Dynamic models can also calculate the target load (TL) which is specific to a particular point of time, such as for example to protect fish species richness in the year 2100. This may require emissions reductions to achieve that goal by 2100 for already-damaged resources or to prevent damage from occurring by 2100 for resources as yet undamaged.

The CL of S, N, or acidity to protect against soil or water acidification can be calculated by simple steady-state models, empirical models, or process-based dynamic models. Such calculations are available at locations throughout much of the United States. Minnesota established a state-wide deposition standard of 3.7 kg S/ha/yr in 1982, which is higher than the estimated total S deposition on FCPC lands (Table 2). The intention with the Minnesota standard was to protect lakes having acid neutralizing capacity (ANC) < 50 $\mu\text{eq/L}$ from acidification damage. In an area where there are multiple proposed new sources or where resources are already impaired, the DAT approach may not be completely protective, and the CL approach may be more appropriate. In considering the potential for adverse impact, the land manager considers the nature of the sensitive AQRVs, CLs, ambient deposition levels, and/or expected impacts due to the proposed source. Where adequate information is not available, the FLAG

report (U.S. Forest Service et al. 2010) recommended use of data from nearby, or ecologically similar, areas.

Critical and target loads have not yet been calculated in a formal manner for protecting lake or stream resources on FCPC lands against damages caused by acidic deposition inputs. Nevertheless, the fact that water ANC below 25 $\mu\text{eq/L}$ has been measured in Devils Lake indicates that the short-term CL to protect against water acidification has already been exceeded in the Devils Lake watershed.

3.3. Critical Load to Protect Terrestrial Resources

Pardo et al. (2011b) reviewed available empirical CL data for ecoregions across the United States to protect against adverse impacts to terrestrial resources attributable to atmospheric N deposition. They concluded that the CL for both hardwood and conifer forest soils in the Northern Forests ecoregion (which includes FCPC lands) could be set at 8 kg N/ha/yr to protect against NO_3^- leaching, which can contribute to surface water and soil acidification (Aber et al. 2003). The empirical CL to protect against negative tree growth response in this ecoregion was estimated by Pardo et al. (2011b) to be 3 kg N/ha/yr, based on research findings of Thomas et al. (2010) documenting effects on tree growth. On this basis, the CL to protect against NO_3^- leaching in the FCPC study area has not been exceeded, but the CL to protect against decreased tree growth has been exceeded by ambient N deposition (Table 2).

4. ESTABLISHMENT OF BACKGROUND LEVELS AND THRESHOLD EFFECTS LEVELS

Proposed emissions sources are assessed by FLMs for evaluation of acidification and nutrient enrichment effects at locations near (within 50 km) and distant (> 50 km) from Class I areas. The U.S. EPA introduced a screening criterion as part of the Regional Haze Rule (RHR), based on the annual emissions strength of a proposed new emissions source and its distance from a Class I area. The FLAG Phase I report (U.S. Forest Service et al. 2010) adopted a similar approach. A new source location > 50 km from a Class I area is deemed to have negligible impact on the acid or nutrient status of Class I AQRVs if its total sulfur dioxide, nitrogen oxides (NO_x), particulate matter, and sulfuric acid annual emissions (in tons/yr based on 24 hr maximum allowable emissions), divided by the distance (in km) from the Class I area (Q/D), is 10 or less. Such a screening criterion may not be appropriate for evaluation of potential effects of

Hg and/or S deposition on Hg methylation, especially because the tipping point for protecting against Hg methylation is already exceeded in many FCPC lakes and because of the toxic and bioaccumulative nature of MeHg.

Recent Federal Land Managers' Air Quality Related Values Work Group (FLAG) documents (U.S. Forest Service et al. 2010, 2011) provided the following guidance regarding determination of potential acidification or nutrient enrichment effects from a new or modified emissions source in a Class I area where the CL has not been exceeded:

1. If the source is further than 50 km from the Class I area, effects are considered negligible if annual emissions in tons/yr (based on 24-hr maximum allowable emissions, Q) divided by distance (D) in km from the Class I area is 10 or less ($Q/D < 10$).
2. If $Q/D > 10$, but the source's predicted contribution to deposition in kg/ha/yr in the Class I area is less than the Deposition Analysis Threshold (DAT; defined below), the source is presumed to be negligible.
3. If a refined analysis suggests that the source's inputs would not cause or contribute to harm in the Class I area (i.e., CL not exceeded) or that the source would provide some ancillary environmental benefits, as described in the FLAG Phase I report (U.S. Forest Service et al. 2010), the impacts are presumed to not be adverse.
4. If there are mitigation strategies that could alleviate the potential adverse impact, a more detailed process could be applied (U.S. Forest Service et al. 2010).

If a source's contribution to increased deposition in a Class I area is predicted to exceed the DAT, a refined analysis may be warranted. This refined analysis would address questions such as the following:

1. Are AQRVs in the Class I area sensitive to deposition?
2. Are AQRVs currently impacted by deposition?
3. Have critical or target loads been developed?
4. If so, does the modeled deposition exceed the critical or target load?

Each of these questions is examined below with respect to potential impacts of S, N, and Hg deposition on the Potawatomi tribal lands that are the subject of this report.

Are AQRVs sensitive to deposition?

As outlined in the previous sections of this report, AQRVs associated with aquatic resources on FCPC lands are sensitive to atmospheric deposition of S, Hg, and potentially N.

This sensitivity is supported by extensive scientific research conducted throughout the northern Great Lakes region, including within the FCPC area that is the focus of this report.

Are AQRVs currently impacted by deposition?

AQRVs on FCPC lands are clearly impacted by recent and current levels of S and Hg deposition. This is reflected in the high concentrations of Hg in fish collected from FCPC lakes and to a lesser extent in the low ANC of Devils Lake. Lake ANC in this lake is influenced by both moderate lakewater SO_4^{2-} concentration contributed by atmospheric S deposition and naturally occurring high DOC.

Have critical or target loads been developed?

Critical and target loads have not been formally developed for atmospheric deposition of S, Hg, or N to watersheds on FCPC lands. Nevertheless, the demonstrated impacts reflected in high Hg concentrations in fish and low lake ANC indicate that CLs have been exceeded, at least in the short term.

Does current deposition exceed the critical or target load?

Current deposition exceeds the critical or target load for protecting aquatic ecosystems on FCPC lands against Hg methylation and at least on occasion also the critical or target load for protecting aquatic ecosystems against acidification. It is not known if critical or target loads are exceeded for protection of aquatic, terrestrial, or transitional ecosystems against nutrient enrichment. Such impacts are possible for the most sensitive ecosystems.

4.1. Sulfur and Mercury Deposition as Related to Hg Methylation

Various indicators can be used to evaluate whether atmospheric deposition is high enough to trigger adverse eutrophication, acidification or nutrient enrichment effects. The NPS and U.S. Fish and Wildlife Service (USFWS) developed DATs for S and N in response to requests by permitting authorities and permit applicants to develop consistent, predictable permit review processes. A DAT is defined as *the additional amount of S or N deposition within a Class I area below which estimated impacts from a proposed new or modified source are considered insignificant*. The DAT represents total (wet plus dry) deposition. It is compared with the

estimated amount of additional deposition resulting from the proposed source, as modeled using an atmospheric transport model.

A DAT for Hg has not been developed on the basis of assumed background Hg deposition by the Federal Land Managers, probably in part because background Hg deposition is less well known. Nevertheless, an alternative DAT for wet deposition can also be expressed on the basis of the minimum concentration that can be detected in precipitation in the analytical laboratory. Using this approach, an increase in wet deposition is considered significant if the increase in precipitation concentration, and associated wet deposition, is measurable in the deposition monitoring network. This approach for specifying a DAT based on what is measurable in the analytical laboratory has been followed by USFS (2000) for the James River Face Class I Wilderness. It constitutes an alternate approach for calculating the DAT for Hg and S for protecting against Hg methylation in view of the lack of data regarding background Hg deposition.

Using this approach, the minimum detectable level of an increase in S or Hg wet deposition to FCPC lands to protect against Hg methylation can be calculated using the following equation:

$$\text{MDI} = C \times P \times \text{UCF} \quad (1)$$

where MDI is the estimated minimum detectable increase in wet deposition; C is the laboratory detection limit for S or Hg measured in water from wet deposition collectors; P is annual precipitation; and UCF is the unit conversion factor value needed to express MDI in the selected units (e.g., kg/ha/yr for S or $\mu\text{g}/\text{m}^2/\text{yr}$ for Hg).

The MDIs with respect to the concentrations of S and Hg wet deposition were calculated for the NADP/NTN monitoring station WI10 for the five-year period 2006 through 2010 based on annual average precipitation and the current analytical method detection limits (MDL) in the NADP/NTN and Mercury Deposition Network (MDN) monitoring programs. Annual average precipitation for 2006-2010 at WI10 was 75.3 cm (range 68.9 to 92.5 cm/yr). The current MDL for SO_4^{2-} wet deposition is 0.01 mg/L SO_4^{2-} (0.0033 mg/L as S). Any measured concentration below the MDL is removed from the published NADP database and replaced with the MDL value (Robert Larson, Data and Information Manager, NADP, pers. comm., March 26, 2012). The current MDL for Hg wet deposition ranges from 0.06 to 0.101 ng/L for the three Hg

analyzers used in the MDN laboratory (Robert Larson, NADP, pers. comm., March 15, 2012). For the purposes of the calculations reported here, the average of that deposition range is used (0.0805 ng/L).

The minimum detectable increase (MDI) in wet Hg deposition for site WI10 is calculated as:

$$\text{MDI}_{\text{Hg}} = \frac{0.0805 \text{ ng}}{\text{L}} \times \frac{75.3 \text{ cm}}{\text{yr}} \times \frac{1}{100} = 0.0606 \text{ } \mu\text{g}/\text{m}^2/\text{yr} \quad (2)$$

Similarly, the MDI in wet S deposition can be calculated as:

$$\text{MDI}_{\text{S}} = \frac{0.0033 \text{ mg}}{\text{L}} \times \frac{75.3 \text{ cm}}{\text{yr}} \times \frac{1}{10} = 0.025 \text{ kg S}/\text{ha}/\text{yr} \quad (3)$$

where the values of $\frac{1}{100}$ (for Hg) and $\frac{1}{10}$ (for S) are unit conversion factors.

For the reasons outlined above, the recommended wet deposition threshold effects levels for protecting water quality and aquatic system AQRVs against harm caused by Hg methylation on FCPC tribal lands can be calculated based on the levels of Hg and S that are detectable in the wet Hg deposition monitoring program:

0.025 kg S/ha/yr for S (1.36% of ambient annual average wet S deposition), and
 0.0606 $\mu\text{g Hg}/\text{m}^2/\text{yr}$ for Hg (0.94% of ambient annual average wet Hg deposition)

Applying the dry:wet conversion factors, total (wet plus dry) minimum detectable deposition increase thresholds are:

$$\text{MDI}_{\text{TS}} = 0.031 \text{ kg S}/\text{ha}/\text{yr} \text{ for S (1.37\% of ambient annual average total S deposition)} \quad (4)$$

$$\text{MDI}_{\text{THg}} = 0.0788 \text{ } \mu\text{g Hg}/\text{m}^2/\text{yr} \text{ for Hg (0.95\% of ambient annual average total Hg deposition)} \quad (5)$$

The DATs for protecting against Hg methylation, based on laboratory detection limits, can then be calculated from the minimum detectable increase in deposition of total Hg (MDI_{THg}) and total S (MDI_{TS}). These equations incorporate the Cumulative Factor ($C = 0.04$) developed by the FLMs for protecting against acidification and nutrient enrichment (see Equation 8 in Section 4.2; Table 6):

$$\begin{aligned} DAT_{Me-Hg} &= 0.0788 \mu\text{g Hg/m}^2/\text{yr} \times C \\ &= 0.0788 \mu\text{g Hg/m}^2/\text{yr} \times 0.04 = 0.0035 \mu\text{g Hg/m}^2/\text{yr} \end{aligned} \quad (6)$$

$$\begin{aligned} DAT_{Me-S} &= 0.031 \text{ kg S/ha/yr} \times C \\ &= 0.031 \text{ kg S/ha/yr} \times 0.04 = 0.0012 \text{ kg S/ha/yr} \end{aligned} \quad (7)$$

Table 6. Deposition analysis thresholds (DATs) for FCPC lands.

Atmospherically Deposited Pollutant	Purpose of DAT	DAT Value
N or S	Protect against water acidification or eutrophication	0.01 kg/ha/yr
S	Protect against Hg methylation	0.0012 kg/ha/yr
Hg	Protect against Hg methylation	0.0035 $\mu\text{g/m}^2/\text{yr}$

The rationale for inclusion of the Cumulative Factor (C) into calculation of the DATs to protect against Hg methylation (DAT_{Me-Hg} and DAT_{Me-S}) is the same as the rationale given by the FLMs for including C in calculation of the DATs to protect against acidification and nutrient enrichment described in Section 4.2 below. Over time, one might expect 25 or fewer new emissions sources to increase deposition to a Class I area. The 0.04 represents a safety factor to protect Class I areas from cumulative impacts.

For the purpose of protecting FCPC resources against Hg methylation in response to increased Hg or S deposition, this report recommends following the approach described above using the DATs given in Table 6. Model estimates of either increased S or Hg deposition above these thresholds due to a new or modified existing emissions source would be considered to be above the threshold effects level for protecting against Hg methylation on FCPC lands and would therefore be unacceptable.

Class I land managers must take into consideration site-specific information regarding each Class I area, not just in relation to the DAT or the MDI, but also in relation to other factors, including whether adverse impacts have been documented, or are suspected (National Park Service and U.S. Fish and Wildlife Service 2002). Because MeHg concentrations are unacceptably high in fish from FCPC lakes, posing risk to humans who consume fish and to piscivorous wildlife, it is clear that the CLs of S and Hg deposition to protect against Hg methylation are exceeded in the short term, based on ambient conditions. For that reason, and because of the toxic and bioaccumulative nature of MeHg, DATs based on the MDI approach are more appropriate than DATs based on assumed background deposition. This is because the aquatic ecosystems are already damaged and no additional S or Hg input to these damaged aquatic ecosystems is acceptable. Further increase in MeHg will increase toxicity to people and wildlife who consume fish.

Biomagnification of Hg is also an important DAT issue. An increase in S and/or Hg deposition that is too small to detect in a precipitation sample may nevertheless biomagnify to a detectable level in fish. I do not have a basis for quantifying that effect. Nevertheless, this complication provides additional support for the need to be conservative in setting DATs for protecting against Hg methylation.

4.2. Sulfur and Nitrogen Deposition as Related to Acidification and Fertilization of Water and Soils

The DAT calculation reported by FLAG (U.S. Forest Service et al. 2011) relies on an estimate of Natural Background Deposition, B, developed from the peer-reviewed scientific literature and expert judgment. This natural background estimate is modified by including a Variability factor, V and a Cumulative Factor, C:

$$\text{DAT} = B \times V \times C \quad (8)$$

Values of B equal to 0.5 kg/ha/yr for the East and 0.25 kg/ha/yr for the West were selected by NPS and USFWS (National Park Service and U.S. Fish and Wildlife Service 2002) for both S and N as representative of the low end of the regional range of reasonable estimated natural background deposition values.

The variability factor, $V = 0.5$, was selected with the reasoning that a total increase in deposition from all sources by an amount greater than 50% of natural background deposition would trigger management concerns. Thus, an additional 50% of natural background deposition could be added to existing natural plus anthropogenic deposition and still be considered insignificant. The FLMs made the determination that it was appropriate to consider that all combined anthropogenic sources could contribute up to 50% of natural background deposition without triggering concerns regarding resource impacts because the estimates of natural background S and N deposition were very conservative and the range of natural variability in deposition could be plus or minus 50% (U.S. Forest Service et al. 2011). For specifying a DAT for Hg using this or a related approach, the estimate of background Hg deposition or alternative parameter should be similarly conservative.

The Cumulative Factor (C) adjusts for the concern that cumulative effects from multiple relatively small sources could produce impacts even if a single source, on its own, would not. The federal land managers (FLMs) applied the 4% value used by EPA in determining Class I increment levels of significant impact. Thus, $C = 0.04$, and C represents a 4% safety factor to protect Class I areas from cumulative deposition impacts. This assumes that, over time, one might expect 25 or fewer new sources to increase deposition in a Class I area. The FLMs reserved the right to modify the DAT in the event that the number of existing and proposed new sources near a Class I area exceeded 25 (U.S. Forest Service et al. 2011).

For Class I parks and refuges east of the Mississippi, the DAT is calculated as:

$$\text{DAT} = 0.5 \text{ kg/ha/yr} \times 0.5 \times 0.04 = 0.01 \text{ kg/ha/yr of N or S} \quad (9)$$

The FLMs adopted this approach and resulting values in the *Federal Land Managers' Interagency Guidance for Nitrogen and Sulfur Deposition Analyses* (U.S. Forest Service et al. 2011). On this basis, the U.S. Forest Service has adopted DATS for both S and N deposition to protect against water acidification equal to 0.01 kg/ha/yr of S or N (http://www.fs.fed.us/air/technical/class_1/wilds.php?recordID=6) at Boundary Waters Canoe Area Wilderness, MN and at Rainbow Lakes Wilderness, WI.

For the purposes of protecting FCPC resources against acidification damage or fertilization of water, this report recommends following the approach used by the NPS and U.S. Forest Service and using a DAT of 0.01 kg/ha/yr of N or S (Table 6).

The DAT, MDI, and CL approaches provide different tools that can be used to evaluate ecosystem protection issues. Any or all of these tools can be used to screen proposed emissions sources to determine the likelihood of adverse impact.

5. SUMMARY

This report identifies appropriate threshold effects levels for water quality and aquatic systems AQRVs on FCPC lands. The primary concern for protection of these AQRVs is the potential for increases in MeHg in Devils Lake and other water bodies on FCPC Class I lands that could be caused by increased atmospheric deposition of either sulfur (S) or mercury (Hg). The most important route of entry of Hg into remote ecosystems in the region that includes FCPC lands is atmospheric deposition of inorganic Hg. However, the most important source of the neurotoxic MeHg to aquatic ecosystems is via microbially mediated methylation of inorganic Hg.

Atmospheric pollutant thresholds are addressed in the Federal Land Managers Air Quality Related Values Work Group (FLAG) Phase I report, initially drafted in 2000 and more recently revised in 2010 and 2011. The purpose of the FLAG Phase I report is to assist permit applicants, state authorities, and land managers to develop more consistent and objective approaches to the evaluation of AQRVs. This is needed for evaluation of air pollution effects, caused by new or modified emissions sources, on AQRVs in Class I areas under the New Source Review program, especially in the review of Prevention of Significant Deterioration air quality permit applications. The FLAG Phase I report and its modifications provide background information on effects levels and screening tools to determine if effects are expected to be negligible based on the planned quantity of emissions, the distance between the proposed source and the Class I area, and dose/response relationships. Among the primary issues of concern to Federal Land Managers (FLMs) and tribal land managers are the effects of pollutant deposition (including S, N, and Hg) on soils and surface waters.

The FCPC has designated Water Quality and Aquatic Systems as AQRVs to be protected against air pollution degradation. These AQRVs can be adversely impacted by atmospheric deposition of Hg, S, and N. Important effects include Hg methylation (which is related to both Hg and S deposition), acidification (which is related to both S and N deposition), and fertilization of water and soils (which is related to N deposition). In order for the FCPC to

determine whether any future emissions source would adversely impact the designated AQRVs, The FCPC must establish threshold effects levels, which represent the amount of additional (above ambient) deposition from a proposed new, or modified existing, source below which an adverse effect will not occur. Such thresholds are proposed herein. Once atmospheric deposition of S, N, or Hg is high enough that a deposition concern threshold has been reached, any further increase in emissions that would contribute additional deposition is considered unlikely to be acceptable.

Studies of FCPC surface waters and surface waters elsewhere in Wisconsin and the broader northern Great Lakes region have documented that many water bodies are contaminated with high levels of Hg. This is especially true for Devils Lake, which is of great cultural, recreational, and spiritual importance to the Potawatomi Tribe. As a consequence of high levels of Hg found in fish tissue tested from Devils Lake, the Tribe issued a fish advisory to limit human consumption of fish from Devils Lake. Because Hg is already at unsafe levels in Devils Lake and other water bodies in and near the FCPC reservation, the collective CLs of S and Hg deposition for protecting these impacted watersheds against Hg methylation have already been exceeded. Additional sources of either S or Hg are considered unacceptable because of the likelihood of enhanced Hg methylation that this would cause. Additional inputs of either S or Hg to FCPC lakes would contribute to further diminishment of the water bodies' significance to the Tribe, impair the natural ecosystem, and impair the quality of Tribal members' and visitor's experiences. Based on guidance from the FLMs provided in the FLAG Phase I Report, such effects are unacceptable.

Devils Lake and other water bodies on and near the FCPC Class I area contain levels of Hg that are considered to be unsafe for protecting human health and fish-eating wildlife. Any increase in the atmospheric deposition of either S or Hg would likely increase MeHg production within the Class I area, thereby causing additional Hg contamination of fish and wildlife. For that reason, the threshold effects level for additional atmospheric deposition of both S and Hg should be set at zero in order to prevent further Hg methylation and bioaccumulation.

Reported fish Hg content risk thresholds for protecting piscivorous wildlife against sublethal effects range between about 0.2 and 0.3 ppm (wet weight) in fish fillets or whole fish. Measured values in fish collected from several of the sampled lakes on FCPC lands and elsewhere within the study area were substantially higher than these threshold effects levels.

Furthermore, concentrations of Hg measured in fish from Devils Lake were substantially higher than the median Hg concentrations reported for yellow perch collected throughout the northern Great Lakes region (0.14 ppm, wet weight [ww]).

The U.S. Forest Service (USFS) designated concern thresholds for Hg in fish in two federal Class I areas in reasonable proximity to the FCPC lands that are the subject of the analysis reported here: the Boundary Waters Canoe Area Wilderness, MN and Rainbow Lakes Wilderness, WI. The concern threshold for Hg in fish tissue was established as 0.05 ppm (ww, skin-on fillet) in both wilderness areas. This level is exceeded in all monitored FCPC lakes.

The USFS also designated concern thresholds for surface water acid neutralizing capacity (ANC) in the Boundary Waters Canoe Area Wilderness and the Rainbow Lakes Wilderness. The threshold for maintaining healthy biological functioning in lakes and streams requires chronic $ANC \geq 25 \mu\text{eq/L}$. Thus, FLMs have adopted a screening level for lake ANC equal to $25 \mu\text{eq/L}$ to evaluate the likelihood of adverse impacts of acidification on lake ecosystems. This screening level is applied here to measured data for lakes on FCPC lands. Water chemistry has been measured by FCPC in 4 lakes and 12 streams over the last decade. Published surface water ANC has been below $25 \mu\text{eq/L}$ in Devils Lake and its inlet stream.

It is not known to what extent surface waters on FCPC lands are N-limited and therefore susceptible to nutrient enrichment impacts due to atmospheric N deposition. It is also not known whether atmospheric inputs of N have been sufficient to cause changes in plant species distributions or abundance due to nutrient enrichment. Current atmospheric N deposition to FCPC lands in the range of 4 kg N/ha/yr is at the low end of the range suggested to protect peatlands in the United States from effects associated with changes in productivity. Peatlands are among the plant community types expected to be most sensitive to nutrient enrichment. Thus, fertilization effects on FCPC lands are possible for the most sensitive plant communities under current N loading.

Potawatomi tribal members have a close link to the natural environment, including its water quality and wildlife. Elements of the natural environment are important to religious rituals, ceremonies, and medicines. Contamination of tribal waters, including entry of Hg into the food web, jeopardizes the tribe's cultural and religious beliefs and practices and threatens public health. The FCPC designated water quality and aquatic systems as AQRVs because:

- tribal members rely on fish as part of their traditional subsistence way of life,

- pure water is important to the tribe's cultural and religious practices, and
- recreational fishing is important to the tourism economy in and around the FCPC reservation, upon which the FCPC depends.

The traditional subsistence way of life followed by many FCPC tribal members relies heavily on fish, clean water, and other natural resources.

AQRVs associated with aquatic resources on FCPC lands are sensitive to atmospheric deposition of S, Hg, and potentially N. As documented in this report, this sensitivity is supported by extensive scientific research conducted throughout the northern Great Lakes region. AQRVs on FCPC lands are clearly impacted by recent and current levels of S and Hg deposition. This is reflected in the high concentrations of Hg in fish collected from FCPC lakes and to a lesser extent in the low ANC of Devils Lake. The ANC of this lake is influenced by both moderate lakewater sulfate concentration, likely contributed by atmospheric S deposition, and naturally occurring high dissolved organic carbon associated with wetlands within the watershed.

Critical and target loads have not been formally developed for atmospheric deposition of S, Hg, or N to watersheds on FCPC lands. Nevertheless, the demonstrated impacts reflected in high Hg concentrations in fish and low lake ANC indicate that CLs have been exceeded, at least in the short term.

Current deposition exceeds the critical or target load for protecting aquatic ecosystems on FCPC lands against Hg methylation and at least on occasion also the critical or target load for protecting aquatic ecosystems against acidification. It is not known if critical or target loads are exceeded for protection of aquatic, terrestrial, or transitional ecosystems against nutrient enrichment. Such impacts are possible for the most sensitive ecosystems.

Various indicators can be used to evaluate whether atmospheric deposition is high enough to trigger adverse acidification or nutrient enrichment effects. The National Park Service and U.S. Fish and Wildlife Service developed deposition analysis thresholds (DATs) for S and N in response to requests by permitting authorities and permit applicants to develop consistent, predictable permit review processes. A DAT is defined as *the additional amount of S or N deposition within a Class I area below which estimated impacts from a proposed new or modified source are considered insignificant*. The DAT represents total (wet plus dry) deposition. It is compared with the estimated amount of additional deposition resulting from the proposed new or modified existing source, as modeled using an atmospheric transport model.

The DAT calculation reported by FLAG in 2011 relies on an estimate of Natural Background Deposition developed from the peer-reviewed scientific literature and expert judgment. This natural background estimate is modified in calculating the DAT by including a Variability factor and a Cumulative Factor.

The variability factor ($V = 0.5$) was selected with the reasoning that a total increase in deposition from all sources by an amount greater than 50% of natural background deposition would trigger management concerns. Thus, an additional 50% of natural background deposition could be added to existing natural plus anthropogenic deposition and still be considered insignificant. The Cumulative Factor ($C = 0.04$) adjusts for the concern that cumulative effects from multiple relatively small sources could produce impacts even if a single source, on its own, would not. The FLMs applied the 4% value used by EPA in determining Class I increment levels of significant impact. Thus, C represents a 4% safety factor to protect Class I areas from cumulative deposition impacts. This assumes that, over time, one might expect 25 or fewer new sources to increase deposition in a Class I area. The FLMs reserved the right to modify the DAT in the event that the number of existing and proposed new sources near a Class I area exceeded 25.

For Class I parks and refuges east of the Mississippi, the DAT is calculated as:

$$\text{DAT} = 0.5 \text{ kg/ha/yr} \times 0.5 \times 0.04 = 0.01 \text{ kg/ha/yr of N or S}$$

The FLMs adopted this approach and resulting values in the *Federal Land Managers' Interagency Guidance for Nitrogen and Sulfur Deposition Analyses*. On this basis, the USFS has adopted DATS for both S and N deposition to protect against water acidification equal to 0.01 kg/ha/yr of S or N at Boundary Waters Canoe Area Wilderness, MN and at Rainbow Lakes Wilderness, WI. For the purposes of protecting FCPC resources against acidification damage or fertilization of water, this report recommends following the approach used by the NPS and U.S. Forest Service and using a DAT of 0.01 kg/ha/yr of N or S.

A DAT for Hg has not been developed on the basis of assumed background Hg deposition by the FLMs, probably in part because background Hg deposition is less well known. Nevertheless, an alternative DAT for wet deposition can be expressed on the basis of the minimum concentration that can be detected in precipitation in the analytical laboratory. Using this approach, an increase in wet deposition is considered significant if the increase in

precipitation concentration, and associated wet deposition, is measurable in the deposition monitoring network. This approach for specifying a DAT based on what is measurable in the analytical laboratory has been followed by USFS for the James River Face Class I Wilderness. It constitutes an alternate approach for calculating the DATs for Hg methylation in view of the lack of information regarding background Hg deposition.

Using this approach, the minimum detectable level of an increase in S or Hg wet deposition to FCPC lands to protect against Hg methylation can be calculated using the following equation:

$$\text{MDI} = C \times P \times \text{UCF}$$

where MDI for the estimated minimum detectable increase in wet deposition; C is the laboratory detection limit for S or Hg measured in water from wet deposition collectors; P is annual precipitation; and UCF is the unit conversion factor value needed to express MDI in the selected units (e.g., kg/ha/yr for S or $\mu\text{g}/\text{m}^2/\text{yr}$ for Hg).

The MDIs with respect to the concentrations of S and Hg wet deposition were calculated for the NADP/NTN monitoring station W110 located within the Potawatomi study area for the five-year period 2006 through 2010 based on annual average precipitation and the current analytical method detection limits (MDL) in the NADP/NTN and Mercury Deposition Network (MDN) monitoring programs.

The DATs for protecting against Hg methylation, based on laboratory detection limits, can then be calculated from the minimum detectable increase in deposition of total Hg (MDI_{THg}) and total S (MDI_{TS}). These equations incorporate the Cumulative Factor ($C = 0.04$) developed by the FLMs for protecting against acidification and nutrient enrichment:

$$\begin{aligned} \text{DAT}_{\text{Me-Hg}} &= 0.088 \mu\text{g Hg}/\text{m}^2/\text{yr} \times C \\ &= 0.088 \mu\text{g Hg}/\text{m}^2/\text{yr} \times 0.04 = 0.0035 \mu\text{g Hg}/\text{m}^2/\text{yr} \end{aligned}$$

$$\begin{aligned} \text{DAT}_{\text{Me-S}} &= 0.031 \text{ kg S}/\text{ha}/\text{yr} \times C \\ &= 0.031 \text{ kg S}/\text{ha}/\text{yr} \times 0.04 = 0.0012 \text{ kg S}/\text{ha}/\text{yr} \end{aligned}$$

The rationale for inclusion of the Cumulative Factor (C) into calculation of the DATs to protect against Hg methylation (DAT_{Me-Hg} and DAT_{Me-S}) is the same as the rationale given by the FLMs for including C in calculation of the DATs to protect against acidification and nutrient enrichment. Over time, one might expect 25 or fewer new emissions sources to increase deposition to a Class I area. The factor C represents a safety factor to protect Class I areas from cumulative impacts.

The CLs, DATs, and MDIs constitute tools that can be used individually or collectively to evaluate the likelihood that a proposed new, or modified existing, emissions source might adversely impact AQRVs in the Class I area. These tools can be used to screen proposed emissions sources for more in-depth analyses to evaluate potential adverse impacts.

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