Decreased atmospheric nitrogen deposition in eastern North America: Predicted responses of forest ecosystems

Frank S. Gilliam, Douglas A. Burns, Charles T. Driscoll, Serita D. Frey, Gary M. Lovette, Shaun A. Watmough

1 Department of Biology, University of West Florida, Pensacola, FL 32514, United States
2 U.S. Geological Survey New York Water Science Center, 425 Jordan Road, Troy, NY 12180, United States
3 Department of Civil and Environmental Engineering, Syracuse University, Syracuse, NY 13244, United States
4 Department of Natural Resources and the Environment, University of New Hampshire, Durham, NH 03824, United States
5 Cary Institute of Ecosystem Studies, PO Box AB, Millbrook, NY 12546, United States
6 School of the Environment, Trent University, 1600 West Bank Drive, Peterborough, ON K9L 0G2, Canada

ABSTRACT

Historical increases in emissions and atmospheric deposition of oxidized and reduced nitrogen (N) provided the impetus for extensive, global-scale research investigating the effects of excess N in terrestrial and aquatic ecosystems, with several regions within the Eastern Deciduous Forest of the United States found to be susceptible to negative effects of excess N. The Clean Air Act and associated rules have led to decreases in emissions and deposition of oxidized N, especially in eastern U.S., representing a research challenge and opportunity for ecosystem ecologists and biogeochemists. The purpose of this paper is to predict changes in the structure and function of North American forest ecosystems in a future of decreased N deposition. Hysteresis is a property of a system wherein output is not a strict function of corresponding input, incorporating lag, delay, or history dependence, particularly when the response to decreasing input is different from the response to increasing input. We suggest a conceptual hysteretic model predicting varying lag times in recovery of soil acidity, plant biodiversity, soil microbial communities, forest carbon (C) and N cycling, and surface water chemistry toward pre-N impact conditions. Nearly all of these can potentially respond strongly to reductions in N deposition. Most responses are expected to show some degree of hysteresis, with the greatest delays in response occurring in processes most tightly linked to "slow pools" of N in wood and soil organic matter. Because experimental studies of declines in N loads in forests of North America are lacking and because of the expected hysteresis, it is difficult to generalize from experimental results to patterns expected from declining N deposition. These will likely be long-term phenomena, difficult to distinguish from other, concurrent environmental changes, including elevated atmospheric CO2, climate change, reductions in acidity, invasions of new species, and long-term vegetation responses to past disturbance.

© 2018 Elsevier Ltd. All rights reserved.

1. Introduction

Gaseous N2 comprising 78% of the atmosphere (non-reactive N—N2) neither undergoes photochemical transformations nor enters metabolic pathways in organisms, other than through fixation by a small group of prokaryotes and archaea. By contrast, there are numerous forms of reactive N (N) (e.g., NH3, NH4+, NO, NO2, NO3, N2O5, and HNO3—Horii et al., 2005), which are all capable of undergoing transformations in the atmosphere and entering into biogeochemical cycles of terrestrial and aquatic ecosystems. In addition to emissions of N via internal combustion and power production, ever-increasing amounts of N2 are converted to N-based fertilizer via the Haber-Bosch (H-B) industrial process; currently, fixation via H-B exceeds natural N fixation by >20% (Canfield et al., 2010).

Historical awareness of increasing rates of atmospheric deposition of N2, along with projections by some models of further increases in the future, provided the initial impetus toward research
on excess N in forest ecosystems (Lamarque et al., 2013). Based on estimates from Galloway et al. (2004), total global atmospheric deposition of NH₄⁺ and NO₃⁻ in terrestrial ecosystems has increased from 17 Tg N yr⁻¹ in 1860 to 64 Tg N yr⁻¹ in the early 1990s, with projected further increases to 125 Tg N yr⁻¹ by 2050, suggesting that atmospheric deposition of N will have increased by over 7-fold during this 190-year period. Bobbink et al. (2010) predicted similar increases in N deposition by 2030.

Much of the initial work on effects of elevated atmospheric N deposition on terrestrial ecosystems in North America focused on biogeochemical responses, with many studies examining spatial variation in stream/surface water chemistry. Findings generally indicated that with increases in N deposition, leaching rates and stream concentrations of NO₃⁻ and cations (Ca²⁺, Mg²⁺, K⁺, Al³⁺, H⁺) coupled to NO₃⁻ mobility also increased (Murdoch and Stoddard, 1992; Aber et al., 2002; Adams et al., 2006). Large areas of temperate forests in North America, especially those of the eastern U.S., have been shown to be sensitive to N saturation, a phenomenon wherein the rate of atmospheric N deposition exceeds the ability of an ecosystem to retain it (Lovett and Goodale, 2011). More recent work has emphasized the effects of excess N on plant biodiversity, microbial responses, and health of temperate forests (Lloret and Valiela, 2016), with broad consensus indicating that excess N has decreased biodiversity and compromised the health of some forests of the eastern U.S. (Clark et al., 2013).

The biogeochemistry of N cycling has received extensive research emphasis (Gorham and Kelly, 2018), especially regarding effects of excess N on ecosystem biogeochemistry and biotic communities. Beginning in the 19th century, with the initiation of the Park Grass Experiment in Rothamsted, England, there is a rich history of experimental studies investigating the effects of N on plant species diversity (Richardson, 1938; Silvertown et al., 2006). As the longest on-going N experiment, Park Grass offered the first evidence that excess N can reduce plant species diversity, results confirmed in the U.S. by ongoing investigations at the Cedar Creek Natural Area in Minnesota (e.g., Clark and Tilman, 2008). Research was initially directed primarily toward herb-dominated communities, but increasingly has since focused on forested ecosystems (Gilliam, 2006; Sutton et al., 2014; Clark et al., 2013; Gilliam et al., 2016).

Empirical data show that emissions of N from vehicles and power plants in the U.S. have decreased >50% since the 1980s (Lloret and Valiela, 2016), largely the result of the Clean Air Act (CAA). Because the focus of the CAA has been on oxidized, rather than reduced, forms of N, temporal patterns of atmospheric deposition have differed among N forms. Data collected by the National Atmospheric Deposition Program (http://nadp.srh.wisc.edu/) and by the Clean Air Status and Trends Network (https://www.epa.gov/castnet) indicate that both wet and dry deposition of oxidized N increased from the 1980s, peaked in the mid-to late-1990s, and then decreased notably toward recent years (Du, 2016; Lloret and Valiela, 2016). By contrast, absolute and relative rates of deposition of reduced N have either increased or remained level at most sites (Warner et al., 2017), with increases driven primarily by decreases in SO₂ and NOₓ emissions (due to decreases in scavenging of atmospheric NH₄⁺ by SO₂⁻ and NO₃⁻) and increases in soil temperature (Fig. 1; Du et al., 2014).

Spatial patterns of total deposition of oxidized versus reduced N over time reveal even sharper contrasts. Whereas total oxidized N deposition reached uniformly high values throughout the eastern U.S. in 2000, by 2014 elevated deposition was confined to major metropolitan areas (Fig. 2a and b). Deposition of reduced N shows a patchy distribution throughout the U.S., driven mainly by agricultural practices, including animal husbandry (Fig. 3a and b)
N deposition has increased over time, then reflects the decrease in excess N as N deposition decreases (i.e., decreasing in North America after the mid-1990s—Fig. 1). Our conceptual hysteretic model predicts varying lag times in recovery of soil acidification, plant biodiversity, soil microbial communities, forest C and N cycling, and surface water chemistry toward pre-N impact conditions as deposition of N continues to decline.

Fig. 2. Total deposition of NOx for 2000 (a) and 2014 (b). Maps were taken from the National Atmospheric Deposition Program (http://nadp.slh.wisc.edu/).
2. Soil acidification

The CAA targets several criteria pollutants, among them the oxides of S and N that contribute to acid deposition, a phenomenon that has been shown to accelerate soil and water acidification. Although soil acidification is a natural phenomenon, occurring in regions where precipitation quantity exceeds evapotranspiration (van Breeman et al., 1983; Jones et al., 2012), acid deposition can accelerate natural soil acidification with adverse consequences for acid-sensitive ecosystems. Acid sensitive regions typically are...
regions by depleting available Ca and Mg from soil exchange sites needed for ecosystems to recover from acid deposition. Acid inputs of strong acids, provide poorer growing conditions for those that have been altered by acid deposition are less able to neutralize occur in extremely Ca-poor ecosystems (Bailey et al., 1996). For exceptions to this pattern at atmospheric deposition. Although this pathway is generally small Calcium and other nutrient cations may also enter forests through surface waters.

Understanding the cycling of Ca, Mg, and potassium in forest ecosystems involves consideration of the inputs and losses of these important nutrient cations. For many forest ecosystems, the primary supply of Ca and other nutrient cations largely occurs by weathering (i.e., the breakdown of rocks and minerals in soil). Calcium and other nutrient cations may also enter forests through atmospheric deposition. Although this pathway is generally small compared with chemical weathering, exceptions to this pattern occur in extremely Ca-poor ecosystems (Bailey et al., 1996). For example, natural sources such as airborne soil particles originating from arid regions contribute to high atmospheric Ca deposition in Northeast Asia, and human emissions of particulate matter from cement production and other industrial activities result in high atmospheric Ca deposition in South Asia (Larsen and Carmichael, 2000; Zhao et al., 2011).

Ecosystem losses of Ca occur largely by vegetation uptake and subsequent harvesting, as well as by leaching in drainage waters. The soil available pool or soil cation exchange complex provides an important and readily available source of Ca and other nutrient cations. Plants are generally able to utilize these nutrients from solution in equilibrium with soil exchange surfaces. Forest ecosystems that are naturally sensitive to acid deposition are often characterized by low rates of weathering and low quantities of available nutrient cations. Under conditions of elevated acid deposition and subsequent transport of SO4 and NO3 in drainage waters, nutrient cations are displaced from available pools and leached from soil (Ruess and Johnson, 1986). This condition is not problematic in areas with high weathering rates and abundant pools of available nutrient cations. However, in acid-sensitive areas with shallow soil characterized by minerals with high resistance to weathering, enhanced loss of Ca and other nutrient cations can result in depletion of soil available pools. Both NO3 and SO4 contribute to this process but with different seasonal patterns (Ruess and Johnson, 1986). Sulfate leaching generally occurs chronically throughout the annual cycle, due to the fact that it is largely controlled by abiotic processes (i.e., atmospheric deposition and soil adsorption). In contrast, NO3 leaching often shows strong seasonal patterns driven by biotic processes and is strongly dependent on hydrologic conditions (Stoddard, 1994).

Over the last century, acid deposition and associated leaching by strong acid anions has accelerated the loss of relatively large amounts of available Ca and Mg from soil in acid-sensitive areas (Likens et al., 1996; Warby et al., 2009). This depletion occurs when nutrient cations are displaced from the soil by acid deposition at a rate faster than they can be replenished by the slow breakdown of rocks and minerals or deposition from the atmosphere. Depletion of nutrient cations can fundamentally alter soil development, compromise the nutrition and health of sensitive tree species, influence adjacent surface waters, and limit the capacity of sensitive soils to recover. For example, based on a hindcast analysis, more than half of the available Ca has been lost from soil at the Hubbard Brook Experimental Forest, New Hampshire over the past 70 years (Likens et al., 1996). Note that while acid deposition to acid-sensitive areas is decreasing and there is some associated recovery of the acid neutralizing capacity (ANC) of surface waters, it appears that forest soils continue to exhibit depletion of exchangeable nutrient cations or at best are recovering soil pools of available nutrient cations at a very slow rate (Warby et al., 2009; Fakhræ et al., 2014; Lawrence et al., 2011).

Aluminum (Al) can be released from soil to soil water, lakes, and streams in forested regions with high acid deposition, low stores of available Ca, and acidic soil (Driscoll and Postek, 1995; Driscoll et al., 2001). One of the most significant ecological effects of acid deposition is the mobilization of Al from soil and a shift in the form of Al in water from non-toxic organic forms to toxic Al3+. Concentrations of Al increase markedly with decreases in pH below 6, particularly Al3+. Note that because of short-term temporal variations in NO3 concentrations, seasonal and episodic variations in Al generally closely track concentrations of NO3 in chronically and episodically acidic waters (Driscoll and Van Dreason, 1993; Driscoll and Postek, 1995). High concentrations of Al3+ can be toxic to plants, fish, and other organisms (Gensemer and Playle, 1999). Concentrations of Al3+ in acid-impacted surface waters in eastern North America and Europe are above levels considered toxic to fish (~2 μmol/L) and much greater than concentrations observed in forest watersheds that receive low inputs of acidic deposition (Driscoll and Postek, 1995; Driscoll et al., 2001).

Empirical evidence suggests that the temporal pattern of soil available nutrient cations in acid sensitive soils likely follows a hysteretic pattern as described herein for N (Fig. 5a). During the acidification phase, increases in deposition and leaching of strong acid anions facilitate the depletion of exchangeable nutrient cations. Due to low weathering rates, decreases in acid deposition and NO3 and SO42− leaching will not concurrently result in parallel increases in soil base saturation resulting in slowly recovering soil fertility. In contrast, soil exchangeable Al and concentrations of Al3+ in surface waters are highly reversible in response to changes in acid deposition and leaching of strong acid anions in soils, even those with low base saturation (Cronan and Schofield, 1990; Fig. 5b; Lawrence et al., 2011). When soil base saturation is below a critical level (~12−15%), increases in concentrations of strong acid anions associated with increases in acid deposition will result in increases in concentrations of Al3+, because the mobilization/immobilization of Al3+ are controlled by fast processes (dissolution/precipitation of secondary Al minerals, adsorption/desorption of Al on soil surfaces).
The emerging issue of ‘browning of surface waters’ is associated with the recovery of forested watersheds in North America and Europe from the effects of acid deposition as manifested by increases in the mobilization of dissolved organic matter (Monteith et al. 2007). This process has implications for the recovery of ecosystems from acid and N deposition. First, a component of naturally occurring organic matter functions as organic acids (Fakhraei and Driscoll, 2015). As a result, elevated leaching of dissolved organic matter can also contribute to cation leaching and the acidification of soil and water, albeit as a natural acidification process. These organic acids can also facilitate the mobilization of Al (Driscoll and Postek, 1995; Fakhraei and Driscoll, 2015). However, in this case the form is dissolved organic Al, which is less toxic due to organic complexation. Finally, spatial stream surveys have shown an inverse relationship between concentrations of dissolved organic carbon and NO$_3^-$ (Goodale et al., 2005; Taylor and Townsend, 2010). The mechanism for this empirical relationship is unclear, but may be due to the role of dissolved organic matter as an electron donor during denitrification, or to microbial immobilization of NO$_3^-$ associated with the processing of dissolved organic matter as an energy source. Regardless, if this empirical relationship reflects a biogeochemical mechanism, then the enhanced mobilization of dissolved organic matter following decreases in acid deposition could accelerate the retention or denitrification of NO$_3^-$, an additional process that may affect ecosystem recovery.

An unresolved question remains—to what extent will the temporal shift in chemical speciation of atmospheric N deposition observed in the eastern U.S. affect N cycling processes and ecosystem recovery? Ammonium increased from <40% of wet-deposited N in the 1980s to >60% currently (Fig. 1). Does such a change have implications for forest ecosystem response? Certainly, these shifts could impact the recovery of the acid–base chemistry of soil and surface waters. In theory, NH$_4^+$ deposition followed by nitrification and leaching of NO$_3^-$ results in greater soil acidification than atmospheric NO$_3^-$ deposition and resulting leaching of NO$_3^-$. Both oxidation and biotic uptake of NH$_4^+$ are acidifying processes (van Breeman et al. 1983). In contrast, ecosystem uptake of NO$_3^-$ is an alkalization process. We know of no studies that have quantified shifts in soil acidification associated with decreasing atmospheric N deposition and a shift in its speciation. Nevertheless, the overall decreases in atmospheric N deposition and associated decreases in NO$_3^-$ leaching should diminish soil and surface water acidification. However, because this decrease has largely occurred due to decreases in oxidized N deposition, residual N deposition is relatively enriched in NH$_4^+$ and will have a relatively greater acidification potential (per mol of N deposition) than historical N deposition.

3. Plant biodiversity

In addition to its biogeochemical implications, soil acidification substantially alters availability of essential nutrients for uptake by plants. Thus, there is a strong potential for related changes in forest community structure and diversity. Excess N can have a direct fertilizer effect on plant communities, especially those of the most sensitive stratum of forest vegetation—the herbaceous layer (Gilliam, 2007).

Evidence is ubiquitous that chronically increased N deposition has decreased biodiversity of herb strata in essentially all forest types (Sutton et al., 2014), Clark et al. (2013) estimated, via hindcast modeling, N-mediated species loss for this region of up to 30% from 1985 to 2010. Using data from >15,000 sites, Simkin et al. (2016) demonstrated that negative relationships between atmospheric N deposition and plant species richness are quite common. Gilliam et al. (2016) found a pronounced shift in herb layer composition in response to 25 years of experimental N additions to a central Appalachian hardwood forest. This change arose from increases in a nitrophilic species (Rubus allegheniensis) that competitively eliminated numerous N-efficient herbaceous species, resulting in a loss of plant diversity.

Less work has focused on effects of excess N on forest overstory communities. Thomas et al. (2010) modeled the potential effects of N on the sequestration of tree biomass C in temperate forests, along with survivorship of tree seedlings. They found that chronically-elevated N deposition enhanced C storage, but decreased survivorship in eight of 11 common temperate tree species. Not considered in Thomas et al. (2010) is the degree to which herbaceous species can determine the success of tree seedling emergence out of the herb layer, often referred to as an ‘ecological filter’ (George and Bazazz, 2014). Thus, N-mediated changes in forest herb communities can represent an indirect effect of N on tree species.

Although few published data directly assess forest diversity response to decreased N deposition, there are studies—virtually all on the forest herb communities—that allow us to address this question to hypothesize on how N-impacted forests may respond to decreased N. Boxman et al. (1998) reported on results of transparent roof studies—the NITREX project in Europe. This networked study integrated results from seven sites throughout Europe wherein high ambient N deposition was excluded experimentally.
by roofs to simulate decreased pollutant N inputs. Results from the Netherlands showed that above-ground biomass of nitrophilous species decreased rapidly under the roof that excluded N to the forest floor over a 5 year period (Boxman et al., 1998), suggesting a short-term recovery of the forest herb community to decreased N inputs.

Strengbom et al. (2001) employed two separate forest fertilization experiments in northern Sweden assessing legacy effects of increased N loading following cessation of added N. The first experiment added NH₄NO₃ at rates up to 108 kg N ha⁻¹·yr⁻¹ from 1971 to 1990 (nine years prior to the Strengbom et al. study). Herb layer composition varied significantly between control and treatment areas, with herb diversity significantly lower on treatment areas, arising primarily from sharp declines in ericaceous species and increases in the nitrophilous grass, Deschampsia flexuosa. Strengbom et al. (2001) concluded that effects of increased N deposition on the herb layer can be long lived.

In the grass-dominated communities of the Park Grass Experiment, Storkey et al. (2015) demonstrated a positive response of biodiversity to decreases in N from either atmospheric deposition or fertilizers. However, Stevens (2016) reviewed studies from throughout Europe and concluded that plant species composition is likely to follow recovery during this period with the period of decreasing atmospheric N deposition. In contrast, she concluded that soil N dynamics are potentially more responsive to such decreases. She further identified possible barriers to recovery, including continued critical load exceedance and lack of either a seed bank or local seed source. Thus, the potential exists for plant communities to reach an alternative stable state wherein species loss from excess N may not allow recovery to pre-impact plant community composition.

That all these studies are from Europe represents a challenge, and a research opportunity, for forests of the eastern U.S. Among these challenges is the wide range of micro-environmental conditions that precludes broad generalizations. The response time of forest herb communities to increases in N load is typically inversely related to ambient N deposition and the antecedent (pre-N enhanced) N status of the forest. Sites with low N availability respond more quickly to increasing N than high N sites (Gilliam et al., 2016). Working in upstate New York and Massachusetts, which receive low levels of N deposition (relative to other regions of the eastern US; Fig. 2), Hurd et al. (1998) and Rainey et al. (1999), respectively, found loss of herb diversity to added N 1–2 years following initiation of treatment. By contrast, Gilliam et al. (2006) found no similar response following 5 year of N additions in a central hardwood forest of West Virginia, a region experiencing chronically high levels of N deposition.

Similar to the effect on soil acidity, the increase in ratio of NH₄ to NO₃ in atmospheric deposition (Fig. 1) previously discussed (see Soil acidification) has potential implications for future responses of plant species. Most plant species exhibit little discrimination in form of N for uptake, with the ability to take up both forms of mineral N in soil. In fact, most plant species perform best with a form of N for uptake, with the ability to take up both forms (often called calcifuges) show a notable preference for NH₄, whereas those adapted to neutral to alkaline soils (calcicoles) prefer NO₃ (Marschner, 1995; Nordin et al., 2001). Thus, potential exists for future plant communities to change toward calcifuge dominance, especially those of the herbaceous stratum.

We hypothesize that the degree of hysteresis (i.e., the distance between lines, Fig. 4) will vary directly with the N status of the impacted forest ecosystem. The results of Boxman et al. (1998) for the NITREX study and Storkey et al. (2015) for the Park Grass Experiment suggest that herb diversity can return to pre-impact levels in a decadal time frame. Although the greater physiognomical complexity of forests (compared to grasslands) should increase recovery time for the forest herb communities (up to 40 yr—Strengbom et al., 2001), we anticipate recovery of biodiversity of such communities under future decreases in N deposition. On the other hand, conclusions of Stevens (2016) are convincing that the composition may be quite different from the pre-impact condition, given alteration of seed pools and sources of dispersal. That is, the number of species may recover, but species composition may remain quite different, relative to pre-impact levels.

4. Soil microbial communities

Excess N deposition that alters both tree and herb communities of impacted forests also affects soil microbiota, both directly in forest floor and mineral soil, and indirectly through linkages between tree and herb strata and soil microbial communities (Gilliam et al., 2014). To predict how these communities in temperate forests may respond to a regional decrease in atmospheric N deposition, it is first important to understand microbial responses to decades-long soil N enrichment and the concomitant changes in soil properties (e.g., pH, nutrient concentrations) and plant communities. Considerable research has documented microbial responses to ambient and simulated N deposition, with general consensus that chronic soil N enrichment reduces total soil microbial biomass, active biomass, and the fungal:bacterial biomass ratio (Frey et al., 2004; Wallenstein et al., 2006; Frey et al., 2014); reduces microbial growth (Kamble et al., 2013) and the activities of extracellular enzymes, especially those associated with lignin breakdown (Carreiro et al., 2000; Sinsabaugh et al., 2002; Deforest et al., 2004; Frey et al., 2004); and reduces the capacity of the microbial community to utilize a suite of C substrates typically found in soil (Frey et al., 2004; Kamble et al., 2013; van Diepen et al., 2017). The mycorrhizal component of the microbial community is particularly sensitive to excess soil N, showing reductions in sporocarp production, root colonization, and mycorrhizal species richness (Egerton-Warburton and Allen, 2000; Lilleshov et al., 2002; Treseder, 2004; Morrison et al., 2016; Carrara et al., 2018).

Recent molecular analyses also indicate significant N-induced shifts in the diversity and composition of both the bacterial (Fierer et al., 2012; Turlapati et al., 2012) and total fungal communities (Entwistle et al., 2013; Mueller et al., 2014; Freedman et al., 2015; Hesse et al., 2015; Morrison et al., 2016), with increased relative abundances of copiotrophic bacteria (Fierer et al., 2012), weak fungal decomposers (e.g., yeasts, cellulolytic ascomycetes; Morrison et al., 2016, 2018), and nitrophilic taxa (Morrison et al., 2016). Thus, chronic soil N enrichment appears to favor the growth and competitive ability of fast-growing, N-tolerant copiotrophs, taxa that may have a lower capacity for efficient organic matter degradation. These shifts in temperate forest microbial communities in response to chronic soil N enrichment are consistently associated with reduced rates of leaf litter (Magill and Aber, 1998; Knorr et al., 2005) and organic matter decomposition (Pregitzer et al., 2008; Liu and Greaver, 2010; Zak et al., 2011; Lovett et al., 2013; Frey et al., 2014), leading to an accumulation of soil organic matter with altered chemistry (Pregitzer et al., 2008; Zak et al., 2008; Lovett et al., 2013; Frey et al., 2014).

It is a reasonable assumption that changes in microbial communities resulting from long-term N additions are reversible if decreases in N deposition are coupled with recovery in soil N cycling rates, pH, base cation and trace nutrient concentrations, and plant community structure. However, a limited number of studies have examined the impact of decreases in N deposition on soil microorganisms. Available data are equivocal, with some evidence for rapid recovery and other results suggesting that microbial
community diversity and at least some microbially-mediated processes may remain impacted for many years. Mycorrhizal diversity and fruiting body abundance has been shown to significantly increase ~5–15 years of N after fertilization ceased (Högberg et al., 2014; Stevens, 2016) and references therein), though this result is inconsistent across studies. For example, in a long-term N addition and recovery experiment, mycorrhizal spore abundance remained significantly different on formerly N-enriched plots compared to untreated controls following more than four decades of recovery (Strengbom et al., 2001). Additionally, differences in mycorrhizal species composition remained with spore-crop production of N-sensitive mycorrhizal fungi lower in formerly N-treated plots. Bacterial community composition is also slow to recover (Högberg et al., 2014). Microbial community reassembly and soil processes (e.g., decomposition) may be slower to recover due to legacy effects resulting from the hysteretic recovery of soil pH and macro- and micronutrient concentrations (e.g., Ca, Mg, P, Mn). Note that, if some aspects of the microbial community recover relatively quickly (e.g., species richness, mycorrhizal spore abundance), the microbial community, as a whole, may not return to the same state (i.e., composition) as prior to the onset of high N deposition rates.

One of the surprises of N deposition research has been the high retention for added N in northeastern U.S. forests (Aber and Magill, 2004). Even systems exposed for decades to high N fertilization rates show retention of greater than 70% of total N inputs (Aber et al., 1989; Aber et al., 1998), with most of the added N being retained in the forest floor and to a lesser extent in the mineral soil (Nadelhoffer et al., 1999, 2004; Templer et al., 2012). So, although soil inorganic N pools may recover to background levels within a few years (Stevens, 2016), total soil N and its overall availability may remain elevated for many years, resulting in cascading effects on other soil properties (e.g., pH, macro- and micronutrient concentrations). As already discussed (see Plant biodiversity), plant communities may also show slow recovery, with nitrophilic species self-perpetuating in the understory through mechanisms such as allelopathy, shading of neighbors, and via litter accumulation. Microbial community recovery to its pre-N enriched state may be also be impeded or delayed if the composition of the microbial spore bank was significantly altered (Glassman et al., 2016) or if microbes lost their capacity, over decades of soil N enrichment, to perform particular functions (van Diepen et al., 2017).

5. Forest carbon and nitrogen cycling

Clearly, both increases and decreases in N supply to terrestrial ecosystems can have a profound influence on soil microbial dynamics, which in turn influence the cycling of C and N in impacted forest ecosystems. In addition, two complexities make predicting responses of forest C and N cycling to declining N deposition especially challenging. First, declines in N deposition can cause multiple, sometimes offsetting, responses. For instance, if N is retained by vegetation, it can have a fertilization effect, increasing productivity, whereas if NO₃⁻ leaches from the ecosystem it can acidify soils and reduce productivity (Lovett and Goodale, 2011) (Fig. 6). Similarly, low levels of N addition can increase plant tissue N concentrations, stimulating N mineralization, but high levels of N addition can decrease decomposition, inhibiting N mineralization rates (Fig. 6). Thus, we may expect different responses at different positions along the trajectory of response to decreases in N deposition.

Second, forest ecosystem pools and fluxes related to C and N cycling are likely to vary widely in the rate at which they respond to decreases in N deposition. These processes that respond slowly are associated with the slow-turnover pools of C and N in the ecosystem, principally soil organic matter (SOM) and wood. Faster-responding processes are associated with pools of N that often have a shorter turnover time such as soil mineral (inorganic) N, microbial biomass, and sometimes foliage and fine roots. Lags in response of the slow pools can produce a hysteresis by slowly feeding N to the fast-response components of the system. In addition, the slow pools can sometimes increase in N content more quickly than they decrease; for example, N in microbial exudates can quickly be incorporated into stable SOM which may take decades or more to turn over. We hypothesize that the SOM and wood pools are likely to respond slowly to declines in N deposition and exhibit pronounced hysteresis, whereas fast pools such as soil mineral N will respond rapidly and exhibit less hysteresis. Further, we predict that the response of ecosystem N cycling processes will vary depending on the extent to which they depend on these slow or fast pools. For example, when the forest C and N cycling model Spe-CN (Crowley et al., 2016; Crowley and Lovett, 2017) is used to simulate the response of a second-growth forest to an increase followed by a decrease in N deposition, net primary production (NPP) responds more slowly and shows more hysteresis than N leaching (Fig. 7a and b). This difference occurs because NPP depends strongly on N availability which is primarily controlled by N mineralization from the slow-turnover pools (e.g., N mineralization during the dormant season) is controlled much more directly by deposition levels acting through the fast-turnover soil mineral N pools.

These hypotheses are largely consistent with results of field studies of decreases in N deposition in European forests (Stevens, 2016). In the Netherlands and Germany, soil inorganic N fluxes declined by over 50% in coniferous forest plots within 2 years after roofs were placed over the soil to exclude N deposition (Boxman et al., 1995, Bredemeier et al. 1998). Tree-associated variables have changed more slowly in these studies. Tree growth increased significantly in the Dutch N-exclusion plots after 6 years of treatment (Boxman et al., 1998), indicating that the trees had been experiencing negative effects of soil acidification that outweighed the benefits of N fertilization. In Scots pine stands in Sweden, foliar %N and ectomycorrhizal function recovered to control plot levels within 6–15 years after the termination of a 20-year N addition experiment (Högberg et al., 2011).

Although similar N exclusion experiments have not been reported for forests of eastern North America, N addition experiments have shown that soil NO₃⁻ and NH₄⁺ concentrations, soil solution NO₃⁻, and stream NO₃ generally show relatively fast (i.e., 1–2 years) responses to N addition (e.g., Gilliam et al., 2001; Edwards et al., 2006; Lovett and Goodale, 2011; Fatemi et al., 2012), indicating that those variables are governed by fast-turnover pools of N and are likely to decrease rapidly in response to decreases in N deposition.

In contrast, net N mineralization rates often exhibit no observable change in response to experimental N additions (Wang and Fernandez, 1999; Lovett and Goodale, 2011; Lovett et al., 2013) or changes may develop slowly (Jefts et al., 2004). Gilliam et al. (2018) found that the temporal pattern of net N mineralization and nitrification at the Fernow Experimental Forest in West Virginia was determined more by year-to-year changes in temperature than by experimental N additions through 25 years treatment. Forest growth showed mixed responses to experimental increases in N inputs, with some sites exhibiting increases in vegetation growth (e.g., Fowler et al., 2015; Magill et al. 2004), some showing no change (e.g. Lovett et al., 2013), some showing declines in growth and slow tree mortality (e.g., Magill et al. 2004; Wallace et al., 2007; McNulty et al., 2005). Some sites show increases in tree growth after N fertilization, followed by later decreases, presumably because acidification effects eventually overwhelms fertilization effects (e.g., DeWalle et al., 2006). Differences in the direction
and timing of tree growth response can probably be attributed to differences in forest age, species composition, and the sensitivity of the site to soil acidification.

Another approach to evaluating patterns of response is to track C and N cycling properties of a forest as ambient N deposition declines. For example, in the Netherlands, decreases in ambient N deposition by about 25% (from ~60 to ~45 kg N ha\(^{-1}\) y\(^{-1}\)) led to rapid declines in inorganic N concentrations in soil solution and soil leachate (Boxman et al., 2008).

Mathias and Thomas (2018) studied red spruce forests of the Central Appalachian region using a multiproxy approach with 75-year tree ring chronologies of basal area growth, C isotope discrimination (\(\Delta^{13}C\), a proxy for leaf gas exchange), and \(\Delta^{15}N\) (a proxy for ecosystem N status) to examine individual tree and ecosystem-level responses to global change, including atmospheric CO\(_2\) concentrations, and acid and N deposition. They documented a notable recovery of red spruce ecosystems—including increases in basal area increment (BAI), photosynthesis, stomatal conductance, and water use efficiency—beginning in 1989, which they ascribed to the efficacy of the Clean Air Act and its reduction in several pollutants, including N, but especially acid deposition. Although improvements in soil acidity exhibited the best correlation with recent increases in tree growth, decreases in NO\(_x\) emissions from 1989 to the present were significantly correlated with increased BAI in red spruce during this period. Similarly, Wason et al. (2017) found that red spruce in the Adirondack region of New York are showing recent increases in growth rates attributed largely to declines in atmospheric deposition of SO\(_2\) and NO\(_x\) as well as a minor contribution from a warming climate.

At the Hubbard Brook Experimental Forest in NH, USA, one of the best-studied forest research sites in North America, bulk N deposition has declined by 50% from about 8 kg N ha\(^{-1}\) y\(^{-1}\) in 1990 to about 4 kg N ha\(^{-1}\) y\(^{-1}\) currently (Rosi-Marshall et al., 2016). During this period there has also been a decline in potential net N mineralization, nitrification and soil inorganic N pools, which Duran et al. (2016) attributed primarily to climate change. There has been a slight decline in aboveground biomass in the reference watershed, attributed primarily to soil acidification (Battles et al., 2014). Soil solution and stream water N export has remained at the lowest levels in the 50-year record at this site (notwithstanding several spikes likely due to disturbance events); the mechanism for these low levels of export is still unknown (Yanai et al., 2013; Fuss et al., 2015), but has been attributed to successional processes (Bernal et al., 2012). All of these trends are consistent with what is expected from declining N deposition, yet in each case the authors attribute them primarily to other causes, illustrating the complexity of distinguishing forest responses to N deposition from responses to other concurrent environmental drivers.

In summary, the evidence presented here leads us to hypothesize that as N deposition declines in eastern North America the processes of N mineralization, nitrification and soil inorganic N pools, which Duran et al. (2016) attributed primarily to climate change. There has been a slight decline in aboveground biomass in the reference watershed, attributed primarily to soil acidification (Battles et al., 2014). Soil solution and stream water N export has remained at the lowest levels in the 50-year record at this site (notwithstanding several spikes likely due to disturbance events); the mechanism for these low levels of export is still unknown (Yanai et al., 2013; Fuss et al., 2015), but has been attributed to successional processes (Bernal et al., 2012). All of these trends are consistent with what is expected from declining N deposition, yet in each case the authors attribute them primarily to other causes, illustrating the complexity of distinguishing forest responses to N deposition from responses to other concurrent environmental drivers.

In summary, the evidence presented here leads us to hypothesize that as N deposition declines in eastern North America the processes of N mineralization, soil C storage and forest NPP will require decades to respond because they rely on the soil N pool, much of which turns over quite slowly. However, we hypothesize that the soil inorganic N pools and NO\(_3\) leaching will track that decline with a lag of a year or two, although these pools will remain somewhat elevated for decades as N is mineralized from the soil N pool.

Observing this sort of long-term response in forests will be difficult because other factors that influence forest ecosystem processes are changing simultaneously. The C and N cycling of forests in the Northeast is affected by ongoing climate change and increases in atmospheric CO\(_2\) as well as many types of disturbance, including wind storms, ice storms, and attacks by insects and diseases, all of which can recur on time scales shorter than the

Fig. 6. Conceptual model of the fates of N added to an ecosystem. N can simultaneously move to two internal sinks (plant biomass and detritus/SOM) and two loss mechanisms (leaching and gaseous loss). The fate of the added N determines the response of the ecosystems (boxes below the horizontal line). Reduction in N deposition should cause reduced losses and reduced rate of accumulation in the sinks. Because there are multiple fates of N, there can be multiple effects, potentially offsetting one another, such as increased productivity from N fertilization and decreased productivity from soil acidification. Diagram from Lovett and Goodale (2011).
Fig. 7. Simulation of hysteresis in responses of net primary productivity (a) and N leaching (b) to changing N deposition. This simulation uses the Spe-CN model (Crowley and Lovett, 2017). Simulation is for a mixed forest of Acer saccharum, Betula alleghaniensis and Fagus grandifolia, harvested in 1915 and allowed to regrow as a second-growth forest. N deposition increases from 0.2 g N m⁻² yr⁻¹ in 1940 to 1.1 in 1990 and then decreases to 0.2 in 2040 (50 years of increase followed by 50 years of decrease). Response variables are plotted from 1940 to 2040, and the arrows indicate the direction of the temporal sequence. The forest ages from 25 to 125 years old during this simulation, and climate is assumed not to change.

recovery from N deposition. Thus, the signal of a long-term N deposition response will need to be observed through the noise resulting from multiple disturbances. Further, acid deposition is declining concurrently with N deposition, and the de-acidification of the forest can have profound effects on the N cycle (Rosi-Marshall et al., 2016).

6. Surface water chemistry

Also related to effects of de-acidification and decreased atmospheric N loading on N cycling are the direct effects of these declines on the chemistry of surface waters. Four forms of N are generally present in surface waters: NO₃, dissolved organic N (DON), NH₄, and particulate N (PN). Particulate N, which is generally associated with organic colloids, is usually a minor component of surface water N in minimally-disturbed catchments, except during high flow, when PN can be dominant (Inamdar et al., 2015). Dissolved organic N, an operational term applied to a complex mix of organic N-bearing constituents, is often the dominant form of dissolved N in many surface waters (Berman and Bronk, 2003). Although DON is infrequently measured in surface water monitoring programs, DON loads increase with increasing atmospheric N deposition (Brookshire et al., 2007), indicating potential importance as a response indicator. Transformations among DON and dissolved inorganic N (DIN) are observed, with DON serving as both nutrient and energy source in aquatic ecosystems (Wymore et al., 2015). In this section we will largely focus on DIN, of which NO₃ is typically the dominant form and the most frequently studied and modeled.

In catchments of the northeastern U.S. and southeastern Canada minimally affected by direct human land use, atmospheric N deposition is generally the dominant source of N to watersheds. Two other potential sources—bedrock and N-fixation—may be important in some settings, such as catchments underlain by metasedimentary bedrock and those with significant N-fixing species often associated with wetlands (Willard et al., 2005; Morford et al., 2011).

Elevated NO₃ leaching from forested catchments in the eastern United States and elsewhere is usually only apparent above an atmospheric N deposition threshold of approximately 5 kg N ha⁻¹ yr⁻¹ (Aber et al., 2003; Watmough et al., 2005). Above this threshold, which encompasses much of the eastern U.S., there is considerable variability in NO₃ export such that adjacent catchments receiving the same N deposition can exhibit up to a ten-fold difference in stream NO₃ export (Aber et al., 2003). There are numerous factors that contribute to this variation in NO₃ leaching such as those that affect NO₃ production and consumption in forest soils and NO₃ transport from soils to surface waters. The balance between NO₃ production and consumption is primarily influenced by stand age, tree species composition, land use history, and soil physical and chemical characteristics (Lovett and Mitchell, 2004; Lovett et al., 2010; Aber et al., 2003; Lovett et al., 2002). Hence, factors that may lead to changes in tree species composition, e.g., pest or pathogen outbreaks, forest management (silviculture), or climate change, can potentially lead to changes in stream NO₃ export irrespective of changes in atmospheric N deposition (Crowley and Lovett, 2017).

Factors that alter soil nitrification and plant uptake can greatly alter NO₃ export, leading to considerable spatial variability. Forest disturbance, including timber harvesting, ice storm damage, soil freezing, and pest or pathogen outbreaks, can lead to differences in NO₃ export among forested catchments (Aber et al., 2002; Burns and Murdoch, 2005; Eshleman et al., 1998; Nielsen et al., 2001). Increased NO₃ export following disturbance events is attributable to a combination of reduced plant uptake as well as changes in soil properties (moisture, temperature, fine root death) and these impacts can persist for many years (Compton and Boone, 2000; Goodale et al., 2000).

Spatial variation in NO₃ export within and among forested catchments may also originate from a variety of other factors including the mix of tree species, differences in flow paths, and the presence of a riparian zone (Burns et al., 1998; Creed and Band, 1998; Lovett et al., 2002). Given the extremely large inherent spatial variability in stream NO₃ export among forested catchments it is not surprising that the response of surface waters to reductions in atmospheric NO₃ deposition has been mixed (Argerich et al., 2013). In some cases, a clear response to reductions in atmospheric NO₃ deposition has been observed (Kothawala et al., 2011; Eshleman et al., 2013; Driscoll et al., 2016), whereas in some cases, no response is evident (Lawrence et al., 2011; McHale et al., 2017) (Fig. 8a and b). Kothawala et al. (2011) reported significant decreases in stream NO₃ export in response to decreases in N deposition, but only in streams draining catchments with a low wetland fraction. The explanation is that NO₃ accumulated in the winter
snowpack was more readily flushed in the spring in catchments with small riparian zones and hence there was a stronger connection to the decline in atmospheric NO$_3^-$ deposition. This strong connection however can be masked by climate change that will likely alter snowpack dynamics and rain-on-snow events that have been shown to increase NO$_3^-$ export (Casson et al., 2014; Crossman et al., 2016). However, as the climate warms further and snowpack depth and duration decrease there will likely be less opportunity for rain-on-snow events and the spring NO$_3^-$ flush may diminish leading to a decline in NO$_3^-$ export (Crossman et al., 2016).

Other factors such as temporal changes in air temperature and hydrology, and the response to anthropogenic (e.g., forest harvesting) and natural disturbances (e.g., invasive insects) are believed to affect the long-term trajectory of surface water NO$_3^-$ export and to cause an apparent decoupling of the relationship with declining atmospheric N deposition (Argerich et al., 2013). Soil freezing is another contributor to the variability in NO$_3^-$ export from forested catchments (Mitchell et al., 1996; Campbell et al., 2014), which has serious implications in light of ongoing climate change. Snow manipulation studies have suggested that removal of snow can lead to greater soil freezing and more root damage, which can lead to enhanced NO$_3^-$ export in the spring (Campbell et al., 2014). Beyond the influence of changing snow cover and winter dynamics, the response of stream chemistry to decreases in N deposition is complex.

Most studies suggest that N will continue to accumulate in forests in eastern North America even with reductions in atmospheric N deposition (Crossman et al., 2016) and ultimately factors that alter the relative rates of soil NO$_3^-$ production and uptake (e.g., tree species change, climate, soil OM) and transport (altered riparian zone, hydrology) will have considerable influence on future stream NO$_3^-$ export. Hence, as much as stream NO$_3^-$ concentrations have varied in response to increasing N deposition, we expect the response of stream chemistry to decreasing N to be just as variable, with only some sites showing clear relationships with atmospheric N deposition.

7. Synthesis and conclusions: forest response to decreased N deposition in a time of rapid environmental change

The components of forest ecosystem structure and function discussed herein recall the classic quote from Frank Egler (Egler, 1977):

“Ecosystems are not only more complex than we think, but more complex than we can think.”

Furthermore, such discussion exemplifies the words of Frank Golley that an ecosystem is “... more than the sum of its parts” (Golley, 1996). Although we have addressed several aspects of N-impacted forests of the eastern U.S. separately, it is clear that their responses to decreases in atmospheric deposition of N are highly intermeshed, highlighting the complexity inherent in such forest ecosystems.

As N deposition has declined, ecologists are challenged to understand how N limitation and N excess can exist simultaneously, and how forest ecosystems respond to decreases in available N while still experiencing legacy effects of excess N deposition. A case in point is the Hubbard Brook Experimental Forest, which is undergoing what Groffman et al. (2018) describe as “N oligotrophication.” They suggest that C-based reductions in soil N availability could limit forest productivity, a response they see as further enhanced by climate change, including enhanced assimilation caused by increasing atmospheric CO$_2$ and changes in phenology and length of growing season. Ultimately, they conclude, as we do here, with an urgent call for further research on N dynamics, emphasizing the need to re-evaluate the nature and extent of N cycling in temperate forests. This will lead to a better understanding of how oligotrophication will influence forest ecosystem response in the context of global change (Groffman et al., 2018).

We reach several overarching conclusions from our examination of processes and review of the literature on responses to declining N deposition:

1) Nearly all the processes and conditions that we examined, including soil acidification, microbial populations and processes, C and N cycling, NPP, plant species composition, and stream chemistry, can potentially respond strongly to reductions in N deposition. Similar to the period of increasing atmospheric N deposition, the responses will likely vary considerably from site to site due to variation in forest age, soil type, N status, canopy and understory species composition, and hydrology.
2) Most responses are expected to show some degree of hysteresis, with the greatest delays in response occurring in the processes most tightly linked to the “slowly changing pools” of N in wood and soil organic matter.

3) Experimental studies of declines in N loads in forests of North America are lacking, so it is necessary to infer responses from studies in Europe (where deposition levels have been higher) or from studies in which N loads were increased experimentally (e.g., Oulehle et al., 2011). Because of the expected hysteresis, it is difficult to generalize from experimental increases to patterns expected from declining N deposition. Good data are available from North America tracking stream chemistry as N deposition has declined, and recent studies have used retrospective analysis of tree rings to better understand tree physiological responses.

4) The response to declining N deposition will most certainly be a long-term phenomenon, and will be difficult to distinguish from other, concurrent environmental changes, including elevated atmospheric CO2, climate change, reductions in acidity, invasions of new species, and long-term vegetation responses to past disturbance. These other environmental factors will govern in part whether measures of ecosystem responses (e.g. soil exchangeable Ca, stream NO3 concentrations, etc.) will eventually return to an initial condition that existed prior to the advent of anthropogenically-enhanced atmospheric N deposition (Fig. 4), or will reach an alternate stable state, and thus never close the hysteresis loop.

Despite numerous and increasing multi-factorial global change experiments, most understanding of N deposition effects and recovery comes from single-factor fertilization experiments and observational gradient studies. Well-documented single factor responses need to be supplemented with multi-factorial observations on how chronic N additions and their cessation interact with other global change stressors. Indeed, the hypothetical nature of much of our discussion illustrates the need for data from impact forests to address the question of recovery from N deposition. These field studies should include both long-term monitoring of ecosystems under declining ambient N deposition, and experimental studies in which current N fertilizations are gradually decreased and ecosystem responses are tracked. Such studies will augment understanding and improve modeling of impacted ecosystems, enhancing evaluation of air pollution policy and prediction of forest responses to future environmental change.

Sir Arthur Tansley famously described ecological succession as “a variable approaching a variable, not a constant” to emphasize the dynamic nature of ecosystem response to an ever-changing environment (Gilliam, 2016). Anthropogenic climate change has created an unprecedented degree of uncertainty in how these forests will respond to future environmental change, including the current trends of decreasing inputs of N from atmospheric deposition. What is certain is that the ambient CO2 and temperature conditions following any return to pre-impact N status will not be the same as they were prior to historical increases in N deposition.

Acknowledgements

Support for this analysis was provided by the National Science Foundation through the Hubbard Brook and Harvard Forest Long-Term Ecological Research Programs (grants DEB-1637685 and DEB-1237491), the U.S. Geological Survey National Water Quality Program, and from the New York State Energy Research and Development Environmental Monitoring, Evaluation and Protection program. We thank Heather Flaherty for invaluable assistance with graphics. Any use of trade, firm, or product names is for descriptive purposes only and does not imply endorsement by the U.S. Government.

Appendix A. Supplementary data

Supplementary data related to this article can be found at https://doi.org/10.1016/j.envpol.2018.09.135.

References

Aber, J.D., 1992. Nitrogen cycling and nitrogen saturation in temperate forest eco-

Aber, J.D., Goodale, C.L., Ollinger, S.V., Smith, M.L., Magill, A.H., Martin, M.E.,

Aber, J.D., Magill, A.H., 2004. Chronic nitrogen additions at the Harvard Forest
196, 1–5.

Aber, J.D., McDowell, W.H., Nadelhoffer, K.J., Magill, A., Benntson, G., Kamakea, M.,
temperate forest ecosystems: hypotheses revisited. Bioscience 48, 921–934.

northern forest ecosystems—hypotheses and implications. Bioscience 39,
379–386.

Aber, J.D., Ollinger, S.V., Driscoll, C.T., Likens, G.E., Holmes, R.T., Freuder, R.J.,
Goodale, C.L., 2002. Inorganic nitrogen losses from a forested ecosystem in
response to physical, chemical, biotic, and climatic perturbations. Ecosystems 5,
648–658.


Pollut. 54, 185–197.

Argerich, A., Johnson, S.L., Sebestyen, S.D., Rhodeas, C.C., Greatheath, E.,
Knoopp, J.D., Adams, M.B., Likens, G.E., Campbell, J.L., McDowell, W.H.,
Scatena, F., 2013. Trends in stream nitrogen concentrations for forested
reference catchments across the USA. Environ. Res. Lett. 9, 014035.

Bailey, S.W., Hornbeek, J.W., Driscoll, C.T., Gaudette, H.E., 1996. Calcium inputs and
transport in a base-poor forest ecosystem as interpreted by Sr isotope. Water

Battles, J.L., Kafye, T.J., Driscoll, C.T., Blum, J.D., Johnson, C.E., 2014. Restoring soil


the forest nitrogen cycle to climate change. Proc. Natl. Acad. Sci. Unit. States Am.
109, 3406–3411.

Bobbink, R., Hicks, K., Galloway, J., Sprunger, T., Alkemade, R., Ashmore, M.,
Bustamante, M., Ciderby, S., Davidson, E., Dentener, F., Emmett, B., Erismann,
J-W., Finn, M., Gilliam, F., Nordin, A., Pardo, L., de Vries, W., 2010. Global
assessment of nitrogen deposition effects on terrestrial plant diversity effects of

and S throughfall deposition and effects on soil solution chemistry in a Scots

decrease in nitrogen input to a Scots pine stand at Ysselsteyn, The Netherlands.

Ecosystem responses to reduced nitrogen and sulfur inputs into 2 coniferous

Bredemeier, M., Blanck, K., Xu, Y.J., Tietema, A., Boxman, A.W., Emmett, B.,

deposition increases organic N loss from temperate forests. Ecosystems 10,
378–386.

and N mineralization in a northern hardwood forest, Catskill Mountains, New
York, USA. Biogeochemistry 72, 123–146.

springs on NO3– concentrations during summer in Catskill Mountain streams.

Campbell, J.L., Reimann, A.B., Templer, P.H., 2014. Soil freezing effects on sources of


2018. Interactions among plants, bacteria, and fungi reduce extracellular
enzyme activities under long-term N fertilization. Global Change Biol. 24,

enzyme shifts explain litter decay responses to simulated nitrogen deposition.


Wason, J.W., Dovciak, M., Beier, C.M., Battles, J.J., 2017. Tree growth is more sensitive than species distributions to recent changes in climate and acidic deposition in the northeastern United States. J. Appl. Ecol. 54, 1648–1657.


