

NADP released *Nitrogen in the Nation's Rain* in the year 2000 as a non-technical summary of our understanding of nitrogen (N) deposition, its consequences and NADP's efforts to measure it. This booklet was designed as an informative source for policy makers, students and others interested in N deposition and its impacts on people and the environment. Since its publication over 15 years ago, our understanding of the role of nitrogen has grown and a new, more comprehensive booklet titled *Nitrogen from the Atmosphere* is now available. We summarize here some of the new information included in this booklet. **Please help yourself to a booklet if you or your colleagues will find it useful.**

SOME HIGHLIGHTS (An Overall Summary):

Nitrogen is a fundamental constituent of life and the most abundant gas (N₂) in the atmosphere. However N₂ gas is generally non reactive (N_{n,r}). It is only when it is converted to a reactive (N_r) form of nitrogen that it becomes available to most organisms. Organisms such as nitrogen fixing bacteria convert N₂ to N_r and in the early 1900's humans learned to convert N₂ to N_r, in the form of NH₃ (the Haber-Bosch process). Thus N-fertilizer (and N based explosives!) became readily available.

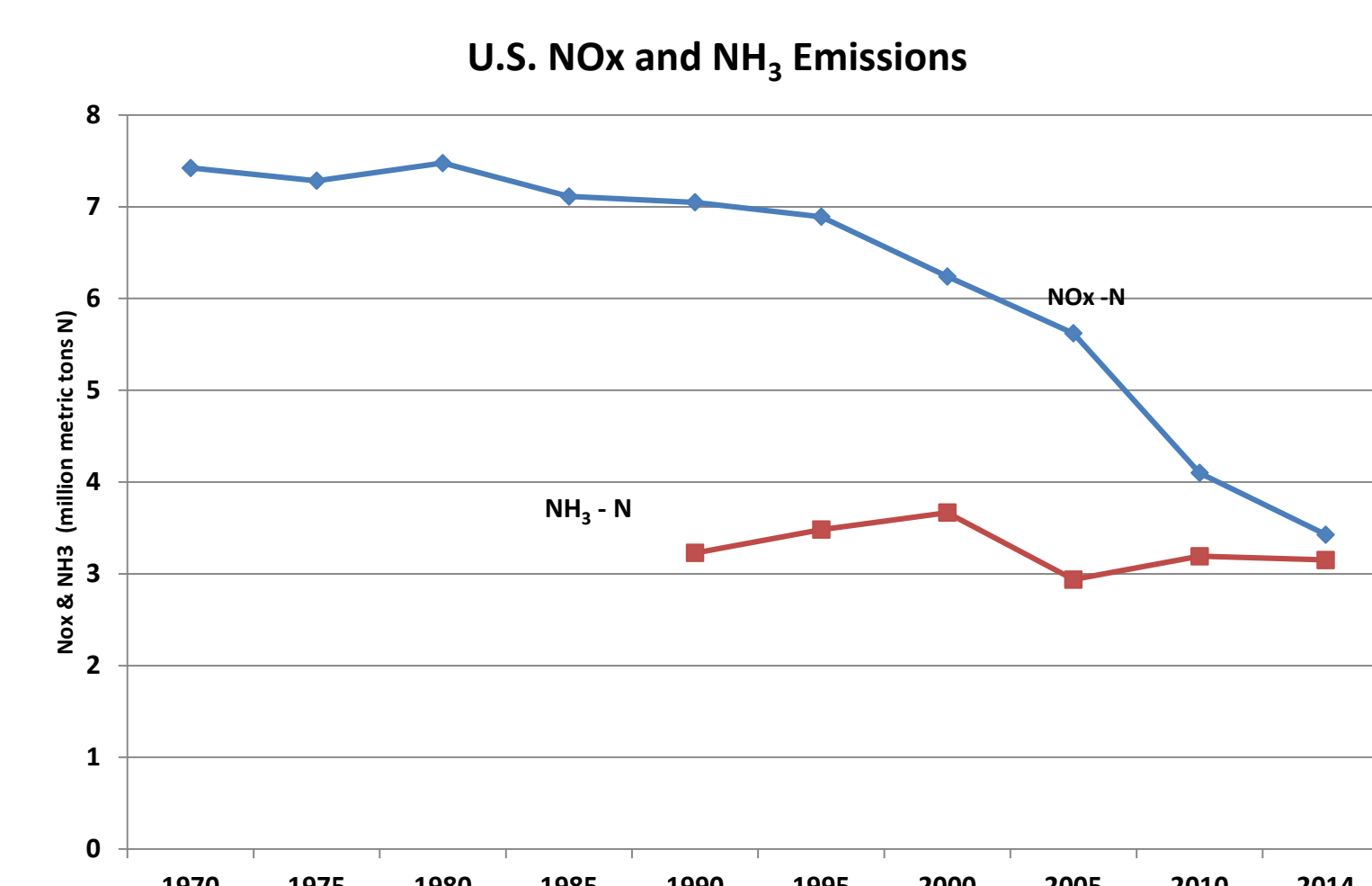
N_r causes a cascade of effects as it moves through ecosystems and forms different N containing compounds in the process.

NADP measures N_r in rain in reduced (ammonium (NH₄⁺)) and oxidized (NO₃⁻ (nitrate)) form. These are the most abundant forms of N_r in precipitation but not the only forms (e.g. organic nitrogen). NADP, since 2007, also measures ammonia gas, NH₃, which is both deposited and emitted in the landscape.

On a global basis we have doubled the N_r circulating in the biosphere and this has both positive and negative consequences. Therefore it is important to measure and monitor trends in N_r derived from the atmosphere. This is an important contribution that NADP participates in.

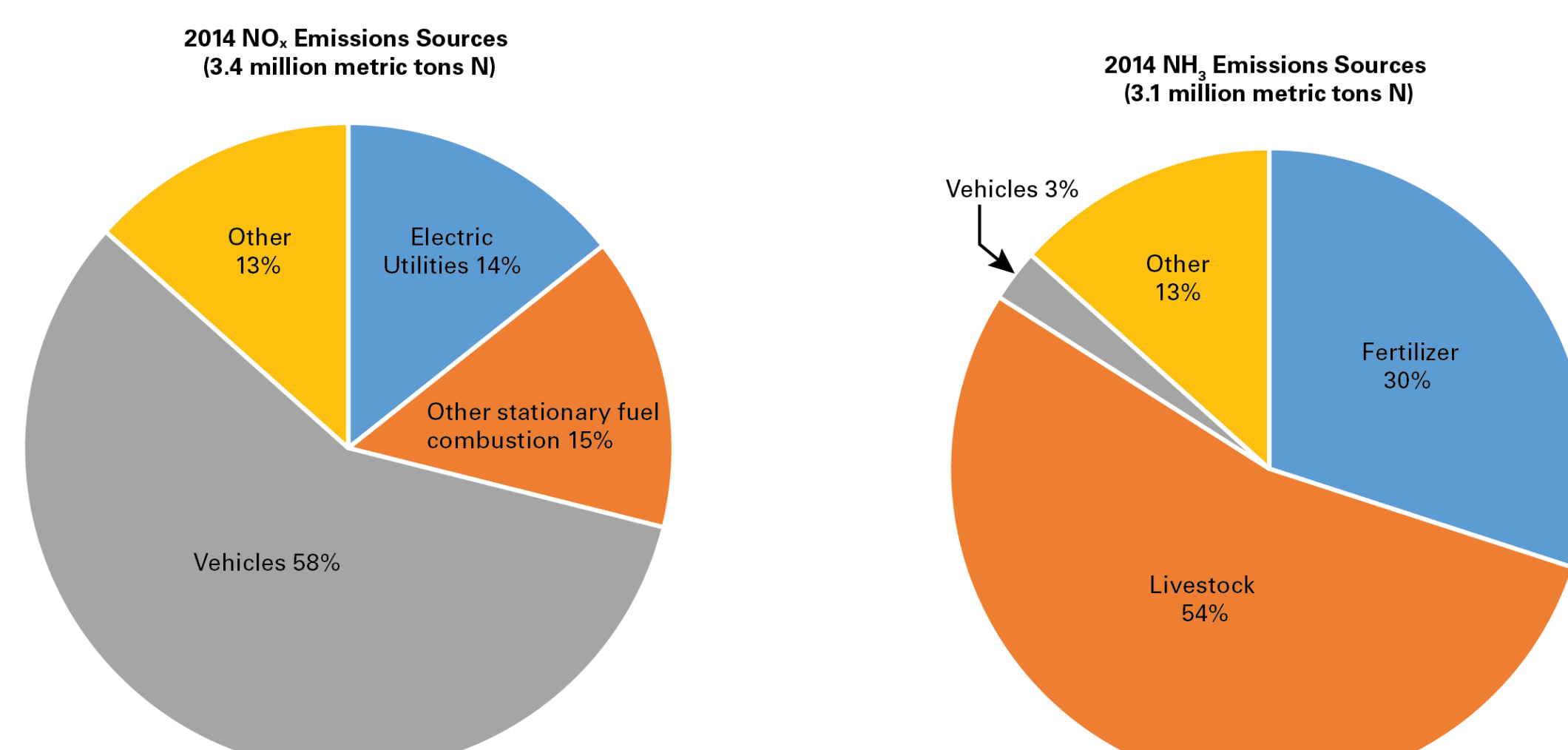
WHICH HUMAN ACTIVITIES CONTRIBUTE NITROGEN?

Reactive N in the atmosphere is derived from NO_x emissions and NH₃ emissions, both from largely different sources. In the past most N_r emissions in the USA were from NO_x, but now nearly equivalent amounts of N_r are derived from NH₃ emissions. In most of the USA

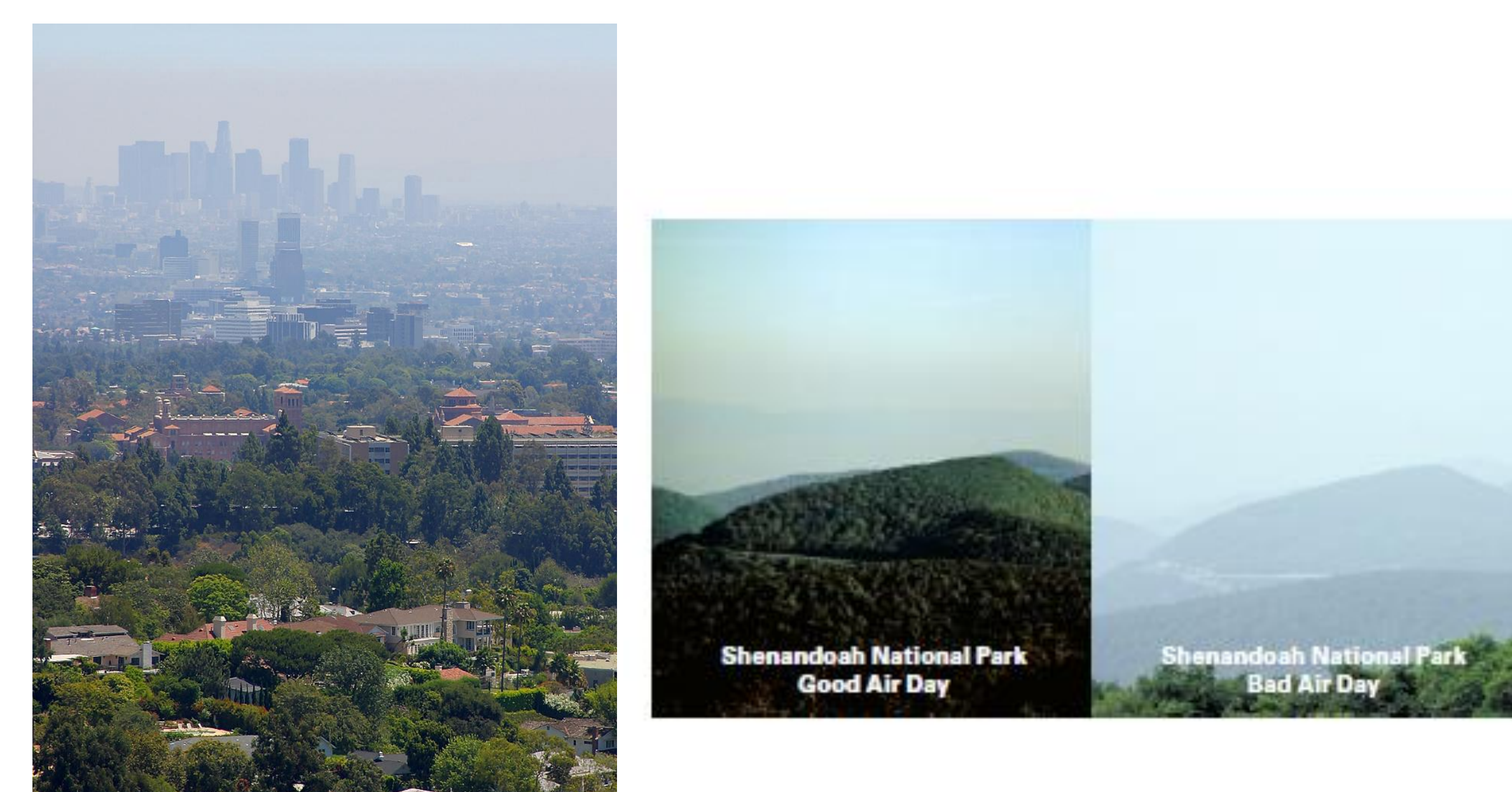


NO_x is derived mainly from fossil fuel combustion. Vehicle emissions are the main contributor (58% of the total). Electric utilities have become relatively less important due to clean air legislation.

Over 80% of NH₃ emissions are from the agricultural sector. Livestock production is the largest source (54%) followed by fertilizer application (30% of total).



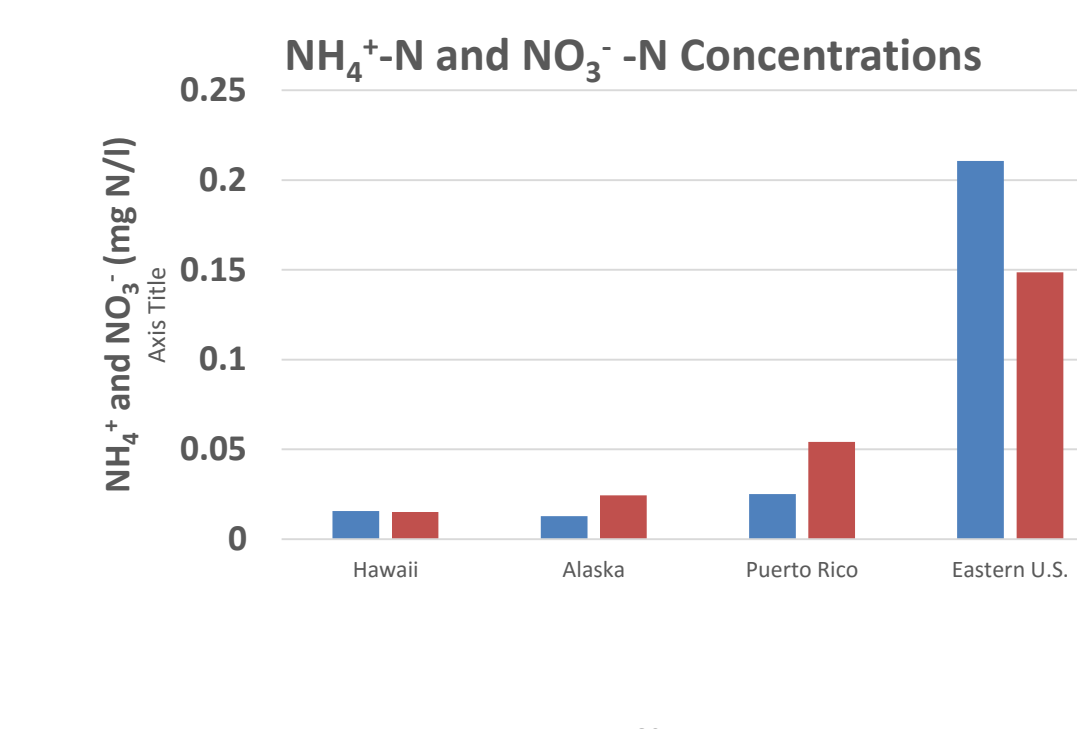
USA 2014 NO_x and NH₃ emissions by source from National Emissions Inventory (NEI).



NO_x emissions (and volatile organic carbon (VOC) emissions) interacting with sunlight, are a major contributor to smog seen here in the Los Angeles Basin (west), and Shenandoah National Park (east).

NADP measurements in precipitation: NO₃⁻ and NH₄⁺

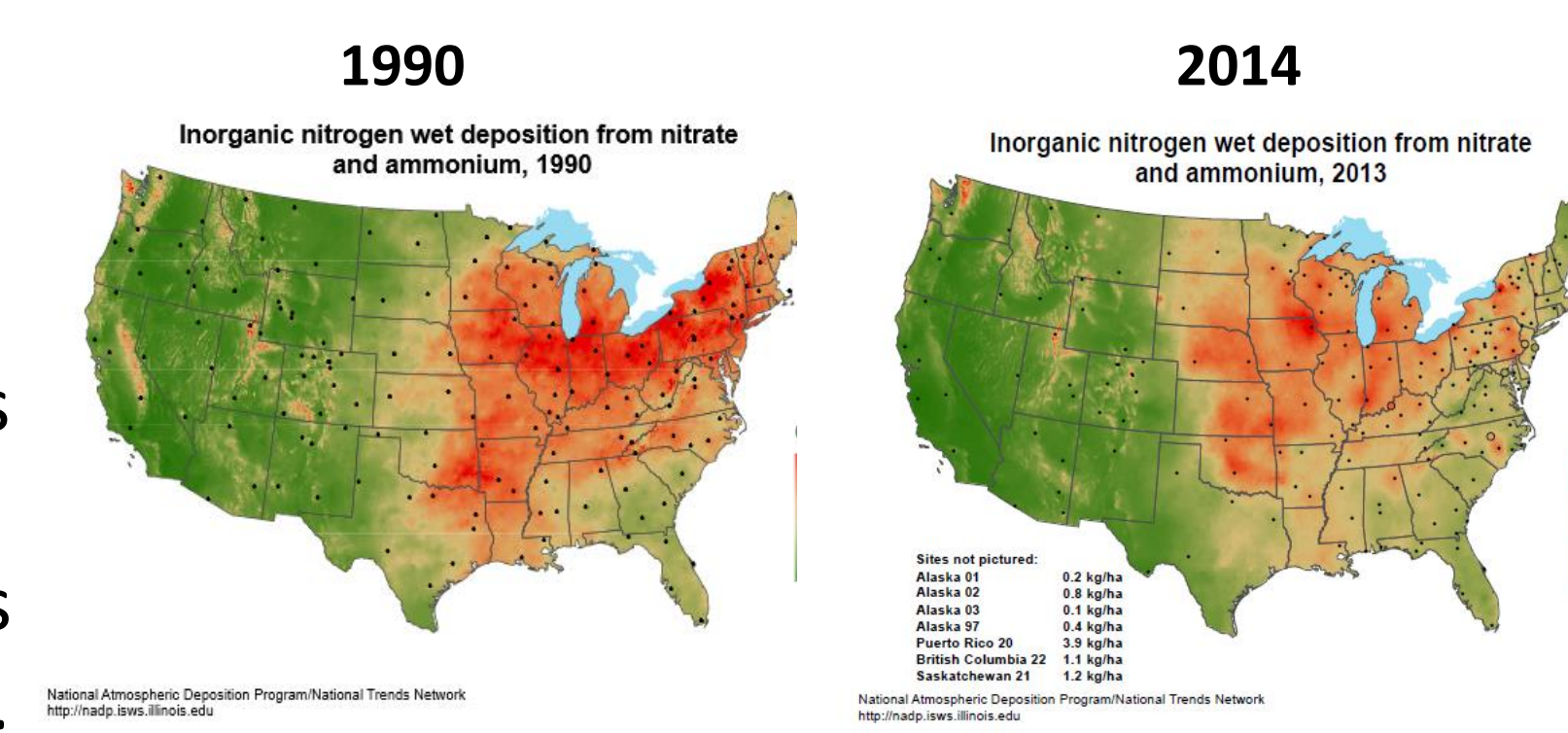
Remote vs populated areas



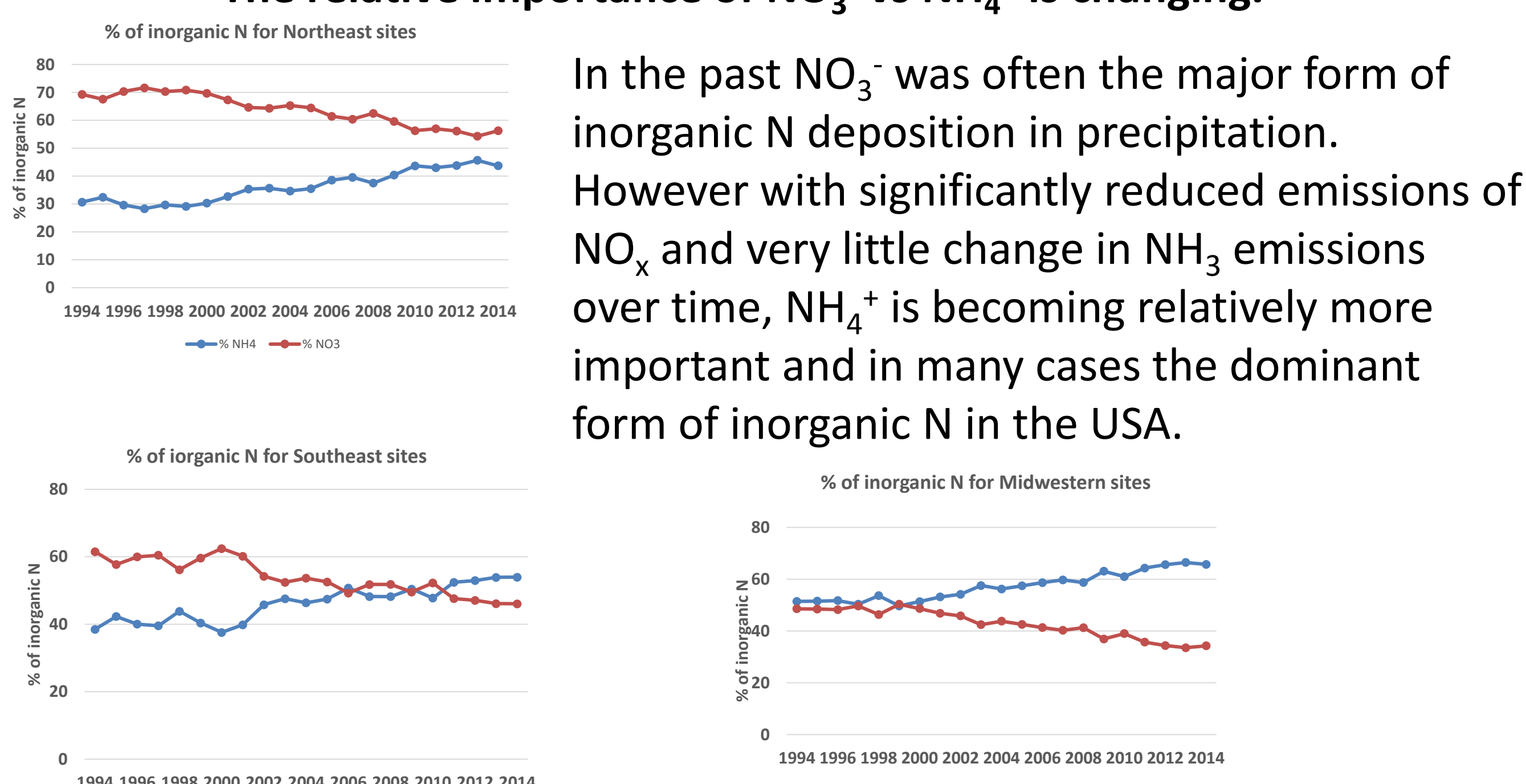
Comparison of NH₄⁺ and NO₃⁻ content (as N) of precipitation in remote (Hawaii, Alaska, Puerto Rico) and more highly populated (eastern USA) areas of the USA. Based on NADP 2012 to 2014 annual volume-weighted mean concentrations (except Hawaii, which is based on 2002 to 2004; site closed in 2005).

Temporal Trends

Inorganic N (NO₃⁻-N + NH₄⁺-N) in precipitation is declining in most areas of the USA as illustrated in these deposition maps comparing 1990 to 2014.



The relative importance of NO₃⁻ vs NH₄⁺ is changing.

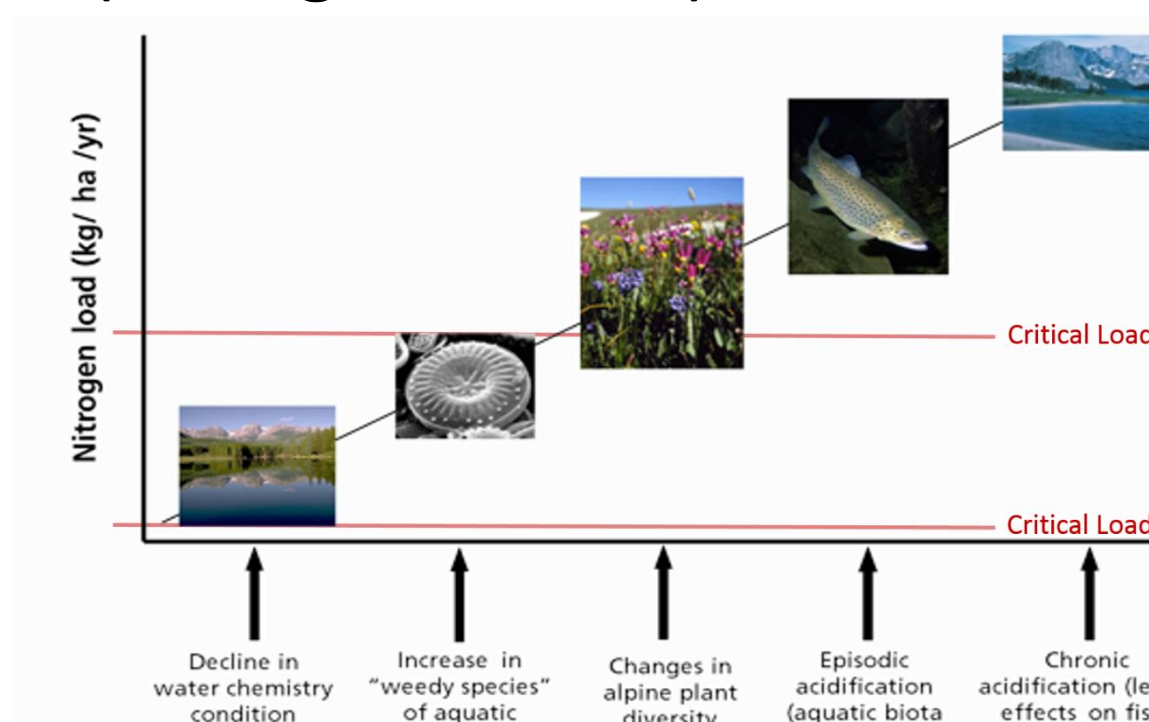


In the past NO₃⁻ was often the major form of inorganic N deposition in precipitation. However with significantly reduced emissions of NO_x and very little change in NH₃ emissions over time, NH₄⁺ is becoming relatively more important and in many cases the dominant form of inorganic N in the USA.

Note: NADP also measures air concentrations of NH₃ at 94 sites as part of an effort to understand deposition of NH₃

CRITICAL LOADS: HOW MUCH N DEPOSITION IS TOO MUCH?

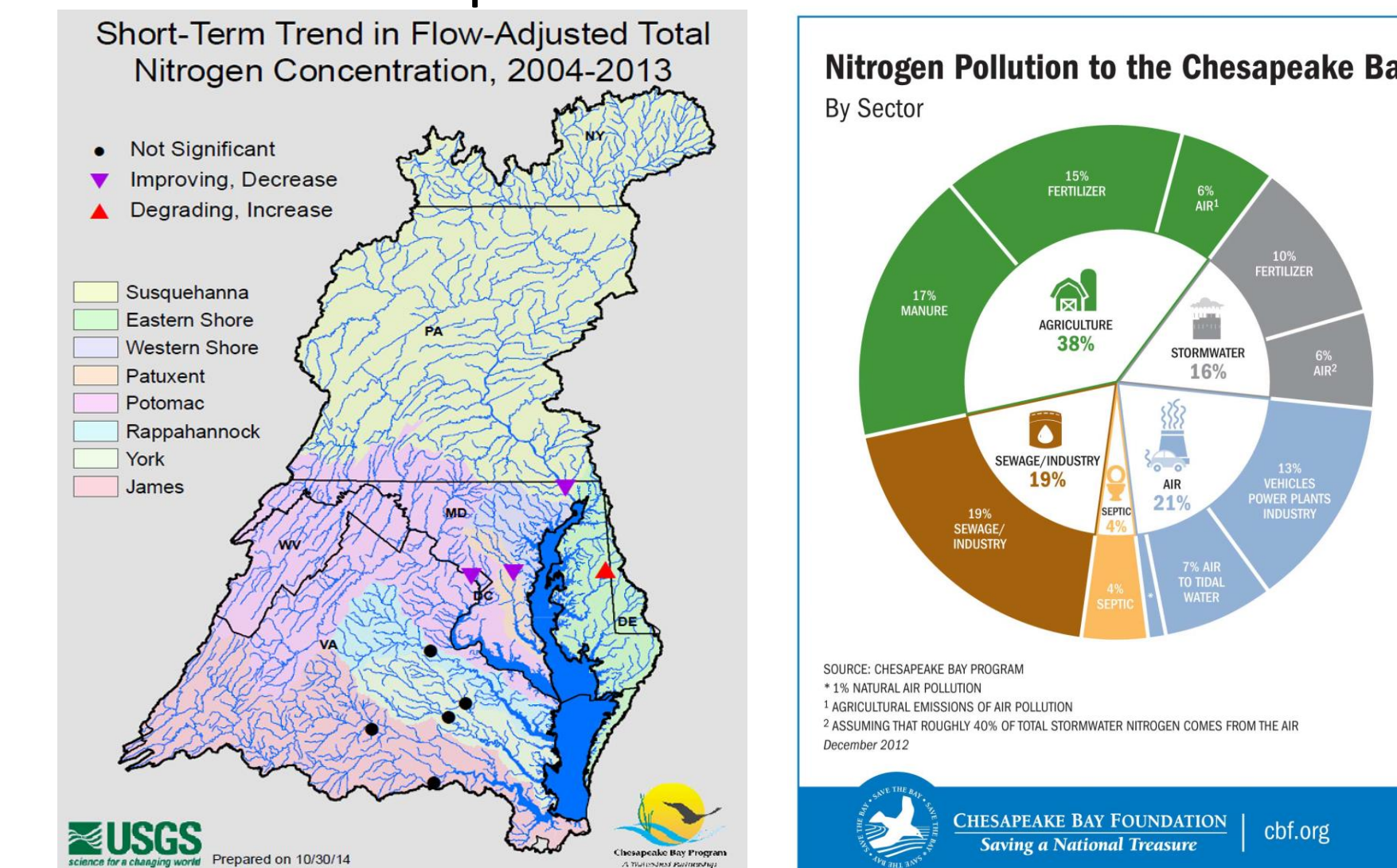
The term critical load is used to describe the threshold of atmospheric deposition that causes harm to sensitive resources in an ecosystem, typically expressed in terms of kilograms per hectare per year (kg/ha/yr) of wet or total (wet + dry) deposition. Critical loads can be developed for a variety of ecosystem responses (see figure below).



When critical loads are exceeded, the environmental effects can extend over great distances. For example, excess nitrogen can change soil and surface water chemistry, which in turn can cause eutrophication of downstream estuaries. The long-term, spatially extensive wet deposition data provided by NADP are instrumental in developing critical loads in the USA, thereby helping to quantify the impacts of air pollution on ecosystems. NADP also provides a format for critical loads development through the **CLAD (Critical Loads and Atmospheric Deposition) Science Committee** (<http://nadp.isws.illinois.edu/committees/clad/>).

THE CHESAPEAKE BAY: A CASE STUDY

The Chesapeake Bay is the largest of 130 estuaries in the nation and is located in coastal Maryland and Virginia. The population within the Chesapeake Bay watershed is approximately 17 million and growing. **One of the largest problems facing the environmental health of the Bay is an overabundance of nutrients, especially N.** This often leads to increased algal production and organic matter, a process known as eutrophication.



Nitrogen is introduced into the Chesapeake Bay as runoff or wastewater from agricultural operations, storm water drainage, sewage and industry, and 1/3 is from deposition from the air.

The Chesapeake Bay Program seeks ways to reduce N entering the Bay, which include programs to promote farming conservation practices to curb drainage and runoff of fertilizers and animal waste. Researchers are using NADP data to compute the amount of N deposited by precipitation in the Bay and its watershed.

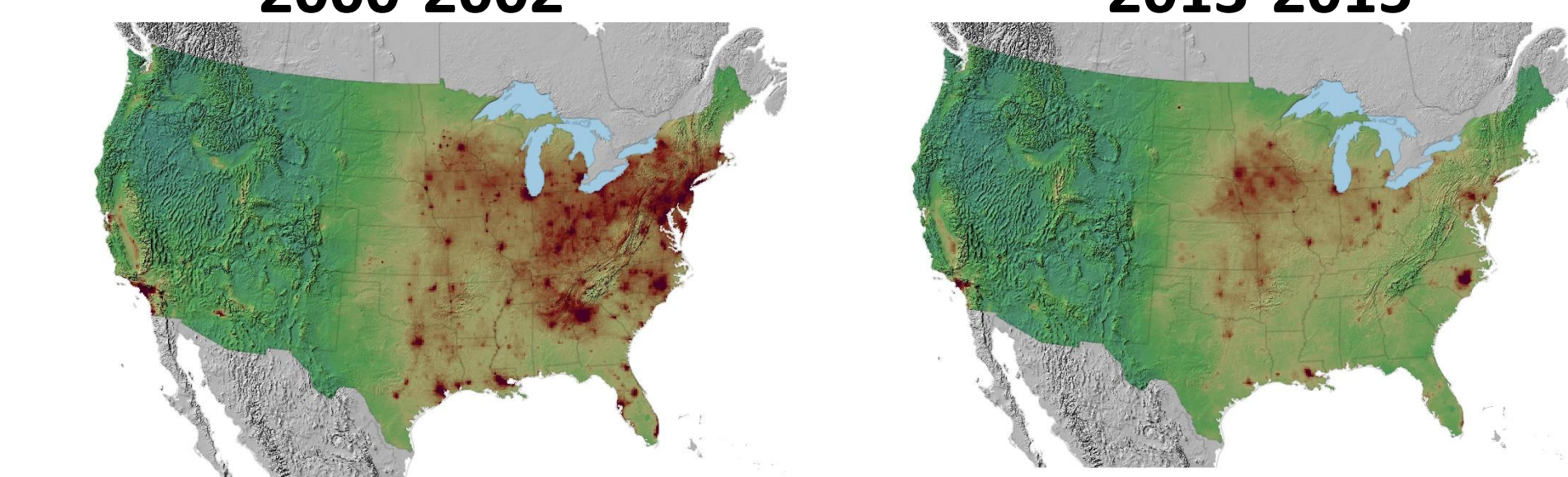


Aerial view of algal blooms in the Elizabeth, Lafayette, and James Rivers, VA. N in the water is causing the explosive growth of algae, which later die, depleting the water of oxygen and leading to hypoxic zones. Source: Chesapeake Bay Foundation.

MEASURING TOTAL DEPOSITION (wet + dry)

NADP has also attempted to better understand total wet and dry nitrogen deposition through the **Total Deposition (TDEP) Science Committee**. Using a "hybrid" approach of combining measured data (e.g., precipitation NO₃⁻ and NH₄⁺, dry deposition of particulate NH₄⁺ and NO₃⁻, and gaseous NH₃) with modeled deposition data of other known, but not measured, nitrogen species (e.g., wet and dry organic N, HONO, NO₂, etc.), estimates of total N deposition are obtained.

Mean annual total N deposition 2000-2002 vs 2013-2015



Such work aids another NADP science committee, the Critical Loads and Atmospheric Deposition (CLAD) Committee in evaluating whether sensitive areas are experiencing too much N deposition.

* The Clean Air Status and Trends Network (**CASTNET**) measures dry species of N (particulate NH₄⁺ & NO₃⁻, as well as HNO₃) and calculates N deposition of these species at 95 sites.

The Nitrogen Cascade

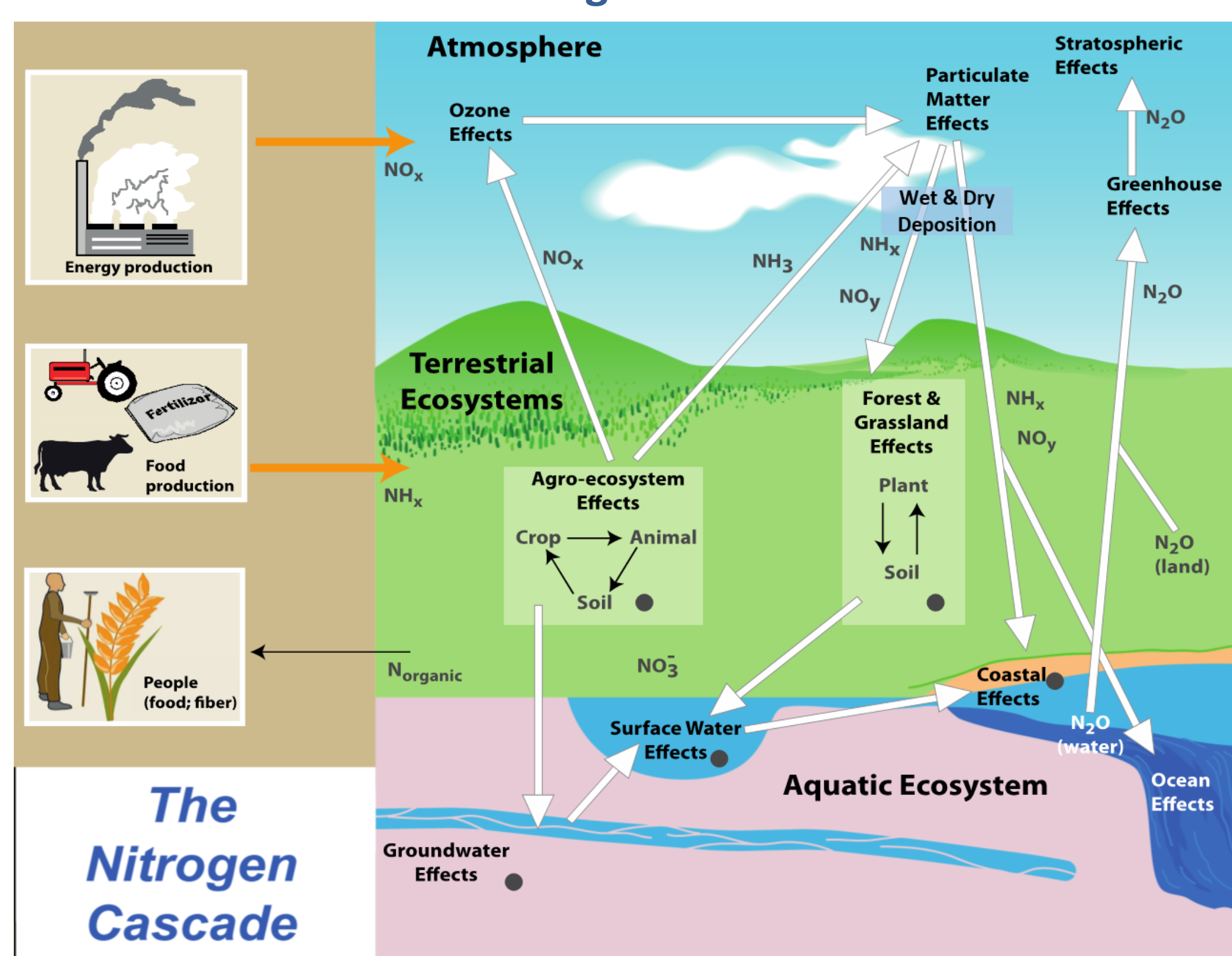


Illustration of the nitrogen cascade showing the sequential effects that a single atom of nitrogen in its various molecular forms can have in various reservoirs after it has been converted from nonreactive N₂ to a reactive form by energy and food production (orange arrows). Once created, the reactive nitrogen has the potential to continue to contribute to impacts until it is converted back to N₂. The small black circle indicates that there is the potential for denitrification to occur within that reservoir. Source: adapted from Ciais et al, 2013 with permission from the GEO Yearbook 2003.