

Pollution of Earth's Troposphere: Acid Rain & Aerosols

AOSC / CHEM 433 & AOSC / CHEM 633

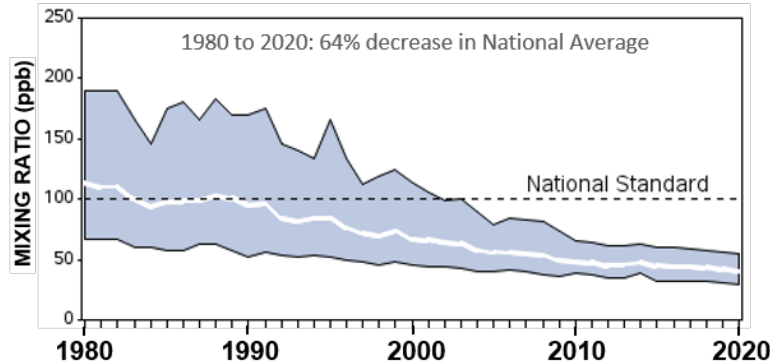
Ross Salawitch

Class Web Sites:

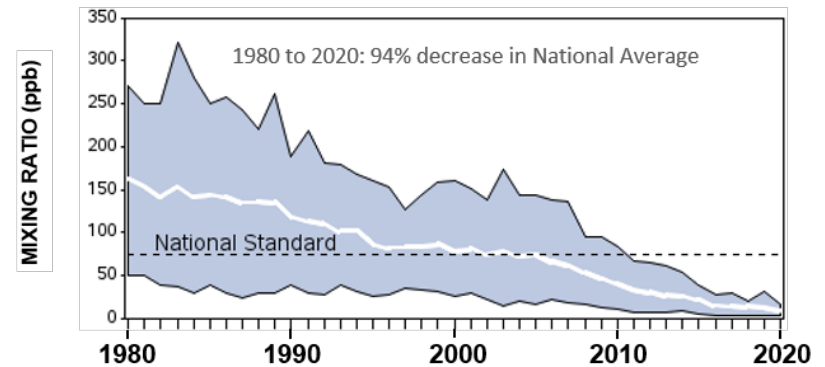
<http://www2.atmos.umd.edu/~rjs/class/spr2022>

<https://myelms.umd.edu/courses/137772>

NO₂ Air Quality, 1980 to 2020
Annual 98th Percentile of Daily Max 1-Hour Average
National Trend based on 20 Sites



SO₂ Air Quality, 1980 to 2020
Annual 99th Percentile of Daily Max 1-Hour Average
National Trend based on 32 Sites



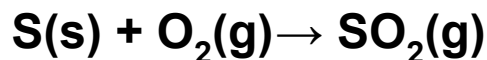
Lecture 14

29 March 2022

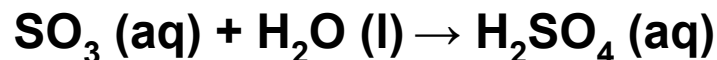
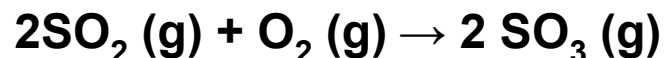
Acid Rain: SO₂

Chemical formula of coal: C₁₃₅H₉₆O₉NS (S varies with coal type)

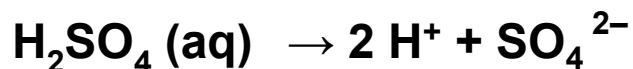
Combustion of leads to release of sulfur dioxide (SO₂)



SO₂ reacts with O₂ to form sulfur trioxide (SO₃)



Followed by:



pH for 200, 400, & 600

My pH is 5.6

My pH is approximately 8.2

Dissolution of these types of rocks tends to maintain the world's oceans in a slightly basic state

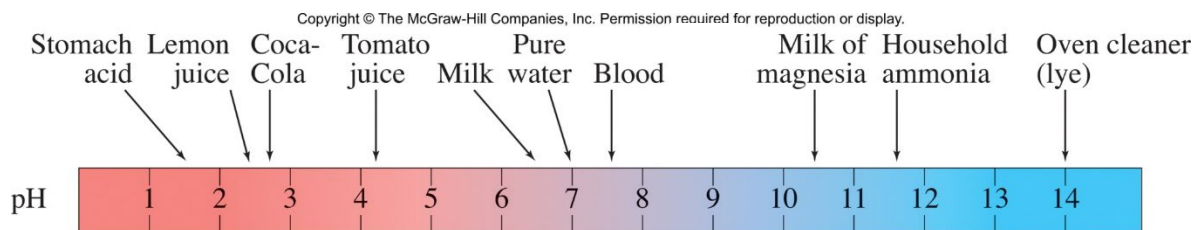


Figure 6.4, Chemistry in Context.

pH for 800 & 1000

Origin of the term pH



Metropolitan area with the most acidic measurement of fog or rain

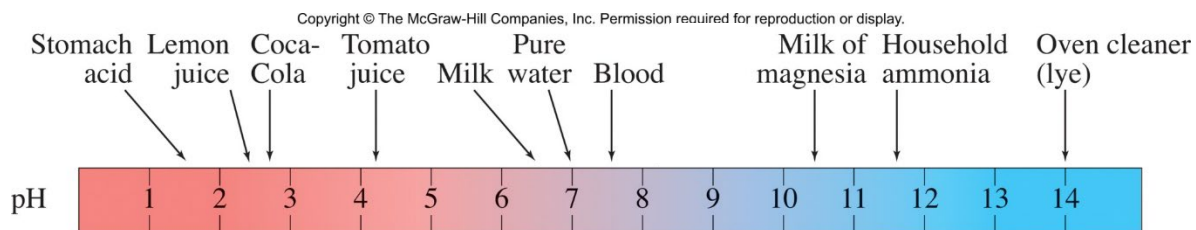
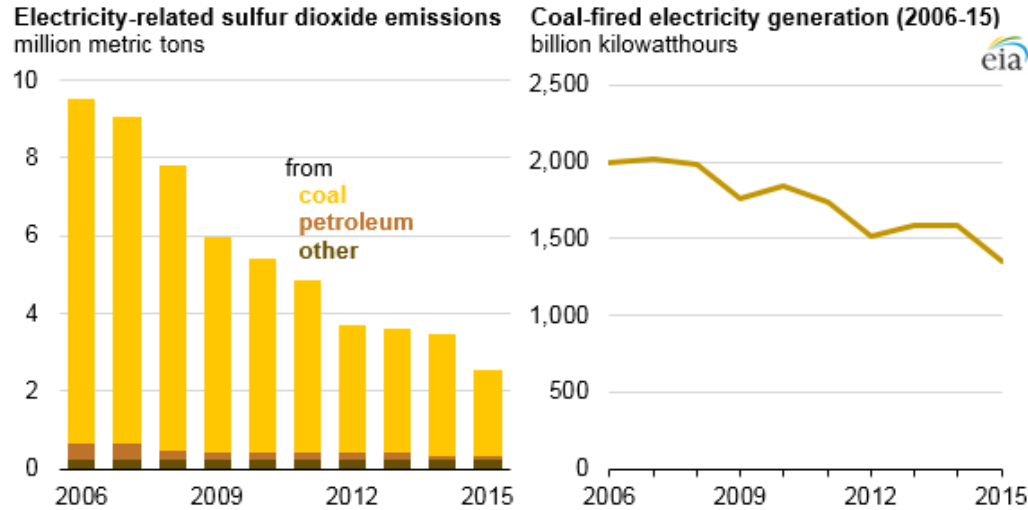


Figure 6.4, Chemistry in Context.

SO₂ Sources (U.S.)



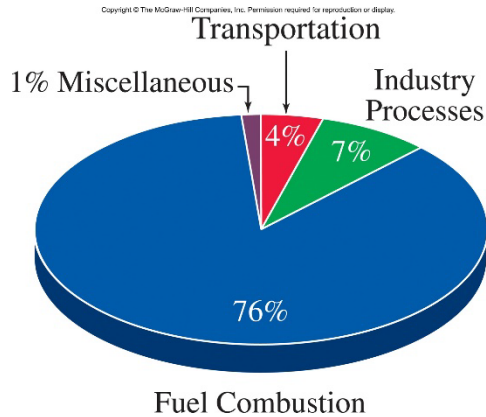
SO₂ emissions from U.S. power plants in the United States declined by 73% from 2006 to 2015, a much larger reduction than the 32% decrease in coal-fired electricity generation over that period.

From 2014 to 2015, SO₂ emissions fell 26%—the largest annual percentage drop in the previous decade.

Nearly all electricity-related SO₂ emissions are associated with coal-fired generation.

<https://www.eia.gov/todayinenergy/detail.php?id=29812>

SO₂ Sources (U.S.)



Primary source of SO₂ is fuel combustion; emissions from this sector are decreasing.

Emissions from transportation are small and largely unchanged.

Figure 6.14, Chemistry in Context.
U.S. SO₂ emission sources, year 2007

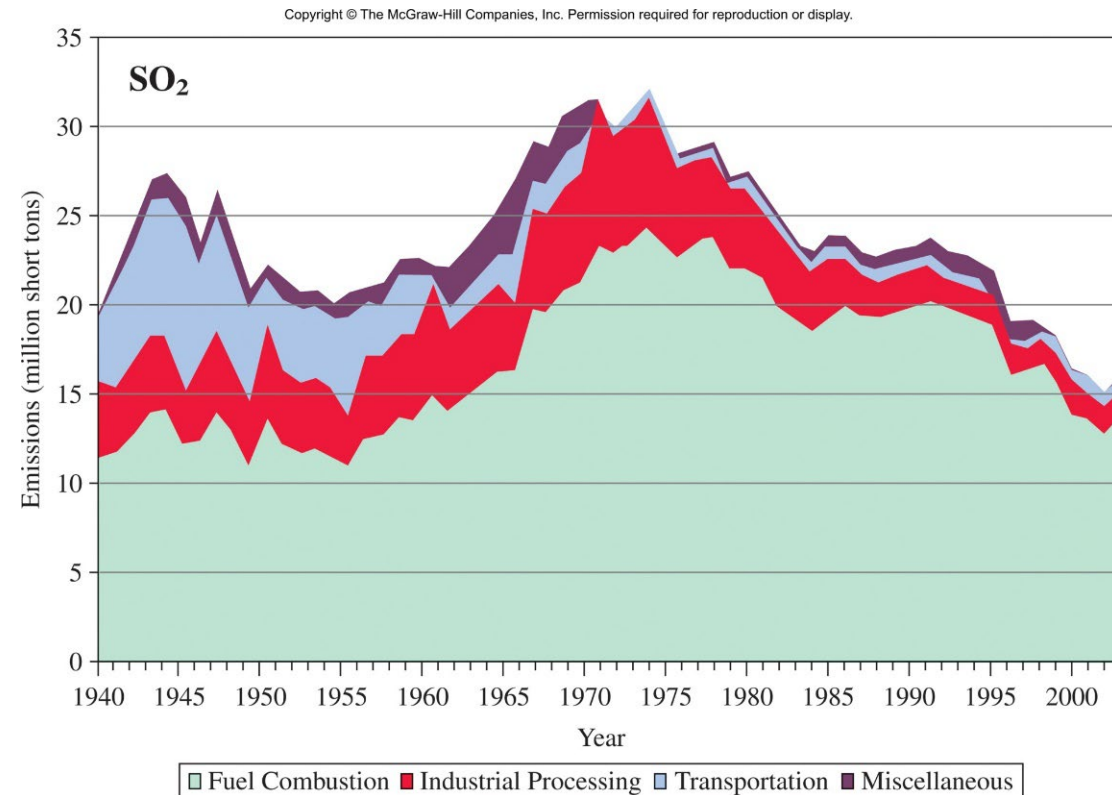
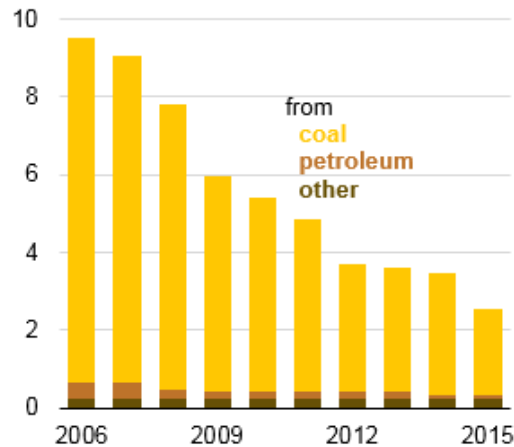


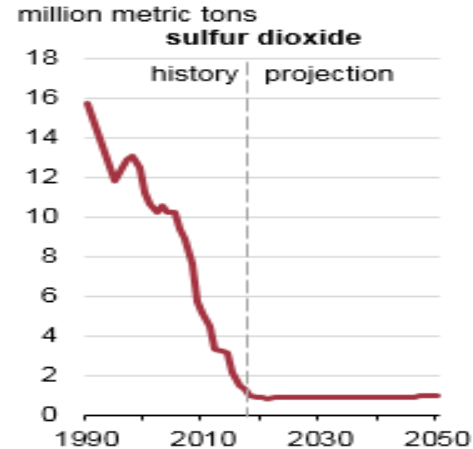
Figure 6.21, Chemistry in Context.
U.S. SO₂ emissions, 1940 to 2003

SO₂ Sources (U.S.)

Electricity-related sulfur dioxide emissions
million metric tons



U.S. electric power sector SO₂ emission
update & projection



Factors that have contributed to lower SO₂ emissions:

1) Changes in the electricity generation mix

Electricity generation from coal fell 14% from 2014 to 2015; mostly offset by an increase in electricity generation from natural gas. Also, over time, coal plants that emitted especially large amounts of SO₂ were used less often than cleaner coal fired power plants.

2) Legislation

The CAAA (Clean Air Act Amendments of 1990) required several regulations that reduced emissions of SO₂ and NO_x. The Acid Rain Program imposed a cap on emissions of SO₂ and NO_x from coal and residual-fuel oil-fired power plants starting in 1995.

The main compliance approach by electric generators for Mercury and Air Toxics Standards (MATS), with an April 2015 initial deadline, was to install flue-gas desulfurization (scrubber) or dry sorbent injection equipment, both of which also remove SO₂ and NO_x in addition to the targeted air pollutants regulated under MATS.

In 2005, the Clean Air Interstate Rule (CAIR) addressed regional interstate transport of contributors to ground-level ozone (smog) by requiring 27 eastern states to file implementation plans to reduce SO₂ and NO_x emissions. CAIR was replaced by the Cross-State Air Pollution Rule (CSAPR) in 2015.

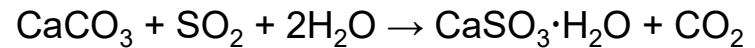
<https://www.eia.gov/todayinenergy/detail.php?id=29812> & <https://www.eia.gov/todayinenergy/detail.php?id=37752>

Removal of SO₂ from Power Plants

SO₂ Control: Flue Gas Desulphurization



Pulverized limestone (CaCO₃) is mixed with water to make a slurry sprayed into flue gas, resulting in:

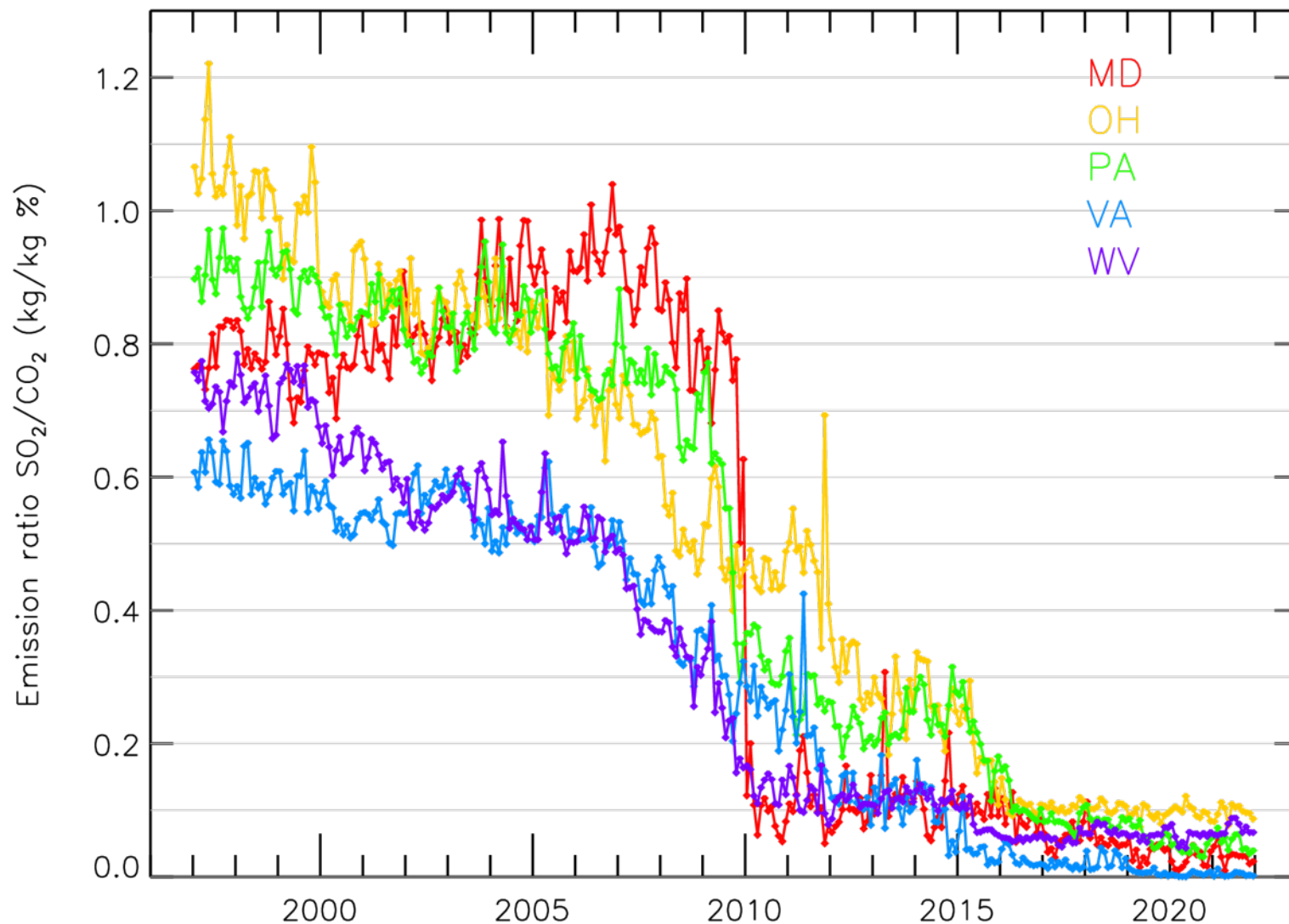


Cost on order \$200 million per unit

Another technology using lime, CaO, exists but is not in widespread use due to high cost of lime

What happens to the CaSO₃·H₂O ?

Trends in power plant emission, region



Thanks to Doyeon Ahn for this wonderful analysis of CEMS (Continuous Emission Monitoring System) Data provided by EPA

Maryland Healthy Air Act

The Maryland Healthy Air Act was developed with the purpose of bringing Maryland into attainment with the National Ambient Air Quality Standards (NAAQS) for ozone and fine particulate matter by the federal deadline of 2010. The act and the subsequent regulations also requires the reduction of **mercury** emissions from coal-fired electric generating units and significantly reduces atmospheric deposition of nitrogen to the Chesapeake Bay and other waters of the State.

The Healthy Air Act is the toughest power plant emission law on the east coast. The HAA requires reductions in nitrogen oxide (**NO_x**), sulfur dioxide (**SO₂**), and **mercury** emissions from large coal burning power plants. The Healthy Air Act also requires that Maryland become involved in the Regional Greenhouse Gas Initiative (RGGI) which is aimed at reducing greenhouse gas emissions.

Which pollutants are covered by this rule and how much pollution will be reduced?

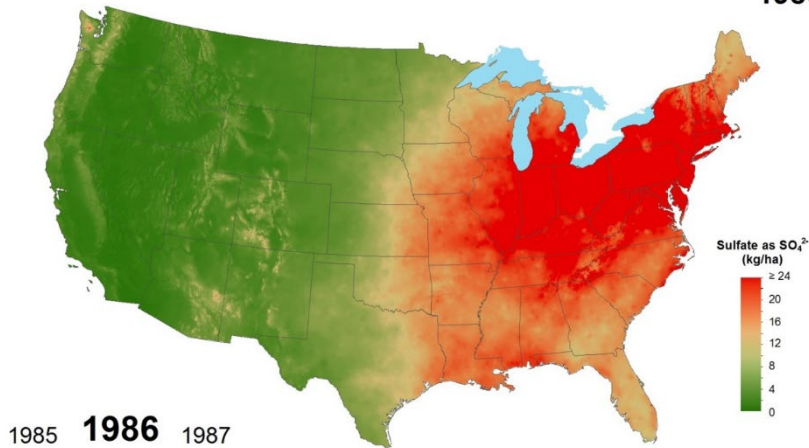
The Healthy Air Act requires year-round emission controls that will significantly reduce nitrogen oxides (**NO_x**), sulfur dioxide (**SO₂**), and **mercury** from power plants located in Maryland. NO_x emissions in Maryland will be reduced almost 70% in 2009. A second phase of NO_x control will reduce emissions by a total of 75% by 2012. SO₂ emissions will be reduced by 80% in 2010 with a second phase of controls in 2013, which will increase the emission reduction to 85%. When the rule is adopted, mercury emissions will be reduced by 80% in 2010. A second phase of controls will reduce mercury emissions by 90% by 2013. All of the above emission reductions are based on a comparison to a 2002 emissions baseline.

http://www.mde.maryland.gov/programs/air/pages/md_haa.aspx

Sulfate Deposition (see Fig 6.12)

1986

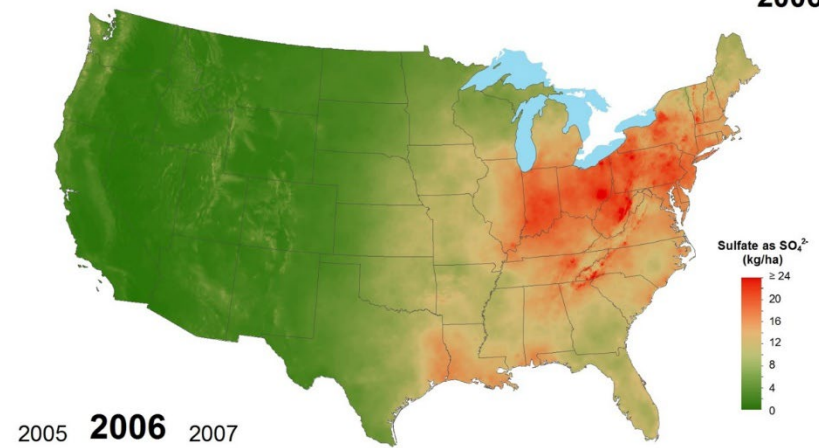
Sulfate ion wet deposition
1986



National Atmospheric Deposition Program/National Trends Network
<http://nadp.isws.illinois.edu>

2006

Sulfate ion wet deposition
2006



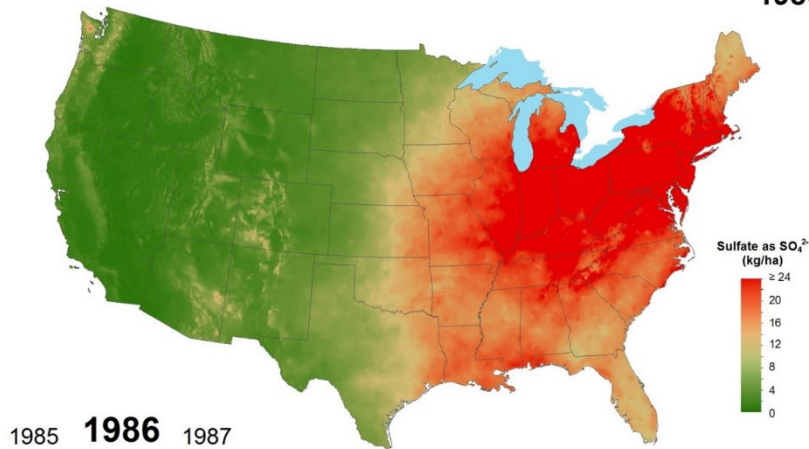
National Atmospheric Deposition Program/National Trends Network
<http://nadp.isws.illinois.edu>

<http://nadp.slh.wisc.edu>

Sulfate Deposition (see Fig 6.12)

1986

Sulfate ion wet deposition
1986



National Atmospheric Deposition Program/National Trends Network
<http://nadp.isws.illinois.edu>

2020

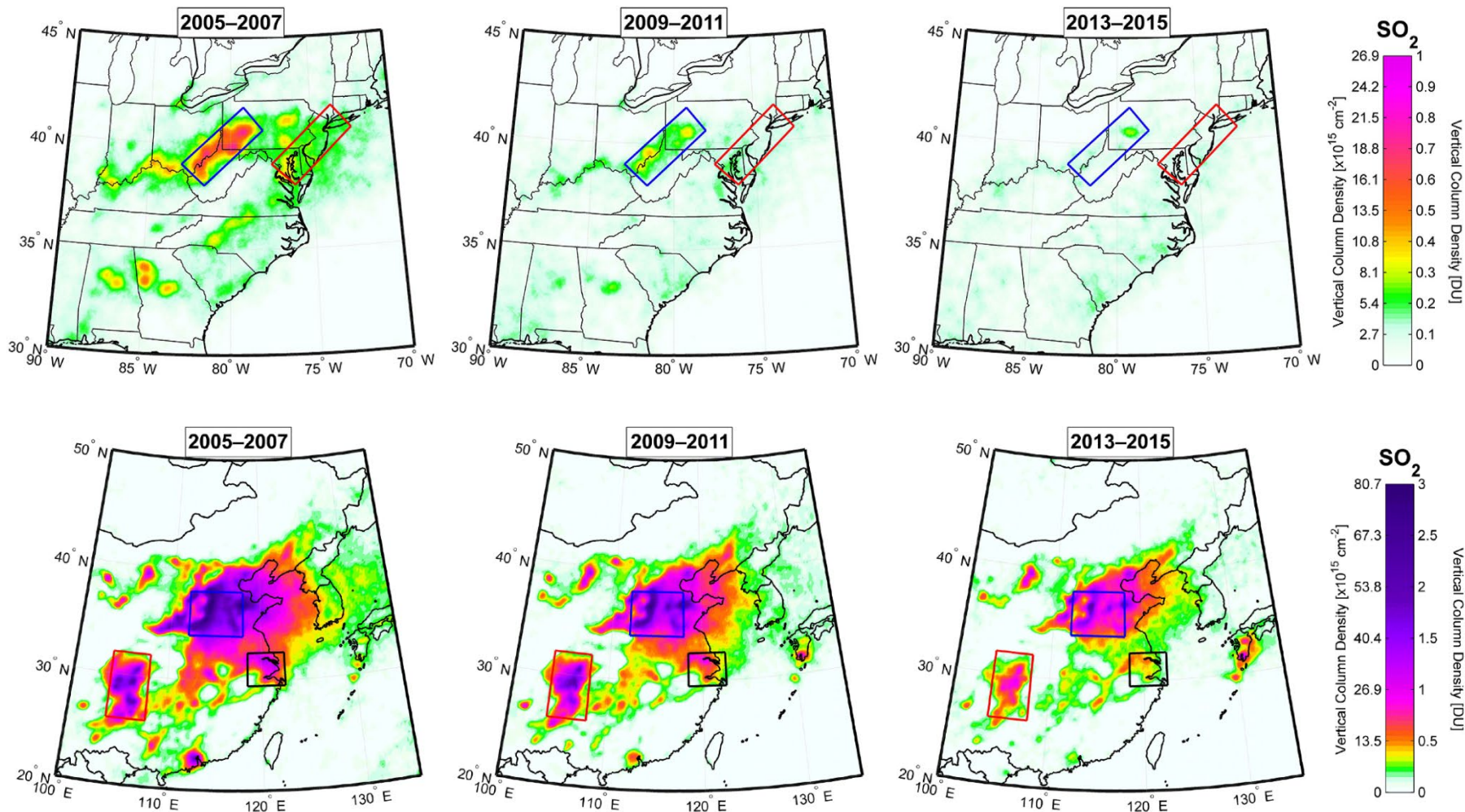
Sulfate ion wet deposition
2020



National Atmospheric Deposition Program/National Trends Network
<http://nadp.slh.wisc.edu>

<http://nadp.slh.wisc.edu>

SO₂ Trends from Space

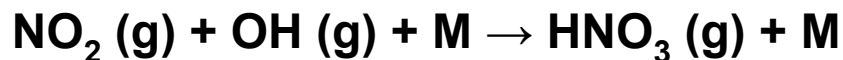


Krotkov *et al.*, ACP, 2016

Acid Rain: NO_x

NO_x plays major role in tropospheric O_3 formation.

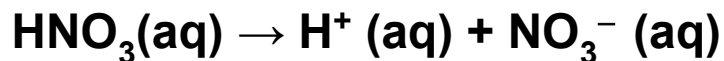
In Lecture 13, we emphasize the critical importance of radical termination:



Nitric acid, HNO_3 , is soluble!

Hence, in the presence of liquid water, $\text{HNO}_3 (\text{g})$, can become $\text{HNO}_3 (\text{aq})$

$\text{HNO}_3 (\text{aq})$ will then dissociate:



and well “oops, we did it again”

NO_x Sources (U.S.)

Primary source of NO₂ is transportation.

The EPA inventory suggests emissions from this sector are holding steady. However, UMD researchers believe mobile NO_x emission in the mid-Atlantic are much lower than estimated by EPA (Anderson *et al.*, Atmos. Envir., 2014)

Figure 6.16, Chemistry in Context.
U.S. NO_x emission sources, year 2007

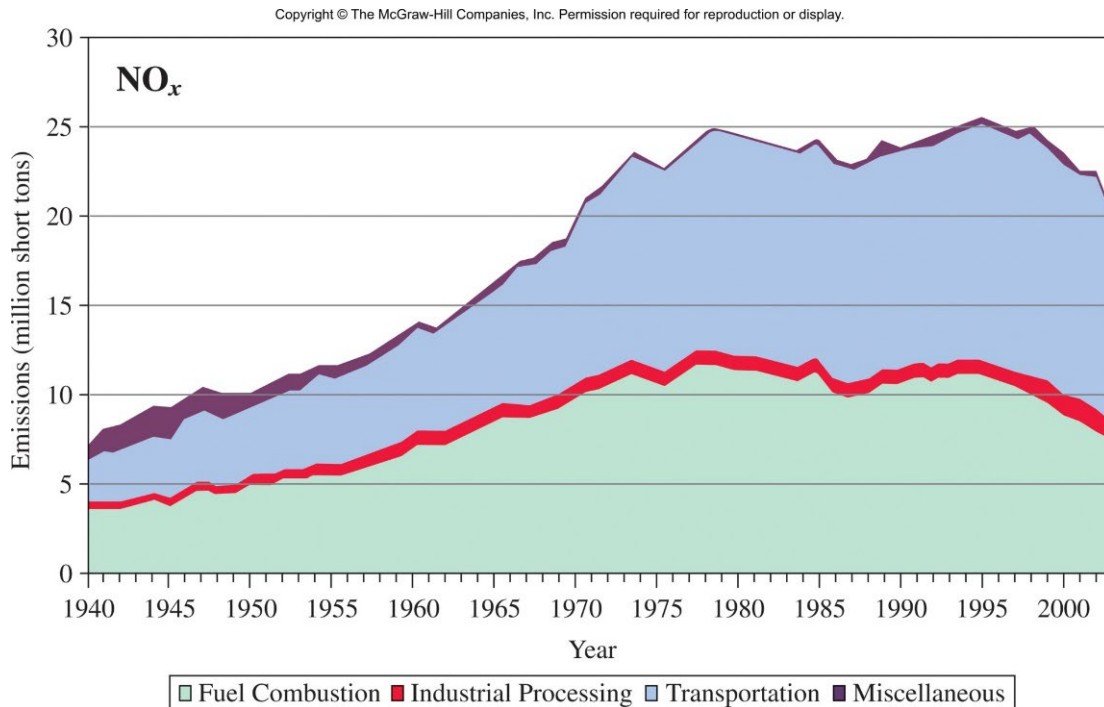
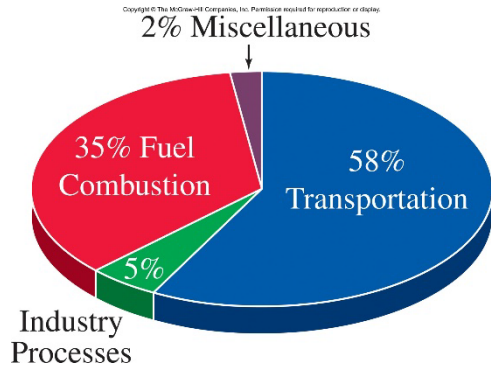
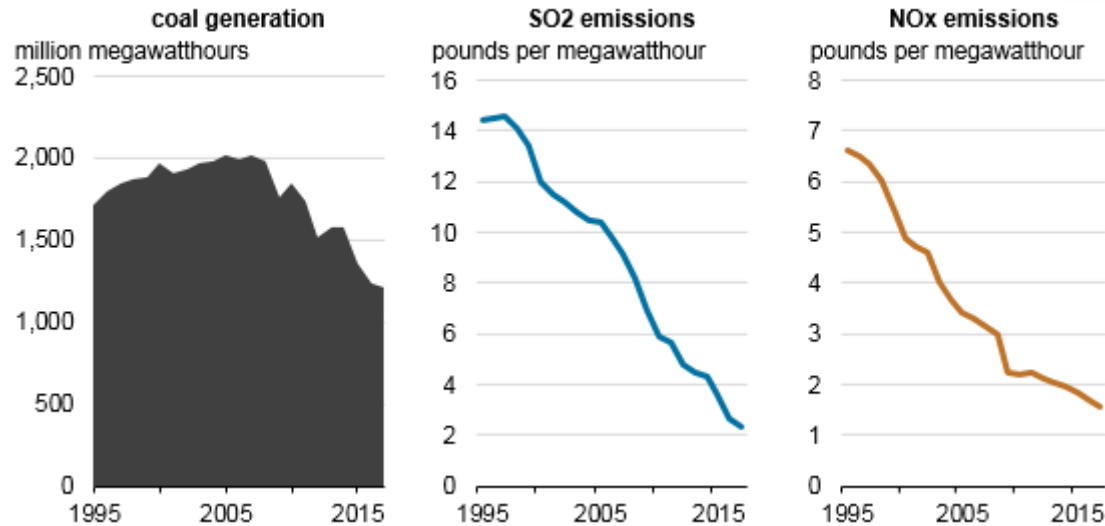


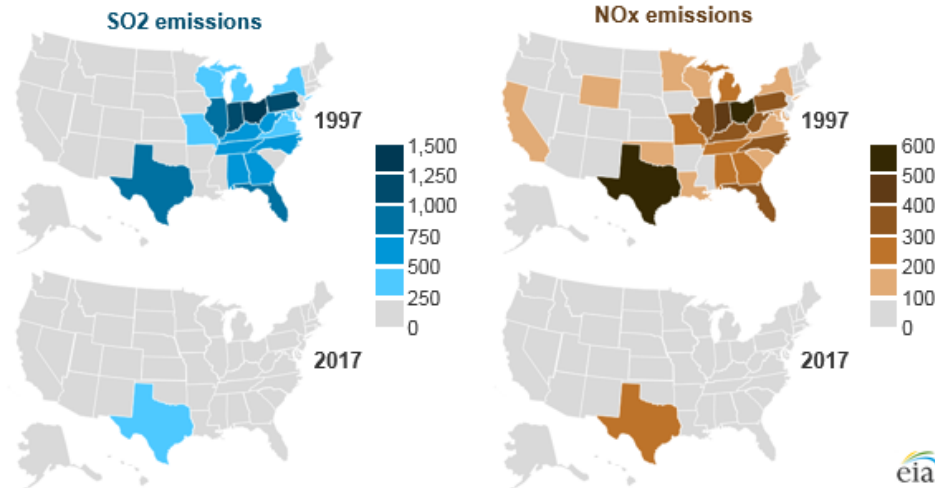
Figure 6.21, Chemistry in Context.
U.S. NO_x emission sources, 1940 to 2003.

Decline of SO₂ and NOx Sources (U.S.)

U.S. coal generation and SO₂ and NOx emission rates (1995-2017)

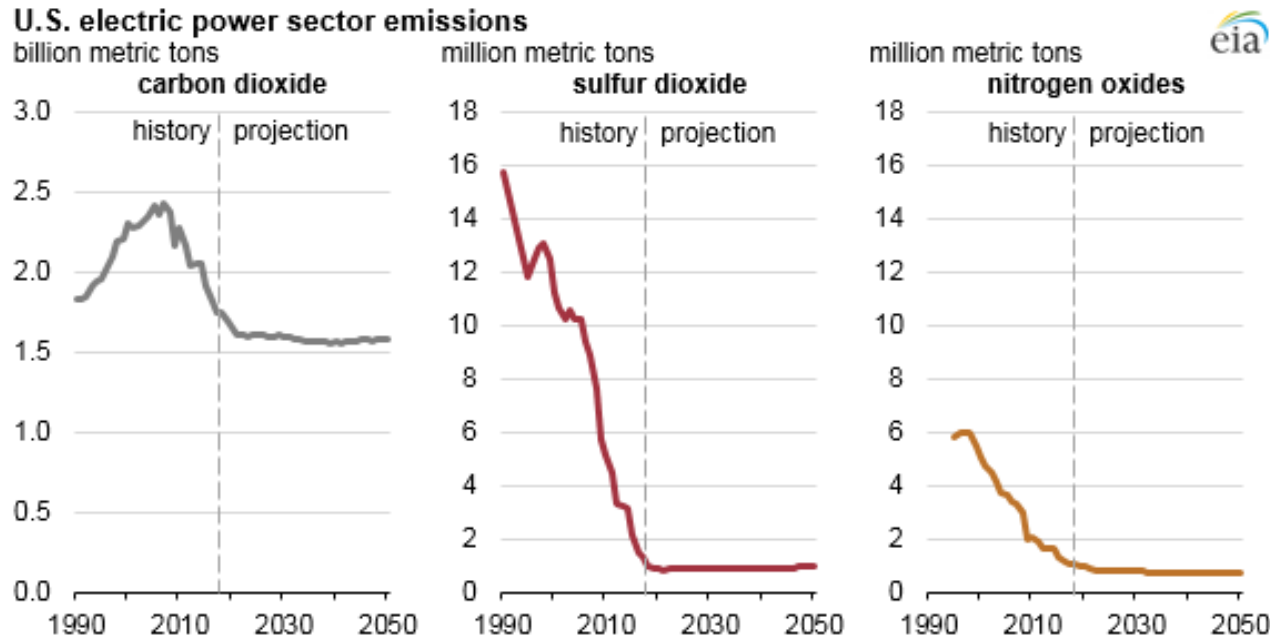


State-level sulfur dioxide (SO₂) and nitrogen oxides (NO_x) emissions (1997 and 2017)
thousand short tons



<https://www.eia.gov/todayinenergy/detail.php?id=37752>

NO_x Sources (U.S.)



EIA Annual Energy Outlook 2019 projects U.S. electric power sector emissions of **SO₂**, **NO_x** and **CO₂** will remain mostly flat through 2050 **assuming no changes to current laws and regulations.**

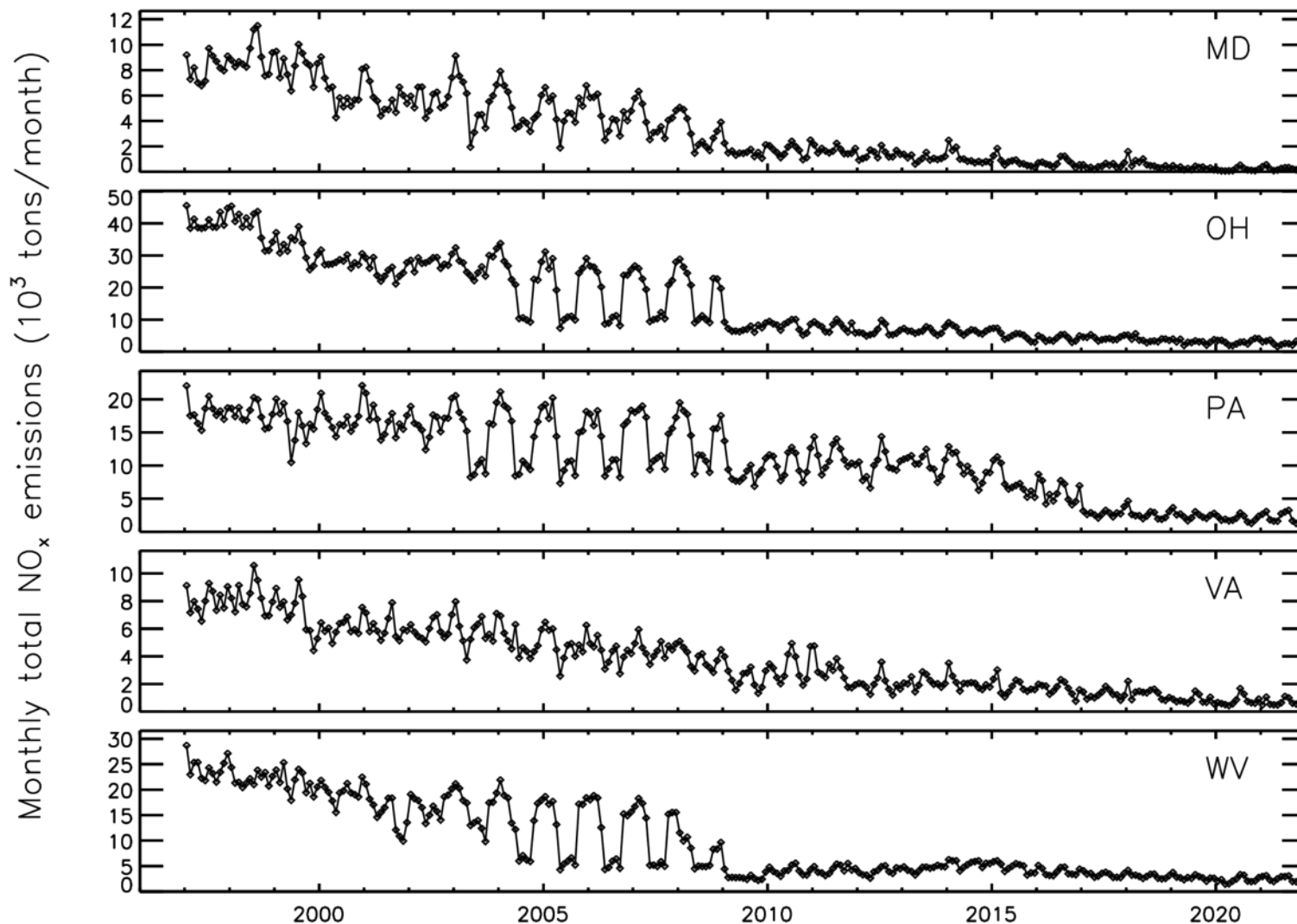
SO₂ and **NO_x** emissions from the electric power sector have declined over the past several decades, largely because of the phased implementation of regulations under the Clean Air Act Amendments of 1990. For SO₂, these regulations include acid rain cap-and-trade program deadlines in 1995 and 2000. One of the main regulations affecting NO_x emissions was the 2003 expansion of the Environmental Protection Agency's (EPA) NO_x Budget Trading Program (Title I) to include most states east of the Mississippi River.

In addition, the EPA's **Mercury** and Air Toxics Standards (MATS), announced in 2011 and implemented in 2015, required power generators to comply with emissions limits for toxic air pollutants that ... also decreased emissions of **SO₂** and **NO_x**.

These programs did not directly target emissions of **CO₂** but they did affect the economics of power plant operation cost as well as retirement decisions. Emissions of CO₂ in the U.S. power sector have been declining since a peak in 2007.

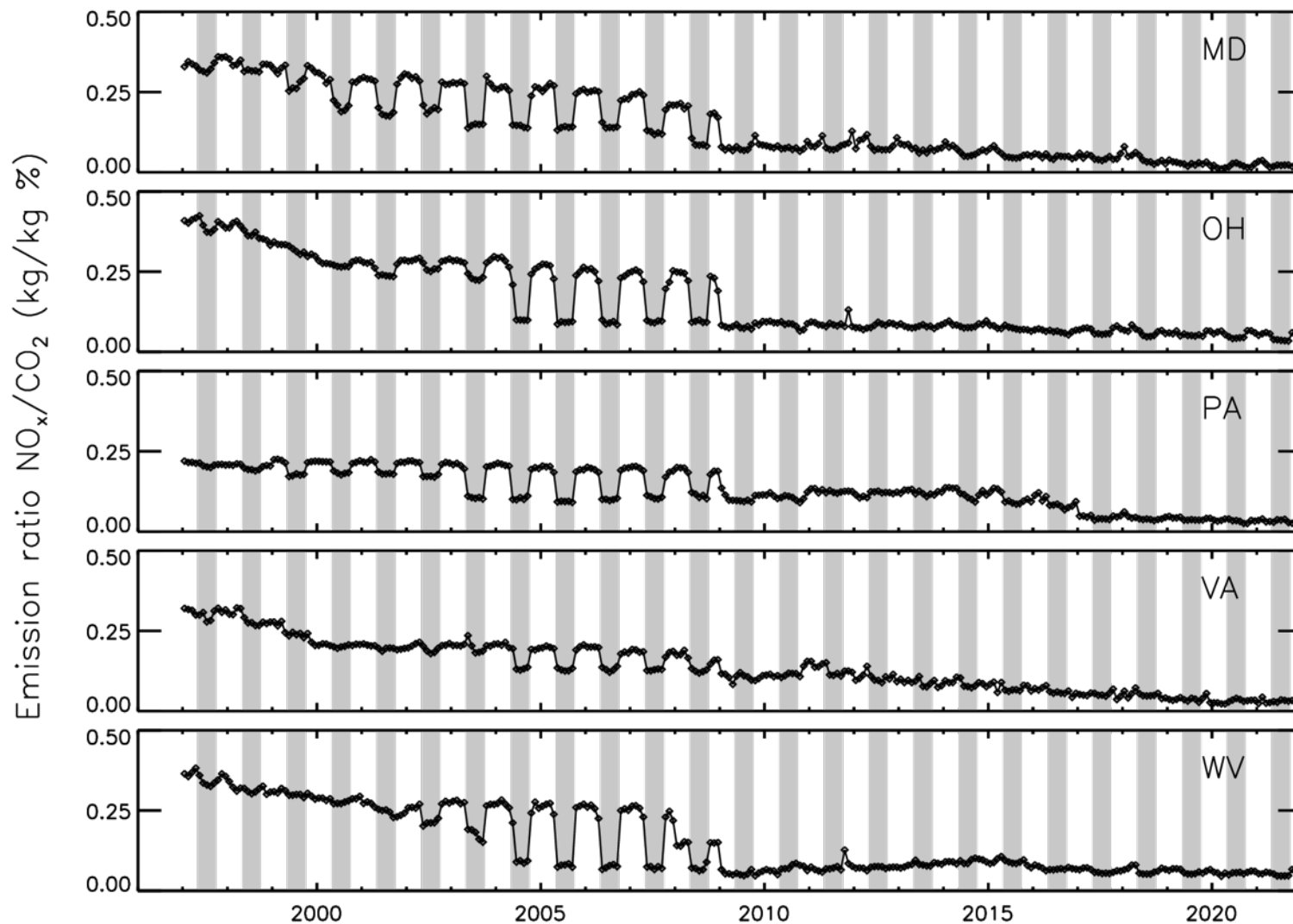
<https://www.eia.gov/todayinenergy/detail.php?id=38293>

Trends in power plant emission, region



Thanks to Doyeon Ahn for this wonderful analysis of CEMS (Continuous Emission Monitoring System) Data provided by EPA

Trends in power plant emission, region

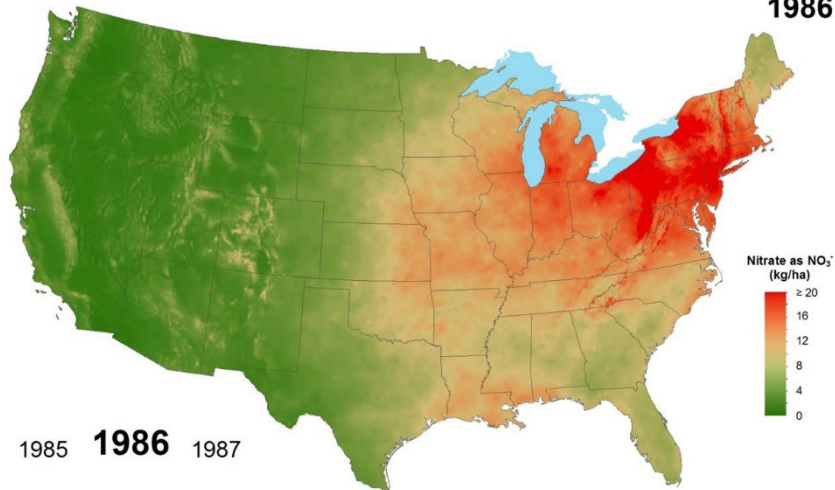


Shading denotes “ozone season”, April to Sept

Nitrate Deposition (see Fig 6.12)

1986

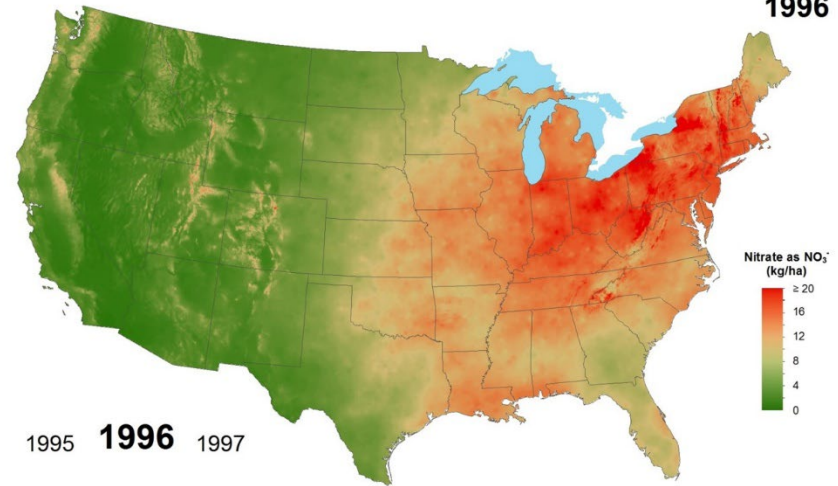
Nitrate ion wet deposition
1986



National Atmospheric Deposition Program/National Trends Network
<http://nadp.isws.illinois.edu>

1996

Nitrate ion wet deposition
1996



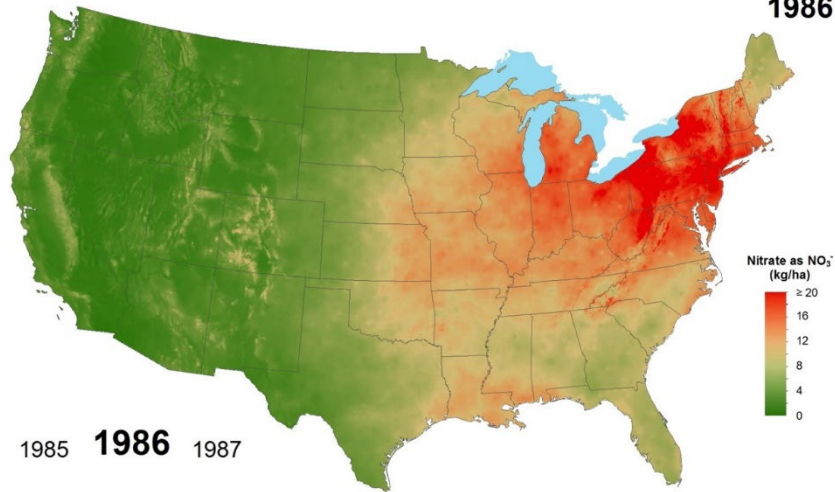
National Atmospheric Deposition Program/National Trends Network
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<http://nadp.slh.wisc.edu>

Nitrate Deposition (see Fig 6.12)

1986

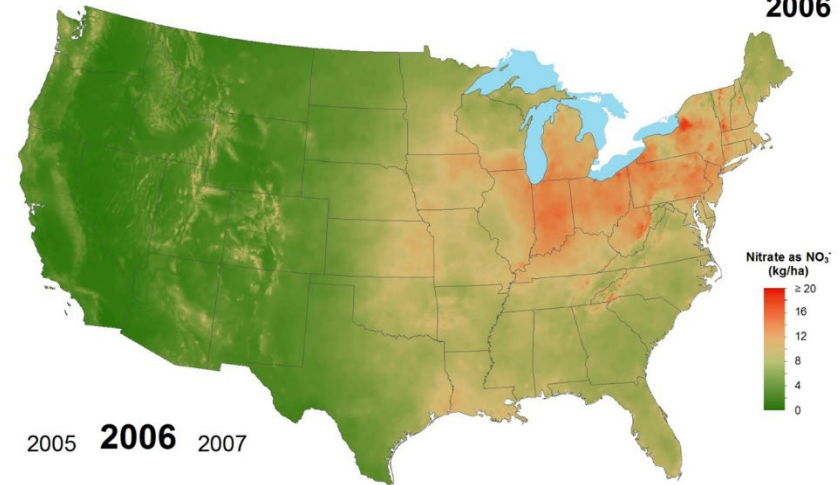
Nitrate ion wet deposition
1986



National Atmospheric Deposition Program/National Trends Network
<http://nadp.isws.illinois.edu>

2006

Nitrate ion wet deposition
2006



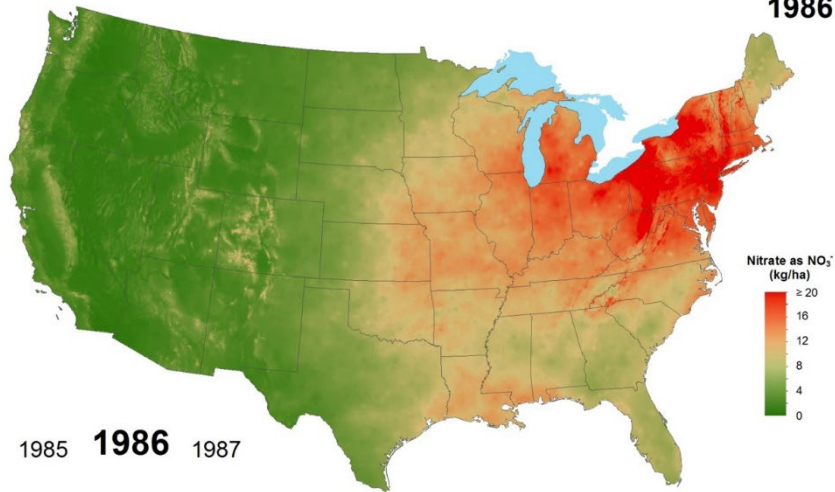
National Atmospheric Deposition Program/National Trends Network
<http://nadp.isws.illinois.edu>

<http://nadp.slh.wisc.edu>

Nitrate Deposition (see Fig 6.12)

1986

Nitrate ion wet deposition
1986



National Atmospheric Deposition Program/National Trends Network
<http://nadp.isws.illinois.edu>

2020

Nitrate ion wet deposition
2020



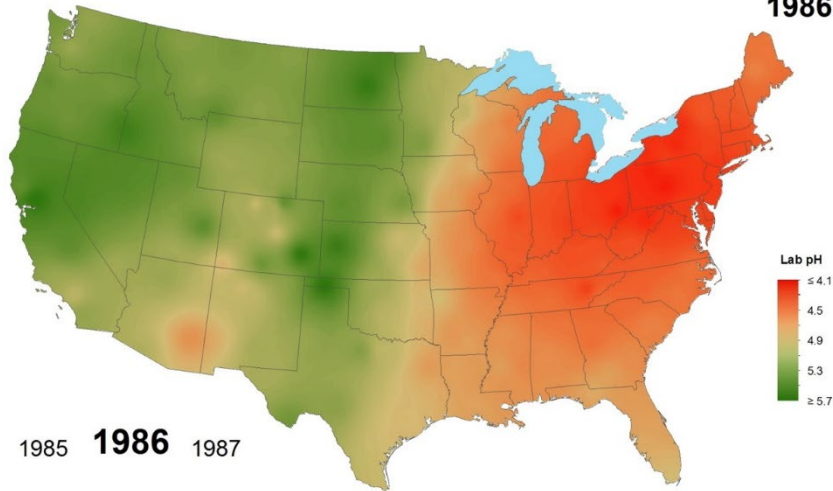
National Atmospheric Deposition Program/National Trends Network
<http://nadp.slh.wisc.edu>

<http://nadp.slh.wisc.edu>

pH of rain samples (see Fig 6.11)

1986

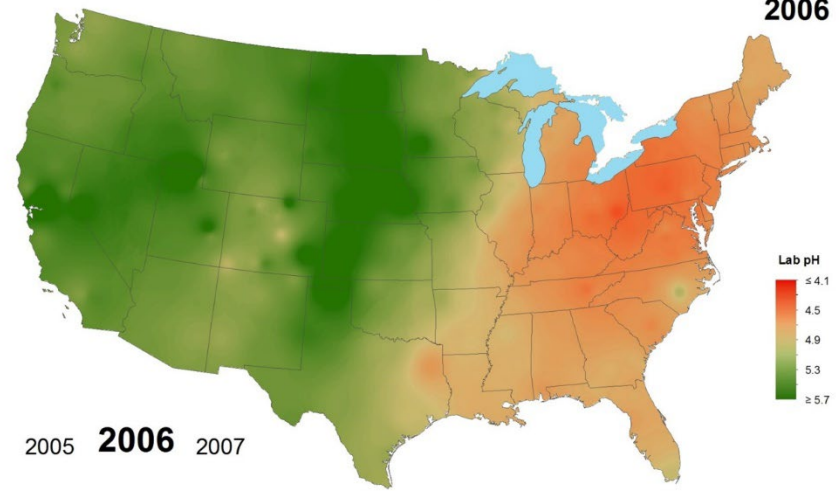
Hydrogen ion concentration as pH
1986



National Atmospheric Deposition Program/National Trends Network
<http://nadp.isws.illinois.edu>

2006

Hydrogen ion concentration as pH
2006



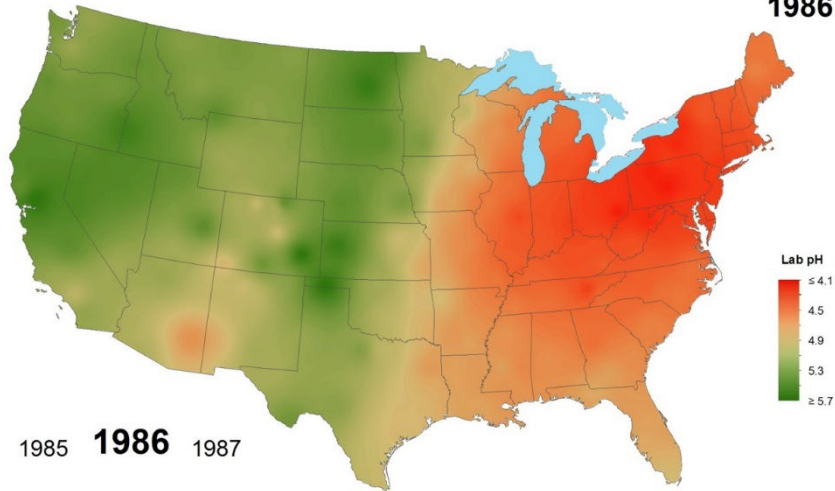
National Atmospheric Deposition Program/National Trends Network
<http://nadp.isws.illinois.edu>

<http://nadp.slh.wisc.edu>

pH of rain samples (see Fig 6.11)

1986

Hydrogen ion concentration as pH
1986

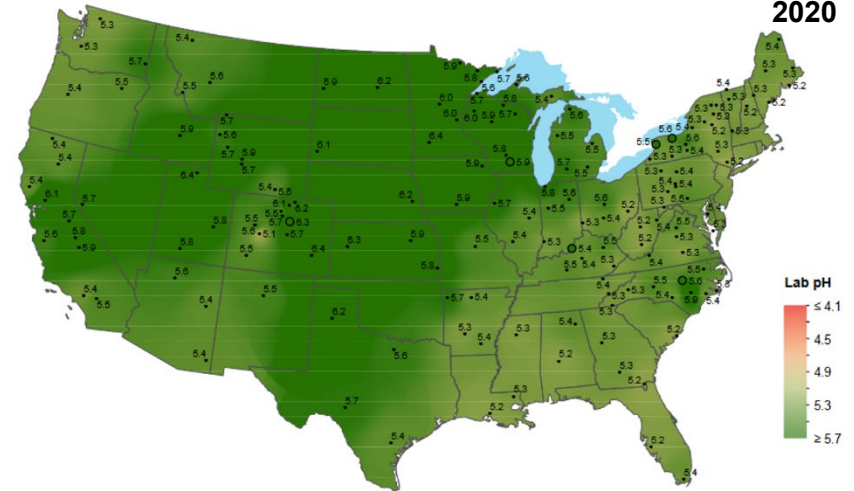


1985 **1986** 1987

National Atmospheric Deposition Program/National Trends Network
<http://nadp.isws.illinois.edu>

2020

Hydrogen ion concentration as pH
2020



National Atmospheric Deposition Program/National Trends Network
<http://nadp.slh.wisc.edu>

<http://nadp.slh.wisc.edu>

Cultural Degradation

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© Kristen Brochmann/Fundamental Photographs NYC

In 1944

At present

Figure 6.22, Chemistry in Context.
Limestone statue of George Washington, NYC

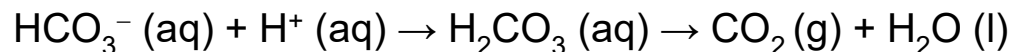
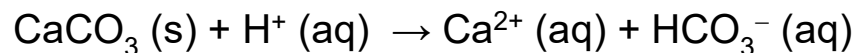
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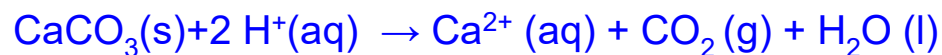
S.A. J. Copley/Visuals Unlimited

Figure 6.24, Chemistry in Context.
Mayan art, Mexico.

Marble limestone, composed mainly of calcium carbonate (CaCO_3), slowly dissolves in the presence of hydrogen ion:



or:



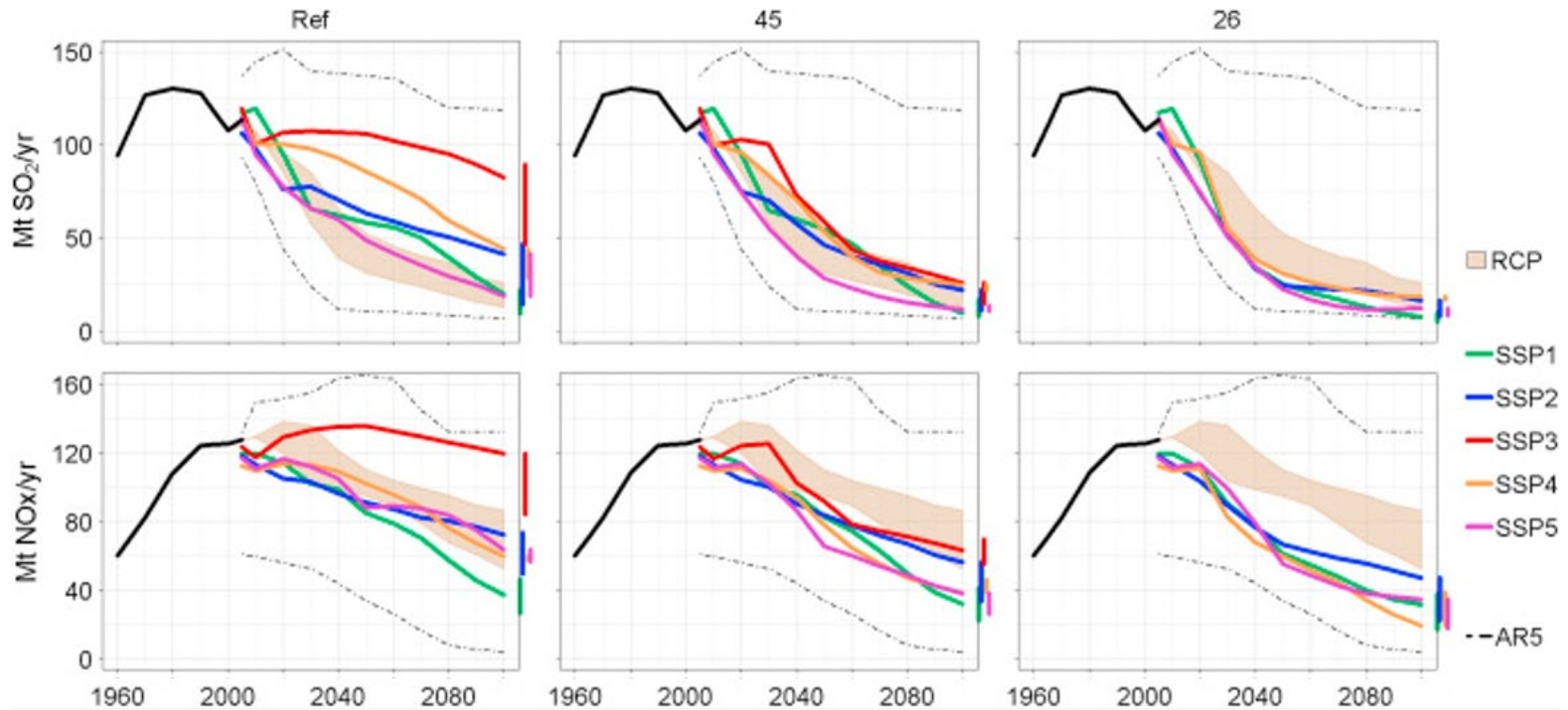
Lake Acidification

Do all lakes respond to atmospheric transport of acidic substances in the same manner?

If so, what remarkable chemical process results in this property?

If not, where are the lakes that are least sensitive to the effects of acid rain located, and what is the process that allows these lakes to be less susceptible?

Future SO₂ (top) and NO₂ (bottom) Trends



Emissions of SO₂, NO_x in SSP baseline (Ref) and 4.5 (labeled as 45) and 2.6 (labeled as 26) W/m² climate mitigation cases. Shaded area indicates range of total emissions from the RCP scenarios. Assessment Report (AR5) range refers to the full range of scenarios reviewed in the Fifth Assessment Report (AR5) of Working Group III of the Intergovernmental Panel on Climate Change (IPCC) <https://tntcat.iiasa.ac.at/AR5DB>. Vertical colored bars indicate the range of uncertainty in 2100.

Rao et al., *Glob. Envir. Change*, 2017

Overview of Aerosols

- Aerosols aka particulate matter (PM)
- Size generally ranges from 0.005 μm to 100 μm diameter
- Can be liquid or solid
- Solids: produced by grinding or crushing operation
- Liquids: formed by condensation of gases on water droplets
- Smoke or soot: carbon particles resulting from incomplete combustion
- SOA: secondary organic aerosol, formed by condensation of decomposition products of VOCs (volatile organic compounds) including isoprene (C_5H_8) which is mainly biogenic and benzene (C_6H_6) which is mainly anthropogenic
- PM can be emitted directly as carbonaceous material (primary pollutant) or formed in atmosphere upon condensation/transformation of gaseous emissions of SO_2 , NO_x , and NH_3

Eastern US: sulfates had dominated due to greater reliance on coal

Western US: carbon and nitrates dominate due to agriculture & transportation

Overview of Aerosols

- Health effects driven by size and chemical composition
- **Smaller** particles most hazardous
- Benzene-like compounds called polycyclic aromatic hydrocarbons (PAH) most hazardous

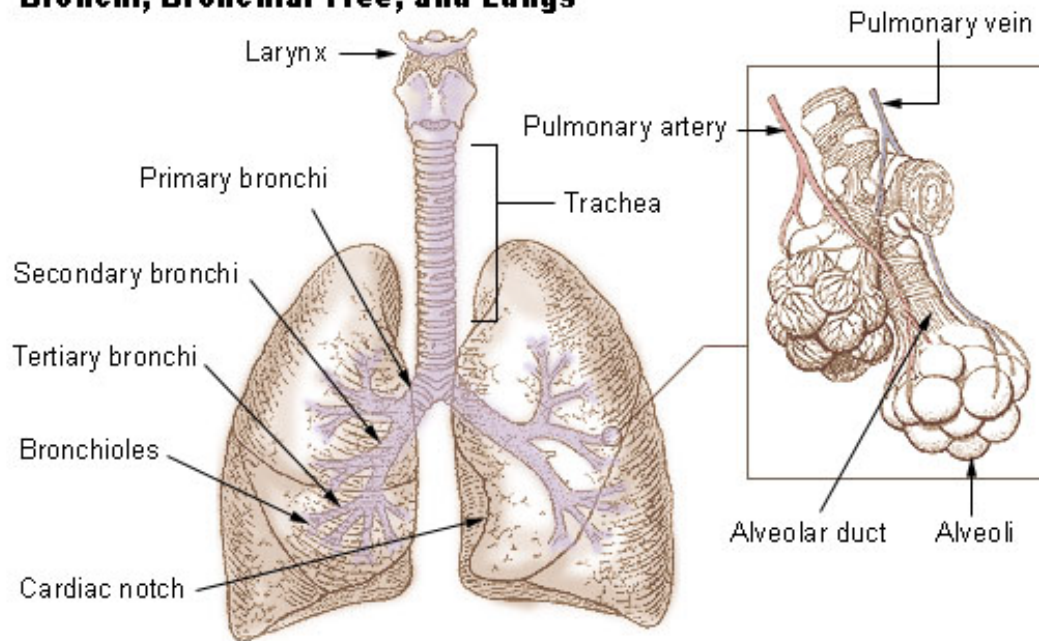


<http://www.barnesandnoble.com/w/polycyclic-aromatic-hydrocarbons-pierre-a-haines>

- Fall speed of aerosols varies as $(\text{diameter})^2$
2 μm diameter particle has **residence time** in 1 km of atmosphere of **2 months**,
if removed by only gravitational settling
 \Rightarrow small particles are suspended in the atmosphere until removed by _____ ?

Health Effects of Aerosols

Bronchi, Bronchial Tree, and Lungs



Our natural defenses help us to cough or sneeze larger particles out of our bodies.

These defenses don't work as well for particles smaller than about 10 microns (or micrometers) in diameter

Small particles get trapped in the lungs (bad) and some pass through the lungs into the bloodstream (really bad).

Exposure to elevated levels of PM leads to increase risk of respiratory illnesses, cardiopulmonary disease, heart disease, and heart attacks.

<https://www.lung.org/our-initiatives/healthy-air/outdoor/air-pollution/particle-pollution.html>

Terminal Velocity as a Function of Particle Diameter

D_p	v_t (cm/s)
100 μm	32
10 μm	0.32
1 μm	0.0032
100 nm	0.000032

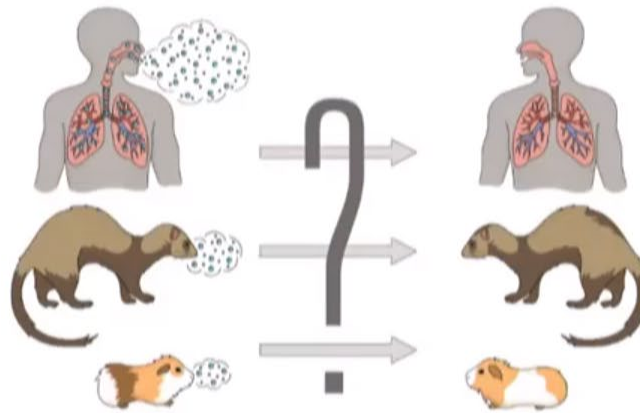
← Falls 1 m
within ~ 3 s

- Large droplets deposit pathogens on surfaces (fomites)
- SARS-CoV 2 can survive 9 DAYS on non-porous surfaces
- Inactivated within ~ 1 minute by
 - Dilute bleach
 - Dilute hydrogen peroxide
 - 62-71% ethanol

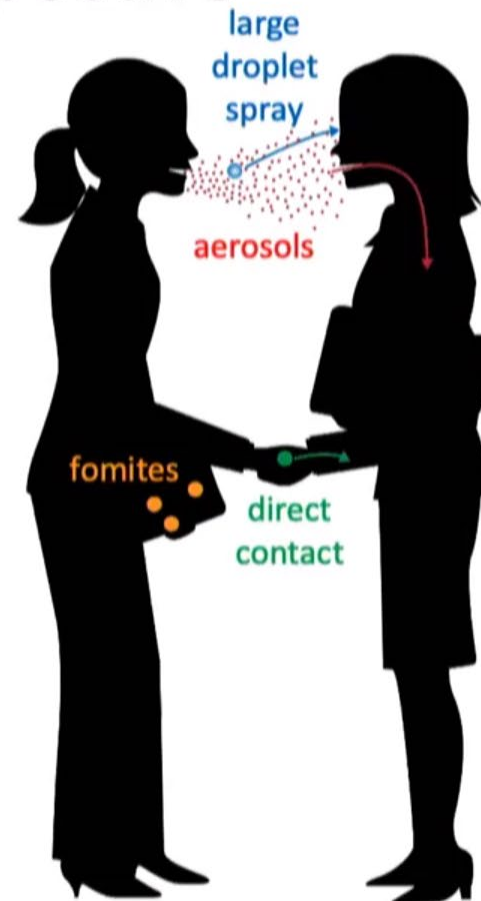
Kampf et al. *J. Hosp. Infect.* 2020 **104**(3) 246-251

<https://www.youtube.com/watch?v=9V9LgdE4W8c>

A Mechanistic Perspective



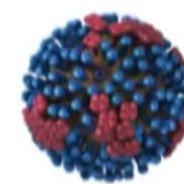
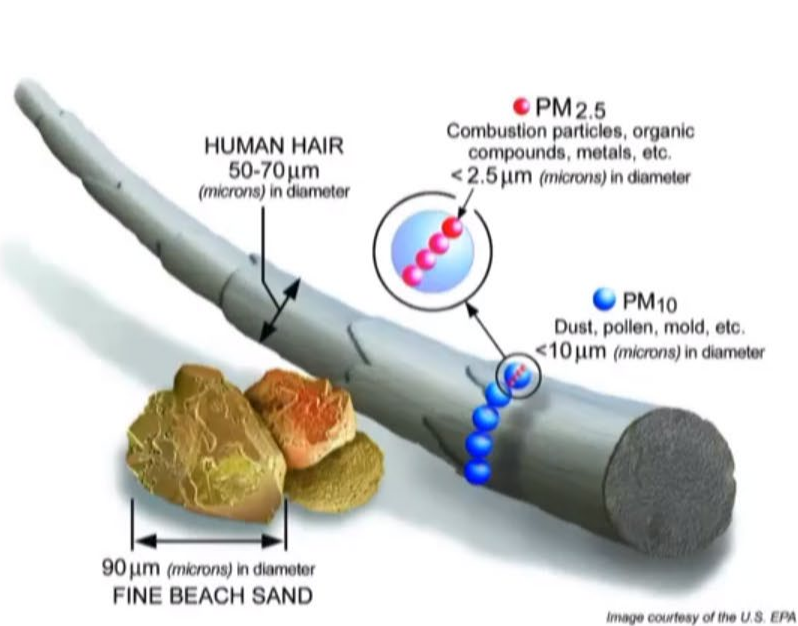
In many studies, a transmission event is observed, but we do not know the path of the virus through the environment.



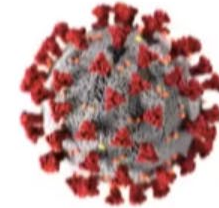
Prof. Linsey Marr, Virginia Tech

<https://www.nationalacademies.org/event/08-26-2020/airborne-transmission-of-sars-cov-2-a-virtual-workshop>

Virus Size



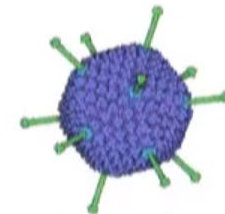
influenza
0.1 μm



SARS-CoV-2
0.12 μm



rhinovirus
0.03 μm



adenovirus
0.1 μm

<https://www.cdc.gov/flu/resource-center/freeresources/graphics/images.htm>, <http://solutionsdesignedforhealthcare.com/rhinovirus>,
<https://phil.cdc.gov/Details.aspx?pid=23312>, <https://pdb101.rcsb.org/motm/132>

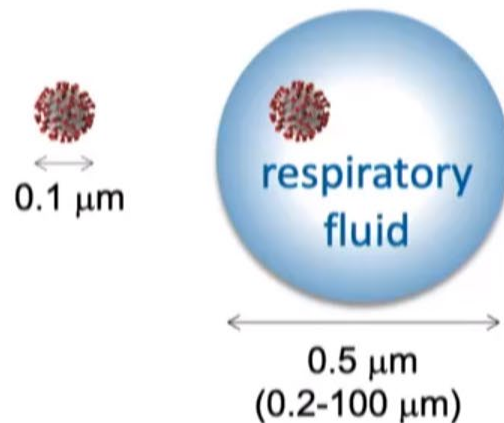


Prof. Linsey Marr, Virginia Tech

<https://www.nationalacademies.org/event/08-26-2020/airborne-transmission-of-sars-cov-2-a-virtual-workshop>

Size of Droplet/Aerosol is Critical

1. Airborne virus is not naked
2. Size of carrier droplet/aerosol defines transport



- How long it stays aloft
- How far it can travel
- How quickly it falls to surfaces
- Where it deposits in the respiratory system
- How efficiently it is removed by masks and filters
- Physics is the same for all viruses

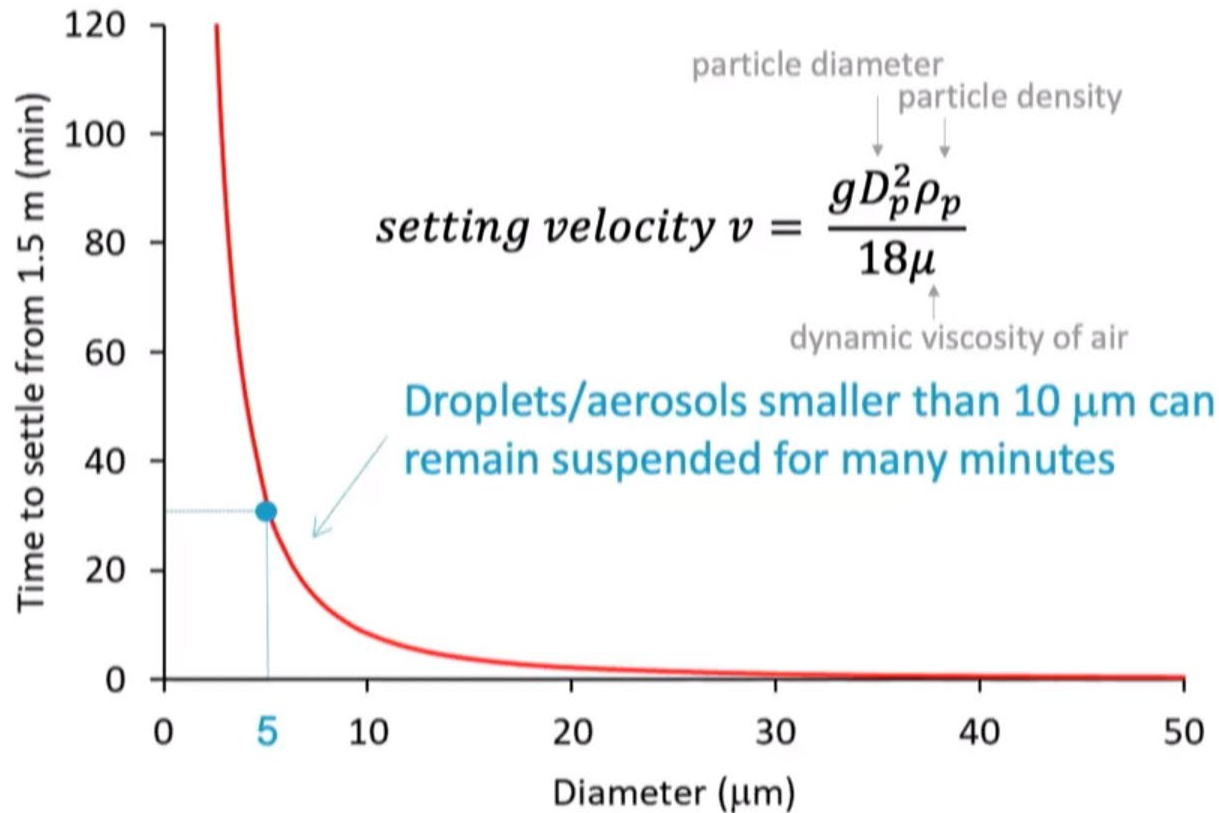
3. SARS-CoV-2 vs. measles vs. other viruses: (1) viral load in different size droplets/aerosols, (2) inactivation rate in droplets/aerosols, (3) location and dose to initiate infection



Prof. Linsey Marr, Virginia Tech

<https://www.nationalacademies.org/event/08-26-2020/airborne-transmission-of-sars-cov-2-a-virtual-workshop>

Settling Velocity and Time



Prof. Linsey Marr, Virginia Tech

<https://www.nationalacademies.org/event/08-26-2020/airborne-transmission-of-sars-cov-2-a-virtual-workshop>

Interventions

1. Source control



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Interventions

1. Source control



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Interventions

1. Source control 2. Ventilation and filtration 3. Distance and PPE 4. Hygiene



Prof. Linsey Marr, Virginia Tech

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International New York Times

Air Pollution Raises Stroke Risk

By NICHOLAS BAKALAR MARCH 24, 2015 4:30 PM 7 Comments



Air pollution — even for just one day — significantly increases the risk of stroke, a large review of studies has found.

Researchers pooled data from 103 studies involving 6.2 million stroke hospitalizations and deaths in 28 countries.

The analysis, [published online in BMJ](#), found that all types of pollution except ozone were associated with increased risk for stroke, and the higher the level of pollution, the more strokes there were.

Daily increases in pollution from nitrogen dioxide, sulfur dioxide, carbon monoxide and particulate matter were associated with corresponding increases in strokes and hospital admissions. The strongest associations were apparent on the day of exposure, but increases in particulate matter had longer-lasting effects.

The exact reason for the effect is unclear, but studies have shown that air pollution can constrict blood vessels, increase blood pressure and increase the risk for blood clots. Other research has tied air pollution to a higher risk of heart attacks, stroke and other ills.

<http://well.blogs.nytimes.com/2015/03/24/air-pollution-raises-stroke-risk>

BMJ: British Medical Journal

Short term exposure to air pollution and stroke: systematic review and meta-analysis

Anoop S V Shah,¹ Kuan Ken Lee,¹ David A McAllister,² Amanda Hunter,¹ Harish Nair,² William Whiteley,³ Jeremy P Langrish,¹ David E Newby,¹ Nicholas L Mills¹

¹BHF/University Centre for Cardiovascular Science, University of Edinburgh, Edinburgh EH16 4SB, UK

²Centre of Population Health Sciences, University of Edinburgh, Edinburgh, UK

³Centre for Clinical Brain Sciences, University of Edinburgh, Edinburgh, UK

Admission to hospital for stroke or mortality from stroke was associated with an increase in concentrations of carbon monoxide (relative risk 1.015 per 1 ppm, 95% confidence interval 1.004 to 1.026), sulphur dioxide (1.019 per 10 ppb, 1.011 to 1.027), and nitrogen dioxide (1.014 per 10 ppb, 1.009 to 1.019). Increases in PM_{2.5} and PM₁₀ concentration were also associated with admission and mortality (1.011 per 10 $\mu\text{g}/\text{m}^3$ (1.011 to 1.012) and 1.003 per 10 $\mu\text{g}/\text{m}^3$ (1.002 to 1.004), respectively).

Gaseous and particulate air pollutants have a marked and close temporal association with admissions to hospital for stroke or mortality from stroke. Public and environmental health policies to reduce air pollution could reduce the burden of stroke.

The lead author, Dr. Anoop Shah, a lecturer in cardiology at the University of Edinburgh, said that there was little an individual can do when air pollution spikes. "If you're elderly, or have co-morbid conditions, you should stay inside," he said. But policies leading to cleaner air would have the greatest impact, he said. "It's a question of getting cities and countries to change."

NOVAXNEXT

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Posted by Caleb Finch and Jiu-Chiuan Chen on
Tue, 28 Feb 2017

[Air Pollution Exposure May Increase Risk of Dementia](#)

We designed this study to answer three broad questions. First, we wanted to know whether older people living in locations with higher levels of outdoor PM_{2.5} have an increased risk for cognitive impairment, especially dementia. We also wanted to know whether people who carry the high-risk gene for Alzheimer's disease, APOE₄, are more sensitive to the damage potentially caused by long-term exposure to PM_{2.5} in the air.

We focused on older women and female mice because APOE₄ confers a greater Alzheimer's disease risk in women than in men.

We found that women exposed to higher levels of PM_{2.5} had faster rates of cognitive decline and a higher risk of developing dementia. Older women living in places where PM_{2.5} levels exceeded the U.S. Environmental Protection Agency's standard had an 81% greater risk of global cognitive decline and were 92% more likely to develop dementia, including Alzheimer's. This environmental risk raised by long-term PM_{2.5} exposure was two to three times higher among older women with two copies of the APOE₄ gene, compared with women who had only the background genetic risk with no APOE₄ gene.

<http://www.pbs.org/wgbh/nova/next/body/air-pollution-exposure-may-increase-risk-of-dementia/>

Cultural Degradation



\$25 million dollar restoration of the Lincoln Memorial began in 2016

Will repair cracks in marble due to 2011 earthquake, install a new roof, and also patch a “baseball-size gouge of the penthouse’s ornate outer wall caused by an errant anti-aircraft bullet in 1942”. During World War II, a gun was set up near a local bridge to defend D.C. against air attack. A soldier accidentally pulled the trigger, hitting the memorial’s east side.

Work should be completed by 2022, in time for the memorial’s centennial.

<https://bangordailynews.com/2018/06/14/news/nation/battered-by-time-nature-and-anti-aircraft-fire-lincoln-memorial-gets-facelift>

<https://www.youtube.com/watch?v=pBo2PSF2Pvg>

Carbon Water Chemistry

Acidity of pure water is 7. This means $[H^+] = 10^{-7}$ moles/liter or 10^{-7} M.

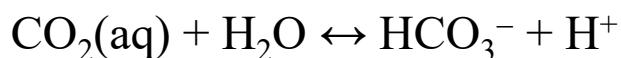
What is acidity of water in equilibrium with atmospheric CO_2 ?

$$[CO_2(aq)] = H_{CO_2} p_{CO_2} = 3.4 \times 10^{-2} \text{ M / atm } p_{CO_2}$$

For $CO_2 = 400$ ppm:

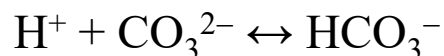
$$[CO_2(aq)] = 3.4 \times 10^{-2} \text{ M / atm } \times 4.0 \times 10^{-4} \text{ atm} = 1.36 \times 10^{-5} \text{ M}$$

First equilibrium between CO_2 , HCO_3^- (bicarbonate), and H^+



$$K_1 = \frac{[HCO_3^-][H^+]}{[CO_2(aq)]} = 4.3 \times 10^{-7} \text{ M (at 298 K)}$$

Second equilibrium between CO_3^{2-} (carbonate), HCO_3^- , and H^+



$$K_2 = \frac{[CO_3^{2-}][H^+]}{[HCO_3^-]} = 4.7 \times 10^{-11} \text{ M (at 298 K)}$$

**Can solve if we assume charge balance: $[H^+] = [HCO_3^-] + 2 [CO_3^{2-}]$
- or – by taking a short-cut (see next slide)**

Carbon Water Chemistry

Acidity of pure water is 7. What is acidity of water in equilibrium with atmospheric CO₂ ?

Shortcut:

$$[\text{CO}_2(\text{aq})] = 1.36 \times 10^{-5} \text{ M for present atmosphere}$$

$$[\text{H}^+] [\text{HCO}_3^-] = K_1 [\text{CO}_2(\text{aq})] = 4.3 \times 10^{-7} \text{ M} \times 1.36 \times 10^{-5} \text{ M} = 5.85 \times 10^{-12} \text{ M}^2$$

Assume charge balance is primarily between [H⁺] and [HCO₃⁻]:

i.e., that [H⁺] ≈ [HCO₃⁻] and that both are >> [CO₃²⁻]

$$[\text{H}^+] [\text{H}^+] = 5.85 \times 10^{-12} \text{ M}^2 \Rightarrow [\text{H}^+] = 2.418 \times 10^{-6} \text{ M}$$

$$\textcolor{blue}{pH} = -\log_{10} [\text{H}^+] = \textcolor{blue}{5.6} \text{ (400 ppm, 298 K)}$$

Is the *assumption* justified? :

$$[\text{CO}_3^{2-}] = K_2 [\text{HCO}_3^-] / [\text{H}^+] \approx 4.7 \times 10^{-11} \text{ M}$$

$$[\text{H}^+] \text{ \& \ } [\text{HCO}_3^-] \text{ are both } \sim 2.4 \times 10^{-6} \text{ M which is } \gg 4.7 \times 10^{-11} \text{ M}$$

As noted in class, the actual ocean is basic. The net charge from a series of **cations** (positively charged ions) and minor **anions** (negatively charged ions) is balanced by the total negative charge of the bicarbonate and carbonate ions. We write:

$$[\text{Alk}] = [\text{HCO}_3^-] + 2 [\text{CO}_3^{2-}] = [\text{Na}^+] + [\text{K}^+] + 2[\text{Mg}^{2+}] + 2[\text{Ca}^{2+}] - [\text{Cl}^-] - [\text{Br}^-] - 2 [\text{SO}_4^{2-}] + \dots$$

where Alk stands for Alkalinity

Henry's Law and the equations for the first and second dissociation constants yield:

$$p\text{CO}_2(\text{vmr}) = \frac{[\text{CO}_2(\text{aq})]}{\alpha} \quad K_1 = \frac{[\text{HCO}_3^-][\text{H}^+]}{[\text{CO}_2(\text{aq})]} \quad K_2 = \frac{[\text{CO}_3^{2-}][\text{H}^+]}{[\text{HCO}_3^-]}$$

The three equations above can be re-arranged to yield: $p\text{CO}_2(\text{vmr}) = \left(\frac{K_2}{\alpha K_1} \right) \frac{[\text{HCO}_3^-]^2}{[\text{CO}_3^{2-}]}$

If we substitute $[\text{HCO}_3^-] = \text{Alk} - 2 [\text{CO}_3^{2-}]$ into the eqn above, we arrive at a quadratic eqn for $[\text{CO}_3^{2-}]$ as a function of $p\text{CO}_2$ and Alk. Note that α , K_1 , and K_2 vary as a function of temperature (T) and ocean salinity (S) (<http://en.wikipedia.org/wiki/Salinity>)

If T, Alk, & S are specified, it is straightforward to solve for $[\text{CO}_3^{2-}]$ from the quadratic eqn.

Values for $[\text{CO}_2(\text{aq})]$, $[\text{HCO}_3^-]$, and $[\text{H}^+]$ are then found from Henry's law & the dissoc eqns.

Finally, Ocean Carbon is found from $[\text{CO}_2(\text{aq})] + [\text{HCO}_3^-] + [\text{CO}_3^{2-}]$.

Numerical values on the slides entitled "Uptake of Atmospheric CO_2 by Oceans" were found in this manner, using Fortran program http://www.atmos.umd.edu/~rjs/class/code/ocean_carbon.f